# Preliminary Information

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Welcome: Texas Medicaid Provider Procedures Manual

This manual is a comprehensive guide for Texas Medicaid providers. It contains information about Texas Medicaid fee-for-service benefits, policies, and procedures including medical, dental, and children’s services benefits.

Refer to: The Medicaid Managed Care Handbook (Vol. 2, Provider Handbooks) for information about the Medicaid Managed Care, which is administered by Texas Health and Human Services Commission (HHSC)-contracted managed care organizations (MCOs), dental managed care organizations, and behavioral health organizations (BHOs) across the state.

The Texas Medicaid Provider Procedures Manual is updated monthly on the TMHP website at www.tmhp.com to include revisions to policies and procedures that went into effect in the prior month. The manual is available in portable document format (PDF) as a complete book and as individual sections and handbooks. A hypertext markup language (HTML) version is also be available.

The current version of the manual always appears prominently on the Texas Medicaid Provider Procedures Manual web page. All previously published annual editions of the Texas Medicaid Provider Procedures Manual have been archived. Users can access the archives through links on the Texas Medicaid Provider Procedures Manual web page.

Providers can determine what has changed each month by following the Release Notes link on the Texas Medicaid Provider Procedures Manual web page. The release notes include the sections and handbooks that have changed for the current month and the nature of the changes. Most changes have been previously announced in news articles on the TMHP website, and, where appropriate, the release notes link to prior website articles.

Publishing the manual monthly has eliminated the need for the Texas Medicaid Bulletin, which was discontinued following the publication of the September/October 2012 Texas Medicaid Bulletin, No. 243. Special bulletins, such as the annual Healthcare Common Procedure Coding System (HCPCS) bulletin, which is published in January of each year, will continue to be published on an as-needed basis.

The Texas Medicaid Provider Procedures Manual is divided into two volumes as follows:

- **Volume I: General Information**
  
  Volume 1 applies to all health-care providers who are enrolled in Texas Medicaid and provide services to Texas Medicaid fee-for-service clients. The sections in Volume 1 include general information for enrolling in the program, receiving appropriate reimbursement, prior authorizations, claim submissions and appeals for services rendered.

- **Volume 2: Provider Handbooks**
  
  Each handbook in Volume 2 covers Medicaid policies, procedures, and claims filing requirements for specific products or services. Volume 2 includes the following handbooks:
  
  - Ambulance Services Handbook
  - Behavioral Health, Rehabilitation, and Case Management Services Handbook
  - Certified Respiratory Care Practitioners (CRCP) Services Handbook
  - Children’s Services Handbook
  - Clinics and Other Outpatient Facility Services Handbook
  - Durable Medical Equipment, Medical Supplies, and Nutritional Products Handbook
  - Gynecological, Obstetrics, and Family Planning Title XIX Services
  - Health and Human Services Commission Family Planning Program Services Handbook
  - Healthy Texas Women Program Handbook
• Home Health Nursing and Private Duty Nursing Services Handbook
• Inpatient and Outpatient Hospital Services Handbook
• Medicaid Managed Care Handbook
• Medical and Nursing Specialists, Physicians, and Physician Assistants Handbook
• Medical Transportation Program Handbook
• Outpatient Drug Services Handbook
• Physical Therapy, Occupational Therapy, and Speech Therapy Services Handbook
• Radiology and Laboratory Services Handbook
• Telecommunications Services Handbook
• Vision and Hearing Services Handbook

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Introduction

Texas Medicaid Administration

The Texas Medical Assistance (Medicaid) Program was implemented on September 1, 1967, under the provisions of Title XIX of the federal Social Security Act and Chapter 32 of the Texas Human Resources Code.

The state of Texas and the federal government share the cost of funding Texas Medicaid. The Health and Human Services Commission (HHSC), the single state Medicaid agency, is responsible for the Title XIX Program. The administration of the program is accomplished through contracts and agreements with the following:

- Medical providers
- Texas Medicaid & Healthcare Partnership (TMHP), the fee-for-service claims administrator
- MAXIMUS, the enrollment broker
• Various managed care organizations (MCOs) and dental managed care organization (dental plans), that administer Medicaid Managed Care benefits.

• The Institute for Child Health Policy (ICHP), the quality monitor

• State agencies

Texas Medicaid providers are reimbursed for services through contracts with health-insuring contractors, fiscal agents, or direct vendor payments.

By signing an HHSC Medicaid Provider Agreement (through the enrollment process) and submitting Medicaid claims, each enrolled provider agrees to abide by the policies and procedures of Medicaid, published regulations, and the information and instructions in manuals, bulletins, and other instructional material furnished to the provider.

Refer to: “Appendix A: State, Federal, and TMHP Contact Information” (Vol. 1, General Information) for addresses and telephone numbers of HHSC and Department of State Health Services (DSHS) regional offices.
SECTION 1: PROVIDER ENROLLMENT AND RESPONSIBILITIES

TEXAS MEDICAID PROVIDER PROCEDURES MANUAL: VOL. 1

MAY 2021
## SECTION 1: PROVIDER ENROLMENT AND RESPONSIBILITIES

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1.1 **Provider Enrollment**

To be eligible for Texas Medicaid reimbursement, a provider of health-care services (including an out-of-state provider) must:

- Meet all applicable eligibility criteria.
- Be approved by the Texas Health and Human Services Commission (HHSC) for enrollment.
- Obtain a National Provider Identifier (NPI) from the National Plan and Provider Enumeration System (NPPES).

**Refer to:** Subsection 1.1.2, “NPI and Taxonomy Codes” in this section.

- File with the Texas Medicaid & Healthcare Partnership (TMHP) the required Texas Medicaid enrollment application ensuring that the application is correct, complete, and includes all required attachments and additional information.
- Provide any additional information requested by TMHP, HHSC, or the HHSC Office of Inspector General (OIG) in connection with the processing of the application.
- Be approved by HHSC for enrollment and enter into a written provider agreement with HHSC.

Providers can use the online provider enrollment on the portal (PEP) tool to enroll electronically through the TMHP website at [www.tmhp.com](http://www.tmhp.com).

**Refer to:** Subsection 1.1.3, “Online Enrollment” in this section.

Paper versions of the enrollment applications are also available for download from the Forms section of the TMHP website.

**Refer to:** Subsection 1.1.4, “Paper Application Enrollment” in this section.

After receipt of all information necessary to process the application, the entire application process can typically take up to 60 days. This may be extended in special circumstances. Requests for exceptions to the enrollment process, risk category, and provider types that require additional state approval may extend the length of the application process.

All providers must be enrolled in Texas Medicaid before enrollment can be approved for any other service or program, including, but not limited to, Medicaid managed care.

Certain provider types are required to enroll in Medicare as a prerequisite for enrolling in Texas Medicaid. During the Texas Medicaid enrollment process, with HHSC approval, the Claims Administrator may waive the mandatory prerequisite for Medicare enrollment for certain providers whose type of practice will never serve Medicare-eligible individuals (e.g., pediatrics, obstetrician/gynecologist [OB/GYN]).

Providers must maintain a valid, current license or certification to be entitled to Texas Medicaid reimbursement. Providers cannot enroll in Texas Medicaid if their license or certification is due to expire within 30 days of application. A current license or certification must be submitted, if applicable.

**Refer to:** Subsection 1.1.10.11, “Copy of License, Temporary License, or Certification” in this section.

A provider identifier is issued when a determination has been made that a provider qualifies for participation.

**Refer to:** Subsection 1.10, “Enrollment Criteria for Out-of-State Providers” in this section for additional criteria that out-of-state providers must meet to enroll in Texas Medicaid.
1.1.1 Provider Enrollment Revalidation Requirements

To remain in compliance with Title 42 Code of Federal Regulations §455.414, providers must complete the revalidation process before the end of their enrollment period, which is also called the “revalidation date.” If a provider’s enrollment period ends before TMHP has received the provider’s revalidation application, TMHP automatically disenrolls providers on the date that their enrollment period ends. Providers can view and confirm their revalidation date and enrollment information on the Provider Information Management System (PIMS). Most providers have an enrollment period of 5 years. Some providers have shorter enrollment periods, which are based on risk categories and other considerations.

Providers can only submit the revalidation application online. Providers that are unable to revalidate using PEP must download and submit the appropriate paper enrollment application.

Some of the fields on the revalidation application cannot be altered. To avoid application processing delays, providers should update the following information before submitting a revalidation application:

- First and last name
- Organization name
- Social Security number
- Date of birth
- Employer’s Tax Identification Number (Tax ID) and legal name

Providers can update this information using PIMS or the Provider Information Change (PIC) Form, which can be found on the TMHP website at www.tmhp.com. Providers will experience delays if they do not submit the updated information before starting the revalidation process.

After the information has been confirmed and updated, providers should submit their revalidation applications as soon as they are available in PEP so that the process can be completed before the enrollment period ends. Providers must allow 30 business days from the date on which TMHP receives the form for the changes to take effect before they can complete a revalidation application.

If providers do not complete the revalidation process before the end of their enrollment period, they will be disenrolled. Their claims will not be paid, and their prior authorization requests will be denied. They will not be eligible to participate as a network provider in Medicaid MCOs.

1.1.2 NPI and Taxonomy Codes

The NPI final rule, Federal Register 45, Code of Federal Regulations (CFR) Part 162, established the NPI as the standard unique identifier for health-care providers and requires covered health-care providers, clearinghouses, and health plans to use this identifier in Health Insurance Portability and Accountability Act (HIPAA)-covered transactions. An NPI is a 10-digit number assigned randomly by the NPPES. An NPI must be obtained before a provider can enroll as a Texas Medicaid provider.

An NPI is not required for enrollment for certain provider types; however, the provider must submit a signed letter on company letterhead that attests that they are not a health-care provider and are unable to obtain an NPI.

The Health Care Provider Taxonomy Code Set is an external, non-medical collection of alphanumeric codes designed to classify health-care providers by provider type and specialty. Providers may have more than one taxonomy code. (Taxonomy codes can be obtained from the Washington Publishing Company website at www.wpc-edi.com).

During the enrollment process, providers must select a primary and, if applicable, secondary taxonomy code associated with their provider type. Providers will be supplied a list of taxonomy codes to choose from that correspond to the services rendered by the type of provider they wish to enroll as. Only the
code will be displayed. Due to copyright laws, TMHP is unable to publish the taxonomy description. Therefore, providers must verify the taxonomy code associated with their provider type and specialty before beginning the online attestation process.

### 1.1.3 Online Enrollment

The following provider types can use PEP to enroll and reenroll:

- Texas Medicaid providers (except medical transportation providers)
- Home and Community Based Services – Adult Mental Health (HCBS-AMH) providers
- Ordering/referring-only providers
- Children with Special Health Care Needs (CSHCN) Services Program providers (except hospice and medical food providers)
- Texas Health Steps (THSteps) dental services provider, including fee-for-service dental providers and managed care
- THSteps medical checkup providers

**Important:** When completing the Texas Medicaid provider application online, qualified providers can choose to opt in as Texas Health Steps Medical checkup providers and CSHCN Services Program providers. Providers who do not opt in when completing the Texas Medicaid provider application online may enroll in the CSHCN Services Program and in Texas Health Steps at a future date using the online PEP tool.

Providers can begin the enrollment process on the [Provider Enrollment](#) page of the TMHP website.

Online enrollment has the following advantages:

- Applications are validated immediately to ensure that all fields have been completed.
- Most of the application can be completed online so that only a few forms need to be printed, completed, and mailed to TMHP.
- Applicants can view both incomplete and complete applications that have been submitted online.
- Some form fields are automatically completed, reducing the amount of information that has to be entered.
- Providers can complete the Provider Information Change (PIC) form online.
- Providers will receive email notifications when messages or deficiency notices about their applications are posted online. The messages can be viewed on the secured access portion of the website. Providers may opt out of email communication and receive messages or deficiency letters by mail.
- Providers can create templates, which make it easier to submit multiple enrollment applications.
- Providers who enroll as a group can assign portions of the application to performing providers to complete. Performing providers can complete their portion of a group application by logging into the online PEP tool with their unique user name and password.
- Providers can navigate to completed sections of the application without having to click through all pages of the application.
- Information that is on file for owners and subcontractors of the applying provider are auto-populated in the application.

Before submitting an application to TMHP for processing, providers are required to review a portable document format (PDF) copy of the application and verify it is complete. Providers are able to edit submitted applications to correct identified deficiencies.
Enrolling online promotes accurate submissions, decreases processing time, and enables immediate feedback on the status of the application.

### 1.1.4 Paper Application Enrollment

As an alternative to applying for enrollment online, a provider may file a paper enrollment application with TMHP. Providers may download the Texas Medicaid Provider Enrollment Application at [www.tmhp.com](http://www.tmhp.com) or request a paper application form by contacting TMHP directly at 1-800-925-9126. Enrollment applications are updated periodically. When an application has been updated, the older version will no longer be accepted and will not be available on the website. It is recommended that the provider check the website regularly for updates and notifications.

A paper enrollment application may also be requested from and must be submitted to the following address:

Texas Medicaid & Healthcare Partnership  
Provider Enrollment  
PO Box 200795  
Austin, TX 78720-0795

**Note:** The Texas Medicaid Provider Enrollment Application must include the accounting/mailing address and the physical address where the provider renders services to Medicaid clients.

**Important:** Qualified providers that want to enroll in Texas Health Steps Medical or CSHCN Services Program must check the appropriate additional program enrollment box on the paper enrollment application. If Medicaid providers choose to enroll in Texas Health Steps Medical or CSHCN Services Program at a later date a separate application must be submitted.

### 1.1.5 Provider Enrollment Identification

Providers are required to identify the type of entity for which they are requesting enrollment. Providers can choose from one of the following on each application they submit (only one per application is allowed):

- **Individual.** This type of enrollment applies to an individual health-care professional who is licensed or certified in Texas, and who is seeking enrollment under the name, and Social Security number or federal tax identification number of the individual. An individual may also enroll as an employee, using the federal tax identification number of the employer. Certain provider types must enroll as individuals, including dieticians, licensed vocational nurses, occupational therapists, registered nurses, and speech-language pathologist.

- **Group.** This type of enrollment applies to health-care items or services provided under the auspices of a legal entity, such as a partnership, corporation, limited liability company, or professional association, and the individuals providing health-care items or services are required to be certified or licensed in Texas. The enrollment is under the name and federal tax identification number of the legal entity. For any group enrollment application other than as a THSteps medical checkup provider group, there must also be at least one enrolling performing provider. THSteps providers are only enrolled at the group level.

During the PEP process, the available taxonomy code list is populated with either taxonomy code 193200000X or 193400000X for a clinic/group practice, depending on which specialty is chosen. The taxonomy codes for clinic/group practice providers are accurate and have been approved by HHSC. The most appropriate taxonomy codes should be selected for any performing providers that will be enrolled according to their specific performing provider type and specialty.

- **Performing provider.** This type of enrollment applies to an individual health care professional who is licensed or certified in Texas, and who is seeking enrollment under a group. The enrollment is under the federal tax identification number of the group, and payment is made to the group. If a health-care professional is required to enroll as an individual, as explained above, but the person is
an employee and payment is to be made to the employer, the health-care professional does not enroll as a performing provider. Instead, the health-care professional enrolls as an individual provider under the federal tax identification number of their employer.

- **Facility.** This type of enrollment applies to situations in which licensure or certification applies to the entity. Although individuals working for or with the entity may be licensed or certified in their individual capacity, the enrollment is based on the licensure or certification of the entity or the supervising licensed practitioner who is assuming responsibility for the facility’s operation. For this reason, facility enrollment does not require enrollment of performing providers. However, certain provider types must enroll as facilities, including the following:
  - Ambulance and air ambulance
  - Ambulatory surgical center (ASC) and hospital-based ambulatory surgical center (HASC)
  - Birthing center
  - Catheterization lab
  - Chemical dependency treatment facility (licensed by the Texas Commission on Alcohol and Drug Abuse)
  - Consumer Directed Services Agency
  - County Indigent Health Care Program
  - Community mental health center
  - Comprehensive health center
  - Comprehensive outpatient rehabilitation facility/outpatient rehabilitation facility
  - Durable medical equipment (DME)
  - Early Childhood Intervention
  - Federally Qualified Health Center (FQHC)
  - Freestanding psychiatric facility
  - Freestanding rehabilitation facility
  - Home and Community Based Services – Adult Mental Health (HCBS-AMH) providers
  - Home Health/Home and community support services agency
  - Hospital/critical access hospital/out-of-state hospital
  - Military hospital
  - Hyperalimentation
  - Independent diagnostic testing facility/physiological lab
  - Indian Health Services
  - Independent laboratory
  - Intellectual or developmental disability (IDD) case management
  - Maternity services clinic
  - Mental health targeted case management and mental health rehabilitative services
  - Milk bank donor
  - Personal care services
  - Pharmacy
• Portable X-ray
• Prescribed Pediatric Extended Care Centers (PPECC)
• Radiation treatment center
• Radiological laboratory
• Renal dialysis facility
• Rural health clinic (RHC)
• School health and related services (SHARS)/non-school SHARS
• Service responsibility option
• Skilled nursing facility
• State Supported Living Center (SSLC)
• Vision medical supplier
• Women, Infant and Children

Providers must submit a separate Texas Medicaid Provider Enrollment Application for each enrollment type that they request unless otherwise approved as a dual enrollment. For example, enrolled hospital providers will be issued a hospital provider identifier that is specific to hospital services and a separate HASC provider identifier that is specific to ambulatory surgical services unless the provider is subject to restricted reimbursement. However, a health-care professional who is already enrolled with Texas Medicaid as an individual with his or her own practice, and who wishes to bill for services provided in connection with a group, must submit a separate enrollment application and be approved as a performing provider with the group. Similarly, a health-care professional who is enrolled as a performing provider with one group, but who wishes to bill for services provided in connection with another group, must submit a separate enrollment application and be approved as a performing provider with the other group.

**Note:** A separate provider identifier is issued for each enrollment type that is approved. The provider is authorized to use the provider identifier only to bill for services provided as indicated in the approved enrollment application. It is a program violation for a provider to use a provider identifier for any purpose other than billing for the types of services, and under the type of enrollment, for which that provider identifier was issued. Improper use of a provider identifier constitutes program abuse and/or fraud.

**Refer to:** Subsection 1.11, “Medicaid Fraud, Waste, and Abuse Policy” in this section for additional information.

### 1.1.5.1 Ordering- or Referring-Only Providers

Individual providers who are not currently enrolled in Texas Medicaid and whose only relationship with Texas Medicaid is to order or refer for supplies or services for Texas Medicaid-eligible clients must enroll in Texas Medicaid as participating providers. This requirement is in accordance with provisions of the Affordable Care Act of 2010 (ACA), 42 CFR §455.410(b), which requires all fee-for-service (FFS) and managed care network ordering or referring physicians or other professionals who order or refer for supplies or services under the Medicaid State plan, or under a waiver of the plan, to enroll in Medicaid as participating providers.

Providers who are out of network for Medicaid managed care organizations (MCOs) do not need to enroll as ordering or referring-only providers. The enrollment requirement includes providers who order or refer for supplies or services for dually eligible clients (i.e., clients who are enrolled in both Medicare and Medicaid), as the client’s claims would be considered Medicaid claims.
These providers can enroll online using the PEP tool by clicking the check box for Ordering/Referring Provider, or they can use the streamlined paper Texas Medicaid Provider Enrollment Application Ordering and Referring Providers Only, which is available for download on the TMHP website at www.tmhp.com.

The ordering or referring-only enrollment application is for individual providers who are not currently enrolled as a billing or performing provider in Texas Medicaid or the CSHCN Services Program and who do not currently have an active Texas Medicaid or CSHCN Services Program billing provider Texas Provider Identifier (TPI).

**Important:** Individual providers who are currently enrolled in Texas Medicaid or the CSHCN Services Program and who currently have an active Texas Medicaid or CSHCN Services Program billing provider TPI can use their current TPI for ordering or referring services and do not need to obtain an ordering and referring-only provider TPI. A current billing provider’s active TPI will be deactivated if the provider enrolls as an ordering or referring-only provider.

Providers who enroll in Texas Medicaid as ordering or referring-only providers receive one TPI that can be used for orders and referrals for both Texas Medicaid clients and CSHCN Services Program clients. Although ordering or referring-only providers do not submit claims to TMHP for rendered services, the ordering or referring-only provider’s NPI is required on claims that are submitted by the billing providers that render the services or provide the supplies or services.

**Refer to:** Subsection 6.4.2.4, “Ordering or Referring Provider NPI” in “Section 6: Claims Filing” (Vol. 1, General Information) for information about filing claims that require an ordering- and referring-only provider NPI.

### 1.1.5.2 Ordering or Referring-Only Providers Participating in Other State Health-Care Programs

Providers who order or refer for supplies or services for Children’s Health Insurance Program (CHIP) clients must enroll as a participating providers with TMHP. Providers who are out of network for CHIP MCOs do not need to enroll as ordering or referring-only providers. Providers who order or refer for supplies or services for Healthy Texas Women (HTW) and CSHCN clients must also enroll as participating providers with TMHP.

### 1.1.5.3 Interns and Medical Residents Who Order, Prescribe, or Refer

Interns and medical residents with only Physician-In-Training (PIT) permits issued by the Texas Medical Board cannot enroll in Texas Medicaid. An intern or resident’s licensed supervising physician must be reported as the ordering or referring provider on claims that are generated from the order or referral of the intern or resident. The NPI of the supervising physician must be listed on orders or referrals written by the interns or residents they supervise. The licensed supervising physician must be enrolled in Texas Medicaid as a billing provider or as an ordering or referring-only provider.

### 1.1.6 Affordable Care Act of 2010 (ACA) Enrollment Requirements

Providers are required to fulfill certain requirements for enrollment in order to comply with the provisions of ACA. Providers that are enrolled in Texas Medicaid and have fulfilled the ACA requirements through their Texas Medicaid enrollment are considered ACA-compliant for all programs in which they are enrolled.

**Refer to:** TMHP website at www.tmhp.com for additional information about ACA requirements.

In accordance with Section 6401 of ACA, the following requirements apply:

- Upon initial enrollment, revalidation, and re-enrollment, all participating providers are screened based on their categorical risk level. (complies with 42 CFR §§455.410 and 455.450)
- All providers are required to revalidate at least every three to five years based on provider type.
• Institutional providers who are enrolling, revalidating, or re-enrolling are required to pay an application fee if one has not already been paid to Medicare or another state’s Medicaid program or Children’s Health Insurance Program (CHIP).

• Ordering and referring-only providers are required to enroll in Texas Medicaid as participating providers.

Refer to: Subsection 1.1.5.1, “Ordering- or Referring-Only Providers” in this section.

1.1.6.1 Provider Screening Requirement
In compliance with ACA, all providers must be screened, which includes:

• Providers who submit a provider enrollment application for new enrollment, a new practice location, or other type of enrollment or re-enrollment.

• Providers who are currently enrolled in Texas Medicaid and are required to revalidate their enrollment in Texas Medicaid.

1.1.6.2 Provider Revalidation
In compliance with 42 CFR §455.414, all providers are required to revalidate at least every three to five years:

• DME providers are required to revalidate enrollment information at least once every three years.

• All other provider types must revalidate their enrollment information at least once every five years.

During revalidation, the provider screening will be repeated.

1.1.6.3 Application Fee
Under ACA, institutional providers are subject to an application fee for applications, including initial applications, applications for new practice locations, revalidation, and re-enrollment applications. Upon completion of the PEP online application, providers will be notified whether they are required to pay an application fee. The amount of the application fee is subject to change every calendar year.

Providers that complete the paper Texas Medicaid Provider Enrollment Application can refer to the TMHP website for the list of provider types that are required to pay the application fee.

Note: Providers that are required to pay the application fee but have already paid the fee to Medicare or another state’s Medicaid program or CHIP have fulfilled the fee requirement and do not have to submit the fee to Texas Medicaid. Proof of payment must be submitted with the application. Providers who are enrolled in Medicare must provide documentation that specifies whether or not they have completed the ACA rescreening process with Medicare.

1.1.6.4 Ordering- or Referring-Only Providers Search on the Online Provider Lookup (OPL)
Providers can verify that an ordering- or referring-only provider is enrolled in Medicaid by using either the basic or advanced provider search function of the OPL.

1.1.7 Surety Bond Enrollment Requirement
All newly enrolling and re-enrolling DME and nongovernment-operated ambulance providers must obtain a surety bond that complies with 1TAC §352.15 as a condition of enrollment and continued participation in Texas Medicaid.

DME providers can refer to subsection 2.1.2, “Surety Bond Requirements” in the Durable Medical Equipment, Medical Supplies, and Nutritional Products Handbook (Vol. 2, Provider Handbooks).
Ambulance providers can refer to subsection 1.1.7.1, “Ambulance Providers” in this handbook.

**Important:** Surety bonds obtained for the purpose of accreditation in the Medicare program, which lists the Centers for Medicare & Medicaid Services (CMS) as obligee, do not fulfill the surety bond requirement for Texas Medicaid.

The surety bond submitted to Texas Medicaid must meet the following requirements:

- A bond in an amount of no less than $50,000 must be provided for each enrolled location.
- The bond must be submitted on the State of Texas Medicaid Provider Surety Bond Form. No other form will be accepted. The use of this form designates HHSC as the sole obligee of the bond. Instructions are included with the form.
- The bond must be issued for a term of 12 months. Bonds for longer or shorter terms are not acceptable.
- The bond must be in effect on the date that the provider enrollment application is submitted to TMHP for consideration. The effective date stated on the bond must be:
  - No later than the date that the provider enrollment application is submitted.
  - No earlier than 12 months before the date that the provider enrollment application is submitted.
- The bond must be a continuous bond. A continuous bond remains in full force and effect from term to term unless the bond is canceled.

**Important:** An annual bond that specifies effective and expiration dates for the bond is not acceptable.

At the time of enrollment, revalidation, or re-enrollment, providers must submit the surety bond form with original signatures and a copy of the Power of Attorney document from the surety company that issued the bond.

**Note:** Surety companies may refer to Texas Department of Insurance (TDI) file #9212562912 or TDI link #132456 when filing the bond.

DME and non-government-operated ambulance providers must maintain a current surety bond to continue participation in Texas Medicaid. To avoid losing Medicaid enrollment status, providers must submit proof of continuation to TMHP Provider Enrollment before the expiration date of the bond currently on file. The completed proof of continuation document must include the bond number, original signatures of the authorized corporate representative of the DME or ambulance provider (principal), the attorney-in-fact of the surety company, date of the original bond, and new “good through” date. Providers may submit a copy of the proof of continuation (scan, fax, photocopy) pending the submission of the original document.

**Refer to:** The State of Texas Medicaid Provider Surety Bond Form in the Forms section of the TMHP website at www.tmhp.com.

### 1.1.7.1 Ambulance Providers

Ambulance providers that participate in Texas Medicaid fee-for-service, managed care programs, or the CHIP must, as a condition of emergency medical services (EMS) provider license renewal, obtain a surety bond that complies with 1 TAC §352.15 and submit the bond to TMHP according to the requirements listed above. A copy of the bond must be included with their application to the Department of State Health Services (DSHS) to renew their emergency services provider license.

Providers can refer to the DSHS website for additional information.

Ambulance providers that are directly operated by a governmental entity are exempt from the surety bond requirement.
1.1.8 Provider Enrollment Application Determinations

An application for provider enrollment may be approved with conditions, or denied. The provider applicant is issued a notice of the enrollment determination.

Refer to: Subsection 1.1.6, “Affordable Care Act of 2010 (ACA) Enrollment Requirements” in this section for additional information about the ACA 3- to 5-year revalidation requirement.

When an application for enrollment is approved with conditions, the applicant has no right of appeal or administrative review of the enrollment determination. The types of conditional enrollment include, among other things:

- An application may be approved for time-limited enrollment, meaning the provider is granted a contract to participate in Medicaid for a specific period of time. In this case, the provider is sent a notice that includes the deactivation date of the contract. It is the provider’s responsibility, if the provider chooses to seek continued Medicaid participation, to file a complete and correct reenrollment application before the deactivation date of the provider’s current contract. It is recommended that the provider submit a reenrollment application at least 60 days before the current contract deactivation date, to ensure that the reenrollment application is complete and correct before the deactivation date. This may avoid a lapse between the provider’s current contract and the new contract, if a new contract is granted.

- An application may be approved subject to restricted reimbursement, meaning the provider is eligible to have only certain types of claims paid. This includes, among other things, reimbursement of only Medicare crossover claims (i.e., claims with respect to “dual eligible” recipients who are covered by both Medicare and Medicaid).

An application might be denied. If an application is denied, TMHP will send the provider a denial notice that explains the basis for the denial. The notice also explains the provider’s right to make a written request for an informal desk review of the denial decision and the procedures for filing such a request. The administrative rules governing the informal desk review of the denial decision are found in 1 TAC §371.1015. Any informal desk review request must be received within 20 business days of the date on the letter and filed in accordance with the instructions provided in the denial notice. HHSC will conduct the informal desk review and render a final enrollment determination. HHSC’s final determination is not subject to further administrative review or reconsideration.

The enrollment date is the day on which a new TPI was issued. This date impacts claims filing deadlines.

Refer to: Subsection 6.1.4.2, “Claims for Newly Enrolled Providers” in “Section 6: Claims Filing” (Vol. 1, General Information) for timely-filing guidelines for newly enrolled providers.

HHSC determines the enrollment effective dates for providers that choose to enroll in Medicaid and Texas State Health-Care Services. Enrollment notification letters that contain the enrollment information for newly enrolling providers are printed the following business day and mailed to the physical address that is listed on the application. Revalidation notifications that contain the updated revalidation due date are printed the following business day and mailed to the physical address listed on the application.

1.1.9 Enrollment in Medicaid Managed Care Programs

To be reimbursed for services rendered to Medicaid managed care clients, providers must be enrolled in Texas Medicaid and then must enroll with the client’s health plan to be eligible for reimbursement for services rendered.

Refer to: Subsection 2.2, “Provider Enrollment and Responsibilities” in the Medicaid Managed Care Handbook (Vol. 2, Provider Handbooks).
1.1.10 **Required Enrollment Forms**

To enroll in Texas Medicaid, providers must complete and submit the appropriate Texas Medicaid enrollment application, including all required forms as indicated in the application.

*Note:* All paper documents must be signed by the person who is applying for enrollment. If the applicant is an entity, a principal of the entity must sign the application.

Whether they are completing the online application or a paper application, providers can refer to the checklist in the paper Texas Medicaid Provider Enrollment Application for information about required forms and other documentation. This checklist explains, by provider type, the documents and information that must be provided with the application. Applications must be complete in order to process and issue a provider identifier.

*Note:* If enrolled in Medicare, the provider must submit a copy of the Medicare enrollment letter to enroll in Texas Medicaid. Otherwise the enrollment application will be considered incomplete.

When prompted to enter a tax identification number (tax ID) on either a paper or electronic copy of an enrollment application, the applicant should list the entity’s nine-digit federal tax identification number.

Providers can call the TMHP Contact Center at 1-800-925-9126 for help with completing the application. Providers should retain a copy of the original application for future reference.

All pages of the application (excluding instructions) must be present even if the forms are left blank because they are not pertinent to the provider’s situation. Providers will be notified of incomplete applications and will have 30 business days to provide the requested missing information. If the information is not provided within 30 business days, TMHP will terminate the enrollment process. If the provider wants to enroll at a later date, a new enrollment application must be submitted. Providers are required to review their enrollment application for correctness and completeness before submitting it to TMHP.

By signing the HHSC Medicaid Provider Agreement, a provider is certifying that all information submitted in connection with the application for enrollment is complete and correct. Any false, misleading, or incomplete information submitted in connection with an enrollment application constitutes a Medicaid program violation, and may result in administrative, civil, or criminal liability.

*Refer to:* Subsection 1.1.11, “Medicaid Fraud, Waste, and Abuse Policy” in this section.

1.1.10.1 **Application Payment Form**

All providers who are required to pay an application fee to participate in the Medicaid program must submit an Application Payment Form. The application cannot be processed if the application fee is required and is not submitted with the application.

*Refer to:* Subsection 1.1.6.3, “Application Fee” in this section.

1.1.10.2 **HHSC Medicaid Provider Agreement**

The HHSC Medicaid Provider Agreement must be submitted by all providers who enroll in Texas Medicaid and must be signed by the provider who is applying for enrollment. If the applicant is an entity, a principal of the entity who has the authority to bind the entity to the requirements of the HHSC Provider Agreement must sign the agreement. “Principal” is defined in the following section.

*Refer to:* Subsection 1.1.10.8, “Corporate Board of Directors Resolution” in this section for information about corporations.)
If the provider is city- or government-owned, the agreement must be signed by a person who is authorized under the city or government charter. This form is an agreement between HHSC and the provider performing services under the State Plan wherein the provider agrees to certain provisions as a condition of participation.

**Note:** The person who signs the HHSC Medicaid Provider Agreement is certifying that all of the information in the application packet, including every completed Provider Information Form (PIF-1) and Principal Information Form (PIF-2), is complete and correct. This includes a certification that every person who is required to complete a PIF-2 has done so, and all required PIF-2s are included with the application.

TMHP must receive all of the pages of the HHSC Medicaid Provider Agreement in a single submission for a valid contract. If corrections are required on any page within the agreement, a new agreement with an original signature and date is required.

### 1.1.10.3 Provider Information Form (PIF-1)

The PIF-1 must be completed by, or on behalf of, the provider that is applying for enrollment. If the provider is an entity, the PIF-1 must be completed on behalf of the entity.

### 1.1.10.4 Principal Information Form (PIF-2)

A PIF-2 must be completed by each principal/creditor, subcontractor, and creditor of the provider that is applying for enrollment with the following exceptions:

- Performing providers who are applying to join a group that is already enrolled
- THSteps provider applications that are received within one year of the TPI enrollment date
- Individuals who enrolled using their own Social Security number and an entity type of Individual/Sole Proprietorship

Principals of the provider include all of the following:

- An owner with a direct or indirect ownership or control interest of 5 percent or more
- Corporate officers and directors
- Managing employees or agents who exercise operational or managerial control, or who directly or indirectly manage the conduct of day-to-day operations
- Limited or nonlimited partners
- Shareholders of a professional corporation, professional association, limited liability company, or other legally designated entity
- Any employee of the provider who exercises operational or managerial control over the entity or who directly or indirectly conducts the day-to-day operations of the entity
- All individuals, companies, firms, corporations, employees, independent contractors, entities, or associations that have been expressly granted the authority to act for or on behalf of the provider
- All individuals who are able to act on behalf of the provider because their authority is apparent
- An individual or entity with a security interest in a debt that is owed by the provider if the creditor’s security interest is protected by at least 5 percent of property listed in Section III(c) of the Disclosure of Ownership

A subcontractor of the provider is defined as follows:

- An individual, agency, or organization to which a disclosing entity has contracted or delegated some of its management functions or responsibilities of providing medical care to its patients; or
• An individual, agency, or organization with which a fiscal agent has entered into a contract, agreement, purchase order, or lease (or leases of real property) to obtain space, supplies

Note: This includes the on-site manager, supervising licensed practitioner, or medical director for each physical location of the provider in Texas.

1.1.10.5 Disclosure of Ownership and Control Interest Statement
All providers must submit the Disclosure of Ownership and Control Interest Statement as part of the enrollment application, with the following exceptions:

• Performing providers who are applying to join a group that is already enrolled
• Texas Health Steps Medical or CSHCN Services Program providers whose applications are received within one year of the TPI enrollment date
• ECI, MHMR, MHR, Blind Children’s Vocational Discovery & Development Program, YES Waiver, TB Clinic, Military Hospital, State Hospital or SHARS providers

This form also contains questions that must be answered under federal law. Failure to provide complete and accurate information as instructed on this form will constitute an incomplete application, which may result in denial of enrollment. Incomplete or inaccurate information on this form constitutes a violation of the rules of Medicaid and may also result in administrative, civil, or criminal liability.

Refer to: Subsection 1.11, “Medicaid Fraud, Waste, and Abuse Policy” in this section.

Note: Providers are required to submit any change in ownership, corporate officers, or directors to TMHP Provider Enrollment within 30 calendar days of the change.

Refer to: Subsection 1.7.2, “Maintenance of Provider Information” in this section.

1.1.10.6 Internal Revenue Service (IRS) W-9 Form
The IRS W-9 Form must be completed and submitted for all types of enrollment, except in the case of performing providers seeking to join an already enrolled group.

1.1.10.7 Medicaid Audit Information Form
The Medicaid Audit Information Form is required by facilities that file cost reports such as hospitals, home health agencies, FQHCs, RHCs, and dialysis facilities.

1.1.10.8 Corporate Board of Directors Resolution
All providers who indicate that they are a corporation on the Disclosure of Ownership and Control Interest Statement are required to submit the Corporate Board of Directors Resolution. This form indicates the individual (by name) who is authorized by the corporation to sign the agreement forms. The secretary of the corporation must sign the Corporate Board of Directors Resolution and have it notarized. If a business is city or government-owned, this form is not required.

1.1.10.9 Franchise Tax Account Status Page
When enrolling as a “Corporation” type of entity, providers must submit a Franchise Tax Account Status Page. This information can be obtained from the Texas State Comptroller’s Office website at https://comptroller.texas.gov/taxinfo/coasintr.html.

Providers who have a 501(c)(3) Internal Revenue Exemption are not required to submit the Franchise Tax Account Status Page, but they must submit the IRS exemption letter.

1.1.10.10 Certificate of Formation or Certificate of Filing/Articles or Certificate of Incorporation/Certificate of Fact
When enrolling as a “Corporation” type of entity, providers must submit the Certificate of Formation or Certificate of Filing form. Obtain the form from the Office of the Secretary of State of Texas. The name on this form must exactly match the legal name shown on the W-9 form.
The following certificates also apply for corporations:

- For corporations formed prior to January 1, 2006, Articles or Certificate of Incorporation/Certificate of Authority/Certificate of Fact
- For corporations formed on or after January 1, 2006, Certificates of Formation or Certificate of Filing
- For corporations registered in a state other than Texas, Certificate of Authority or Certificate of Filing

The Certificate and any required certifications to provide certain services in Texas must be submitted when a corporation is registered in a state other than Texas. The form identifies the legal name of the corporation and is proof that the corporation is registered to do business in Texas.

Note: Out-of-state providers that do not provide services in the state of Texas are exempt from submitting this form.

1.1.10.11 Copy of License, Temporary License, or Certification

Providers cannot enroll in Texas Medicaid if their license is due to expire within 30 days. During the enrollment process, TMHP verifies licensure using available resources. If TMHP cannot verify a license at the time of enrollment, it is the providers’ responsibility to provide a copy of the active license to TMHP. Psychologists and facilities must submit a copy of their license since these licenses cannot be verified online.

TMHP will notify the provider by letter if a copy has not been submitted and the license cannot be verified.

Once a provider is enrolled in Texas Medicaid the license or certification must be kept current. A reminder letter for renewal will be sent to the provider 60 days before the provider’s license expires.

TMHP directly obtains licensure information from the following licensing boards:

- Texas Medical Board (TMB) (for physicians only)
- Texas Board of Nursing (BON)
- Texas State Board of Dental Examiners (TSBDE)
- National Board of Certification and Recertification for Nurse Anesthetists (NBCRNA)
- The Executive Council of Physical Therapy and Occupational Therapy Examiners (ECPTOTE) (Physical Therapist only)
- Texas Optometry Board (TOB)

If a license cannot be verified due to a delay in obtaining the board licensing information, providers must request a letter from the licensing board for their individual provider information and submit it to TMHP by the deadline indicated in the reminder letter. The letter must contain the provider’s specific identification information, license number, and licensure period.

All other licenses and certifications that are not issued by TMB, BON, or TSBDE must be submitted to TMHP upon renewal.

Important: Providers are also required to submit to TMHP, within 10 days of occurrence, notice that the provider’s license or certification has been partially or completely suspended, revoked, or retired. Not abiding by this license and certification update requirement may impact a provider’s qualification to continued participation in Texas Medicaid.

Refer to: Subsection 1.1.10.14, “Licensure Renewal” in this section.
1.1.10.12 Federally Qualified Health Center Affiliation Affidavit
All FQHC must identify and attest that all contractual affiliation agreements with contracted providers have been submitted to and approved by the Bureau of Primary Health Care (BPHC).

- Texas Medicaid defines an affiliate agreement as a contract between an FQHC and another provider for the provision of FQHC services for which the FQHC will bill Medicaid under the FQHC prospective payment system (PPS).
- Affiliations do not include contracts for the direct employment of providers or staff.

Refer to: Federally Qualified Health Center Affiliation Affidavit on the TMHP website at www.tmhp.com.

1.1.10.13 Physician’s Letter of Agreement
Upon initial enrollment and upon revalidation, Certified Nurse Midwife (CNM) or Licensed Midwife (LM) providers must complete and submit to TMHP with the Medicaid provider enrollment application the Physician’s Letter of Agreement affirming the CNM’s supervising physician arrangement or the LM’s referring or consulting physician arrangement.

- According to HHSC rules, 1 TAC §354.1253(c), and 1 TAC §354.1252(3), CNM providers and LM providers are required to inform HHSC in writing of the identity of a licensed physician or group of physicians with whom the CNM or LM has arranged for referral and consultation in the event of medical complications. For purposes of this rule, “consultation” means discussion of patient status, care, and management.
- A separate agreement must be submitted for each physician with whom an arrangement is made.

This agreement must be signed by the CNM or LM and the physician.

A new agreement must also be completed and submitted to TMHP when a new arrangement is made and when changes to an arrangement are made.

Refer to: Subsection 1.4.2, “Provider Status (Individual, Group, Performing Provider, or Facility)” in this section.

1.1.10.14 Licensure Renewal
Not abiding by the license and certification update requirement may impact a provider’s qualification for continued participation in Texas Medicaid. If a provider’s license has expired, a deactivation letter will be sent to the provider, and all claims filed on and after the expiration date will be denied.

To have claims payments resumed, updated information must be sent to the applicable licensing board to renew the license. Payment will be considered for dates of service on or after the date of license renewal. Claims denied due to an inactive license may be appealed, and payment will be considered for dates of service on or after the date of return to active license status. Payment deadline rules for the fiscal agent arrangement must be met.

Refer to: Subsection 6.1.5, “HHSC Payment Deadline” in “Section 6: Claims Filing” (Vol. 1, General Information).

1.1.10.15 Medicare Participation
Under federal law, Medicaid is the payor of last resort, so Medicare-covered services must first be billed to and paid by Medicare. Therefore, in order to be eligible to enroll in Texas Medicaid, a provider must be a Medicare participating provider. Certain types of providers, however, are not required to meet the Medicare participation requirement, including:

- Pediatric providers
- Family planning providers
- Case Management for Children and Pregnant Women program providers
• CCP providers
• Early Childhood Intervention (ECI) providers
• Licensed professional counselors (LPCs)
• Licensed marriage and family therapists (LMFTs)
• OB/GYN providers
• THSteps medical and dental services providers

Some provider types may apply for a waiver of the Medicare certification requirement of the application process if they do not serve Medicare-eligible individuals. The following provider types are eligible to apply for this waiver:

• Audiologist
• Dentist (D.D.S. or D.M.D.)
• Nurse practitioner/clinical nurse specialist (NP/CNS)
• Optometrist (OD)
• Orthotists
• Physician (DO)
• Physician (MD)
• Physician assistant (PA)
• Prosthetist

All providers that are required to participate in Medicare must include a valid and current Medicare number and a copy of the provider’s notice of Medicare participation as part of the Texas Medicaid Provider Enrollment Application.

Each group and each performing provider of a Medicare group must have a current Medicare number. The group enrollment application must include the current and valid Medicare number for the group and for each performing provider in the group, as well as a copy of the notice of Medicare enrollment for the group and for each performing provider in the group.

Each group enrolling as a Medicaid-only group does not need to submit a current Medicare number for the group. Performing providers added to this Medicaid-only group also do not require a current Medicare number.

1.1.10.16 Group Information Changes

If additions or changes occur in a group’s enrollment information (for example, a performing provider leaves or enters the group, changes the physical address or the accounting/mailing address, or a provider is no longer licensed) after the enrollment process is completed, the provider group must notify Texas Medicaid in writing within 90 calendar days of occurrence of the changes. Failure to provide this information may lead to administrative action by HHSC. Filing claims and receiving payment without having followed this requirement constitutes a program violation and may also result in administrative, civil, or criminal liability.

Refer to: Subsection 1.11, “Medicaid Fraud, Waste, and Abuse Policy” in this section for additional information.
1.2 Payment Information

Texas Medicaid reimbursements are available to all enrolled providers by check or electronic funds transfer (EFT). Providers are strongly encouraged to utilize EFT, which allows for more rapid reimbursement.

1.2.1 Using EFT

HHSC recommends that all Texas Medicaid providers receive payment by EFT. EFT is a method for directly depositing funds into a designated bank account. EFT does not require special software, and providers can enroll immediately.

1.2.2 Advantages of EFT

Advantages of EFT include:

- Electronically-deposited funds are available more quickly than with paper checks.
- Providers do not have to worry about lost or stolen checks.
- TMHP includes provider and Remittance and Status (R&S) Report numbers with each transaction submitted. If the bank’s processing software captures and displays the information, both numbers would appear on the banking statement.

1.2.3 EFT Enrollment Procedures

The Electronic Funds Transfer (EFT) Notification can be found on the TMHP website at www.tmhp.com. Completed EFT forms can be uploaded to the Provider Information Management System (PIMS), faxed to 1-512-514-4214, or mailed to:

Texas Medicaid & Healthcare Partnership  
Attn: Provider Enrollment  
PO Box 200795  
Austin, TX 78720-0795

To enroll for EFT, providers must submit a completed Electronic Funds Transfer (EFT) Authorization Agreement to TMHP. A voided check or letter on bank letterhead, containing the bank routing and account information, must be attached to the enrollment form. One completed form must be filled out for each billing provider identifier, including an original signature of the provider.

After the Electronic Funds Transfer (EFT) Authorization Agreement has been processed, TMHP issues a prenotification transaction during the next cycle directly to the provider’s bank account. This transaction serves as a checkpoint to verify EFT is working correctly.

If the bank returns the prenotification without errors, the provider will begin receiving EFT transactions with the third cycle following the enrollment form processing. Providers will continue to receive paper checks until they begin to receive EFT transactions.

If the provider changes bank accounts, the provider must submit a new Electronic Funds Transfer (EFT) Authorization Agreement to TMHP Provider Enrollment. The prenotification process is repeated and, once completed, the EFT transaction is deposited to the new bank account.

Refer to:  
1.2.4 Receiving Paper Checks
Providers must have a current physical and accounting/mailing address and telephone number on file so that they can promptly receive reimbursement checks and other TMHP correspondence. Providers must send all changes to addresses and telephone numbers to:

Texas Medicaid & Healthcare Partnership
Provider Enrollment
PO Box 200795
Austin, TX 78720-0795
1-800-925-9126
Fax: 1-512-514-4214

1.2.5 Stale-Dated Checks
Stale-dated checks (i.e., checks that are older than 180 days) that have not been cashed are voided and applied to either IRS levies or outstanding accounts receivable. Once a check has been voided, the associated claims may not be payable, and the transaction will be finalized after 24 months. Providers may submit a voided check appeal to TMHP Cash Financial at the following address:

Texas Medicaid & Healthcare Partnership
Attn: Cash Financial
12357B Riata Trace Parkway
Austin, TX 78727

TMHP encourages providers to receive payment via EFT to eliminate stale-dating issues. EFT ensures that providers receive payments through direct deposit in a bank account of their designation.

Refer to: Subsection 1.2.3, "EFT Enrollment Procedures" in this section.

1.3 Provider Deactivation/Disenrollment
Payment denial codes are applied to a TPI that has had no claim activity for a period of 24 months or more. The TPI will be considered inactive and will not be able to be used to submit claims.

A courtesy letter will be sent to all providers whose TPIs have been identified as not having any claims activity over the previous 18 months. Providers will have six months to submit claims and prevent the TPI from being deactivated. If the provider is enrolled in both Medicaid and the CSHCN Services Program, the provider identifiers for both programs will be examined to determine whether claims activity has occurred.

After 24 months without claim activity, providers will be sent a deactivation letter, and a payment denial code will be applied to their provider identifier. If a provider’s Medicaid TPI is deactivated, any enrollments associated with the inactive TPI with the CSHCN Services Program will also be deactivated.

Claims that are submitted for a deactivated TPI after the payment denial code has been applied will be denied.

To have the payment denial code removed from a provider identifier, providers must submit a completed application for the state health-care program in which they wish to enroll, and the application must be approved. The information on this application must match exactly the information currently on the provider’s file for the payment denial code to be removed.

1.3.1 Excluded Entities and Providers
The United States Health and Human Services (HHS)-OIG and the HHSC-OIG exclude certain individuals and entities from participation in all federal or state health-care programs. The exclusions restrict individuals from receiving any reimbursement for items or services furnished, ordered, or prescribed.
All current providers and providers who are applying to participate in state health-care programs must screen their employees and contractors every month to determine whether they are excluded individuals or entities. These screenings are a condition of the provider’s enrollment, revalidation, or re-enrollment into state health-care programs.

Providers can determine whether an individual or entity is excluded by searching the List of Excluded Individuals/Entities (LEIE) website at www.oig.hhs.gov/fraud/exclusions.asp. A downloadable version of the database is available but it does not include Social Security Numbers (SSNs) or Employer Identification numbers (EINs). The Texas HHSC-OIG website is found at https://oig.hhsc.texas.gov/exclusions. If a name matches a name on the exclusion list, it can be verified online with a SSN or EIN.

Providers must search the LEIE website monthly to capture any exclusions or reinstatements that have occurred since the last search. Providers must immediately report to HHS-OIG any exclusion information they discover when searching the LEIE database.

CFR section 1003.102(a)(2), states that civil monetary penalties may be imposed against Medicaid providers and managed care entities (MCEs) that employ or enter into contracts with excluded individuals or entities to provide items or services to Medicaid clients. In addition, no Medicaid payments can be made for any items or services directed or prescribed by an excluded provider or other authorized person when the individual or entity furnishing the services either knew or should have known of the exclusion. This prohibition applies even when the Medicaid payment itself is made to another provider, practitioner, or supplier that is not excluded.

1.4 Provider Reenrollment

Providers must submit a new application and a new provider identifier must be issued when there are changes in the Medicare number.

The new application may be submitted electronically using PEP or by submitting a completed paper Texas Medicaid Provider Enrollment Application. A new application is required when one of the following changes:

1.4.1 Medicare Number

If Medicare has issued a new Medicare number, the provider must complete and submit a Texas Medicaid Provider Enrollment Application in order to enroll with the new information.

1.4.2 Provider Status (Individual, Group, Performing Provider, or Facility)

Providers leaving group practices must send a signed letter or a Provider Information Change Form to TMHP that states the date of deactivation. The letter should include the provider identifier, effective date of deactivation, and the group’s provider identifier. The letter should be signed by an authorized representative of the group or the individual provider leaving the group. If the provider is joining a new group practice has changed from a group practice to an individual enrollment or vice versa, the provider must complete and submit a new Texas Medicaid Provider Enrollment Application to request enrollment with the new status information.

1.4.3 Physical Address

If a provider moves or has an address change and the new address is within the same Medicare locality, the provider must update their address information within 90 days of the change. Providers should update their address information through My Account on www.tmhp.com. Alternately, providers may also update their address information by submitting a completed Provider Information Change Form, which is available on the Forms section of www.tmhp.com. If address information updates are submitted on the Provider Information Change Form, the provider must also submit a Form W-9.
If the new address is not within the previous Medicare locality and Medicare has issued the provider a new Medicare number, the provider must submit a new enrollment through My Account or a completed Texas Medicaid Provider Enrollment Application. Texas Health Steps Medical and Dental providers must complete an online enrollment application or submit a paper application for each new practice location.

**Note:** Providers that are not enrolled with Medicare must submit a new enrollment application to enroll additional physical address locations.

### 1.4.4 Change in Principal Information

As defined in subsection 1.1.10.4, “Principal Information Form (PIF-2)” in this section, change in principal information includes a change in corporate officers or directors, professional association membership, and managing employees. The change must be reported to TMHP within 30 calendar days of when it occurs.

**Refer to:** Subsection 1.7.2.3, “Online Provider Lookup (OPL)” in this section.

Providers must contact the Electronic Data Interchange (EDI) help desk directly and request an Electronic Remittance & Status (ER&S) Report each time a new provider identifier is issued to the provider. This form must be completed and returned to EDI with unique identifying information related to the new provider identifier to ensure there is no suspension in the provider’s ability to access their ER&S statement on the secure provider portal through [www.tmhp.com](http://www.tmhp.com).

Providers must also contact any third party EDI vendors with whom they are contracted to add any new provider identifiers to their ER&S Report. To obtain a PDF copy of the ER&S Report on the TMHP Home Page, the provider must create an administrator account for each provider identifier belonging to them.

Providers that have been issued a new provider identifier through the TMHP enrollment or re-enrollment process must ensure that any prior authorizations affected have been updated to reflect the new provider identifier.

### 1.5 Change of Ownership Requirements

The new owner must do the following:

- Obtain recertification as a Title XVIII (Medicare) facility under the new ownership
- Submit CMS Acknowledgement of Change of Ownership Letter
- Provide TMHP with a copy of the Contract of Sale (specifically, a signed agreement that includes the identification of previous and current owners in language that specifies who is liable for overpayments that were identified subsequent to the change of ownership, that includes dates of service before the change of ownership)
- Provide a separate change of ownership and Texas Medicaid provider enrollment application for all of the provider identifiers affected by the change of ownership
- Submit any new enrollment application relating to a change of ownership to TMHP Provider Enrollment within 30 calendar days of the change

When the change of ownership has been processed, the original TPI used by the provider to bill claims will be deactivated, and the provider will lose the ability to download R&S Reports from the TMHP portal as well as the ability to verify client eligibility online. Claims status inquiries through the TMHP portal will also be unavailable. After a TPI has been deactivated, the provider can call the contact center to check on client eligibility and the status of claims. Paper R&S Reports can be printed by the TMHP Contact Center, and delivered to providers, up to 30 days from the date the TPI is deactivated.
1.6 Claims Filing During Enrollment

Providers must adhere to claim filing deadlines throughout the enrollment, re-enrollment, and revalidation processes. Claims submitted by newly enrolled providers must be received within 95 days of the date that the new provider identifier is issued and within 365 days of the date of service.

Providers that are enrolling in Texas Medicaid for the first time or that are making a change that requires the issuance of a new TPI can submit claims within 95 days of the date on which their TPI is issued as long as claims are submitted within 365 days of the date of service.

Providers that are revalidating an existing enrollment can continue to file claims while they are completing the revalidation process. TMHP must receive claims within 95 days of the date of service.

For clients who have retroactive eligibility, the 95-day deadline is based on the date of service or the date on which the client eligibility information is added to the TMHP eligibility file, whichever is later. For clients who have dual Medicare and Medicaid eligibility, when a service is a benefit of both Medicare and Medicaid, the claim must be filed with Medicare first. In these cases, the 95-day deadline is based on the date of Medicare disposition.


1.7 Provider Responsibilities

1.7.1 Compliance with Texas Family Code

1.7.1.1 Child Support

The Texas Family Code 231.006 places certain restrictions on child support obligors. Texas Family Code §231.006(d) requires a person who applies for, bids on, or contracts for state funds to submit a statement that the person is not delinquent in paying child support. This law applies to an individual whose business is a sole proprietorship, partnership, or corporation in which the individual has an ownership interest of at least 25 percent of the business entity. This law does not apply to contracts/agreements with governmental entities or nonprofit corporations.

The required statement has been incorporated into the Texas Medicaid Provider Agreement.

The law also requires that payments be stopped when notified that the contractor/provider is more than 30 days delinquent in paying child support. Medicaid payments are placed on hold when it is discovered that a currently enrolled provider is delinquent in paying child support. A provider application may be denied or terminated if the provider is delinquent in paying child support.

1.7.1.2 Reporting Child Abuse or Neglect

Title 5 Texas Family Code (TFC) §261.101 states: “(a) A person having cause to believe that a child’s physical or mental health or welfare has been adversely affected by abuse or neglect by any person shall immediately make a report as provided by this subchapter; (b) If a professional has cause to believe that a child has been abused or neglected, or may be abused or neglected, or that a child is a victim of an offense under section 21.11, Penal Code, and the professional has cause to believe that the child has been abused as defined by section 261.001 or 261.401, the professional shall make a report no later than the 48th hour after the hour the professional first suspects that the child has been, or may be abused or neglected, or is a victim of an offense under section 21.11, Penal Code.” A professional may not delegate to or rely on another person to make the report. In this subsection, professional means an individual who is licensed or certified by the state or who is an employee of a facility licensed, certified, or operated by the state and who, in the normal course of official duties or duties for which a license or certification is required, has direct contact with children. The term includes teachers, nurses, doctors, day-care employees, employees of a clinic or health-care facility that provides reproductive services, juvenile probation officers, and juvenile detention or correctional officers.
All Medicaid providers shall comply with the provisions of state law as set forth in Chapter 261 of the Texas Family Code relating to investigations of reports of child abuse and neglect and the provisions of HHSC policy. Reimbursement shall only be made to providers who have demonstrated a good faith effort to comply with child abuse reporting guidelines and requirements in Chapter 261 and HHSC policy. Provider staff shall respond to disclosures or suspicions of abuse or neglect of minors, by reporting to the appropriate agencies as required by law.

All providers shall adopt this policy as their own, report suspected sexual abuse of a child as described in this policy and as required by law, and develop internal policies and procedures that describe how to determine, document, and report instances of sexual or nonsexual abuse.

This information is also available on the HHSC and TMHP websites at [https://hhs.texas.gov/laws-regulations/handbooks/fpp/section-3000-abuse-neglect-reporting](https://hhs.texas.gov/laws-regulations/handbooks/fpp/section-3000-abuse-neglect-reporting) and [www.tmhp.com](http://www.tmhp.com).

**1.7.1.3 Procedures for Reporting Abuse or Neglect**

Professionals as defined in the law are required to report no later than the 48th hour after the hour the professional first has cause to believe the child has been or may be abused or is the victim of the offense of indecency with a child.

Nonprofessionals shall immediately make a report after the nonprofessional has cause to believe that the child’s physical or mental health or welfare has been adversely affected by abuse.

A report shall be made regardless of whether the provider staff suspect that a report may have previously been made.

Reports of abuse or indecency with a child must be made to one of the following:

- Department of Family and Protective Services (DFPS) if the alleged or suspected abuse involves a person responsible for the care, custody, or welfare of the child (the DFPS Texas Abuse/Neglect Hotline, at 1-800-252-5400, operated 24 hours a day, 7 days a week)
- Any local or state law enforcement agency
- The state agency that operates, licenses, certifies, or registers the facility in which the alleged abuse or neglect occurred
- The agency designated by the court to be responsible for the protection of children

The law requires the report to include the following information if known:

- The name and address of the minor
- The name and address of the minor’s parent or the person responsible for the care, custody, or welfare of the child if not the parent
- Any other pertinent information concerning the alleged or suspected abuse

Reports can be made anonymously.

A provider may not reveal whether the child has been tested or diagnosed with human immunodeficiency virus (HIV) or acquired immunodeficiency syndrome (AIDS).

If the minor’s identity is unknown (e.g., the minor is at the provider’s office anonymously to receive testing for HIV or a sexually transmitted disease [STD]), no report is required.

**1.7.1.4 Procedures for Reporting Suspected Sexual Abuse**

All providers shall ensure that their employees, volunteers, or other staff report a victim of abuse who is a minor 14 years of age or younger who has engaged in sexual activity with any individual to whom the minor is not married. Sexual activity would be indicated if the minor is pregnant or has a confirmed STD acquired in a manner other than through perinatal transmission.
Sexual activity may include, but is not limited to, the actions described in Penal Code §21.11(a) relating to indecency with a child; §21.01(2) defining sexual contact; §43.01(1) or (3)-(5) defining various sexual activities; §22.011(a)(2) relating to sexual assault of a child; or §22.021(a)(2) relating to aggravated sexual assault of a child.

Providers may voluntarily use the HHSC checklist for monitoring all clients who are 14 years of age or younger, unmarried, and sexually active. The checklist, if used, as well as any report of child abuse, shall be retained as part of the client’s record by each provider and made available during any monitoring conducted by HHSC.

Refer to: Texas Medicaid - Title XIX Acknowledgment of Hysterectomy Information on the TMHP website at www.tmhp.com.

1.7.1.5 Training
All providers must develop training for all staff on the policies and procedures in regard to reporting child abuse. New staff must receive this training as part of their initial training/orientation. Training must be documented. As part of the training, staff must be informed that the staff person who conducts the screening and has cause to suspect abuse has occurred is legally responsible for reporting. A joint report may be made with the supervisor.

1.7.1.6 Reporting Abuse and Neglect of the Elderly or Disabled
Title 2 Human Resources Code (HRC) §48.051 states: “(a) Except as prescribed by Subsection (b), a person having cause to believe that an elderly or disabled person is in the state of abuse, neglect, or exploitation, including a disabled person receiving services as described by Section 48.252, shall report the information required by Subsection (d) immediately to the Department of Health and Human Services and Department of Protective and Regulatory Services. (b) If a person has cause to believe that an elderly or disabled person, other than a disabled person receiving services as described by Section 48.252, has been abused, neglected, or exploited in a facility operated, licensed, certified, or registered by a state agency, the person shall report the information to the state agency that operates, licenses, certifies, or registers the facility for investigation by that agency. (c) The duty imposed by Subsections (a) and (b) applies without exception to a person whose knowledge concerning possible abuse, neglect, or exploitation is obtained during the scope of the person’s employment or whose professional communications are generally confidential, including an attorney, clergy member, medical practitioner, social worker, and mental health professional.”

(d) The report may be made orally or in writing. It shall include:

- The name, age, and address of the elderly or disabled person;
- The name and address of any person responsible for the elderly or disabled person’s care;
- The nature and extent of the elderly or disabled person’s condition;
- The basis of the reporter’s knowledge; and
- Any other relevant information.

(e) If a person who makes a report under this section chooses to give self-identifying information, the caseworker who investigates the report shall contact the person if necessary to obtain any additional information required to assist the person who is the subject of the report.”
1.7.1.7 Procedures for Reporting Abuse or Neglect of the Elderly or Disabled

Title 2 HRC §48.151 states: "(a) Not later than 24 hours after the department receives a report of an allegation of abuse, neglect, or exploitation under Section 48.051, the Department of Health and Human Services and Department of Protective and Regulatory Services shall initiate a prompt and thorough investigation as needed to evaluate the accuracy of the report and to assess the need for protective services, unless the department determines that the report:

- Is frivolous or patently without a factual basis; or
- Does not concern abuse, neglect, or exploitation, as those terms are defined by rules adopted by the executive commissioner under Section 48.002(c), except that if the executive commissioner has not adopted applicable rules under that section, the statutory definitions of those terms under Section 48.002(a) shall be used.

(b) The Department of Health and Human Services and Department of Protective and Regulatory Services have adopted rules for conducting investigations under this chapter.

(c) The Department of Human Services and Department of Protective and Regulatory Services by rule may assign priorities and prescribe investigative procedures for conducting investigations according to the degree of severity and immediacy of the alleged harm to the individual. Notwithstanding Subsection (a), the department’s priorities and procedures may provide that an investigation is not required to be initiated within 24 hours in all cases.

(d) The Department of Human Services and Department of Protective and Regulatory Services shall prepare and keep on file a report of each investigation conducted by the department.

(e) This section does not apply to investigations conducted under Subchapter F or H."

1.7.2 Maintenance of Provider Information

Within 90 calendar days of occurrence, providers must report changes in address (physical location or accounting), telephone number, name, federal tax ID, and any other information that pertains to the structure of the provider’s organization (for example, performing providers). Changes in address, office telephone or fax number, and email address should be updated online using the OPL update page. Alternatively, providers may update their address information using the PIC Form, which is available on the TMHP website. A W9 is required if the provider is changing the mailing address. A copy of the Medicare approval letter listing the additional location or site must be submitted when adding alternate physical addresses. Failure to provide this information may lead to administrative action by HHSC.

Refer to: Subsection 1.7.2.3, “Online Provider Lookup (OPL)” in this section.

Refer to: Subsection 1.4.1, “Medicare Number” in this section.

Providers are notified when they have an invalid address on file with TMHP. Account administrators who log onto their accounts through the TMHP website at www.tmhp.com are notified when they have an invalid address on file for any of the TPIs associated with their NPI.

The Check Status Amount Search screen on the provider’s secure homepage of the TMHP website will alert providers when payments are pending because of inaccurate or incomplete provider information. R&S Reports that are viewed on the TMHP website also notify the provider of pending payments.

Pending payments are released in the financial cycle of the following week after the address information has been updated. Payments that are pending for more than 180 days will be voided.
Other changes (in name, ownership status, federal tax ID, etc.) must be reported in writing to TMHP Provider Enrollment. Failure to notify TMHP of changes affects accurate processing and timely claims payment. In addition, failure to timely report such changes is a violation of the rules of Medicaid, and may result in administrative, civil, or criminal liability.

Refer to: Subsection 1.11, “Medicaid Fraud, Waste, and Abuse Policy” in this section.

Providers will be prompted to verify their address(es) and make necessary changes at least once a year. After the PIC form has been completed, it can be uploaded to PIMS, faxed to 1-512-514-4214, Attn: Provider Enrollment, or mailed to:

Texas Medicaid & Healthcare Partnership
Provider Enrollment
PO Box 200795
Austin, TX 78720-0795

Providers should keep a copy of the completed form for their records.

Providers that have a moderate or high risk category cannot render or submit claims for services at a new practice location until it has been approved and added to the enrollment record. Providers are encouraged to check PIMS for verification that the practice location has been approved prior to rendering or submitting claims for services.

Refer to: The Affordable Care Act (ACA) Provider Enrollment Frequently Asked Questions on the TMHP website at www.tmhp.com for more information on risk category screening requirements.

1.7.2.1 Contracted Provider Groups

Contracted provider groups, independent physician groups, and physician management companies that have a hospital-based practice location must ensure that the organization’s contact information is listed as the accounting or mailing address rather than the hospital’s information. Providers can check this information in Provider Information Management System (PIMS).

1.7.2.2 NPI Verification

TMHP verifies NPIs with NPPES to ensure that the NPI is active. If the NPI is shown by NPPES to be inactive, TMHP will notify the provider by letter.

The provider will be allowed a 60-day grace period to contact NPPES and resolve their NPI status. If the inactive NPI has not been reinstated within the 60-day grace period, TMHP will disenroll all TPIs associated with the inactive NPI.

1.7.2.3 Online Provider Lookup (OPL)

The OPL is available on the public access portion of the TMHP website at www.tmhp.com. Provider information can be viewed by providers, clients, and anyone who accesses the TMHP website.

Providers with certain provider types must verify and update key demographic information every six months in the Provider Information Management System (PIMS) to ensure their information is correct in the OPL. Affected provider types include, but are not limited to, physicians, nurses, dentists, and durable medical equipment providers.

If more than six months have elapsed since the required demographic information in the OPL was verified, access to the secure provider portal is blocked until the verification takes place. Upon logging into their accounts, users with administrative rights see a list of NPIs that require verification and update. After addressing each NPI listed on the page, administrative providers are able to proceed to their accounts.
If access to the secure portal has been blocked because of needed verification, nonadministrative users are not able to perform work functions on NPIs listed on the Review Required page. Nonadministrative users are advised to notify users with administrative rights so that they can verify demographic information and remove the block.

The My Account page has a link to the Provider Demographic Update web page. Current information will be displayed with a button to allow editable fields to be changed. Demographic information may be updated only by authorized administrators. This authorization is controlled through the Permissions Management link, also located on the My Account page. Fields that can be updated online include the following:

- Primary physical address:
  - Street address lines 1 and 2
  - City, state, ZIP Code
  - County
- Telephone numbers
- Email address
- Office hours
- Accepting new clients, current clients only, or not accepting new clients
- Additional sites where services are provided
- Languages spoken
- Additional services offered
- Medicaid waiver programs
- Client age or gender limitations
- Counties served

The following enhancements have also been made to the OPL to increase overall functionality:

- Clients are able to search for providers in up to 5 counties in a single search.
- Doing business as (DBA) names appear for providers or provider groups.
- The State of Texas Access Reform (STAR) Health program has been added as a searchable health plan.
- The default ZIP Code radius for provider search has been increased to 10 miles from 5 miles.
- Providers who make address updates may receive a confirmation email from TMHP after the address has been verified and if their email address has been provided.
- Users will be able to search for providers within a ZIP Code that crosses multiple counties.

Each provider specialty and subspecialty listed in the OPL now has a corresponding definition. Users can view the definitions by clicking “more information” on either the basic or advanced search page or by hovering over the specialty on the results page. The definitions have been added to help clients locate the correct type of provider.

Providers are able to self-declare as many as three subspecialties to identify the services they offer. Providers may declare only subspecialties that are within the scope of their practice. Users are able to search for a provider on the OPL using these subspecialties.
Clients using the OPL will use drop-down boxes to select search criteria. An initial list will display all providers that meet the specified search criteria. Clicking on any name in that list will display the provider’s specific information, including a map of the office location.

Links to health maintenance organization (HMO) websites are also provided, enabling clients to search each HMO’s network of participating providers. The OPL supports both English and Spanish language users, and search results can be printed.

### 1.7.2.4 Updating NPI and Taxonomy Codes

Providers are required to provide their NPI in the enrollment application. During the enrollment process, providers must also select a primary and, if applicable, secondary taxonomy codes associated with their provider type. Due to copyright restrictions, TMHP is unable to publish the taxonomy descriptions. Providers must verify the taxonomy codes associated with their provider type and specialty before beginning the enrollment process.

**Refer to:** Subsection 1.1.2, “NPI and Taxonomy Codes” in this section.

Providers must maintain and update their NPI and/or taxonomy code information with Texas Medicaid. The available taxonomy code selections are auto populated according to the provider type and specialty associated with the TPI entered. The taxonomy code options may not match the taxonomy code listed in the confirmation letter received from NPPES. Providers must contact the TMHP Contact Center at 1-800-925-9126 to validate their provider type and specialty associated with their TPIs.

**Refer to:** Subsection 1.7.2.5, “Updating Provider Specialty” in this section if a taxonomy code that you want to use is not available for the enrolled provider type or specialty.

**Important:** The taxonomy code that is included in electronic transactions must match a taxonomy code that is included in the attestation record. Secondary taxonomy codes included during the attestation process are used as additional matching criteria for claims and authorization processing.

### 1.7.2.5 Updating Provider Specialty

Providers that have made a change in their specialty must submit their updated specialty information to Texas Medicaid. The forms that must be submitted to Texas Medicaid depend on the provider’s enrollment, as follows:

- Medicare-enrolled providers whose Medicare number has not changed must submit a copy of the Medicare letter listing the updated specialty along with a PIC Form to Texas Medicaid. The PIC form and specialty letter may be uploaded to PIMS, faxed to TMHP (Attention Provider Enrollment) at 1-512-514-4214, or mailed to:
  
  Texas Medicaid & Healthcare Partnership  
  Provider Enrollment  
  PO Box 200795  
  Austin, TX 78720-0795

- Providers that are not enrolled in Medicare or whose Medicare number has changed must submit a new application

**Refer to:** [Provider Information Change Form](https://www.tmhp.com) on the TMHP website at [www.tmhp.com](http://www.tmhp.com).

**Refer to:** Subsection 1.4, “Provider Reenrollment” in this section for more information about provider reenrollment in Texas Medicaid.

### 1.7.3 Retention of Records and Access to Records and Premises

The provider must maintain and retain all necessary documentation, records, R&S Reports, and claims to fully document the services and supplies provided and delivered to a client with Texas Medicaid coverage, the medical necessity of those services and supplies, costs included in cost reports or other
documents used to determine a payment rate or fee, and records or documents necessary to determine whether payment for those items or services was due and was properly made for full disclosure to HHSC and its designee. A copy of the claim or R&S Reports without additional documentation will not meet this requirement.

The documentation includes the following, without limitation:

- Patient clinical health records
- Other records pertaining to the patient
- Any other records of services, items, equipment, or supplies provided to the patient and payments made for those services
- Diagnostic tests
- Documents related to diagnosis
- Charting
- Billing records
- Invoices
- Treatments
- Services
- Laboratory results
- X-rays
- Documentation of delivery of items, equipment, and supplies

Accessible information must include information that is necessary for the agencies specified in this section to perform statutory functions.

**Note:** The required information may also include, without limitation, business and accounting records with backup support documentation, statistical documentation, computer records and data, and patient sign-in sheets and schedules. Additionally, it includes all requirements and elements described in 1 TAC §§371.1607 and 371.1667 (definition of “failure to grant immediate access”).

The provider is required to submit original documents, records, and accompanying business records affidavits to representatives of the organizations listed in this section. These records should also be provided to any agents and contractors related to the organizations. At the discretion of the requestor, the provider may be permitted to instead provide copies notarized with the required business records affidavit. Requested records must be provided promptly and at no cost to the state or federal agency. If the provider was originally requested to provide original documents and subsequent requests for copies of these records are made by the provider, any and all costs associated with copying or reproducing any portion of the original records will be at the expense of the provider. This applies to any request for copies made by the provider at any point in the investigative process until such time as the agency deems the investigation to be finalized. A method of payment for the copying charge, approved by the agency, would be used to pay for the copying of the records. If copies of records are requested from the provider initially, the provider must submit copies of such records at no cost to the requestor’s organization.

The provider must provide immediate access to the provider’s premises and records for purposes of reviewing, examining, and securing custody of records, documents, electronic data, equipment, or other requested items, as determined necessary by the requestor to perform statutory functions. Nothing in this section will in any way limit access otherwise authorized under state or federal law. If, in the opinion of the Inspector General or other requestor, the documents may be provided at the time of the request or in less than 24 hours or the Inspector General or other requestor suspects the requested documents or other requested items may be altered or destroyed, the response to the request must be completed by
the provider at the time of the request or in less than 24 hours as allowed by the requestor. If, in the opinion of the Inspector General or other requestor, the requested documents and other items requested cannot be completely provided on the day of the request, the Inspector General or requestor may set the deadline for production at 24 hours from the time of the original request.

Failure to supply the requested documents and other items, within the time frame specified, may result in payment hold to the provider’s Medicaid payments, recoupment of payments for all claims related to the missing records, contract cancellation, and/or exclusion from Texas Medicaid.

As directed by the requestor, the provider or person will relinquish custody of the requested documents and other items and the requestor will take custody of the records, removing them from the premises. If the requestor should allow longer than “at the time of the request” to produce the records, the provider will be required to produce all records completed, at the time of the completion or at the end of each day of production, as directed by the requestor who will take custody of the requested items.

If the provider places the required information in another legal entity’s records, such as a hospital, the provider is responsible for obtaining a copy of these requested records for use by the requesting state and federal agencies.

These documents and claims must be retained for a minimum period of five years from the date of service or until all audit questions, appeal hearings, investigations, or court cases are resolved. Freestanding RHCs must retain their records for a minimum of six years, and hospital-based RHCs must retain their records for a minimum of ten years. These records must be made available immediately at the time of the request to employees, agents, or contractors of HHSC OIG, the Office of the Attorney General (OAG) Medicaid Fraud Control Unit (MFCU) or Antitrust and Civil Medicaid Fraud Section, TMHP, DFPS, HHSC, DSHS, Texas Workforce Commission (TWC), U.S. Department of Health and Human Services (HHS) representative, any state or federal agency authorized to conduct compliance, regulatory, or program integrity functions on the provider, person, or the services rendered by the provider or person, or any agent, contractor, or consultant of any agency or division delineated above. In addition, the provider must meet all requirements of 1 TAC §371.1667.

The records must be available as requested by each of these entities, during any investigation or study of the appropriateness of the Medicaid claims submitted by the provider.

1.7.3.1 Payment Error Rate Measurement (PERM) Process

CMS assesses Texas Medicaid using the PERM process to measure improper payments in Texas Medicaid. Providers will be required to provide medical record documentation to support the medical reviews that the federal review contractor will conduct for Texas Medicaid fee-for-service and Primary Care Case Management Medicaid and State Children’s Health Insurance Program (SCHIP) claims.

Under the PERM process, if a claim is selected in a sample for a service that a provider rendered to a Medicaid client, the provider will be contacted to submit a copy of the medical records that support the medical review of the claim. All providers should check the TMHP system to ensure their current telephone number and addresses are correct in the system. If the information is incorrect or incomplete, providers must request a change immediately to ensure the PERM medical record request can be delivered. Client authorization for release of this information is not required.

Once a provider receives the request for medical records, the provider must submit the information electronically or in hard copy within 60-calendar days. It is important that providers cooperate by submitting all requested documentation in a timely manner because no response or insufficient documentation will count against the state as an error. This can ultimately negatively impact the amount of federal funding received by Texas for Medicaid.

1.7.4 Medicare Overpayment

Title 42 CFR §447.30 provides for withholding the federal share of Medicaid payments to recover Medicare overpayments from providers in a coordinated effort with CMS.
1.7.5 Credit Balance and Recovery Vendor
Trend Health Partners helps Texas Medicaid resolve credit balances and recover overpayments. Trend Health Partners reviews the credit balances of all current accounts with claims that received a primary or secondary payment from both TMHP and a health insurance carrier, but the health insurance carrier was liable for payment before Medicaid.

1.7.6 Release of Confidential Information
Information regarding the diagnosis, evaluation, or treatment of a client with Texas Medicaid coverage by a person licensed or certified to diagnose, evaluate, or treat any medical, dental, mental/emotional disorder, or drug abuse, is confidential information that the provider may disclose only to authorized persons. Family planning information is sensitive, and confidentiality must be ensured for all clients, especially minors.

Only the client may give written permission for release of any pertinent information before client information can be released, and confidentiality must be maintained in all other respects. If a client’s medical records are requested by a licensed Texas health-care provider or a provider licensed by any state, territory, or insular possession of the United States or any State or province of Canada, for purposes of emergency or acute medical care, a provider must furnish such records at no cost to the requesting provider. This includes records received from another physician or health-care provider involved in the care or treatment of the patient. If the records are requested for purposes other than for emergency or acute medical care, the provider may charge the requesting provider a reasonable fee and retain the requested information until payment is received.

The client’s signature is not required on the claim form for payment of a claim, but HHSC recommends the provider obtain written authorization from the client before releasing confidential medical information. A release may be obtained by having the client sign the indicated block on the claim form after the client has read the statement of release of information that is printed on the back of the form. The client’s authorization for release of such information is not required when the release is requested by and made to HHSC, DSHS, TMHP, DFPS, TWC, HHSC OIG, the MFCU or Antitrust and Civil Fraud Division, or HHS.

1.7.7 Compliance with Federal Legislation
HHSC complies with HHS regulations that protect against discrimination. All contractors must agree to comply with the following:

- Title VI of the Civil Rights Act of 1964 (Public Law 88-352), section 504 of the Rehabilitation Act of 1973 (Public Law 93-112), the Americans with Disabilities Act of 1990 (Public Law 101-336), Title 40, Chapter 73, of the TAC, all amendments to each, and all requirements imposed by the regulations issued pursuant to these acts. The laws provide in part that no persons in the United States (U.S.) shall, on the grounds of race, color, national origin, age, sex, disability, political beliefs, or religion, be excluded from participation in or denied any aid, care, service, or other benefits provided by federal and/or state funding, or otherwise be subjected to any discrimination.
• Health and Safety Code 85.113 as described in “Model Workplace Guidelines for Businesses, State Agencies, and State Contractors” on page G-2 (relating to workplace and confidentiality guidelines on AIDS and HIV)

Exception: In the case of minors receiving family planning services, only the client may consent to release of health-care information. Providers must comply with the laws and regulations concerning discrimination. Payments for services and supplies are not authorized unless the services and supplies are provided without discrimination on the basis of race, color, sex, national origin, age, or disability. Send written complaints of noncompliance to the following address:

Executive Commissioner
1100 West 49th Street
Austin, TX 78756-3172

Reminder: Each provider must furnish covered Medicaid services to eligible clients in the same manner, to the same extent, and of the same quality as services provided to other patients. Services made available to other patients must be made available to Texas Medicaid clients if the services are benefits of Texas Medicaid.

1.7.8 Tamper-Resistant Prescription Pads

Providers are required by federal law (Public Law 110-28) to use a tamper-resistant prescription pad when writing a prescription for any drug for Medicaid clients.

Providers must take necessary steps to ensure that tamper-resistant pads are used for all written prescriptions provided to Medicaid clients. Providers may also use compliant, non-written alternatives for transmitting prescriptions such as by telephone, fax, or electronic submittal. Pharmacies are required to ensure that all written Medicaid prescriptions submitted for payment to the Vendor Drug Program are written on a compliant tamper-resistant pad.

If a prescription is not submitted on a tamper-resistant prescription form, a pharmacy may fill the prescription and obtain a compliant prescription by fax, electronic prescription, or re-written on tamper-resistant paper within 72 hours after the date the prescription was filled.

Providers may purchase tamper-resistant prescription pads from the vendor of their choice.

Special copy-resistant paper is not a requirement for prescriptions printed from electronic health records (EHRs) or ePrescribing generated prescriptions. These prescriptions may be printed on plain paper and will be fully compliant with all three categories of the tamper-resistant regulations, provided they contain at least one feature from each of the three following categories:

• Prevents unauthorized copying of completed or blank prescription forms.
• Prevents erasure or modification of information written on the prescription form.
• Prevents the use of counterfeit prescription forms.

1.7.9 Utilization Control — General Provisions

Title XIX of the Social Security Act, sections 1902 and 1903, mandates utilization control of all Texas Medicaid services under regulations found at Title 42 CFR, Part 456. Utilization review activities required by Texas Medicaid are completed through a series of monitoring systems developed to ensure the quality of services provided, and that all services are both medically necessary and billed appropriately. Both clients and providers are subject to utilization review monitoring. Utilization control procedures safeguard against the delivery of unnecessary services, monitor quality, and ensure payments are appropriate and according to Texas Medicaid policies, rules, and regulations. All providers identified as a result of utilization control activities are presented to HHSC-OIG to determine any and all subsequent actions.
The primary goal of utilization control activity is to identify providers with practice patterns inconsistent with the federal requirements and Texas Medicaid scope of benefits, policies, and procedures. The use of utilization control monitoring systems allows for identification of providers whose patterns of practice and use of services fall outside of the norm for their peer groups. Providers identified as exceptional are subject to an in-depth review of all Texas Medicaid billings. These review findings are presented to the HHSC-OIG to determine any necessary action. Medical records may be requested from the provider to substantiate the medical necessity and appropriateness of services billed to Texas Medicaid. Inappropriate service utilization may result in recoupment of overpayments and/or sanctions, or other administrative actions deemed appropriate by the HHSC-OIG. There are instances when a training specialist may be directed to communicate with the provider to offer assistance with the technical or administrative aspects of Texas Medicaid.

At the direction of the HHSC-OIG, a provider’s claims may be manually reviewed before payment. Parameters are developed for prepayment review based on the specific areas of concern identified in each case. As part of the prepayment review process, providers are required to submit paper claims, rather than electronic claims, along with supporting medical record documentation (e.g., clinical notes, progress notes, diagnostic testing results, other reports, superbills, X-rays, and any related medical record documentation) attached to each claim for all services billed. This documentation is used to ascertain that the services billed were medically necessary, billed appropriately, and according to Texas Medicaid requirements and policies. Services inconsistent with Texas Medicaid requirements and policies are adjudicated accordingly. Claims submitted initially without the supporting medical record documentation will be denied. Additional medical record documentation submitted by the provider for claims denied as a result of the prepayment review process is not considered at a later time. A provider is removed from prepayment review only when determined appropriate by the HHSC-OIG. Once removed from prepayment review, a follow-up assessment of the provider’s subsequent practice patterns is performed to monitor and ensure continued appropriate use of resources. Noncompliant providers are subject to administrative sanctions up to and including exclusion and contract cancellation, as deemed appropriate by the HHSC-OIG as defined in the rules in 1TAC §§371.1701, 371.1703, 371.1705, 371.1707, 371.1709, 371.1711, 371.1713, and 371.1715. Providers placed on prepayment review must submit all paper claims and supporting medical record documentation to the following address:

Texas Medicaid & Healthcare Partnership
Attention: Prepayment Review MC-A11 SURS
PO Box 203638
Austin, Texas 78720-3638

1.7.10 Provider Certification/Assignment

Texas Medicaid service providers are required to certify compliance with or agree to various provisions of state and federal laws and regulations. After submitting a signed claim to TMHP, the provider certifies the following:

- Services were personally rendered by the billing provider or under supervision of the billing provider, if allowed for that provider type, or under a substitute arrangement.
- The information on the claim form is true, accurate, and complete.
- All services, supplies, or items billed were medically necessary for the client’s diagnosis or treatment. Exception is allowed for special preventive and screening programs (for example, family planning and THSteps).
- Health records document all services billed and the medical necessity of those services.
- All billed charges are usual and customary for the services provided. The charges must not be higher than the fees charged to private-pay patients.
• The provider will not bill Texas Medicaid for services that are provided or offered to non-Medicaid patients, without charge, discounted or reduced in any fashion including, but not limited to, sliding scales or advertised specials. Any reduced, discounted, free, or special fee advertised to the public must also be offered to Texas Medicaid clients.

• Services were provided without regard to race, color, sex, national origin, age, or handicap.

• The provider of health care and services files a claim with Texas Medicaid agreeing to accept the Medicaid reimbursement as payment in full for those services covered under Texas Medicaid. In accordance with 1 TAC §354.1005, the reimbursement for services covers the costs for a covered service, and any function incidental to the provision of a covered service (refer to subsection 1.7.11, “Billing Clients” for more information). The client with Medicaid coverage, or others on their behalf, must not be billed for the amount above that which is paid on allowed services or for services denied or reduced as a result of errors made in claims filing, claims preparation, missed filing deadlines, or failure to follow the appropriate appeal process. However, the client may be billed for noncovered services for which Texas Medicaid does not make any payment. Before providing services, providers should always inform clients of their liability for services that are not a benefit of Texas Medicaid, including use of the Client Acknowledgment Statement.

• The provider understands that endorsing or depositing a Texas Medicaid check is accepting money from federal and state funds and that any falsification or concealment of material fact related to payment may be grounds for prosecution under federal and state laws.

Providers must not bill for, and agree not to bill for, any service provided for which the client bears no liability to pay (i.e. free services). The only exceptions to this ban on billing for services that are free to the user are:

• Services offered by or through the Title V agency when the service is a benefit of Texas Medicaid and rendered to an eligible client

• Services included in the Texas Medicaid client’s individualized education plan (IEP) or individualized family service plan (IFSP) if the services are covered under the Title XIX state plan, even though they are free to the users of the services


Subsection 1.7.10.1, “Delegation of Signature Authority” in this section.

1.7.10.1 Delegation of Signature Authority

A provider delegating signatory authority to a member of the office staff or to a billing service remains responsible for the accuracy of all information on a claim submitted for payment. A provider’s employees or a billing service and its employees are equally responsible for any false billings in which they participated or directed.

If the claim is prepared by a billing service or printed by data processing equipment, it is permissible to print “Signature on File” in place of the provider’s signature. When claims are prepared by a billing service, the billing service must obtain and keep a letter on file that is signed by the provider authorizing claim submission.

1.7.11 Billing Clients

A provider cannot require a down payment before providing Medicaid-allowable services to eligible clients, bill, nor take recourse against eligible clients for denied or reduced claims for services that are within the amount, duration, and scope of benefits of Texas Medicaid if the action is the result of any of the following provider-attributable errors:

• Failure to submit a claim, including claims not received by TMHP
• Failure to submit a claim to TMHP for initial processing within the 95-day filing deadline (or the initial 365-day deadline, if applicable)
• Submission of an unsigned or otherwise incomplete claim such as omission of the Hysterectomy Acknowledgment Statement or Sterilization Consent Form with claims for these procedures
• Filing an incorrect claim
• Failure to resubmit a corrected claim or rejected electronic media claim within the 120-day resubmittal period
• Failure to appeal a claim within the 120-day appeal period. Errors made in claims preparation, claims submission, or appeal process
• Failure to submit a claim to TMHP within 95 days of a denial by the DSHS Family Planning Program for family planning services
• Failure to submit a claim within 95 days from the disposition date from Medicare or a primary third party insurance resource
• Failure to obtain prior authorization for services that require prior authorization under Texas Medicaid

Providers must certify that they will accept the reimbursement paid by Texas Medicaid for covered services and will not bill an eligible client for covered services. Federal regulations prohibit providers from charging clients a fee for completing or filing Medicaid claim forms. Providers are not allowed to charge TMHP for filing claims. The cost of claims filing is part of the usual and customary rate for doing business.

Medicaid reimbursement is considered as payment in full for those services covered under Texas Medicaid. In accordance with 1 TAC §354.1005, the reimbursement for services is intended to cover the costs for a covered service, or any function incidental to the provision of a covered service, including, but not limited to:

• Signing, completing, or providing a copy of a health assessment form, such as a physical examination form required for the eligible client’s enrollment in school or participation in school or other activities;
• Providing a copy of a medical record requested:
  • By or on behalf of any health care practitioner for purposes of medical care or treatment of the eligible client;
  • As a supplement to a health assessment form or other form provided incidental to a covered service; or
  • By an eligible client, for any reason, for the first time in a one-year period; and
• Providing a copy of any subsequent amendment, supplement, or correction to a medical record requested by or on behalf of the eligible client.
• If the provider has already provided the eligible client a free copy of the medical record within a one-year period, the provider is required to provide only the amended, supplemented, or corrected portion of the record, if requested, without having to copy the entire record.

Note: A provider may bill or otherwise charge a client a reasonable fee for providing a paper copy of a medical record outside of the above scenarios. A reasonable fee for providing a paper copy of the requested records shall be a charge of no more than $25.00 for the first twenty pages and $.50 per page for every copy thereafter per 22 TAC §165.2.
Completion of required forms submitted by a nursing facility to the physician for signature is also considered incidental to a covered service. It is not acceptable for the physician to charge Texas Medicaid clients, their family, or the nursing facility for telephone calls, telephone consultations, or signing forms.

In accordance with current federal policy, Texas Medicaid and Texas Medicaid clients cannot be charged for the client’s failure to keep an appointment. Only billings for services provided are considered for payment. Clients may not be billed for the completion of a claim form, even if it is a provider’s office policy.

Letters of inquiry about client billing are sometimes sent to providers in lieu of telephone calls from TMHP representatives. In either case, it is mandatory that the questions be answered with the requested pertinent information. Upon receipt, TMHP forwards these letters to HHSC. HHSC uses the information to resolve client billing/liability issues. It is mandatory that these letters be signed, dated, and returned within ten business days.

**Refer to:** The *Inpatient and Outpatient Hospital Services Handbook (Vol. 2, Provider Handbooks)* for more information about spell-of-illness.

Subsection 4.5, “Medically Needy Program (MNP)” in “Section 4: Client Eligibility” (Vol. 1, General Information).

Private Pay Agreement on the TMHP website at [www.tmhp.com](http://www.tmhp.com).

### 1.7.11.1 Client Acknowledgment Statement

Texas Medicaid only reimburses services that are medically necessary or benefits of special preventive and screening programs such as family planning and THSteps. Hospital admissions denied by the Texas Medical Review Program (TMRP) also apply under this policy.

The provider may bill the client only if:

- A specific service or item is provided at the client’s request.
- The provider has obtained and kept a written Client Acknowledgment Statement signed by the client that states:
  - “I understand that, in the opinion of (provider’s name), the services or items that I have requested to be provided to me on (dates of service) may not be covered under the Texas Medical Assistance Program as being reasonable and medically necessary for my care. I understand that the HHSC or its health insuring agent determines the medical necessity of the services or items that I request and receive. I also understand that I am responsible for payment of the services or items I request and receive if these services or items are determined not to be reasonable and medically necessary for my care.”
  - “Comprendo que, según la opinión del (nombre del proveedor), es posible que Medicaid no cubra los servicios o las provisiones que solicité (fecha del servicio) por no considerarlos razonables ni médicamente necesarios para mi salud. Comprendo que el Departamento de Salud de Texas o su agente de seguros de salud determina la necesidad médica de los servicios o de las provisiones que el cliente solicite o reciba. También comprendo que tengo la responsabilidad de pagar los servicios o provisiones que solicite y que reciba si después se determina que esos servicios y provisiones no son razonables ni médicamente necesarios para mi salud.”

A provider is allowed to bill the following to a client without obtaining a signed Client Acknowledgment Statement:

- Any service that is not a benefit of Texas Medicaid (for example, cellular therapy).
- All services incurred on noncovered days because of eligibility or spell of illness limitation. Total client liability is determined by reviewing the itemized statement and identifying specific charges incurred on the noncovered days. Spell of illness limitations do not apply to medically necessary stays for Medicaid clients who are 20 years of age and younger.
• The reduction in payment that is due to the Medically Needy Program (MNP) is limited to children who are 18 years of age and younger and pregnant women. The client’s potential liability would be equal to the amount of total charges applied to the spend down. Charges to clients for services provided on ineligible days must not exceed the charges applied to spend down.

• All services provided as a private pay patient. If the provider accepts the client as a private pay patient, the provider must advise clients that they are accepted as private pay patients at the time the service is provided and responsible for paying for all services received. In this situation, HHSC strongly encourages the provider to ensure that the client signs written notification so there is no question how the client was accepted. Without written, signed documentation that the Texas Medicaid client has been properly notified of the private pay status, the provider cannot seek payment from an eligible Texas Medicaid client.

• The client is accepted as a private pay patient pending Texas Medicaid eligibility determination and does not become eligible for Medicaid retroactively. The provider is allowed to bill the client as a private pay patient if retroactive eligibility is not granted. If the client becomes eligible retroactively, the client notifies the provider of the change in status. Ultimately, the provider is responsible for filing timely Texas Medicaid claims. If the client becomes eligible, the provider must refund any money paid by the client and file Medicaid claims for all services rendered.

A provider attempting to bill or recover money from a client in violation of the above conditions may be subject to exclusion from Texas Medicaid.

**Important:** Ancillary services must be coordinated and pertinent eligibility information must be shared. The primary care provider is responsible for sharing eligibility information with others (e.g., emergency room staff, laboratory staff, and pediatricians).

### 1.7.12 General Medical Record Documentation Requirements

The Administrative Simplification Act of HIPAA mandates the use of national coding and transaction standards. HIPAA requires that the American Medical Association’s (AMA) Current Procedural Terminology (CPT) system or the American Dental Association’s (ADA) Current Dental Terminology (CDT) system be used to report professional services, including physician and dental services. Correct use of CPT and CDT coding requires using the most specific procedure code that matches the services provided based on the procedure code’s description. Providers must pay special attention to the standard CPT descriptions for the evaluation and management services. The medical record must document the specific elements necessary to satisfy the criteria for the level of service as described in CPT. Reimbursement may be recouped when the medical record does not document that the level of service provided accurately matches the level of service claimed. Furthermore, the level of service provided and documented must be medically necessary based on the clinical situation and needs of the patient.

HHSC and TMHP routinely perform retrospective reviews of all providers. HHSC ultimately is responsible for Texas Medicaid utilization review activities. This review includes comparing services billed to the client’s clinical record. The following requirements are general requirements for all providers. Any mandatory requirement not present in the client’s medical record subjects the associated services to recoupment.

**Note:** This list is not all-inclusive. Additional and more specific requirements may apply to special services areas.

**Note:** Health-care documentation that is maintained by a provider in a client’s record can be maintained in a language other than English; however, when TMHP, HHSC, or any other state/federal agency requests a written record or conducts a documentation review, this health-care documentation must be provided in English and in a timely manner.

• (Mandatory) All entries are legible to individuals other than the author, dated (month, day, and year), and signed by the performing provider.
• (Mandatory) Medicaid-enrolled providers must submit claims with their own TPI except when under the agreement of a substitute provider or *locum tenens*.

• (Mandatory) Each page of the medical record documents the patient’s name and Texas Medicaid number.

• (Mandatory) A copy of the actual authorization from HHSC or its designee (e.g., TMHP) is maintained in the medical record for any item or service that requires prior authorization.

• (Mandatory) Allergies and adverse reactions (including immunization reactions) are prominently noted in the record.

• (Mandatory) The selection of evaluation and management codes (levels of service) is supported by the client’s clinical record documentation. Providers must follow either the 1995 or 1997 Documentation Guidelines for Evaluation and Management Services published by CMS, when selecting the level of service provided.

• (Mandatory) The history and physical documents the presenting complaint with appropriate subjective and objective information.

• (Mandatory) The services provided are clearly documented in the medical record with all pertinent information regarding the patient’s condition to substantiate the need and medical necessity for the services.

• (Mandatory) Medically necessary diagnostic lab and X-ray results are included in the medical record and abnormal findings have an explicit notation of follow-up plans.

• (Mandatory) Necessary follow-up visits specify time of return by at least the week or month.

• (Mandatory) Unresolved problems are noted in the record.

• (Desirable) Immunizations are noted in the record as complete or up-to-date.

• (Desirable) Personal data includes address, employer, home/work telephone numbers, sex, marital status, and emergency contacts.

  **Note:** An unenrolled provider that renders services and attempts to use the TPI of a provider who is enrolled in Medicaid will not be reimbursed for the services. During retrospective review, any services that were rendered by a provider that was not enrolled in Texas Medicaid and were billed using the provider identifier of a Medicaid-enrolled provider are subject to recoupment.

### 1.7.13 Informing Pregnant Clients About CHIP Benefits

Section 24, S.B. 1188, 79th Legislature, Regular Session, 2005, requires that Medicaid providers rendering services to a pregnant Medicaid client must inform the client of the health benefits for which the client or the client’s child may be eligible under the CHIP.

CHIP is available to children whose families have low to moderate income, who earn too much money to qualify for Texas Medicaid, and who do not have private insurance. Some clients may have to pay an enrollment fee.

To qualify for CHIP, a child must be:

• A Texas resident

• 18 years of age or younger

• A citizen or legal permanent resident of the United States

• Must meet all income and resource guidelines

CHIP benefits include:

• Physician, hospital, X-ray, and lab services
• Well-baby and well-child visits
• Immunizations
• Prescription drugs
• Dental services
• DME
• Prosthetic devices (with a $20,000 limit per 12-month period)
• Case coordination and enhanced services for children with special health-care needs and children with disabilities
• Physical, speech, and occupational therapy
• Home health services
• Transplants
• Mental health services
• Vision services
• Chiropractic services

Individuals may apply for CHIP by downloading and completing the application found on the CHIP Apply Now page of the HHSC website or by calling the toll-free CHIP number at 1-800-647-6558.

1.7.14 Home Health Providers

To enroll in Texas Medicaid as a provider of home health services, Home Health Services and Home and Community Support Services Agency (HCSSA) providers must complete the Texas Medicaid Provider Enrollment Application. Medicare certification is required for providers that are licensed as a Licensed and Certified Home Health Agency. Providers that are licensed as a Licensed Home Health Agency are not required to enroll in Medicare as a prerequisite to enrollment with Texas Medicaid.

Licensed and Certified Home Health agencies that are enrolled as Medicaid providers can provide personal care services (PCS) using their existing provider identifier. PCS for clients who are 20 years of age and younger will be provided by the Texas Health and Human Services Commission (HHSC) under the PCS benefit.

Refer to: Subsection 2.11, “Personal Care Services (PCS) (CCP)” in the Children’s Services Handbook (Vol. 2, Provider Handbooks).

To provide CCP services, HCSSA providers must follow the enrollment procedures in subsection 5.2, “Enrollment” in the Children’s Services Handbook (Vol. 2, Provider Handbooks).

Providers may download the Texas Medicaid Provider Enrollment Application at www.tmhp.com or request a paper application form by contacting TMHP directly at 1-800-925-9126.

Providers may also obtain the application by writing to the following address:

Texas Medicaid & Healthcare Partnership
Provider Enrollment
PO Box 200795
Austin, TX 78720-0795
1-800-925-9126
Fax: 1-512-514-4214
Providers may request prior authorization for home health services by contacting:

Texas Medicaid & Healthcare Partnership
Home Health Services
PO Box 202977
Austin, TX 78720-2977
1-800-925-8957
Fax: 1-512-514-4209

1.7.14.1 **Home Health Skilled Nursing and Home Health Aide (HHA) Services Provider Responsibilities**

Providers must be a licensed and certified home health agency, enrolled in Texas Medicaid, and must comply with all applicable federal, state, and local laws and regulations and Texas Medicaid policies and procedures. All providers must maintain written policies and procedures:

- That meet the standards of the Texas Family Code, Chapter 32 for obtaining consent for the medical treatment of clients in the absence of the primary caregiver.
- For obtaining physician signatures for all telephone orders within 14 calendar days of receipt of the order.

Providers must only accept clients on the basis of a reasonable expectation that the client’s needs can be adequately met in the place of service (POS). The essential elements of safe and effective home health SN or HHA services include a trained parent, guardian, or caregiver, a primary physician, competent providers, and an environment that supports the client’s health and safety needs.

Necessary primary and back-up utility, communication, and fire safety systems must be available.

*Note:* A parent or guardian, primary caregiver, or alternate caregiver may not provide SN or HHA services to their family member even if he or she is an enrolled provider or employed by an enrolled provider.

1.7.15 **Private Duty Nursing (PDN) Providers**

Home health agencies may enroll to provide PDN under the Comprehensive Care Program (CCP). RNs and licensed vocational nurses (LVNs) may enroll independently to provide PDN under CCP.

Home health agencies must do all of the following:

- Comply with provider participation requirements for home health agencies that participate in Texas Medicaid
- Comply with mandatory reporting of suspected abuse and neglect of children or adults
- Maintain written policies and procedures for obtaining consent for medical treatment for clients in the absence of the parent or guardian
- Comply with all requirements in this manual

Independently-enrolled RNs and LVNs must be enrolled as providers in CCP and comply with all of the following:

- The terms of the Texas Medicaid Provider Agreement
- All state and federal regulations and rules relating to Texas Medicaid
- The requirements of this manual, all handbooks, standards, and guidelines published by HHSC

Independently enrolled RNs and LVNs must also:

- Provide at least 30 days’ written notice to clients of their intent voluntarily to terminate services except in situations of potential threat to the nurse’s personal safety.
- Comply with mandatory reporting of suspected abuse and neglect of children.
• Maintain written policies and procedures for obtaining consent for medical treatment for clients in the absence of the parent or guardian.

Independently enrolled RNs must:

• Hold a current license from the Texas Board of Nursing (BON) or another compact state to practice as an RN.
• Agree to provide services in compliance with all applicable federal, state, and local laws and regulations, including the Texas Nursing Practice Act.
• Comply with accepted professional standards and principles of nursing practice.

Independently enrolled LVNs must:

• Hold a current license from the Texas BON to practice as an LVN.
• Agree to provide services in compliance with all applicable federal, state, and local laws and regulations, including the Texas Nursing Practice Act.
• Comply with accepted standards and principles of vocational nursing practice.
• Be supervised by an RN once per month. The supervision must occur when the LVN is present and be documented in the client’s medical record.

1.7.16 Certified Respiratory Care Practitioner (CRCP) Services

To enroll in Texas Medicaid, a CRCP must be certified by the Texas Medical Board to practice under the Texas Occupations Code, Chapter 604. For CRCPs, Medicare certification is not a prerequisite for Medicaid enrollment. A provider cannot be enrolled if his license is due to expire within 30 days; a current license must be submitted. CRCPs must enroll as individual providers and comply with all applicable federal, state, and local laws and regulations.

1.7.17 Physical, Occupational, and Speech Therapy Providers

Physical therapists, occupational therapists, and speech-language pathologists must be enrolled in Texas Medicaid according to their specific licensure in order to be reimbursed for services rendered to Texas Medicaid clients.

Occupational therapists, physical therapist, physical therapy assistants, or occupational therapy assistants must be registered and licensed by the Executive Council of Physical Therapy and Occupational Therapy Examiners.

**Note:** Auxiliary (aide, orderly, student, or technician), a licensed therapy assistant, and a licensed speech-language pathology intern (Clinical Fellow) are not eligible to enroll as therapy providers in Texas Medicaid.

**Refer to:** Subsection 1.1, “Provider Enrollment” in this section for information about enrollment procedures.


1.7.17.1 CCP Enrollment for Children’s Services—20 Years of Age and Younger

Children’s therapy services are provided under the Comprehensive Care Program (CCP). Physical, occupational, and speech therapy providers must meet Medicaid and Health and Human Services Commission (HHSC) participation standards to enroll in CCP. All CCP providers must be enrolled in Texas Medicaid to be reimbursed for services.
The following facilities or organizations may also enroll in Texas Medicaid to provide CCP therapy services:

- Comprehensive Outpatient Rehabilitation Facilities (CORFs) and Outpatient Rehabilitation Facilities (ORFs)
- Home Health Services and Home and Community Support Services Agency (HCSSA) providers
- Early Childhood Intervention (ECI) providers
- School Health and Related Services (SHARS) providers

Refer to: Subsection 2.1.2, "Enrollment" in the Children’s Services Handbook (Vol. 2, Provider Handbooks) for more information about ECI enrollment.

Subsection 3.2, “Enrollment” in the Children’s Services Handbook (Vol. 2, Provider Handbooks) for more information about SHARS enrollment.

1.7.18 Children’s Services Comprehensive Care Program (CCP)

CCP providers must meet Medicaid and Health and Human Services Commission (HHSC) participation standards to enroll in the program. All CCP providers must be enrolled in Texas Medicaid to be reimbursed for services. Provider enrollment inquiries and application requests must be sent to the TMHP Provider Enrollment department at:

Provider Enrollment
Texas Medicaid & Healthcare Partnership
PO Box 200555
Austin, TX 78720-0555

Home and community support services agencies (HCSSAs) that want to provide CCP private-duty nursing (PDN), home telemonitoring, occupational therapy, physical therapy, or speech therapy services under the licensed-only home health (LHH) category must first enroll with TMHP. To enroll with TMHP in the LHH category, an HCSSA must:

- Complete a provider enrollment form, which can be found on the TMHP website at www.tmhp.com, provide its license information, and check the “Only CCP services” box on the form.
- Obtain a Texas Provider Identifier (TPI) for CCP services.

1.8 Electronic Health Record Incentive/Promoting Interoperability Program

The Texas Medicaid Electronic Health Record (EHR) Incentive/ Promoting Interoperability (PI) Program was authorized under the federal American Recovery and Reinvestment Act of 2009 and was implemented in Texas in 2011. The program provides incentive payments to eligible Medicaid providers and hospitals if they adopt and meaningfully use certified electronic health record technology. The program encourages Texas Medicaid providers and hospitals to make the transition to electronic health records and help build a statewide health information network where patient records are shared among offices electronically.

Individual providers and hospitals can qualify for incentive payments by adopting certified electronic health record technology that meets federal standards and using that technology to improve quality, safety, and effectiveness of patient care.

Enrollment of new providers into the program ended in 2016; however, individual providers that received at least one previous incentive payment can continue receiving up to six annual payments, if they meet the federally defined criteria for meaningful use and promoting interoperability. The program has a statutory end date of December 31, 2021, and no incentive payments will be disbursed after that date.
To learn more about the program, providers can visit https://healthit.hhsc.texas.gov and www.cms.gov/Regulations-and-Guidance/Legislation/EHRIncentivePrograms/index.html for the latest program news and resource documents.

For more information about the Texas Medicaid EHR Incentive/PI Program, providers can:

- Email support@tmhp-mi.com or HealthIT@tmhp.com.
- Call the Medicaid EHR Incentive/PI Program at 1-855-831-6112.
- Call the TMHP Contact Center at 1-800-925-9126.

Providers can find a list of program forms on the Forms page of the Health Information Technology website.

1.8.1 **Attesting to Meaningful Use-Required Documentation for Texas Medicaid Electronic Health Record Incentive/Promoting Interoperability Program**

Providers must maintain auditable records to support attestations that are made by eligible professionals (EPs) in the Texas Medicaid EHR Incentive/PI Program from Adopt/Implement/Upgrade (AIU) to Meaningful Use (MU). To receive incentives, EPs must show that they are meaningfully using certified EHR technology by reporting on federally defined MU, PI, and clinical quality measures. Providers must attest to these measures in the Medicaid EHR Incentive/PI Program portal, which is also known as Medicaid Incentives (MI) 360, that is available through My Account on www.tmhp.com.

During the submission process, providers are guided through the portal step-by-step, including prompts to upload required documents to support the submission or attestation. Documentation requirements may vary depending on the provider’s status and attestation. The broad categories of required documents include:

- **Numerator and Denominator Measures:** For the numerator and denominator measures, providers must upload an EHR-generated summary MU and PI report that shows the numerator and denominator and calculated value for each measure reported. If some measures are not included in the summary report, providers must generate separate reports or other auditable documentation. For example, screenshots showing an MU and PI dashboard with reported measures and values might be acceptable. Providers must ensure that the report date matches the MU and PI reporting period and the report data values are for the attesting provider only.

- **Yes and No Measures:** For the yes and no measures, providers must demonstrate compliance with the following measures:
  - Drug and drug and allergy interaction checks are enabled—Providers must include documentation, such as a screenshot or configuration page, that shows this functionality has been turned on during the entire MU and PI reporting period.
  - Clinical Decision Support (CDS) rules implemented—providers must include documentation, such as a screenshot, that shows CDS rules have been configured.
  - Security risk analysis—Provide must include a copy or documentation of a security risk analysis completed or reviewed each year.
  - Public health registry reporting—For immunization, syndromic surveillance, and other registry reporting, providers must include documentation, such as an email or letter, from the public health agency (PHA) or registry that acknowledges the provider’s active engagement with the PHA or registry. For immunization reporting, the provider must include the ImmTrac2 ID number.
Clinical Quality Measures (CQMs): EPs must report on CQMs and may upload documentation for reported CQMs; however, it is not required during attestation.

Note: Personal health information (PHI) must be removed from all documentation before submission.

1.8.2 Electronic Health Record Incentive/Promoting Interoperability Program Audits and Appeals

HHSC contracts with an independent audit firm to conduct EHR Incentive/PI audits and follows policies and procedures defined in 1 TAC §354.1450 for audits and appeals related to overpayments of Texas Medicaid EHR Incentive/PI payments. Detailed information on the audit and appeal process can be found on the program website at https://healthit.hhsc.texas.gov/policies-and-procedures/audits-and-appeals.

1.8.3 How to Return an Electronic Health Record Incentive/Promoting Interoperability Payment

To ensure that a returned payment is credited properly, providers that have received an EHR incentive/PI payment that must be returned due to an audit or determination of ineligibility must complete the following steps:

1) Download and print the Texas Medicaid Refund Information Remittance Form.
2) Fill out the provider name, Texas Provider Identifier (TPI), and Taxpayer Identification Number (TIN) that correspond to the payee (recipient) of the incentive payment, even if the recipient is different than the attesting provider.
3) Fill out the additional information requested on the form.
4) Write a comment at the bottom of the form designating the check as an “EHR Incentive Program payment refund.”
5) Mail the completed form and the check to the following address:
   Texas Medicaid & Healthcare Partnership
   Accounts Receivable
   PO Box 202948
   Austin, TX 78720-2948

1.9 Computer-Based Training Courses for Providers

TMHP has created a web page to simplify the process of accessing the computer-based training courses that are available on the TMHP Learning Management System (LMS).

The new Computer-Based Training (CBT) web page lists all of the CBT courses that TMHP makes available free of charge to all providers. The web page arranges the CBT titles by the programs they support. Each CBT title has a link to a description of the curriculum and a link to the CBT itself, which resides on the TMHP LMS. Clicking one of the View CBT Now links takes the provider to the LMS where, after the provider logs in, the CBT starts immediately.

The procedure that providers follow to access a CBT has changed to the following:

1) Click Provider Education in the left navigation panel of any provider web page. The Provider Education home page appears.
2) Click Computer-Based Training in the left navigation panel. The CBT web page appears.
3) Click View CBT Now in the same row as the name of the CBT that you want to view. The LMS appears in a new window.
4) Log in to an existing account or create a new account. New visitors to the LMS can take courses immediately after they register. The selected CBT begins as soon as the login is complete.
Additional Improvement to Training Experience

TMHP is taking additional steps to make it easier for providers to get answers to their questions more quickly, including the following:

- TMHP has started to create single-topic "training snapshots" to provide quick training to answer specific needs. Currently, there is only one training snapshot, but more are planned.
- As CBT courses are revised, TMHP is converting many of them to a series of single-topic modules, which are selected from the Main Menu slide for the CBT course. The modular structure makes it easier for providers to get the specific training that they need.

**Note:** Even if a CBT course has not yet been converted to modules, providers can jump to a specific section after the course begins by clicking on the Outline tab and selecting a slide from the list that appears.

### 1.10 Enrollment Criteria for Out-of-State Providers

Texas Medicaid covers medical assistance services provided to eligible Texas Medicaid clients while in a state other than Texas, as long as the client does not leave Texas to receive out-of-state healthcare that can be received in Texas. Services provided outside the state are covered to the same extent medical assistance is furnished and covered in Texas.

**Note:** Border state providers (providers rendering services within 50 miles driving distance of the Texas border) are considered in-state providers.

The administrative rules governing the enrollment of out-of-state providers are found in 1 TAC §352.17. The rule provides that a Medicaid applicant or re-enrolling provider is considered out-of-state if any of the following criteria are met:

- The physical address where services are or will be rendered is located outside the Texas state border and within the United States.
- The physical address where the services or products originate or will originate is located outside the Texas state border and within the United States when providing services, products, equipment, or supplies to a Medicaid recipient in the state of Texas.
- The physical address where services are or will be rendered is located within the Texas state border, but:
  - The applicant or re-enrolling provider maintains all patient records, billing records, or both, outside the Texas state border and
  - The applicant or re-enrolling provider is unable to produce the originals or exact copies of the patient records or billing records, or both, from the location within the Texas state border where services are rendered.

An applicant or re-enrolling provider that is considered out-of-state is ineligible to participate in Medicaid unless HHSC or its designee approves the enrollment on the basis that the applicant has provided, is providing, or will provide services under one or more of the following criteria:

- The services are medically necessary emergency services to a recipient who is located outside of the state.

**Note:** An out-of-state provider seeking enrollment under this criterion must include with the enrollment application a copy of the claim that contains the diagnosis that indicates emergency care or medical record documentation. The documentation must demonstrate that emergency care was provided to a Texas Medicaid client. Providers enrolled under this criterion will be given a time limited enrollment not to exceed one year.
• The services are medically necessary to a recipient who is located outside of the state, and in the expert opinion of the recipient’s attending or other provider, the recipient’s health would be or would have been endangered if the recipient were required to travel to Texas.

  **Note:** An out-of-state provider seeking enrollment under this criterion must include supporting clinical records, signed by the attending provider, explaining why the client’s health would be or would have been endangered if the client had been required to travel to Texas. Providers enrolled under this criterion will be given a time limited enrollment not to exceed one year.

• The services are medically necessary and more readily available to a recipient in the state where the recipient is located.

  **Note:** An out-of-state provider that seeks enrollment under this criterion must include supporting clinical records, signed by the attending provider, explaining why the services are more readily available in the state where the client is located. Providers that are enrolled under this criterion may be enrolled for a limited period of time.

• The services are medically necessary to a recipient who is eligible on the basis of participation in an adoption assistance or foster care program administered by the Texas Department of Family and Protective Services under Title IV-E of the Social Security Act.

  **Note:** An out-of-state provider that seeks enrollment under this criterion must include documentation showing that the client is an adopted child or is in a foster care program and/or is receiving adoption subsidies through the programs listed in this criterion. Providers that are enrolled under this criterion may be enrolled for a limited period of time.

• The services are medically necessary and have been prior authorized by HHSC or its designee, and documented medical justification indicating the reasons the recipient must obtain medical care outside Texas is furnished to HHSC or its designee before providing the services and before payment.

  **Note:** An out-of-state provider that seeks enrollment under this criterion must include documentation showing that the service has been prior authorized by HHSC or its designee (TMHP, or MCO), or supporting clinical documentation (signed by the attending provider) indicating the reasons why the recipient must obtain medical care outside of Texas. Providers that are enrolled under this criterion may be enrolled for a limited period of time.

• The services are medically necessary and it is the customary or general practice of Medicaid recipients in a particular locality within Texas to obtain services from the out-of-state provider, if the provider is located in the United States and within 50 miles driving distance from the Texas state border, or as otherwise demonstrated on a case-by-case basis.

  **Note:** An out-of-state provider does not meet the criterion in this paragraph merely on the basis of having established business relationships with one or more providers that participate in Medicaid. Attach signed letter from the provider stating why it is customary or general practice of clients in a particular locality within Texas to obtain services from the out-of-state provider. Providers that are enrolled under this criterion may be enrolled for a limited period of time.

• The services are medically necessary and the nature of the service is such that providers for this service are limited or not readily available within the state of Texas.

  **Note:** An out-of-state provider that seeks enrollment under the criterion must include documentation showing that the services provided by the applicant are medically necessary and are limited or not readily available within the state of Texas.
• The services are medically necessary services to one or more dually eligible recipients (i.e., recipients who are enrolled in both Medicare and Medicaid) and the out-of-state provider may be considered for reimbursement of co-payments, deductibles, and co-insurance, in which case the enrollment will be restricted to receiving reimbursement only for the Medicaid-covered portion of Medicare crossover claims.

  Note: An out-of-state provider that seeks enrollment under this criterion must include documentation for why this criterion applies, Medicare EOB or MRAN, with documented medical justification as well as any additional information requested by HHSC or its designee. Providers that are enrolled under this criterion may be enrolled for a limited period of time.

• The services are provided by a pharmacy that is a distributor of a drug that is classified by the U.S. Food and Drug Administration (FDA) as a limited distribution drug.

  Note: An out-of-state provider that seeks enrollment under this criterion must include with the enrollment application documentation for why this criterion applies with documented medical justification as well as any additional information requested by HHSC or its designee. Attach signed letter from the provider stating that the enrolling pharmacy is a distributor of a drug that is classified by FDA as a limited distribution drug, include a letter from the FDA stating that the aforementioned drug is considered a limited distribution drug. Providers that are enrolled under this criterion may be enrolled for a limited period of time.

• The services are medically necessary and one or more of the following exceptions for good cause exist and can be documented:
  • Texas Medicaid enrolled providers rely on the services provided by the applicant.
  • Applicant maintains existing agreements as a participating provider through one or more Medicaid managed care organizations (MCO) and enrollment of the applicant leads to more cost-effective delivery of Medicaid services.

• A laboratory may participate as an in-state provider under any program administered by a health and human services agency, including HHSC, that involves laboratory services, regardless of the location where any specific service is performed or where the laboratory’s facilities are located if:
  • The laboratory or an entity that is a parent, subsidiary, or other affiliate of the laboratory maintains laboratory operations in Texas;
  • The laboratory and each entity that is a parent, subsidiary, or other affiliate of the laboratory, individually or collectively, employ at least 1,000 persons at places of employment located in this state; and
  • The laboratory is otherwise qualified to provide the services under the program and is not prohibited from participating as a provider under any benefits programs administered by a health and human services agency, including HHSC, based on conduct that constitutes fraud, waste, or abuse.

Out-of-state providers that seek enrollment under one or more of the above criteria must submit an enrollment application and be approved for enrollment.

TMHP must receive claims from out-of-state providers within 365 days from the date of service.


1.11 Medicaid Fraud, Waste, and Abuse Policy

The OIG has the responsibility to identify and investigate cases of suspected fraud, waste, and abuse in Medicaid and other health and human services programs. This responsibility, granted through state and federal law, gives the OIG the authority to pursue administrative sanctions and to refer cases to prose-
cutors, licensure and certification boards, and other agencies. Additionally, Texas Medicaid is required to disenroll or exclude any provider who has been disenrolled or excluded from Medicare or any other state health-care program.

Anyone participating in Texas Medicaid must understand the requirements for participation. Available methods both to learn and stay up to date on program requirements include the following:

- **Provider education.** Attendance at educational workshops and training sessions. Regular training opportunities are offered by TMHP.
- **Texas Medicaid publications.** These include the Texas Medicaid Provider Procedures Manual and banner messages, which are included in R&S Reports.
- **All adopted agency rules.** These include those related to fraud, waste, and abuse contained in 1 TAC Chapter 371.
- **State and federal law.** Statutes and other law pertinent to Texas Medicaid and fraud, waste, and abuse within Texas Medicaid.

In addition, providers are responsible for the delivery of health-care items and services to Medicaid clients in accordance with all applicable licensure and certification requirements and accepted health care professionals’ community standards. Such standards include those related to medical record and claims filing practices, documentation requirements, and records maintenance. The TAC requires providers to follow these standards. For more information, consult 1 TAC §371.1659.

Texas Medicaid providers must follow the coding and billing requirements of the Texas Medicaid Provider Procedures Manual (TMPPM). However, if coding and billing requirements for a particular service are not addressed in the TMPPM, and if coding and billing requirements are not otherwise specified in program policy (such as in provider bulletins or banners), then providers must follow the most current coding guidelines. These include:

- CPT as set forth in the American Medical Association’s most recently published “CPT books”, “CPT Assistant” monthly newsletters, and other publications resulting from the collaborative efforts of American Medical Association with the medical societies.
- Healthcare Common Procedure Coding System (HCPCS) as developed and maintained by the federal government.
- National Correct Coding Initiative (NCCI), as set forth by the CMS and as explained in the NCCI Policy and Medicare Claims Processing Manuals. NCCI consists of procedure code combinations that a provider must not bill together. One of the codes in the pair is considered a part of the primary procedure and not reimbursable to the same provider on the same date of service.

**Exception:** NCCI outlines use of modifiers some of which are not currently recognized by Texas Medicaid. See the list of modifiers utilized by Texas Medicaid in subsection 6.3.5, “Modifiers” in “Section 6: Claims Filing” (Vol. 1, General Information).

- Current Dental Terminology (CDT) as published by the American Dental Association (ADA).
- Other publications resulting from the collaborative efforts of the ADA with dental societies.
- International Classification of Diseases, 10th Revision, Clinical Modification (ICD-10-CM).
- Diagnostic and Statistical Manual of Mental Disorders (DSM).

Failure to comply with the guidelines provided in these publications may result in a provider being found to have engaged in one or more program violations listed in 1 TAC §371.1659.

All providers are held responsible for any claims preparation or other activities that may be performed under the provider’s authority. For example, providers are held responsible for any omissions and the accuracy of submitted information, even if those actions are performed by office staff, contractors, or...
billing services. This, however, does not absolve these other individuals for their participation in any documents provided to the state or designee with false, inaccurate, or misleading information; or pertinent omissions.

HHSC-OIG may impose one or any combination of administrative actions or administrative sanctions on Texas Medicaid providers or other persons when fraud, waste, or abuse is determined. Those who may be sanctioned include:

- Those furnishing services or items directly or indirectly.
- Those billing for services.
- Those violating any of the provisions delineated in this section.
- Affiliates of a provider or person violating any of the provisions delineated in this section.

Administrative sanctions include, without limitation:

- Exclusion from program participation for a specified period of time, permanently, or indefinitely. Anyone excluded from Texas Medicaid is also automatically excluded from all programs under Titles V and XX of the Social Security Act.
- Suspension of Medicaid payments (payment hold) to a provider.
- Recoupment of Medicaid overpayments, including any overpayments determined through statistical sampling and extrapolation.
- Restricted Medicaid reimbursement (specific services will not be reimbursed to an individual provider during the time the provider is on restricted reimbursement; however, reimbursement for other services may continue).
- Cancellation of the Medicaid provider agreement (however, a deactivation in accordance with the agreement itself is not considered a sanction).
- Exclusion or suspension under the authority of the CFR.

Administrative actions include:

- Amending a provider agreement so that it will deactivate on a specific date.
- Granting an agreement or transferring a provider to an agreement with special terms or conditions, including a probationary agreement.
- Required attendance at provider education sessions.
- Prior authorization of selected services.
- Pre-payment review.
- Post-payment review.
- Required attendance at informal or formal provider corrective action meetings.
- Submission of additional documentation or justification that is not normally required to accompany submitted claims. (Failure to submit legible documentation or justification requested will result in denial of the claim.)
- Oral, written, or personal educational contact with the provider.
- Posting of a surety bond or providing a letter of credit.
- Having a subpoena served to compel an appearance for testimony or the production of relevant evidence, as determined by the HHSC-OIG.
Anyone facing an administrative sanction has a right to formal due process. This formal due process may include a hearing before an administrative law judge. Conversely, anyone facing an administrative action is not entitled to formal due process. People who induce, solicit, receive, offer, or pay any remuneration (including, but not limited to, bribes, kickbacks, or rebates) directly or indirectly in relation to referrals, purchases, leases, or arrangements of services covered by Medicare or Texas Medicaid may be in violation of state statutes and guilty of a federal felony offense. State law also allows for the suspension of providers convicted of a criminal offense related to Medicare or Texas Medicaid. The commission of a felony in Medicaid or Medicare programs may include fines or imprisonment ranging from five years to life in prison. Examples of inducements include a service, cash in any amount, entertainment, or any item of value.

As stated in 1 TAC §§371.1651-371.1669, following is a nonexclusive list of grounds or criteria for the Inspector General’s administrative enforcement and/or referral for criminal, civil, or licensure or certification investigation and judicial action regarding program violations by any provider or person. Violations result from a provider or person who knew or should have known the following were violations. The headings of each of the following groups are provided solely for organization and convenience and are not elements of any program violation.

1) Claims and Billing.
   a) Submitting or causing to be submitted a false statement or misrepresentation, or omitting pertinent facts when claiming payment under the Texas Medicaid or other HHS program or when supplying information used to determine the right to payment under the Texas Medicaid or other HHS program;
   b) Submitting or causing to be submitted a false statement, information or misrepresentation, or omitting pertinent facts to obtain greater compensation than the provider is legally entitled to;
   c) Submitting or causing to be submitted a false statement, information or misrepresentation, or omitting pertinent facts to meet prior authorization requirements;
   d) Submitting or causing to be submitted under Title XVIII (Medicare) or a state health-care program claims or requests for payment containing unjustified charges or costs for items or services that substantially exceed the person’s usual and customary charges or costs for those items or services to the public or the private pay patients unless otherwise authorized by law;
   e) Submitting or causing to be submitted claims with a pattern of inappropriate coding or billing that results in excessive costs to the Texas Medicaid or other HHS program;
   f) Billing or causing claims to be filed for services or merchandise that were not provided to the recipient;
   g) Submitting or causing to be submitted a false statement or misrepresentation that, if used, has the potential of increasing any individual or state provider payment rate or fee;
   h) Submitting or causing to be submitted to the Texas Medicaid or other HHS program a cost report containing costs not associated with Texas Medicaid or other HHS program or not permitted by Texas Medicaid or other HHS program policies;
   i) Presenting or causing to be presented to an operating agency or its agent a claim that contains a statement or representation that the person knows or should have known to be false;
   j) Billing or causing claims to be submitted to the Texas Medicaid or other HHS program for services or items furnished personally by, at the medical direction of, or on the prescription or order of a person who is excluded from Texas Medicaid, other HHS program, or Medicare or has been excluded from and not reinstated within Texas Medicaid, other HHS program, or Medicare;
   k) Billing or causing claims to be submitted to the Texas Medicaid or other HHS program for services or items that are not reimbursable by the Texas Medicaid or other HHS program;
l) Billing or causing claims to be submitted to the Texas Medicaid or other HHS program for a service or item which requires a prior order or prescription by a licensed health-care practitioner when such order or prescription has not been obtained;

m) Billing or causing claims to be submitted to the Texas Medicaid or other HHS program for an item or service substituted without authorization for the item or service ordered, prescribed or otherwise designated by the Texas Medicaid or other HHS program;

n) Billing or causing claims to be submitted to the Texas Medicaid or other HHS program by a provider or person who is owned or controlled, directly or indirectly, by an excluded person; and

o) Billing or causing claims to be submitted to the Texas Medicaid or other HHS program by a provider or person for charges in which the provider discounted the same services for any other type of patient.

2) Records and Documentation.

a) Failing to maintain for the period of time required by the rules relevant to the provider in question records and other documentation that the provider is required by federal or state law or regulation or by contract to maintain in order to participate in the Texas Medicaid or other HHS program or to provide records or documents upon request for any records or documents determined necessary by the Inspector General to complete their statutory functions related to a fraud and abuse investigation. Such records and documentation include, without limitation, those necessary:

i) To verify specific deliveries, medical necessity, medical appropriateness, and adequate written documentation of items or services furnished under Title XIX or Title XX;

ii) To determine in accordance with established rates appropriate payment for those items or services delivered;

iii) To confirm the eligibility of the provider to participate in the Texas Medicaid or other HHS program; e.g., medical records (including, without limitation, X-rays, laboratory and test results, and other documents related to diagnosis), billing and claims records, cost reports, managed care encounter data, financial data necessary to demonstrate solvency of risk-bearing providers, and documentation (including, without limitation, ownership disclosure statements, articles of incorporation, by-laws, and corporate minutes) necessary to demonstrate ownership of corporate entities; and

iv) To verify the purchase and actual cost of products;

b) Failing to disclose fully and accurately or completely information required by the Social Security Act and by 42 CFR Part 455, Subpart B; 42 CFR Part 420, Subpart C; 42 CFR §1001.1101; and 42 CFR Part 431;

c) Failing to provide immediate access, upon request by a requesting agency, to the premises or to any records, documents, and other items or equipment the provider is required by federal or state law or regulation or by contract to maintain in order to participate in the Texas Medicaid or other HHS program (see subparagraphs (a) and (b) of this paragraph), or failing to provide records, documents, and other items or equipment upon request that are determined necessary by the Inspector General to complete their statutory functions related to a fraud and abuse investigation, including without limitation all requirements specified in 1 TAC §§371.1701, 371.1703, 371.1705, 371.1707, 371.1709, 371.1711, 371.1713, and 371.1715 of this subchapter. “Immediate access” is deemed to be within 24 hours of receiving a request, unless the requesting agency has reason to suspect fraud or abuse or to believe that requested records, documents, or other items or equipment are about to be altered or destroyed, thereby necessitating access at the actual time the request is presented or, in the opinion of the Inspector General, the request may be completed at the time of the request and/or in less than 24 hours;
d) Developing false source documents or failing to sign source documents or to retain supporting
documentation or to comply with the provisions or requirements of the operating agency or its
agents pertaining to electronic claims submittal; and
e) Failing as a provider, whether individual, group, facility, managed care or other entity, to
include within any subcontracts for services or items to be delivered within Texas Medicaid all
information that is required by 42 CFR §434.10(b).

3) Program-Related Convictions.
   a) Pleading guilty or nolo contendere, agreeing to an order of probation without adjudication of
guilt under deferred adjudication, or being a defendant in a court judgment or finding of guilt
for a violation relating to performance of a provider agreement or program violation of
Medicare, Texas Medicaid, other HHS program, or any other state’s Medicaid program;
b) Pleading guilty or being convicted of a violation of state or federal statutes relating to
dangerous drugs, controlled substances, or any other drug-related offense;
c) Pleading guilty of, being convicted of, or engaging in conduct involving moral turpitude;
d) Pleading guilty or being convicted of a violation of state or federal statutes relating to fraud,
thief, embezzlement, breach of fiduciary responsibility, or other financial misconduct relating
to the delivery of a health-care item or service or relating to any act or omission in a program
operated or financed by any federal, state, or local government agency;
e) Being convicted in connection with the interference with or obstruction of any investigation
into any criminal offense that would support mandatory exclusion under 1 TAC §371.1705 of
this subchapter or any offense listed within paragraph (3) of this subsection regarding
program-related convictions; and
f) Being convicted of any offense that would support mandatory exclusion under 1 TAC
§371.1705 of this subchapter.

4) Provider Eligibility.
   a) Failing to meet standards required for licensure, when such licensure is required by state or
federal law, administrative rule, provider agreement, or provider manual for participation in
the Texas Medicaid or other HHS program;
b) Being excluded, suspended or otherwise sanctioned within any federal program involving the
provision of health care;
c) Being excluded, suspended or otherwise sanctioned under any state health-care program for
reasons bearing on the person’s professional competence, professional performance or financial
integrity;
d) Failing to fully and/or correctly complete a Provider Enrollment Agreement, Provider Re-
enrollment Agreement or other enrollment form prescribed by the relevant operating agency
or its agent for enrollment; and
e) Loss or forfeiture of corporate charter.

5) Program Compliance.
   a) Failing to comply with the terms of the Texas Medicaid or other HHS program contract or
provider agreement, assignment agreement, the provider certification on the Texas Medicaid or
other HHS program claim form, or rules or regulations published by the Commission or a
Medicaid or other HHS operating agency;
b) Violating any provision of the Human Resources Code, Chapter 32 or 36, or any rule or
regulation issued under the Code;
c) Submitting a false statement or misrepresentation or omitting pertinent facts on any application or any documents requested as a prerequisite for the Texas Medicaid or other HHS program participation;

d) Refusing to execute or comply with a provider agreement or amendments when requested;

e) Failing to correct deficiencies in provider operations after receiving written notice of them from an operating agency, the commission or their authorized agents;

f) Failing to abide by applicable federal and state law regarding handicapped individuals or civil rights;

g) Failing to comply with the Texas Medicaid or other HHS program policies, published Texas Medicaid or other HHS program bulletins, policy notification letters, provider policy or procedure manuals, contracts, statutes, rules, regulations, or interpretation previously sent to the provider by an operating agency or the commission regarding any of the authorities listed above, including statutes or standards governing occupations;

h) Failing to fully and accurately make any disclosure required by the Social Security Act, §1124 or §1126;

i) Failing to disclose information about the ownership of a subcontractor with whom the person has had business transactions in an amount exceeding $25,000 during the previous 12 months or about any significant business transactions (as defined by HHS) with any wholly-owned supplier or subcontractor during the previous five years;

j) Failing, as a hospital, to comply substantially with a corrective action required under the Social Security Act, §1886(f)(2)(B);

k) Failing to repay or make arrangements that are satisfactory to the commission to repay identified overpayments or other erroneous payments or assessments identified by the commission or any Texas Medicaid or other HHS program operating agency;

l) Committing an act described as grounds for exclusion in the Social Security Act, §1128A (civil monetary penalties for false claims) or §1128B (criminal liability for health care violations);

m) Defaulting on repayments of scholarship obligations or items relating to health profession education made or secured, in whole or in part, by HHS or the state when they have taken all reasonable steps available to them to secure repayment;

n) Soliciting or causing to be solicited, through offers of transportation or otherwise, Texas Medicaid or other HHS program recipients for the purpose of delivering to those recipients health-care items or services;

o) Marketing, supplying or selling confidential information (e.g., recipient names and other recipient information) for a use that is not expressly authorized by the Texas Medicaid or other HHS program; and

p) Failing to abide by applicable statutes and standards governing providers.

**Important:** Providers must comply with their applicable licensing agency's laws and regulations, including any related to marketing and advertising, and any applicable state and federal laws and regulations, contractual requirements, and other guidance documents. Providers are encouraged to review the "Provider Marketing Guidelines," which are available on the HHs website at [https://hhs.texas.gov/services/health/medicaid-chip/provider-information/texas-medicaid-chip-communications-resources](https://hhs.texas.gov/services/health/medicaid-chip/provider-information/texas-medicaid-chip-communications-resources).
6) Delivery of Health-Care Services.
   a) Failing to provide health-care services or items to Texas Medicaid or other HHS program recipients in accordance with accepted medical community standards or standards required by statute, regulation, or contract, including statutes and standards that govern occupations;
   b) Furnishing or ordering health-care services or items for a recipient-patient under Title XVIII or a state health-care program that substantially exceed the recipient’s needs, are not medically necessary, are not provided economically or are of a quality that fails to meet professionally recognized standards of health care; and
   c) Engaging in any negligent practice that results in death, injury, or substantial probability of death or injury to the provider’s patients.

7) Improper Collection and Misuse of Funds.
   a) Charging recipients for services when payment for the services was recouped by the Texas Medicaid or another HHS program for any reason;
   b) Misapplying, misusing, embezzling, failing to promptly release upon a valid request, or failing to keep detailed receipts of expenditures relating to any funds or other property in trust for a Texas Medicaid or other HHS program recipient;
   c) Failing to notify and reimburse the relevant operating agency or the commission or their agents for services paid by the Texas Medicaid or other HHS programs if the provider also receives reimbursement from a liable third party;
   d) Rebating or accepting a fee or a part of a fee or charge for a Texas Medicaid or other HHS program patient referral;
   e) Requesting from a recipient in payment for services or items delivered within the Texas Medicaid or other HHS program any amount that exceeds the amount the Texas Medicaid or other HHS program paid for such services or items, with the exception of any cost-sharing authorized by the program; and
   f) Requesting from a third party liable for payment of the services or items provided to a recipient under the Texas Medicaid or other HHS program, any payment other than as authorized at 42 CFR §447.20.

8) Licensure Actions.
   a) Having a voluntary or involuntary action taken by a licensing or certification agency or board that requires the provider or employee to comply with professional practice requirements of the board after the board receives evidence of noncompliance with licensing or certification requirements; and
   b) Having its license to provide health care revoked, suspended, or probated by any state licensing or certification authority, or losing a license or certification, because of action based on assessment of the person’s professional competence, professional performance, or financial integrity, non-compliance with Health and Safety Code, statutes governing occupations, or surrendering a license or certification while a formal disciplinary proceeding is pending before licensing or certification authorities when the proceeding concerns the person’s professional competence, professional performance, or financial integrity.

9) MCOs and Persons Providing Services or Items Through Managed Care.

   **Note:** This paragraph includes those program violations that are unique to managed care; paragraphs (1) through (8) and (11) of this section also apply to managed care.
a) Failing, as an MCO, or an association, group or individual health-care provider furnishing services through an MCO, to provide to recipient enrollee a health-care benefit, service or item that the organization is required to provide under its contract with an operating agency;

b) Failing, as an MCO or an association, group or individual health-care provider furnishing services through an MCO, to provide to an individual a health-care benefit, service or item that the organization is required to provide by state or federal law, regulation or program rule;

c) Engaging, as an MCO, in actions that indicate a pattern of wrongful denial or payment for a health-care benefit, service or item that the organization is required to provide under its contract with an operating agency;

d) Engaging, as an MCO, in actions that indicate a pattern of wrongful delay of at least 45 days or a longer period specified in the contract with an operating agency, not to exceed 60 days, in making payment for a health-care benefit, service or item that the organization is required to provide under its contract with an operating agency;

e) Engaging, as an MCO or an association, group or individual health-care provider furnishing services through managed care, in a fraudulent activity in connection with the enrollment in the organization’s managed care plan of an individual eligible for medical assistance or in connection with marketing the organization’s services to an individual eligible for medical assistance;

f) Discriminating against enrollees or prospective enrollees on any basis, including, without limitation, age, gender, ethnic origin or health status;

g) Failing, as an MCO, to comply with any term within a contract with a Texas Medicaid or other HHS program operating agency to provide healthcare services to Texas Medicaid or HHS program recipients; and

h) Failing, as an MCO, reasonably to provide to the relevant operating agency, upon its written request, encounter data and/or other data contractually required to document the services and items delivered by or through the MCO to Texas Medicaid or other HHS program recipients.


a) Reporting noncovered or nonchargeable services as covered items; e.g., incorrectly apportioning or allocating costs on cost reports; including costs of noncovered services, supplies or equipment in allowable costs; arrangements between providers and employees, related parties, independent contractors, suppliers, and others that appear to be designed primarily to overstate the costs to the program through various devices (such as commissions or fee splitting) to siphon-off or conceal illegal profits;

b) Reporting costs not incurred or which were attributable to nonprogram activities, other enterprises or personal expenses;

c) Including unallowable cost items on a cost report;

d) Manipulating or falsifying statistics that result in overstatement of costs or avoidance of recoupment, such as incorrectly reporting square footage, hours worked, revenues received, or units of service delivered;

e) Claiming bad debts without first genuinely attempting to collect payment;

f) Depreciating assets that have been fully depreciated or sold or using an incorrect basis for depreciation; and

g) Reporting costs above the cost to the related party.
11) Kickbacks and Referrals.

   a) Violating any of the provisions specified in 1 TAC §371.1655 (30) of this subchapter relating to kickbacks, bribes, rebates, referrals, inducements, or solicitation;

   b) As a physician, referring a Texas Medicaid or other HHS program patient to an entity with which the physician has a financial relationship for the furnishing of designated health services, payment for which would be denied under Title XVIII (Medicare) pursuant to §1877 and §1903(s) of the Social Security Act (Stark I and II). Neither federal financial participation nor this state’s expenditures for medical assistance under the state Medicaid plan may be used to pay for services or items delivered within the program and within a relationship that violates Stark I or II. The Commission hereby references and incorporates within these rules the federal regulations promulgated pursuant to Stark I and II, and expressly recognizes all exceptions to the prohibitions on referrals established within those rules;

   c) Failing to disclose documentation of financial relationships necessary to establish compliance with Stark I and II, as set forth in subparagraph (b) of this paragraph; and

   d) Offering to pay or agreeing to accept, directly or indirectly, overtly or covertly any remuneration in cash or in kind to or from another for securing or soliciting a patient or patronage for or from a person licensed, certified, or registered or enrolled as a provider or otherwise by a state health-care regulatory or health and human service agency.

   Involvement in any of these practices may result in provider exclusion or suspension from Texas Medicaid. Providers are notified in writing of any actions taken as well as procedures for appeal and reinstatement. The written notification will specify the date on which Medicaid program participation may resume. The reinstated person may then apply for a contract or provider agreement.

   Providers and individuals who have been excluded from Texas Medicaid may be reinstated only by HHSC-OIG. If HHSC-OIG approves an individual’s request for reinstatement, a written notice will be sent to that individual. The provider must first be reinstated into Medicaid and receive written notification specifying the date on which Medicaid program participation may resume. Once the provider has been reinstated into Medicaid, the provider may then apply for a contract or provider agreement.

   Full investigation of criminal Medicaid fraud is the MFCU’s responsibility and may result in a felony or misdemeanor criminal conviction.

1.11.1 Reporting Fraud, Waste, and Abuse

   Anyone with knowledge about suspected Medicaid fraud, waste, and abuse of provider services must report the information to the HHSC-OIG. To report fraud, waste, and abuse, visit https://oig.hhsc.texas.gov/report-fraud-waste-or-abuse and select IG’s Fraud Reporting Form. Fraud, waste, and abuse may also be reported by calling the OIG hotline at 1-800-436-6184. All reports of fraud, waste, and abuse received through either channel remain confidential.

   HHSC-OIG encourages providers to voluntarily investigate and report fraud, waste, abuse, or inappropriate payments of Medicaid funds in their own office. Providers are required to report these activities to HHSC-OIG when identified. HHSC-OIG will work collaboratively with self-reporting providers. More information about provider self-reporting is available on the OIG website at https://oig.hhsc.texas.gov/resources/information-providers.

1.11.2 Suspected Cases of Provider Fraud, Waste, and Abuse

   HHSC-OIG is responsible for minimizing fraud, waste, and abuse by Medicaid providers. HHSC-OIG has established and continues to refine criteria for identifying cases of possible fraud, waste, and abuse and recouping provider overpayments. When HHSC-OIG identifies fraud, waste, and abuse, a case may be referred to the MFCU or Antitrust and Civil Medicaid Fraud Section, or result in administrative enforcement.
1.11.3 Employee Education on False Claims Recovery

Title 42 United States Code (U.S.C.) §1396a(a)(68) requires any entity that receives or makes annual Medicaid payments of at least $5,000,000 to establish written policies that provide detailed information about each employee’s role in preventing and detecting waste, fraud, and abuse in federal health-care programs. These written policies, which must apply to all employees of the entity (including management) and the employees of any contractor or agent of the entity, must address:

- Administrative remedies for false claims and statements as provided in 31 U.S.C. §3802.
- Texas law relating to civil and criminal penalties for false claims (including Chapter 36 of the Human Resources Code; section 35A.02 of the Penal Code; Title 1, Chapter 371, Subchapter G of the TAC; and other applicable law).
- Whistleblower protections under the above laws (including section 36.115 of the Human Resources Code).

In addition, these written policies must include detailed provisions regarding the entity’s policies and procedures for detecting and preventing fraud, waste, and abuse. The entity must also include a specific discussion of the following in all employee handbooks:

- The above laws
- The entity’s policies and procedures for detecting and preventing fraud, waste, and abuse
- The rights of employees to be protected as whistleblowers

TMHP sends a yearly letter to each provider that receives over $5,000,000 in Medicaid payments. This letter requires providers to verify that they have educated their staff on the False Claims Act. Failure to return this letter, signed by the provider, may result in an administrative hold on the provider’s Texas Medicaid payments.

1.11.4 Managed Care Organization (MCO) Special Investigative Unit (SIU)

All MCOs that contract with HHSC to administer managed care benefits to Texas Medicaid clients are required to establish and maintain an SIU that works in cooperation with HHSC-OIG and the OAG.

Refer to: 1 TAC §533.012, §531.113, §531.1131, §353.501-353.505, and 370.501-370.505 for additional information.

The MCO and SIU will do the following:

- The MCO must maintain the SIU within the MCO or contract with another entity for any investigation.
- The established SIU will identify and investigate cases of suspected fraud, waste, and abuse in Texas Medicaid in accordance with Title 1, Chapter 353, Subchapter F of the TAC.
- The MCO and SIU (as applicable) must submit the following:
  - An annual plan that has been adopted by the MCO and approved by HHSC-OIG describing how it will prevent and reduce fraud and abuse in accordance with 1 TAC, §§353.501 and 353.502.
  - A monthly open case list to OIG Medicaid Program Integrity and the MFCU.

The MCO will refer a case to both HHSC-OIG and MFCU in the following situations:

- When fraud, waste, and abuse is discovered in the Medicaid or CHIP programs. (The MCO SIU must immediately notify the HHSC-OIG and MFCU and begin payment recovery efforts, unless HHSC-OIG or MFCU notifies the MCO to stop the recovery effort, as provided in Texas Government Code §531.1131.)
• When possible fraud, waste, and abuse is discovered in the Medicaid or CHIP programs. (The MCO SIU must refer the alleged fraud or abuse to HHSC-OIG within 30 working days of completing a review. The SIU report and referral must completely and accurately detail its findings in accordance with 1 TAC §353.502.)

• When there is reason to believe that a delay in the referral may result in:
  • Harm or death to patients
  • Loss, destruction, or alteration of valuable evidence
  • Significant monetary loss that may not be recoverable
  • Hindrance of an investigation or criminal prosecution of the offense

1.12 Texas Medicaid Limitations and Exclusions

Medicaid pays for services on behalf of clients to the provider of service according to Texas Medicaid’s limitations and procedures. TMHP does not make Medicaid payments directly to clients.

The following services, supplies, procedures, and expenses are not benefits of Texas Medicaid. This list is not all inclusive.

• Autopsies
• Care and treatment related to any condition for which benefits are provided or available under Workers’ Compensation laws
• Cellular therapy
• Chemolase injection (chymodiactin, chymopapain)
• Dentures or endosteal implants for adults
• Ergonovine provocation test
• Excise tax
• Fabric wrapping of abdominal aneurysms
• Hair analysis
• Heart–lung monitoring during surgery
• Histamine therapy–intravenous
• Hyperthermia
• Hysterectomy for infertility
• Immunizations or vaccines unless they are otherwise covered by Texas Medicaid (These limitations do not apply to services provided through the THSteps Program.)
• Immunotherapy for malignant diseases
• Infertility
• Inpatient hospital services to a client in an institution for tuberculosis, mental disease, or a nursing section of public institutions for persons with intellectual disabilities
• Inpatient hospital tests that are not specifically ordered by a provider who is responsible for the diagnosis or treatment of the client’s condition
• Intragastric balloon for obesity
• Joint sclerotherapy
• Keratoprosthesis/refractive keratoplasty
• Laetrile
• Mammoplasty for gynecomastia

More than $200,000 per client per benefit year (November 1 through October 31) for any health-care and remedial care services provided to a hospital inpatient by the hospital (If the $200,000 amount is exceeded because of an admission for an approved organ transplant, the allowed amount for that claim is excluded from the computation. This limitation does not apply to clients eligible for CCP or clients with an organ transplant.)

More than 30 days of inpatient hospital stay per spell of illness (Each spell of illness must be separated by 60 consecutive days during which the client has not been an inpatient in a hospital.)

**Important:** CCP provides medically necessary, federally allowable treatment for Medicaid/THSteps clients who are 20 years of age and younger. Some health-care services that usually would not be covered under Medicaid may be available to CCP-eligible clients. An additional 30-day spell of illness begins with the date of specified covered organ transplant. No spell-of-illness limitation exists for Medicaid THSteps clients who are 20 years of age and younger.

• Obsolete diagnostic tests
• Oral medications, except when claims are submitted by a hospital for services that are provided given in the emergency room or the inpatient setting (Hospital take-home drugs or medications given to the client are not a benefit.)

**Important:** Outpatient prescription medications are covered through the Medicaid Vendor Drug Program. See the Subsection 1.1, “About the Vendor Drug Program” in the Outpatient Drug Services Handbook (Vol. 2, Provider Handbooks) for more information.

• Orthoptics (except CCP)
• Outpatient and nonemergency inpatient services provided by military hospitals
• Outpatient mental health services provided by a psychiatric assistant, psychological assistant (excluding Master’s level LPA), or a licensed chemical dependency counselor
• Oxygen (except CCP and home health)
• Parenting skills
• Payment for eyeglass materials or supplies regardless of cost if they do not meet Texas Medicaid specifications
• Payment to physicians for supplies (All supplies, including anesthetizing agents such as Xylocaine, inhalants, surgical trays, or dressings, are included in the surgical payment.)
• Podiatry, optometric, and hearing aid services in long term care facilities, unless ordered by the attending physician
• Private room facilities except when:
  • A critical or contagious illness exists that results in disturbance to other patients and is documented as such.
  • It is documented that no other rooms are available for an emergency admission.
  • The hospital only has private rooms.
- Procedures and services considered experimental or investigational

  **Note:** While procedures and services that are experimental or investigational are not a benefit of Texas Medicaid, routine patient care costs for individuals enrolled in clinical trials may be covered as medically necessary when those services are current Texas Medicaid benefits. Texas Insurance Code Section 1379.051 defines “routine patient care costs” as “the costs of any medically necessary health care service for which benefits are provided under a health benefit plan, without regard to whether the enrollee is participating in a clinical trial.” Refer to Texas Insurance Code Chapter 1379 for additional information.

- Prosthetic and orthotic devices (except CCP)
- Prosthetic eye or facial quarter

- Psychiatric services:
  - Outpatient behavioral health services for which no prior authorization has been given

**Refer to:** Section 4, “Outpatient Mental Health Services” in the *Behavioral Health and Case Management Services Handbook (Vol. 2, Provider Handbooks).*

- Quest test (infertility)
- Recreational therapy
- Review of old X-ray films
- Routine cardiovascular and pulmonary function monitoring during the course of a surgical procedure under anesthesia
- Separate fees for completing or filing a Medicaid claim form (The cost of claims filing is to be incorporated in the provider’s usual and customary charges to all clients.)
- Services and supplies to any resident or inmate in a public institution
- Services or supplies for which benefits are available under any other contract, policy, or insurance, or which would have been available in the absence of Texas Medicaid
- Services or supplies for which claims were not received within the filing deadline
- Services or supplies that are not reasonable and necessary for diagnosis or treatment
- Services or supplies that are not specifically provided by Texas Medicaid
- Services or supplies provided in connection with cosmetic surgery except:
  - As required for the prompt repair of accidental injury
  - For improvement of the functioning of a malformed body member
  - When prior authorized for specific purposes by TMHP (including removal of keloid scars)
- Services or supplies provided outside of the U.S., except for deductible or coinsurance portions of Medicare benefits as provided for in this manual
- Services or supplies provided to a client after a finding has been made under utilization review procedures that these services or supplies are not medically necessary
- Services or supplies provided to a Texas Medicaid client before the effective date of his or her designation as a client, or after the effective date of his or her denial of eligibility
- Services that are payable by any health, accident, other insurance coverage, or any private or other governmental benefit system, or any legally liable third party
- Services that are provided by an interpreter (except sign language interpreting services requested by a physician)
• Services that are provided by ineligible, suspended, or excluded providers
• Services that are provided by the client’s immediate relative or household member
• Services that are provided by Veterans Administration facilities or U.S. Public Health Service Hospitals
• Sex change operations
• Silicone injections
• Social and educational counseling except for certain health and disability related and counseling services
• Sterilization reversal
• Sterilizations (including vasectomies) unless the client has given informed consent 30 days before surgery, is mentally competent, and is 21 years of age or older at the time of consent (This policy complies with 42 CFR §441.250, Subpart F.)
• Take-home and self-administered drugs except as provided under the Vendor Drug or family planning pharmacy services or for clients being treated for a substance use disorder
• Tattooing (commercial or decorative only)
• Telephone calls with clients or pharmacies (except as allowed for case management)
• Thermogram
• Treatment of flatfoot conditions for solely cosmetic purposes, the prescription of supportive devices (including special shoes), and the treatment of subluxations of the foot

Refer to the applicable handbooks in Volume 2 of this manual for additional information.

1.13 Forms
The following linked forms can also be found on the Forms page of the Provider section of the TMHP website at www.tmhp.com:

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SECTION 2: TEXAS MEDICAID FEE-FOR-SERVICE REIMBURSEMENT

TEXAS MEDICAID PROVIDER PROCEDURES MANUAL: VOL. 1

MAY 2021
# SECTION 2: TEXAS MEDICAID FEE-FOR-SERVICE REIMBURSEMENT

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2.1 Payment Information
Texas Medicaid reimbursements are available to all enrolled providers by check or electronic funds transfer (EFT).

Refer to: Subsection 1.2, “Payment Information” in “Section 1: Provider Enrollment and Responsibilities” (Vol. 1, General Information).

2.2 Fee-for-Service Reimbursement Methodology
Texas Medicaid reimburses providers using several different reimbursement methodologies, including fee schedules, reasonable cost with interim rates, hospital reimbursement methodology, provider-specific encounter rates, reasonable charge payment methodology, and manual pricing. Each Texas Medicaid service describes the appropriate reimbursement for each service area.

Note: If a client is covered by a Medicaid managed care organizations (MCO) or dental plan, providers must contact the client’s MCO or dental plan for reimbursement information. The MCOs and dental plans are not required to follow the Texas Medicaid fee schedules, so there may be some differences in reimbursement based on decisions made by the individual health and dental plans.

2.2.1 Online Fee Lookup (OFL) and Static Fee Schedules
Texas Medicaid reimburses certain providers based on rates published in the OFL and static fee schedules. These rates are uniform statewide and by provider type. According to this type of reimbursement methodology, the provider is paid the lower of the billed charges or the Medicaid rate published in the applicable static fee schedule or OFL.

Providers can obtain fee information using the OFL functionality on the TMHP website at www.tmhp.com. The online OFL can be used to:

- Retrieve real-time fee information.
- Search for procedure code reimbursement rates individually, in a list, or in a range.
- Search and review contracted rates for a specific provider (provider must login).
- Retrieve up to 24 months of history for a procedure code by searching for specific dates of service within that 2-year period.
- Search for benefit limitations for dental and durable medical equipment (DME) procedure codes.

Providers can obtain the static fee schedules as Microsoft Excel® spreadsheets or portable document format (PDF) files from the TMHP website at www.tmhp.com.

Type of service (TOS) codes payable for each procedure code are available on the OFL and the static fee schedules.

The following provider types and services are reimbursed based on rates published with the rates calculated in accordance with the referenced reimbursement methodology as published in the Texas Administrative Code (TAC), Part 1 Administration, Part 15 Texas Health and Human Services Commission (HHSC), and Chapter 355 Reimbursement Rates.

- **Ambulance.** The Medicaid rates for ambulance services are calculated in accordance with 1 TAC §355.8600.
- **Ambulatory Surgical Center (ASC).** The Medicaid rates for ASCs are calculated in accordance with 1 TAC §355.8121.
- **Case Management for Children and Pregnant Women.** The Medicaid rates for this service are calculated in accordance with 1 TAC §355.8401.
• Targeted Case Management for Early Childhood Intervention (ECI). The Medicaid rate for this service is reimbursed in accordance with 1 TAC §355.8421.

• Specialized Skills Training for ECI. The Medicaid rate for this service is reimbursed in accordance with 1 TAC § 355.8422.

• Certified Nurse-Midwife (CNM). The Medicaid rates for CNMs are calculated in accordance with 1 TAC §355.8161.

• Certified Registered Nurse Anesthetist (CRNA). According to 1 TAC §355.8221, the Medicaid rate for CRNAs is 92 percent of the rate reimbursed to a physician anesthesiologist for the same service.

• Certified Respiratory Care Practitioner (CRCP) Services. The Medicaid rate per daily visit for 99503 is calculated in accordance with 1 TAC §355.8089.

• Chemical Dependency Treatment Facility (CDTF). The Medicaid rates for CDTF services are calculated in accordance with 1 TAC §355.8241.

• Chiropractic Services. The Medicaid rates for chiropractic services are calculated in accordance with 1 TAC §355.8085.

• Dental. The Medicaid rates for dentists are calculated as access-based fees in accordance with 1 TAC §355.8085, 1 TAC §355.8441(11), and 1 TAC §355.455(b).

• Durable Medical Equipment, Prostheses, Orthoses and Supplies (DMEPOS). DMEPOS items provided by home health agencies and providers/suppliers of DMEPOS are reimbursed in accordance with 1 TAC §355.8023. DMEPOS items provided by the Comprehensive Care Program (CCP) are reimbursed in the same manner, in accordance with 1 TAC §355.8441.

• Family Planning Services. The Medicaid rates for family planning services are calculated in accordance with 1 TAC §355.8581.

• Genetic Services. The procedure codes and Medicaid rates for genetic services are listed in the OFL or the Physician - Genetics fee schedule on the TMHP website at www.tmhp.com.

• Hearing Aid and Audiometric Evaluations. Hearing screening services for newborns are provided at the birthing facility before discharge and reimbursed in accordance with the reimbursement methodology for the specific type of birthing facility. Outpatient hearing screening and diagnostic testing services for children are provided by physicians and are reimbursed in accordance with the reimbursement methodology for physician services at 1 TAC §355.8085, 1 TAC §355.8141, and 1 TAC §355.8441.

• Texas Medicaid (Title XIX) Home Health Services. The reimbursement methodology for home health nursing and aide services delivered by home health agencies are statewide visit rates calculated in accordance with 1 TAC §355.8021.

• Independent Laboratory. The Medicaid rates for independent laboratories are calculated in accordance with 1 TAC §355.8610, and the Deficit Reduction Act (DEFRA) of 1984. By federal law, Medicaid payments for a clinical laboratory service cannot exceed the Medicare payment for that service. Early Periodic Screening, Diagnosis, and Treatment (EPSDT)/Texas Health Steps medical and newborn screening laboratory services provided by the Department of State Health Services (DSHS) Laboratory are reimbursed based on the Medicare payment for that service.

• Indian Health Services. The reimbursement methodology for services provided in Indian Health Services Facilities operating under the authority of Public Law 93-638 is located at 1 TAC §355.8620.

• In-Home Total Parenteral Nutrition (TPN) Supplier. The Medicaid rates for these providers are calculated in accordance with 1 TAC §355.8087.
• **Licensed Clinical Social Worker (LCSW).** According to 1 TAC §355.8091, the Medicaid rate for LCSWs is 70 percent of the rate paid to a psychiatrist or psychologist for a similar service per 1 TAC §355.8085.

• **Licensed Marriage and Family Therapist (LMFT).** According to 1 TAC §355.8091, the Medicaid rate for LMFTs is 70 percent of the rate paid to a psychiatrist or psychologist for a similar service per 1 TAC §355.8085.

• **Licensed Midwife (LM).** According to 1 TAC §355.8161, covered professional services provided by an LM and billed under the LM’s own provider number are reimbursed the lesser of the LM’s billed charges or 70 percent of the reimbursement for the same professional service paid to a physician (M.D. or D.O.).

• **Licensed Professional Counselor (LPC).** According to 1 TAC §355.8091, the Medicaid rate for LPCs is 70 percent of the rate paid to a psychiatrist or psychologist for a similar service per 1 TAC §355.8085.

• **Maternity Service Clinic (MSC).** The Medicaid rates for these providers are calculated in accordance with 1 TAC §355.8085.

• **Nurse Practitioner (NP) and Clinical Nurse Specialist (CNS).** According to Title 1 TAC §355.8281, the Medicaid rate for NPs and CNSs is 92 percent of the rate paid to a physician (doctor of medicine [MD] or doctor of osteopathy [DO]) for the same service and 100 percent of the rate paid to physicians for laboratory services, X-ray services, and injections.

• **Physical (PT), Occupational (OT), and Speech (ST) Therapy Services.** Covered therapy services provided by home health agencies, comprehensive outpatient rehabilitation facilities or outpatient rehabilitation facilities, independent therapists (including Early Childhood Intervention) and physicians and other practitioners are reimbursed according to 1 TAC §355.8097.

• **Physical Therapists/Independent Practitioners.** The Medicaid rates for these providers are calculated in accordance with 1 TAC §355.8085.

• **Physician.** The Medicaid rates for physicians and other practitioners are calculated in accordance with 1 TAC §355.8085.

• **Physician Assistant (PA).** According to 1 TAC §355.8093, the Medicaid rate for PAs is 92 percent of the rate paid to a physician (MD or DO) for the same service and 100 percent of the rate paid to physicians for laboratory services, X-ray services, and injections.

• **Psychologist.** The Medicaid rates for psychologists are calculated in accordance with 1 TAC §355.8085.

• **Radiological and Physiological Laboratory and Portable X-Ray Supplier.** The Medicaid rates for these providers are calculated in accordance with 1 TAC §355.8085.

• **Renal Dialysis Facility.** The Medicaid rates for these providers are composite rates based on calculations specified by the Centers for Medicare & Medicaid Services (CMS).

• **School Health and Related Services (SHARS).** The Medicaid rates for these providers are calculated in accordance with 1 TAC §355.8443.

• Texas Health Steps reimburses by provider type in accordance with 1 TAC §355.8441. Approved providers enrolled in Texas Medicaid are reimbursed for THSteps services in the same manner as they are reimbursed for other Medicaid services. THSteps CCP reimburses for DME and expendable supplies in accordance with 1 TAC §355.8441(2).

• Telemedicine, telehealth, and home telemonitoring services are reimbursed in accordance with 1 TAC 355.7001.
• **Tuberculosis (TB) Clinics.** The Medicaid rates for these providers are calculated in accordance with 1 TAC §355.8085.

• **Vision Care (Optometrists, Opticians).** The Medicaid rates for these providers are calculated in accordance with 1 TAC §355. 8001 and §355.8085.

### 2.2.1.1 Non-emergent and Non-urgent Evaluation and Management (E/M) Emergency Department Visits

Section 104 of the Tax Equity and Fiscal Responsibility Act (TEFRA) of 1982 requires that Medicare and Medicaid limit reimbursement for those physician services furnished in outpatient hospital settings (e.g., clinics and emergency situations) that are ordinarily furnished in physician offices. The limit is 60 percent of the Medicaid rate for the non-emergency service furnished in physician offices.

Reimbursement for non-emergent and non-urgent services that are rendered by the facility during the emergency room visit will be limited to 125 percent of the adult, physician office visit fee for procedure code 99202. Reimbursement will not be reduced for those services that were rendered to address conditions that meet any of the following criteria:

- Problems of high-severity
- Problems that require urgent evaluation by a physician
- Problems that pose immediate and significant threats to physical or mental function
- Critically ill or critically injured

Non-emergent and non-urgent services that are rendered by rural hospitals will be reimbursed at 65 percent of the allowed rates.

Non-rural hospitals will receive a flat rate which is limited to 125 percent of the adult, physician office visit fee for procedure code 99202.

Diagnostic services, such as laboratory and radiology, will not be reduced by 40 percent.

**Refer to:** Subsection 9.2.56.3.5, “Physician Services Provided in the Emergency Department” in the *Medical and Nursing Specialists, Physicians, and Physician Assistants Handbook (Vol. 2, Provider Handbooks)* for more information about non-emergent and non-urgent services rendered in the emergency department.

Subsection 4.2.2.1, “Emergency Department Payment Reductions” in the *Inpatient and Outpatient Hospital Services Handbook (Vol. 2, Provider Handbooks)* for more information about non-emergent and non-urgent services rendered in the emergency department.

These procedures are designated with note code “1” in the current fee schedule or OFL on the TMHP website at [www.tmhp.com](http://www.tmhp.com).

The following services are excluded from the 60-percent limitation:

- Services furnished in rural health clinics (RHCs)
- Surgical services that are covered ASC/hospital-based ambulatory surgical center (HASC) services
- Anesthesiology and radiology services
- Prenatal services when billed with modifier TH and the appropriate E/M procedure code to the highest level of specificity
- Emergency services provided in a hospital emergency room after the sudden onset of a medical condition manifesting itself by acute symptoms of sufficient severity (including severe pain), such that the absence of immediate medical attention could reasonably be expected to result in one of the following:
  - Serious jeopardy to the client’s health
• Serious impairment to bodily functions
• Serious dysfunction of any bodily organ or part

### 2.2.1.2 Payment Window Reimbursement Guidelines for Services Preceding an Inpatient Admission

According to the three-day and one-day payment window reimbursement guidelines, most professional and outpatient diagnostic and nondiagnostic services that are rendered within the designated timeframe of an inpatient hospital stay and are related to the inpatient hospital admission will not be reimbursed separately from the inpatient hospital stay if the services are rendered by the hospital or an entity that is wholly owned or operated by the hospital.

These reimbursement guidelines do not apply in the following circumstances:

- The professional services are rendered in the inpatient hospital setting.
- The hospital and the physician office or other entity are both owned by a third party, such as a health system.
- The hospital is not the sole or 100-percent owner of the entity.

**Refer to:** Subsection 3.7.4.14, “Payment Window Reimbursement Guidelines” in the Inpatient and Outpatient Hospital Services Handbook (Vol. 2, Provider Handbooks) for additional information about the payment window reimbursement guidelines.

### 2.2.1.3 Drugs and Biologicals

Physician-administered drugs and biologicals are reimbursed under Texas Medicaid as access-based fees under the physician fee schedule in accordance with 1 TAC §355.8085. Physicians and other practitioners are reimbursed for physician-administered drugs and biologicals at the lesser of their usual and customary or billed charges and the Medicaid fee established by the HHSC. The Medicaid fee is an estimate of the provider’s acquisition cost for the specific drug and biological. An invoice must be submitted when it is in the provider’s possession. Submission of an invoice will document that the provider is billing the lesser of the usual and customary charge or the access-based fee.

HHSC reserves the option to use other data sources to determine Medicaid fees for drugs and biologicals when AWP or ASP calculations are determined to be unreasonable or insufficient.

Prescriptions are covered under the Texas Medicaid Vendor Drug Program (VDP). The reimbursement methodology for pharmacy services is located at 1 TAC §§355.8541–355.8551.

### 2.2.2 Cost Reimbursement

Medicaid providers who are cost reimbursed are subject to cost reporting, cost reconciliation, and cost settlement processes, including time study requirements.

The following providers are cost reimbursed in accordance with the noted TAC rules:

- 1 TAC §355.743—Mental health (MH) case management
- 1 TAC §355.746—Mental retardation (MR) service coordination
- 1 TAC §355.781—MH rehabilitative services
- 1 TAC §355.8443—School Health and Related Services (SHARS)
- 1 TAC §355.8061—Outpatient Hospital Reimbursement
- 1 TAC §355.8056—State-Owned Teaching Hospital Reimbursement Methodology
2.2.3 Reasonable Cost and Interim Rates
Outpatient hospital services are reimbursed in accordance with 1 TAC §355.8061. The reimbursement methodology is based on reasonable costs, and providers are reimbursed at an interim rate based on the provider’s most recent Medicaid cost report settlement. To determine the provider’s payable amount, the interim rate is applied to the claim details allowed amount.

2.2.4 Hospitals
Inpatient and outpatient hospital services are reimbursed as follows:

- 1 TAC §355.8052—Inpatient Hospital Reimbursement
- 1 TAC §355.8056—State-Owned Teaching Hospital Reimbursement Methodology
- 1 TAC §355.8058—Inpatient Direct Graduate Medical Education (GME) Reimbursement
- 1 TAC §355.8060—Reimbursement Methodology for Freestanding Psychiatric Facilities
- 1 TAC §355.8061—Outpatient Hospital Reimbursement
- 1 TAC §355.8065—Disproportionate Share Hospital (DSH) Reimbursement Methodology
- 1 TAC §355.8066—Hospital-Specific Limit Methodology

2.2.5 Provider-Specific Visit Rates
Medicaid provider-specific prospective payment system (PPS) visit rates for RHCs are calculated in accordance with 1 TAC §355.8101, and those for federally qualified health centers (FQHCs) are calculated in accordance with 1 TAC §355.8261.

Refer to:
- Section 4, “Federally Qualified Health Center (FQHC)” in the Clinics and Other Outpatient Facility Services Handbook (Vol. 2, Provider Handbooks)
- Section 7, “Rural Health Clinic” in the Clinics and Other Outpatient Facility Services Handbook (Vol. 2, Provider Handbooks).

2.2.6 Manual Pricing
When services or products do not have an established reimbursement amount, the detail or claim is manually reviewed to determine an appropriate reimbursement. The manual pricing methodology for DME and expendable supplies is included with the reimbursement methodology for these products. DME and medical supplies, other than nutritional products, that have no established fee are subject to manual pricing at the documented MSRP less 18 percent or the provider’s documented invoice cost.

2.3 Reimbursement Reductions
Texas Medicaid implemented mandated rate reductions for certain services. The Online Fee Lookup (OFL) and static fee schedules include a column titled “Adjusted Fee” to display the individual fees with all mandated percentage reductions applied. Additional information about rate changes is available on the TMHP website at www.tmhp.com/resources/rate-and-code-updates/rate-changes.

2.4 Using Payouts to Satisfy Accounts Receivables Across Programs and Alternate Provider Identifiers
The TMHP accounts receivable process identifies funds that a provider owes to TMHP and subtracts these funds from payments to the provider. TMHP satisfies outstanding accounts receivables using all available funds from the providers’ Medicaid payouts, as well as managed care payments, until the accounts receivables have been recovered. Outstanding balances are recovered as follows:

- For outstanding fee-for-service accounts receivables, TMHP first recovers funds from any available fee-for-service payments. If there is still an accounts receivable balance for that week’s financial cycle, TMHP recovers funds from any available managed care payments.
• For outstanding managed care accounts receivables, TMHP first recovers funds from any available managed care payments. If there is still an accounts receivable balance for that week’s financial cycle, TMHP will recover funds from any available fee-for-service payments

1099 Reports
Providers receive one 1099 report for each provider identifier. The 1099 report has combined information for both fee-for-service and managed care programs.

Paper Remittance and Status (R&S) Report
The summary page of the R&S report has combined information from the fee-for-service and managed care programs.

The Financial Transactions Sub-Owner Recoupment page has accounts receivable for both programs. A column on the page identifies the program (Medicaid [fee-for-service] or Managed Care) from which the funds were recouped.

The Financial Transactions Accounts Receivable page has the accounts receivable for both programs. A column identifies the program (Medicaid [fee-for-service] or Managed Care) from which the funds were recouped.

The Original Date in the Accounts Receivable section of the R&S Report reflects the date on which the accounts receivable first appeared on the R&S Report.

ER&S Report
The Pending and Non-Pending ER&S Reports have combined information for both programs.

2.4.1 HHSC Recoupment of Accounts Receivables from Alternate Provider Identifiers
HHSC recoups the outstanding accounts receivable balances of all existing Medicaid and managed care Texas Provider Identifiers (TPIs) from alternate TPIs that use the same Tax ID or National Provider Identifier (NPI).

If a Medicaid or managed care provider has a TPI that is no longer active or has been terminated and that TPI has an outstanding accounts receivable balance, the balance is recouped from future payments made to any and all TPIs that have the same Tax ID or NPI. Recoupments are reflected on future R&S Reports.

Note: This process affects only managed care claims that are submitted to TMHP.

Refer to: Subsection 2.2.5, “Accounts Receivable” in the Medicaid Managed Care Handbook (Vol. 2, Provider Handbooks) for additional information about managed care claims and outstanding accounts receivables.

2.4.2 Medicaid Funds May Be Used to Satisfy Children with Special Health Care Needs (CSHCN) Services Program Accounts Receivables
A service that is rendered to a CSHCN Services Program client who receives retroactive Medicaid eligibility may be reimbursed by the CSHCN Services Program or by Medicaid, but not by both.

The CSHCN Services Program is the payer of last resort. The CSHCN Services Program does not supplement a client’s Medicaid benefits. However, services that are not a benefit of Medicaid may be covered by the CSHCN Services Program. If dual Medicaid and CSHCN Services Program eligibility is determined, claims that have already been paid by the CSHCN Services Program will be reprocessed under the appropriate program.
An accounts receivable is created for each CSHCN Services Program claim that is reprocessed and subsequently reimbursed under Medicaid so that the amount the CSHCN Services Program originally reimbursed can be returned to the CSHCN Services Program. If the CSHCN Services Program payout during the week’s financial cycle in which the claim was reprocessed is not sufficient to satisfy the accounts receivable, the provider’s Medicaid claim payouts are used to satisfy the CSHCN Services Program accounts receivable.

Note: The deduction from Medicaid claim payouts does not exceed the amount Medicaid reimbursed the provider when the CSHCN Services Program claim was reprocessed.

2.5 Additional Payments to High-Volume Providers

High volume provider payments are made to outpatient hospitals per 1 TAC §355.8061 and ASCs/HASCs per 1 TAC §355.8121.

Outpatient hospital services are those services provided by outpatient hospitals and ASCs/HASCs. The definition of a high-volume outpatient hospital provider is one that was paid a minimum of $200,000 during the qualifying period.

The reimbursement rate for non-high-volume hospitals is as follows with the application of the hospital specific interim rate:

<table>
<thead>
<tr>
<th>Non-High-Volume Provider</th>
<th>Current Allowable Rate</th>
</tr>
</thead>
<tbody>
<tr>
<td>Children’s hospitals</td>
<td>72.27 percent of the allowable charges</td>
</tr>
<tr>
<td>Rural hospitals</td>
<td>100 percent of the allowable charges</td>
</tr>
<tr>
<td>State-owned teaching hospitals</td>
<td>72.27 percent of the allowable charges</td>
</tr>
<tr>
<td>Other hospitals</td>
<td>68.44 percent of the allowable charges</td>
</tr>
</tbody>
</table>

The reimbursement rate for high-volume hospitals is as follows with the application of the hospital specific interim rate:

<table>
<thead>
<tr>
<th>High-Volume Provider</th>
<th>Current Allowable Rate</th>
</tr>
</thead>
<tbody>
<tr>
<td>Children’s hospitals</td>
<td>76.03 percent of the allowable charges</td>
</tr>
<tr>
<td>Rural hospitals</td>
<td>100 percent of the allowable charges</td>
</tr>
<tr>
<td>State-owned teaching hospitals</td>
<td>76.03 percent of the allowable charges</td>
</tr>
<tr>
<td>Other hospitals</td>
<td>72 percent of the allowable charges</td>
</tr>
<tr>
<td>ASCs/HASCs that qualify as high-volume providers</td>
<td>Additional 5.2 percent increase in payment rates</td>
</tr>
</tbody>
</table>

2.6 Out-of-State Medicaid Providers

Texas Medicaid covers medical assistance services provided to eligible Texas Medicaid clients while in a state other than Texas, as long as the client does not leave Texas to receive out-of-state medical care that can be received in Texas. Services provided outside the state are covered to the same extent medical assistance is furnished and covered in Texas when the service meets one or more requirements of 1 TAC §354.1440 (a). TMHP must receive claims from out-of-state providers within 365 days from the date of service.

Note: Border state providers (providers rendering services within 50 miles of the Texas border) are considered in-state providers for Texas Medicaid.

Refer to: Subsection 1.10, “Enrollment Criteria for Out-of-State Providers” in “Section 1: Provider Enrollment and Responsibilities” (Vol. 1, General Information).
2.7 Medicare Crossover Claim Reimbursement

2.7.1 Part A
Providers must accept Medicare assignment to receive Medicaid payment for any portion of the coinsurance and deductible amounts for services rendered to Qualified Medicare Beneficiary (QMB) and Medicaid Qualified Medicare Beneficiary (MQMB) clients. If a provider has accepted a Medicare assignment, the provider may receive, on behalf of the QMB or MQMB client, payment for deductible or coinsurance according to current payment guidelines.

Any payments made by Medicare and Medicaid must be considered payment in full. Providers that accept Medicare or Medicaid assignment cannot legally require the client to pay the Medicare coinsurance or deductible amounts or any remaining amount after Medicaid payment has been made.

The payment of the Medicare Part A coinsurance and deductibles for Medicaid clients who are Medicare beneficiaries is based on the following:

- If the Medicare payment amount equals or exceeds the Medicaid payment rate, Medicaid does not pay the Medicare Part A coinsurance/deductible on a Medicare crossover claim.
- If the Medicare payment amount is less than the Medicaid payment rate, Medicaid pays the Medicare Part A coinsurance/deductible, but the amount of the payment is limited to the lesser of the coinsurance/deductible or the amount remaining after the Medicare payment amount is subtracted from the Medicaid payment rate.

2.7.2 Part B
Texas Medicaid reimburses coinsurance liability for MQMB clients on valid, assigned Medicare claims that are within the amount, duration, and scope of the Medicaid program and, if Medicare did not exist, would be covered by Medicaid.

For Medicare crossover claims, Texas Medicaid reimburses the lesser of the following:

- The coinsurance and deductible payment
- The amount remaining after the Medicare payment amount is subtracted from the allowed Medicaid fee or encounter rate for the service (If this amount is less than the deductible, then the full deductible is reimbursed instead.)

If the Medicare payment is equal to or exceeds the Medicaid allowed amount or encounter payment for the service, Texas Medicaid does not make a payment for coinsurance.

Important: Medicaid payment of a client’s coinsurance/deductible liabilities satisfies the Medicaid obligation to provide coverage for services that Medicaid would have paid in the absence of Medicare coverage. The client has no liability for any balance or Medicare coinsurance and deductible related to Medicaid-covered services.

2.7.3 Part C: Medicare Advantage Plans (MAPs)

2.7.3.1 Contracted MAPs
HHSC makes a per-client-per-month payment to MAPs that contract with HHSC. The payment to the MAP includes all costs associated with the Medicare coinsurance and deductible for a client who is dually eligible for Medicare and Medicaid. TMHP does not reimburse the coinsurance or deductible amounts for these claims. These costs must be billed to the MAP and must not be billed to TMHP or the Medicaid client.

Refer to: A list of MAPs that are contracted with HHSC is available on the TMHP website at www.tmhp.com/topics/edi/map-contractors. The list is updated as additional plans receive approved contracts.
2.7.3.2 Noncontracted MAPs
Texas Medicaid reimburses professional and outpatient facility crossover claims the lesser of the following:

- The coinsurance and deductible amounts
- The amount remaining after the Medicare payment amount is subtracted from the allowed Medicaid fee or encounter rate for the service

For Medicare Part B cost sharing obligations, all deductible obligations will be reimbursed at 100 percent of the deductible amount owed, even if the cost sharing comparison results in a lower payment. For all other cost sharing obligations (including Medicare Part A and Part B coinsurance, and C), the cost sharing comparison is performed according to current guidelines.

**Exception:** Texas Medicaid will reimburse coinsurance liability for MQMB clients on valid, assigned Medicare claims that are within the amount, duration, and scope of the Medicaid program, and would be covered by Medicaid when the services are provided, if Medicare did not exist.

If the Medicare payment is equal to or exceeds the allowed Medicaid fee or encounter rate for the service, Texas Medicaid will not make a payment for coinsurance and deductible.

**Important:** Medicaid payment of a client’s coinsurance/deductible liabilities satisfies the Medicaid obligation to provide coverage for services that Medicaid would have paid in the absence of Medicare coverage. The client has no liability for any balance or Medicare coinsurance and deductible related to Medicaid-covered services.

2.7.4 Exceptions

2.7.4.1 Full Amount of Part B and Part C Coinsurance and Deductible Reimbursed
Texas Medicaid reimburses the full amount of the Medicare Part B and Part C (noncontracted MAPs only) coinsurance and deductible for the following services:

- All ambulance services
- Services rendered by psychiatrists, psychologists, and licensed clinical social workers
- Procedure codes R0070 and R0075 for services rendered by physicians

2.7.4.2 Nephrology (Hemodialysis, Renal Dialysis) and Renal Dialysis Facility Providers
Texas Medicaid pays the Medicare coinsurance less 5 percent and full Medicare deductible for Medicare crossover claims that are submitted by nephrology (hemodialysis, renal dialysis) and renal dialysis facility providers.

2.8 Home Health Agency Reimbursement
Home health service claims should not be submitted for payment until Medicaid certification is received and a prior authorization number is assigned.

Home Health Agency providers should note the following:

- The client’s primary physician must request professional, SN, and HHA services through a home health agency, and sign and date the POC.
- Claims are approved or denied according to eligibility, prior authorization status, and medical appropriateness.
- Claims must represent a numerical quantity of one-month for medical supplies according to the billing requirements.
• SN, HHA, OT, and PT services must be provided through a Medicaid-enrolled home health agency. These services must be billed using the home health agency's provider identifier. File these services on a UB-04 CMS-1450 claim form.

• OT and PT are always billed as POS 2 (home) and may be prior authorized to be provided in the home of the client or the home of the caregiver/guardian.

Note: Medical social services and speech-language pathology services are available to clients who are 20 years of age and younger and are not a home health services benefit. These services may be considered a benefit for clients who qualify for CCP.

Texas Medicaid does not reimburse separately for associated DME charges, including but not limited to battery disposal fees or state taxes. Reimbursement for any associated charges is included in the reimbursement for a specific piece of equipment.

Refer to: Subsection 2.2, “Fee-for-Service Reimbursement Methodology” in this section.

2.8.1 Pending Agency Certification
Home health agencies submitting claims before the enrollment process is complete or without prior authorization for services issued by the TMHP Home Health Services Prior Authorization Department will not be reimbursed. The effective date of enrollment is when all Texas Medicaid provider enrollment forms are received and approved by TMHP.

Upon the receipt of notice of Texas Medicaid enrollment, the agency must contact the TMHP Home Health Services Prior Authorization Department before serving a Texas Medicaid client for services that require a prior authorization number. Prior authorization cannot be issued before Texas Medicaid enrollment is complete. Regular prior authorization procedures are followed at that time.

Home health agencies that provide laboratory services must comply with the rules and regulations of the Clinical Laboratory Improvement Amendments (CLIA). Providers who do not comply with CLIA will not be reimbursed for laboratory services.

Refer to: Subsection 2.1.1, “Clinical Laboratory Improvement Amendments (CLIA)” in the Radiology and Laboratory Services Handbook (Vol. 2, Provider Handbooks).

2.8.2 Prohibition of Medicaid Payment to Home Health Agencies Based on Ownership
Medicaid denies Home Health Services claims when TMHP records indicate that the physician ordering treatment has a significant ownership interest in, or a significant financial or contractual relationship with, the nongovernmental home health agency billing for the services. Federal regulation Title 42 CFR §424.22 (d) states that “a physician who has a significant financial or contractual relationship with, or a significant ownership in a nongovernmental home health agency may not certify or recertify the need for Home Health Services care services and may not establish or review a plan of treatment.”

A physician is considered to have a significant ownership interest in a home health agency if either of the following conditions apply:

- The physician has a direct or indirect ownership of five percent or more in the capital, stock, or profits of the home health agency.

- The physician has an ownership of five percent or more of any mortgage, deed of trust, or other obligation that is secured by the agency, if that interest equals five percent or more of the agency’s assets.

A physician is considered to have a significant financial or contractual relationship with a home health agency if any of the following conditions apply:

- The physician receives any compensation as an officer or director of the home health agency.
• The physician has indirect business transactions, such as contracts, agreements, purchase orders, or leases to obtain services, medical supplies, DME, space, and salaried employment with the home health agency.

• The physician has direct or indirect business transactions with the home health agency that, in any fiscal year, amount to more than $25,000 or five percent of the agency’s total operating expenses, whichever is less.

Providers must submit claims for CCP services and general home health services on two separate UB-04 CMS-1450 paper claim forms with the appropriate prior authorization number. Claims that are denied because of an ownership conflict will continue to be denied until the home health agency submits documentation that indicates that the ordering physician no longer has a significant ownership interest in, or a significant financial or contractual relationship with, the home health agency that is providing the services. Providers should send documentation to TMHP Provider Enrollment at the address indicated in the “Appendix A: State, Federal, and TMHP Contact Information” (Vol. 1, General Information).

2.9 Federal Medical Assistance Percentage (FMAP)

The Federal Medical Assistance Percentages (FMAPs) are used in determining the amount of Federal matching funds for State expenditures for assistance payments for certain social services and State medical and medical insurance expenditures. The Social Security Act requires the Secretary of Health and Human Services to calculate and publish the FMAPs each year.

The “Federal Medical Assistance Percentages” are for Medicaid. Section 1905(b) of the Act specifies the formula for calculating Federal Medical Assistance Percentages.

“Enhanced Federal Medical Assistance Percentages” are for the State Children’s Health Insurance Program (SCHIP) under Title XXI of the Social Security Act. Section 2105(b) of the Act specifies the formula for calculating Enhanced Federal Medical Assistance Percentages. The FMAPs are subject to change.
# SECTION 3: TMHP ELECTRONIC DATA INTERCHANGE (EDI)

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</table>
3.1 TMHP EDI Overview

The Texas Health and Human Services Commission (HHSC) and the Texas Medicaid & Healthcare Partnership (TMHP) encourage providers to submit claims using electronic methods. Providers can participate in the most efficient and effective method of submitting requests to TMHP by submitting through the TMHP EDI Gateway. TMHP uses the Health Insurance Portability and Accountability Act (HIPAA)-compliant American National Standards Institute (ANSI) ASC X12 5010 file format through secure socket layer (SSL) and virtual private networking (VPN) connections for maximum security. Providers can access TMHP’s electronic services through the TMHP website at www.tmhp.com, TexMedConnect, vendor software, and third party billing agents. Providers may also submit claims using paper forms.

3.1.1 Advantages of Electronic Services

- **It’s fast.** No more waiting by the mailbox or telephone inquiries; know what’s happening to claims in less than 24 hours and receive reimbursement for approved claims within a week. TexMedConnect users can submit individual requests interactively and receive a response immediately.

- **It’s free.** All electronic services offered by TMHP are free, including TexMedConnect and its technical support and training.

- **It’s easy.** TMHP offers computer-based training (CBT) for TexMedConnect, Medicaid billing, and many other topics, including those for the Children with Special Health Care Needs (CSHCN) Services Program, and Long Term Care, as well as a large library of reference materials and manuals on www.tmhp.com.

- **It’s safe.** TMHP EDI services use VPN and SSL connections, just like the United States government, banks, and other financial institutions, for maximum security.

- **It’s accurate.** TexMedConnect and most vendor software programs have features that let providers know when they’ve made a mistake, which means fewer rejected and denied claims. Rejected claims are returned with messages that explain what’s wrong, so the claim can be corrected and resubmitted right away. Denied claims appear on the provider’s Remittance and Status (R&S) Report along with paid and pending claims.

- **It’s there when it’s needed.** Electronic services are available day and night; from home, the office, or anywhere in the world.

- **It makes record keeping and research easy.** Not only can TexMedConnect be used to send and receive claims, it can check client eligibility, retrieve Electronic Remittance and Status (ER&S) Reports, perform claim status inquiries, and archive claims. TexMedConnect can generate and print reports on everything it sends, receives, and archives.

3.1.2 Electronic Services Available

- Eligibility verification
- Claims submission
- Claim status inquiry (CSI)
- ER&S Reports
- Appeals (also known as correction and resubmission)

3.1.3 Paper Remittance and Status (R&S) Reports No Longer Available

TMHP no longer produces or distributes paper R&S Reports. This initiative saves the state of Texas the cost of printing and mailing Paper R&S Reports.
All R&S Reports are now available online through the secure portion of the TMHP website at [www.tmhp.com](http://www.tmhp.com). Providers who receive an ER&S Report with third party software are not affected by this change.

Online R&S Reports are available as a portable document format (PDF) file every Monday morning—four days earlier than paper R&S Reports were available. Providers must have a provider administrator account on the TMHP website to receive R&S Reports. Providers who do not have a provider administrator account should create one to avoid delays or interruptions to business processes.

Providers can follow the instructions in the [TMHP Portal Security Training Guide](http://www.tmhp.com) to setup a provider administrator account.

### 3.2 Electronic Billing

Providers who want to transition from paper billing to electronic billing must decide how they will submit their claims to TMHP. Providers can use TexMedConnect on the TMHP website at [www.tmhp.com](http://www.tmhp.com), vendor software that submits files directly to TMHP, or they may use a third party billing agent (e.g., billing companies and clearinghouses) who submit files on the provider’s behalf.

#### 3.2.1 TexMedConnect

TexMedConnect is a free, web-based, claims submission application provided by TMHP. Technical support and training for TexMedConnect are also available free from TMHP. Providers can submit claims, eligibility requests, claim status inquiries, appeals, and download ER&S Reports (in either PDF or ANSI 835 formats) using TexMedConnect. TexMedConnect can interactively submit individual claims that are processed in seconds. To use TexMedConnect, providers must have:

- An internet service provider (ISP)
- Microsoft® Internet Explorer®
- Google Chrome®
- Mozilla Firefox®

A broadband connection is recommended but not required. Providers that use TexMedConnect can find the online TexMedConnect manuals for Acute Care and Long Term Care on the TexMedConnect Info web page in the EDI section of the TMHP website at [www.tmhp.com/resources/texmedconnect](http://www.tmhp.com/resources/texmedconnect).

#### 3.2.2 Vendor Software

Providers that do not use TexMedConnect must use vendor software to create, submit, and retrieve data files. Providers can use software from any vendor listed on the EDI Vendor Testing List, which is located in the EDI section of the TMHP website at [www.tmhp.com](http://www.tmhp.com). There are hundreds of software vendors that have a wide assortment of services and that have been approved to submit electronic files to TMHP. Providers that plan to access TMHP’s electronic services with vendor software should contact the vendor for details on software requirements. TMHP does not make vendor recommendations or provide any assistance for vendor software. Not all vendor software offers the same features or levels of support. Providers are encouraged to research their software thoroughly to make certain that it meets their needs and that it has completed testing and has been certified with TMHP.

#### 3.2.3 Third Party Billing Agents

Billing agents are companies or individuals who submit electronic files to TMHP on behalf of the provider. Generally, this means that the provider uses a product that sends billing or other information to the billing agent who processes it and transmits it to TMHP and other institutions. A complete list of billing agents who have completed the testing process and been certified by TMHP can be found on the EDI Vendor Testing List, which is located in the EDI section of the TMHP website at [www.tmhp.com](http://www.tmhp.com).
TMHP does not make billing agent recommendations or provide any assistance for billing agents’ software or services. TMHP has no information on the software or other requirements of billing agents. Providers should contact the billing agent to obtain information about their products and processes.

### 3.3 Gaining Access
Providers must setup their software or billing agent services to access the TMHP EDI Gateway. Providers who use billing agents or software vendors should contact those organizations for information about installation, settings, maintenance, and their processes and procedures for exchanging electronic data.

Providers that download the ANSI 835 file through TexMedConnect and providers that use vendor software must request a submitter ID. A submitter ID is necessary for vendor software to access TMHP’s electronic services. It serves as an electronic mailbox for the provider and TMHP to exchange data files. To order a submitter ID, providers must call the EDI Help Desk at 1-888-863-3638, Option 3. Providers that use a billing agent do not need a submitter ID.

Providers may receive an ER&S Report by completing the Electronic Remittance and Status (ER&S) Agreement and submitting it to the EDI Help Desk after setting up access to the TMHP EDI Gateway.

Refer to: Electronic Remittance Advice (ERA) Agreement on the TMHP website at www.tmhp.com.

### 3.4 Training
Providers should contact the TMHP Contact Center at 1-800-925-9126 for billing and training questions. Information about training opportunities is available in the Provider Education section of the TMHP website at www.tmhp.com. Providers may also use the many reference materials and workbooks available on the website. The TMHP EDI Help Desk provides technical assistance and does not provide training.

### 3.5 Electronic Transmission Reports
Providers are required to retain all claim and electronic file transmission records. Providers must verify that all claims submitted to TMHP are received and accepted. Providers must also track claims submissions against their claims payments to detect and correct all claim errors.

Refer to: Subsection 1.7.3, “Retention of Records and Access to Records and Premises” in “Section 1: Provider Enrollment and Responsibilities” (Vol. I, General Information) for more information about provider responsibility and electronic submissions.

If an electronic file transmission record is missing, providers can request that the transmission report file be reset by contacting the TMHP EDI Help Desk at 1-888-863-3638, Option 3. The TMHP EDI Help Desk will then reset the files for the production submitter ID provided. Requests for transmission reports produced in the previous 30 days will be provided at no cost to providers. Requests for transmission reports produced more than 30 days before the request will result in a charge of $500 plus 8.25 percent sales tax of $41.25 for a total charge of $541.25. Providers that hold a tax-exempt certificate will not be assessed the sales tax. This cost is per transmission report.

### 3.6 Provider Check Amounts Available Online
Acute care providers can search, view, and print on the TMHP website at www.tmhp.com all payment amounts issued during the previous year.

The features of the online check amount include:

- The ability to search information up to one year before the date of the search.
- All results are displayed on a single screen.
• All results can be printed on a single report.
• The 52 weeks of reimbursement payment information includes the:
  • Payment date
  • Payee name
  • Payment amount
  • Program for which payment was issued
  • Hold amount
  • Payment status

Providers must have or must create an administrative account to view their payment amounts online. Providers can then grant “View Payment Amounts” security permission to the office staff of their choice. Providers can access their check amounts by logging into their accounts from the TMHP website and then pressing View Payment Amounts.

Provider check amounts are also available through the automated inquiry system (AIS) telephone line and ER&S Reports.

3.7 Third Party Vendor Implementation

TMHP requires all software vendors and billing agents to complete EDI testing before access to the production server is allowed. Vendors that wish to begin testing may either call the EDI Help Desk at 1-888-863-3638, Option 3, or visit the Edifecs testing site at editesting.tmhp.com and use the TMHP Support link. An Edifecs account will be created for the vendor to begin testing EDI formats once they have enrolled for testing. After the successful completion of Edifecs testing and the submission of a Trading Partner Agreement, vendors must then complete end-to-end testing on the TMHP test server. Software vendors and billing agents must be partnered with at least one Texas provider before a test submitter ID can be issued. When end-to-end testing has been completed, the software vendor or billing agent will be added to the EDI Submitter List. Providers and billing agents may then order production submitter IDs for use with the vendor’s software. Companion guides and vendor specifications are available on the EDI page of the TMHP website at www.tmhp.com.

3.7.1 Automated Maintenance Process for All Electronic Submitters

All submitter folders have a maximum limit of 7500 files, and no files can be more than 30 days old. Files that exceed these limits will be purged by TMHP on a daily basis. Providers should review, retrieve, and backup their electronic response files within 30 days. Files not retrieved within the 30-day time period or files that exceed a maximum file count of 7500 will be purged by TMHP. All electronic submitters are responsible for the maintenance of their submitter folders. Files that are submitted using EDI version 5010 are limited to a maximum of 5,000 transactions per file. Files that have more than 5,000 transactions will be rejected.

Requests for transmission reports produced after the 30-day period, or resulting from a purge of over 7500 files will require fees, as outlined in Subsection 3.5, “Electronic Transmission Reports” in this section.
3.7.2 **Supported File Types**

TMHP EDI supports the following electronic HIPAA-compliant ANSI ASC X12 5010 transaction types:

<table>
<thead>
<tr>
<th>Electronic Transaction Types</th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td>270</td>
<td>Eligibility request</td>
</tr>
<tr>
<td>271</td>
<td>Eligibility response</td>
</tr>
<tr>
<td>276</td>
<td>Claim status inquiry</td>
</tr>
<tr>
<td>277</td>
<td>Claim status inquiry response</td>
</tr>
<tr>
<td>835</td>
<td>ER&amp;S Report</td>
</tr>
<tr>
<td>837D</td>
<td>Dental claims</td>
</tr>
<tr>
<td>837I</td>
<td>Institutional claims</td>
</tr>
<tr>
<td>837P</td>
<td>Professional claims</td>
</tr>
</tbody>
</table>

*Note:* Dental providers who submit American National Standards Institute, Accredited Standards Committee X12 (ANSI ASC X12N) 837D transactions through the TMHP Electronic Data Interchange (EDI) are required to include the header date of service (HDOS) to comply with International Classification of Diseases, Tenth Revision, Clinical Modification (ICD-10-CM) claims processing guidelines.

### 3.8 Forms

The following linked forms can also be found on the Forms page of the Provider section of the TMHP website at [www.tmhp.com](http://www.tmhp.com):

<table>
<thead>
<tr>
<th>Forms</th>
</tr>
</thead>
<tbody>
<tr>
<td>Electronic Remittance Advice (ERA) Agreement</td>
</tr>
<tr>
<td>Claim Status Inquiry Authorization</td>
</tr>
</tbody>
</table>
SECTION 4: CLIENT ELIGIBILITY

TEXAS MEDICAID PROVIDER PROCEDURES MANUAL: VOL. 1

MAY 2021
# SECTION 4: CLIENT ELIGIBILITY

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4.1 General Medicaid Eligibility

A person may be eligible for medical assistance through Medicaid if the following conditions are met:

- The applicant must be eligible for medical assistance at the time the service is provided. It is not mandatory that the process of determining eligibility be completed at the time service is provided; the client can receive retroactive eligibility. Services or supplies cannot be paid under Texas Medicaid if they are provided to a client before the effective date of eligibility for Medicaid or after the effective date of denial of eligibility. Having an application in process for Medicaid eligibility does not guarantee the applicant will be eligible.

- The service must be a benefit and determined medically necessary (except for preventive family planning, annual physical exams, and Texas Health Steps [THSteps] medical or dental checkup services) by Texas Medicaid and must be performed by an approved provider of the service.

- Applicants for medical assistance potentially are eligible for Medicaid coverage up to three calendar months before their application for assistance, if they have unpaid or reimbursable Medicaid-covered medical bills and have met all other eligibility criteria during the time the service was provided. The provision also includes deceased individuals when a bona fide agent requests application for services. An application for retroactive eligibility must be filed with the Health and Human Services Commission (HHSC); it is not granted automatically. The applicant must request the prior coverage from an HHSC representative and complete the section of the application about medical bills.

Clients who are not eligible for Medicaid but meet certain income guidelines may receive family planning services through other family planning funding sources. Clients not eligible for Medicaid are referred to a family planning provider. Clients seeking other services may be eligible for state health-care programs, some of which are described in this section.

Refer to: HHSC website at www.healthytexaswomen.org for information about family planning and the locations of family planning clinics that receive HHSC Family Planning Program funding.

4.1.1 Your Texas Benefits Medicaid Card

Clients receive a Your Texas Benefits Medicaid card that can be used to verify the client eligibility for various state-funded programs, including Medicaid.

The front of the card includes the client’s name, member ID, the ID of the agency that issued the card, and the date on which the card was sent.

The back of the card provides:

- An eligibility verification contact number. The number can be used to determine:
  - Program eligibility dates.
  - Retroactive eligibility (when applicable).
  - Eligible services (when applicable).
  - Medicaid managed care eligibility.

- An eligibility website address for clients and non-pharmacy providers.
- A non-managed care pharmacy claims assistance contact number.
- The Medicaid Client Transfer Line at 1-800-252-8263.

Client TPR and other insurance information can also be verified using the benefit card.
4.1.2 Retroactive Eligibility

Medicaid coverage may be assigned retroactively for a client. For claims for an individual who has been approved for Medicaid coverage but has not been assigned a Medicaid client number, the 95-day filing deadline does not begin until the date the notification of eligibility is received from HHSC and added to the TMHP eligibility file.

The date on which the client’s eligibility is added to the TMHP eligibility file is the add date. To ensure 95-day filing deadlines are met and to check current and retroactive client eligibility, providers can log in to My Account and use one of the following resources:

- The TMHP electronic data interchange (EDI)
- The Medicaid Client Portal for Providers

Additionally, providers can call the Automated Inquiry System (AIS) at 1-800-925-9126 to verify a client eligibility.

If a person is not eligible for medical services under Texas Medicaid on the date of service, reimbursement for all care and services provided must be resolved between the provider and the client receiving the services. Providers are not required to accept Medicaid for services provided during the client’s retroactive eligibility period and may continue to bill the client for those services. This guideline does not apply to nursing facilities certified by the Texas Health and Human Services Commission (HHSC).

If it is the provider’s practice not to accept Medicaid for services provided during the client’s retroactive eligibility period, the provider must apply the policy consistently for all clients who receive retroactive eligibility. Providers must inform the client about their policy before rendering services. If providers accept Medicaid assignment for the services provided during the client’s retroactive eligibility period and want to submit a claim for Medicaid-covered services, providers must refund payments received from the client before billing Medicaid for the services.

Examples of Medicaid identification forms are found at the end of this section. Actual Medicaid forms can be identified by a watermark.

---

**Refer to:** Your Texas Benefits Medicaid card (English and Spanish) on the TMHP website at www.tmhp.com.

4.1.3 Expedited Eligibility (Applies to Medicaid-eligible Pregnant Women Throughout the State)

HHSC processes Medicaid applications for pregnant women within 15 business days of receipt. Once eligibility has been certified, a Your Texas Benefits Medicaid card will be issued to verify eligibility and to facilitate provider reimbursement.

4.1.4 Medicaid Buy-In Program for Employed Individuals with Disabilities

The Medicaid Buy-In (MBI) Program allows employed individuals with disabilities to receive Medicaid services by paying a monthly premium. Some MBI participants, based on income requirements, may be determined to have a $0 premium amount and therefore are not required to make a premium payment. Individuals with earnings of less than 250 percent of the federal poverty limits (FPL) may be eligible to participate in the program. Applications for the program are accepted through HHSC’s regular Medicaid application process.
Participants will receive the Your Texas Benefits Medicaid card. MBI participants in urban service areas will be served through Texas Medicaid fee-for-service.

4.1.5 **Newborn Eligibility**

A newborn child may be eligible for Medicaid for up to 1 year if:

- The child’s mother received Medicaid at the time of the child’s birth.
- The child’s mother is eligible for Medicaid or would be eligible if pregnant.
- The child resides in Texas.

If the newborn is eligible for Medicaid coverage, providers must not require a deposit for newborn care from the guardian. The hospital or birthing center must report the birth to HHSC Eligibility Services at the time of the child’s birth.

After the child has been added to the HHSC eligibility file, a Your Texas Benefits Medicaid card will be issued. Newborn clients will receive the Your Texas Benefits Medicaid card within 7 business days of being certified. A temporary Medicaid Eligibility Verification (Form H1027) may be issued by request through a local office or can be downloaded from YourTexasBenefits.com. Form H1027 includes the client’s Medicaid identification number and effective date of coverage.

Providers can verify the client’s add date and retroactive eligibility by:

- Accessing the Medicaid Client Portal for Providers.
- Using TexMedConnect.
- Calling AIS at 1-800-925-9126.

After the newborn becomes a Medicaid client, the website will verify that the client is eligible, even if the card has not been produced yet.

**Note:** Claims submitted for services provided to a newborn eligible for Medicaid must be filed using the newborn client’s Medicaid number. Claims filed with the mother’s Medicaid number cause a delay in reimbursement.

The Medicaid number on the Medicaid Eligibility Verification (Form H1027) may be used to identify newborns eligible for Medicaid.

**Refer to:** Your Texas Benefits Medicaid card (English and Spanish) on the TMHP website at www.tmhp.com.

4.1.6 **Potential Supplemental Security Income (SSI)/Medicaid Eligibility for Premature Infants**

The Supplemental Security Income (SSI) program includes financial and Medicaid benefits for people who are disabled. When determining eligibility for SSI, the Social Security Administration (SSA) must establish that the person meets financial and disability criteria. When determining financial eligibility for a newborn child, SSA does not consider the income and resources of the child’s parents until the month following the month the child leaves the hospital and begins living with the parents. Determinations of disability are made by the state’s Disability Determination Services and may take several months. Federal regulations state that infants with birth weights less than 1,200 grams are considered to meet the SSI disability criteria.

The SSA issued a policy to local SSA offices to make presumptive SSI disability decisions and payments for these children, making it possible for a child to receive SSI and Medicaid benefits while waiting for a final disability determination to be made by Disability Determination Services. The child’s parent or legal guardian must file an SSI application with the SSA. It is in the child’s best interest that the application with the SSA be filed as soon as possible after birth. The SSA accepts a birth certificate with the child’s birth weight or a hospital medical summary as evidence for the presumptive disability decision.
Providers should not change their current newborn referral procedures to HHSC for children who are born to mothers who are eligible for Medicaid as described in this section. However, providers are encouraged to refer parents and guardians of low birth weight newborns to the local SSA office for an SSI application.

### 4.1.7 Foster Care

Most children in the state of Texas foster care program are automatically eligible for Medicaid.

Extended health-care coverage is also available for some former foster care youth clients enrolled in an institution of higher education through the Former Foster Children in Higher Education (FFCHE) program.

To ensure that these children have access to the necessary health-care services for which they are eligible, providers can accept the Medicaid Eligibility Verification (Form H1027) as evidence of Medicaid eligibility. Although this form may not list the client’s Medicaid identification number, it is an official state document that establishes Medicaid eligibility.

Providers should honor the Medicaid Eligibility Verification (Form H1027) as proof of Medicaid eligibility and must bill Texas Medicaid as soon as a Medicaid ID number is assigned. Medicaid ID numbers will be assigned approximately one month from the issue date of the Medicaid Eligibility Verification (Form H1027). The form includes a Department of Family and Protective Services (DFPS) client number that provides an additional means of identification and tracking for children in foster care.

**Note:** The DFPS client number is accepted by Medicaid Vendor Drug Program (VDP)-enrolled pharmacies to obtain outpatient prescribed drug benefits. VDP pharmacies must submit subsequent pharmacy claims with the Medicaid ID number after it has been assigned.

**Reminder:** Adoption agencies/foster parents are not considered third party resources (TPRs). Medicaid is primary in these circumstances.

### 4.1.8 Former Foster Care

HHSC provides Medicaid health-care coverage to former foster care youth who:

- Are 18 through 25 years of age.
- Were in Texas foster care on their 18th birthday or older and were receiving Medicaid when they aged out of Texas foster care.
- Are U.S. citizens or have a qualified alien status (i.e., green card).

### 4.1.9 Medicaid Managed Care Eligibility

All clients who are determined to be eligible for Texas Medicaid are first enrolled as fee-for-service clients. Specific client groups within the Texas Medicaid population are eligible for managed care based on criteria such as age, location, and need. A client who is determined to be eligible for Medicaid managed care is enrolled in the appropriate managed care organization (MCO) or dental plan with a separate eligibility date. In most cases, Medicaid managed care enrollment is not retroactive.

**Refer to:** The Medicaid Managed Care Handbook (Vol. 2, Provider Handbooks) for more information about managed care eligibility and enrollment.

### 4.1.10 Eligibility Verification

To verify a client’s Texas Medicaid eligibility, use the following options:

- Verify electronically through TMHP EDI, TexMedConnect, or the Medicaid Client Portal for Providers. Providers must submit one of the following for each client:
  - Medicaid or Children with Special Health Care Needs (CSHCN) Services Program identification number.
• One of the following combinations: Social Security number and last name; Social Security number and date of birth; or last name, first name, and date of birth. Providers can narrow the search by entering the client’s county code or sex.

• Verify the client’s Medicaid eligibility using the Medicaid Eligibility Verification (Form H1027).

• Contact the TMHP Contact Center or AIS at 1-800-925-9126 or 1-512-335-5986.

• Submit a hard-copy list of clients to TMHP. This service is only used for clients with eligibility that is difficult to verify. A charge of $15 per hour plus $0.20 per page, payable to TMHP, applies to this eligibility verification. The list includes names, gender, and dates of birth if the Social Security and Medicaid ID numbers are unavailable. TMHP can check the client’s eligibility manually, verify eligibility, and provide the Medicaid ID numbers. Mail the lists to the following address:

  Texas Medicaid & Healthcare Partnership Contact Center
  12357-A Riata Trace Parkway
  Suite 100
  Austin, TX 78727

  Note: Providers can obtain client eligibility information for a client who is enrolled in a Medicaid managed care organization (MCO) from the MCO’s web page. Providers can also check the MCO’s web page for submission of electronic claims, prior authorization requests, claim appeals and reconsiderations, exchange of clinical data, and other documentation necessary for prior authorization and claim processing.

4.1.11 Advantages of Electronic Eligibility Verification

Verifying eligibility using TexMedConnect or EDI has the following advantages:

• They are available 24-hours a day 7 days a week.

• Providers can submit up to 5000 verification requests per EDI transmission.

• Providers can create client eligibility lists for verification in TexMedConnect. Each client group can contain up to 250 clients. Providers can create up to 100 groups for each National Provider Identifier (NPI).

The eligibility search function through the Medicaid Client Portal for Providers have the following advantages:

• It displays the client’s:
  • Name, Patient Control Number (PCN), and status.
  • Texas Health Steps information (when applicable).
  • Medicaid health plans and their contact information.
  • Main doctor and the doctor’s contact information.
  • Main dental home.
  • Main dental provider.
  • Medicaid benefits.

• It allows providers to search for details about a client’s eligibility by month and year.

• It allows providers to view and print the client’s:
  • Current and previous Medicaid eligibility information.
  • CHIP, CSHCN Services Program, Medicare, and other insurance plans.
  • Medical, dental, vision, and pharmacy benefits and limitations.
• Medicaid Cards

Electronic eligibility responses contain:
• Eligibility restrictions, such as lock-in, emergency, or women’s health.
• Medicare Part A, B, and C eligibility and effective dates
• Other insurance information, including name and address, and effective dates. EDI transactions also indicate the patient relationship to policy holder.

4.1.12 Contract with Outside Parties
The State Medicaid Manual, Chapter 2, “State Organization,” (Section 2080.18) allows states to contract with outside agents to confirm for providers the eligibility of a Medicaid client. Medicaid providers may contract with these agents for eligibility verification with a cost to the provider. The provider remains responsible for adhering to the claims filing instructions in this manual. The provider, not the agent, is responsible for meeting the 95-day filing deadline and other claims submission criteria.

4.2 Medicaid Identification and Verification
Providers are responsible for requesting and verifying current eligibility information by using the methods listed in subsection 4.1.10, “Eligibility Verification” in this handbook or by asking clients for their Your Texas Benefits Medicaid card (electronic or paper copies) or Medicaid Eligibility Verification form (H1027).

Providers can verify client eligibility electronically by:
• Using TexMedConnect.
• Accessing the Medicaid Client Portal for Providers.
• Calling AIS at 1-800-925-9126.

Providers must accept Your Texas Benefits Medicaid cards and Medicaid Eligibility Verification forms (H1027) as valid proof of eligibility. Providers should retain a copy for their records to ensure the client is eligible for Medicaid when the services are provided. Clients should share eligibility information with their providers.

Providers should request additional identification if they are unsure whether the person presenting the form is the person identified on the form.

Providers should check the Eligibility Date to see whether the client has possible retroactive eligibility for previous bills.

Only those clients listed on the Medicaid Eligibility Verification form or the Your Texas Benefits Medicaid card are eligible for Medicaid. If a person insists he or she is eligible for Medicaid but cannot produce a current Your Texas Benefits Medicaid card or Medicaid Eligibility Verification (Form H1027), providers can verify eligibility through the methods listed in subsection 4.1.10, “Eligibility Verification” or by calling the TMHP Contact Center at 1-800-925-9126. Providers must document this verification in their records and treat these clients as if they had presented a Your Texas Benefits Medicaid card (electronic or printed copy of the card) or Medicaid Eligibility Verification (Form H1027).

When a client’s Your Texas Benefits Medicaid card has been lost or stolen, HHSC issues a temporary Medicaid verification Form H1027. Clients can also request a replacement card by visiting YourTexasBenefits.com’s Medicaid Client Portal or by calling the Texas Medicaid Call Transfer line at 1-800-252-8263. The following is a sample of forms:
• Form H1027-A. Medicaid eligibility verification is used to indicate eligibility for clients who receive regular Medicaid coverage.
• **Form H1027-B.** Medicaid Qualified Medicare Beneficiary (MQMB) is issued to clients eligible for MQMB coverage.

• **Form H1027-C.** Qualified Medicare Beneficiary (QMB) is issued to clients who are eligible for QMB coverage only.

• **Form H1027-F.** Temporary Medicaid identification for clients receiving Former Foster Care in Higher Education (FFCHE) health care.

**Refer to:** Subsection 4.9.1, “QMB/MQMB Identification” in this section.

The Medicaid Eligibility Verification (Form H1027) is acceptable as evidence of eligibility during the eligibility period specified unless the form contains limitations that affect the eligibility for the intended service. Providers must accept any of the documents listed above as valid proof of eligibility. If the client is not eligible for medical assistance or certain benefits, the client is treated as a private-pay patient.

**Refer to:** Subsection 4.1.10, “Eligibility Verification” in this section.

Providers can determine whether a client’s eligibility has limitations by:

• Checking TMHP EDI.

• Accessing the Medicaid Client Portal for Providers.

• Using TexMedConnect.

• Calling AIS at 1-800-925-9126.

Clients may be required to use a designated primary care provider or pharmacy. QMB clients will be limited to Medicaid coverage of the Medicare Part A premiums, if any, Medicare Part B premiums, and Medicare coinsurance or deductible according to current payment guidelines.

If the client is identified as eligible and no other limitations of eligibility affect the intended service, proceed with the service. Eligibility during a previous month does not guarantee eligibility for the current month. The Medicaid Eligibility Verification (Form H1027) and the Your Texas Benefits Medicaid card (electronic or paper) are the only documents that are honored as verification of Medicaid eligibility.

**Refer to:** “Section 8: Third Party Liability (TPL)” (Vol. 1, General Information) for TPL information.

In accordance with current federal policy, Texas Medicaid and Texas Medicaid clients cannot be charged for the client’s failure to keep an appointment. Only claims for services provided are considered for payment. Clients may not be billed for the completion of a claim form, even if it is a provider’s office policy.

### 4.3 Restricted Medicaid Coverage

The following sections are about limitations that may appear on the Your Texas Benefits Medicaid card, indicating that the client’s eligibility is restricted to specific services. Unless “LIMITED” appears on the form, the client is not locked into a single provider.

#### 4.3.1 Emergency Only

The word “EMERGENCY” on the form indicates the client is restricted to coverage for an emergency medical condition. “Emergency medical condition” is defined in subsection 4.3.2.2, “Exceptions to Lock-in Status” in this section.

Certification for emergency Medicaid occurs after the services have been provided. The coverage is retroactive and limited to the specific dates that the client was treated for the emergency medical condition.
Clients limited to emergency medical care are not eligible for family planning, THSteps, or Comprehensive Care Program (CCP) benefits. Only services directly related to the emergency or life-threatening situations are covered.

Undocumented aliens and aliens with a nonqualifying entry status are identified for emergency Medicaid eligibility by the classification of type programs (TPs) 30, 31, 32, 33, 34, 35, and 36. Under Texas Medicaid, undocumented aliens are only eligible for emergency medical services, including emergency labor and delivery.

Any service provided after the emergency medical condition is stabilized is not a benefit.

If a client is not eligible for Medicaid and is seeking family planning services, providers may refer the client to one of the clinics listed on the HHSC website at www.healthytexaswomen.org.

### 4.3.2 Client Lock-in Program

Texas Medicaid fee-for-service clients can be required to use a designated primary care provider and/or a primary care pharmacy.

The client is assigned to a designated provider for access to medical benefits and services when one of the following conditions exists:

- The client received duplicative, excessive, contraindicated, or conflicting health-care services, including drugs.
- A review indicates abuse, misuse, or fraudulent actions related to Medicaid benefits and services.

After analysis through the neural network component of the Medicaid Fraud and Abuse Detection System (MFADS), qualified medical personnel validate the initial identification and determine candidates for lock-in status. The validation process includes consideration of medical necessity. For the lock-in status designation, medical necessity is defined as the need for medical services to the amount and frequency established by accepted standards of medical practice for the preservation of health, life, and the prevention of more impairments.

Except for specialist consultations, services rendered to a client by more than one provider for the same or similar condition during the same time frame may not be considered medically necessary.

#### 4.3.2.1 Lock-in Medicaid Identification

Clients with lock-in status receive the Your Texas Benefits Medicaid card with “Lock-in” printed on the card. The designated provider and pharmacy names are printed on the card under the word “Lock-in.”

When a Texas Medicaid fee-for-service client in the Lock-in Program attempts to obtain nonemergency services from someone other than their designated lock-in primary care provider, the provider must do one of the following:

- Verify the lock-in status by:
  - Accessing the Medicaid Client Portal for Providers.
  - Calling AIS at 1-800-925-9126.
- Attempt to contact the client’s designated lock-in primary care provider for a referral. If the provider is unable to obtain a referral, the provider must inform clients that they are financially responsible for the services.

#### 4.3.2.2 Exceptions to Lock-in Status

Lock-in clients may go to any provider for the following services or items:

- Ambulance services
- Anesthesia
• Annual well-woman checkup
• Assistant surgery
• Case management services
• Chiropractic services
• Counseling services provided by a chemical dependency treatment facility
• Eye exams for refractive errors
• Eyeglasses
• Family planning services (regardless of place of service [POS])
• Genetic services
• Hearing aids
• Home health services
• Laboratory services (including interpretations)
• Licensed clinical social worker (LCSW) services
• Licensed professional counselor (LPC) services
• Mental health rehabilitation services
• Intellectual disability/related condition assessment performed by an intellectual or developmental disability (IDD) provider
• Nursing facility services
• Primary home care
• Psychiatric services
• Radiology services (including interpretations)
• School Health and Related Services (SHARS)
• Comprehensive Care Program (CCP)
• THSteps medical and dental services

For referrals or questions, contact:

HHSC
Office of Inspector General
Lock-in Program - MC 1323
PO Box 85200
Austin, TX 78708
1-800-436-6184

If an emergency medical condition occurs, the lock-in restriction does not apply. The term emergency medical condition is defined as a medical condition manifesting itself by acute symptoms of sufficient severity (including severe pain), such that the absence of immediate medical attention could reasonably be expected to result in:

• Placing the client’s health (or, with respect to a pregnant woman, the health of the woman or her unborn child) in serious jeopardy.
• Serious impairment to bodily functions.
• Serious dysfunction of any bodily organ or part.

Important: A provider who sends in an appeal because a claim was denied with explanation of benefits (EOB) 00066 must include the performing provider identifier, not just a name or group provider identifier. Appeals without a performing provider identifier are denied. The NPI of the designated provider must be entered in the appropriate paper or equivalent electronic field for nonemergency inpatient and outpatient claims to be considered for reimbursement.

Note: Only when the designated provider or designated provider representative has given permission for the client to receive nonemergency inpatient and/or outpatient services, including those provided in an emergency room, can the facility use the designated provider’s NPI for billing.

4.3.2.3 Selection of Designated Provider and Pharmacy
Texas Medicaid fee-for-service clients identified for lock-in status can participate in the selection of one primary care provider, primary care pharmacy, or both from a list of participating Medicaid providers. Eligible providers cannot be under administrative action, sanction, or investigation. In general, the designated primary care provider’s specialty is general practice, family practice, or internal medicine. Other specialty providers may be selected on a case-by-case basis. Primary care providers can include, but are not limited to the following:

• A physician
• Physician assistant
• Physician group
• Advanced practice nurse
• Outpatient clinic
• Rural health clinic (RHC)
• Federally qualified health center (FQHC)

If the client does not select a primary care provider or primary care pharmacy, HHSC selects one for the client.

When a candidate for the designated provider is determined, HHSC contacts the provider by letter. The designated provider receives a confirmation letter from HHSC that verifies the name of the client confirming the name of the client, primary care provider or primary care pharmacy, and the effective date of the lock-in arrangement.

4.3.2.4 Pharmacy services
The primary care pharmacy helps the Lock-in Program ensure that prescriptions that are filled for clients with lock-in status are written either by the primary care provider or other health-care providers to whom the primary care provider has referred the client. HHSC has identified by therapeutic class those medications that require additional monitoring. When a medication that requires additional monitoring is prescribed by an emergency room provider, the primary care pharmacy may be reimbursed for dispensing up to 72 hours or three business days of the prescribed dosage, which allows for holidays and weekends. The primary care pharmacy may dispense the remainder of the medication after receiving approval by the primary care provider or the other providers that HHSC deems to be appropriate.

Some circumstances allow a client to be approved to receive medications from a pharmacy other than the primary care pharmacy. A pharmacy override occurs when the Lock-in Program approves an individual client’s request to obtain medication at an alternate pharmacy other than the lock-in pharmacy. The Lock-in Program is notified when the client or pharmacist calls the HHSC-OIG Hotline telephone number at 1-800-436-6184 to request a pharmacy override.
The Lock-in Program staff refers the client to the notification letter titled “What You Need to Know About the Lock-in Program,” which was sent at initial lock-in. This letter explains the pharmacy override process. The client is instructed to have the alternate pharmacy call the Lock-in Program to request the override.

The following are allowable circumstances for pharmacy override approval:

- The recipient moved out of the geographical area (more than 15 miles from the lock-in pharmacy).
- The lock-in pharmacy does not have the prescribed medication, and the medication will remain unavailable for more than two to three days.
- The lock-in pharmacy is closed for the day, and the recipient needs the medication urgently.
- The lock-in pharmacy does not carry the medication and is either unable to order it or unwilling to stock it.
- The lock-in pharmacy no longer wants to be the designated pharmacy for a particular lock-in client.
- The client has valid complaints against the lock-in pharmacy or its staff.

For questions about pharmacy services for clients that are locked into a primary care pharmacy, contact the Lock-in Program by calling the HHSC OIG Hotline at 1-800-436-6184.

### 4.3.2.5 Duration of Lock-in Status

The Lock-in Program duration of lock-in status is the following:

- Initial lock-in status period—minimum of 36 months.
- Second lock-in status period—additional 60 months.
- Third lock-in status period—will be for the duration of eligibility and all subsequent periods of eligibility.
- Clients who have been arrested for, indicted for, convicted of, or admitted to a crime that is related to Medicaid fraud will be assigned lock-in status for the duration of eligibility and subsequent periods of eligibility.

HHSC uses the same time frames for clients with a lock-in status as noted by the word “LIMITED” on the Your Texas Benefits Medicaid card.

Clients are removed from lock-in status at the end of the specified limitation period if their use of medical services no longer meets the criteria for lock-in status. A medical review also may be initiated at the client’s or provider’s request. Clients or providers can reach the Lock-in Program by calling the HHSC OIG Hotline at 1-800-436-6184 to request this review.

Providers may request to no longer serve as a client’s designated provider at any time during the lock-in period by contacting the Lock-in Program by calling the HHSC OIG Hotline at 1-800-436-6184. Providers are asked to serve or refer the client until another arrangement is made. New arrangements are made as quickly as possible.

### 4.3.2.6 Referral to Other Providers

Texas Medicaid fee-for-service clients with a lock-in status may be referred by their designated provider to other providers. For the referred provider to be paid, the provider identifier of the referring designated provider must be in the referring provider field of the claim form. Claims submitted electronically must have the NPI of the referring designated provider in the Referring Provider Field. Providers must consult with their vendor for the location of this field in the electronic claims format.

**Refer to:** Subsection 6.2, “TMHP Electronic Claims Submission” in “Section 6: Claims Filing” (Vol. 1, General Information)
4.3.2.7 Hospital Services

An inpatient hospital claim for a lock-in Medicaid fee-for-service client is considered for reimbursement if the client meets Medicaid eligibility and admission criteria. Hospital admission personnel should check the name of the client’s designated provider using TexMedConnect or the Medicaid Client Portal for Providers and share the designated provider’s name with the admitting physician if the two are different.

Provider claims for nonemergency inpatient services for lock-in Texas Medicaid fee-for-service clients are considered for payment only when the designated provider identifier appears on the claim form as the billing, performing, or referring physician.

Providers can get information about claim reimbursement for lock-in clients by calling the TMHP Contact Center at 1-800-925-9126.

4.3.2.8 Lock-in Status Claims Payment

Payment for services to a lock-in Medicaid client is made to the designated provider only, unless the services result from a designated provider referral or emergency. An automated review process determines if the claim includes the lock-in primary care provider’s provider identifier as the billing, performing, or referring provider. If the lock-in primary care provider’s provider identifier is not indicated on the claim, the claim is not paid. Exceptions to this rule include emergency care and services that are included in subsection 4.3.2.2, “Exceptions to Lock-in Status” in this section. Appeals for denied claims are submitted to TMHP and must include the designated Medicaid provider identifier for reimbursement consideration.

Claims for provider services for Texas Medicaid fee-for-service clients must include the provider identifier for the designated primary care provider as the billing or performing provider or a referral number in the prior authorization number (PAN) field.

4.3.3 Hospice Program

HHSC manages the Hospice Program through provider enrollment contracts with hospice agencies. These agencies must be licensed by the state and Medicare-certified as hospice agencies. Coverage of services follows the amount, duration, and scope of services specified in the Medicare Hospice Program. Hospice pays for services related to the treatment of the client’s terminal illness and for certain physician services (not the treatments).

Medicaid Hospice provides palliative care to all Medicaid-eligible clients (no age restriction) who sign statements electing hospice services and are certified by physicians to have six months or less to live if their terminal illnesses run their normal courses. Hospice care includes medical and support services designed to keep clients comfortable and without pain during the last weeks and months before death.

Texas Medicaid clients who are 21 years of age and older and who elect hospice coverage waive their rights to all other Medicaid services related to their terminal illness. They do not waive their rights to Medicaid services that are unrelated to their terminal illness.

Texas Medicaid clients who are 20 years of age and younger and who elect hospice care are not required to waive their rights to concurrent hospice care and treatment of the terminal illness. They do not waive their rights to Medicaid services that are unrelated to their terminal illness.

Medicare and Medicaid clients must elect both the Medicare and Medicaid Hospice programs.

Concurrent hospice care and treatment services include:

- Services related or unrelated to the client’s terminal illness
- Hospice care (palliative care and medical and support services related to the terminal illness).
Direct policy questions about the hospice program to HHSC at 1-512-438-3161. Direct all other general questions related to the hospice program, such as billing, claims, rate key issues, and authorizations to HHSC at 1-512-438-2200.

HHSC pays the provider for a variety of services under a per diem rate for any particular hospice day in one of the following categories:

- Routine home care
- Continuous home care
- Respite care
- Inpatient care

### 4.3.3.1 Hospice Medicaid Identification

Individuals who elect hospice care are issued a Your Texas Benefits Medicaid card. Hospice status can be verified by using TexMedConnect or the Medicaid Client Portal for Providers. Clients can cancel their election at any time.

### 4.3.3.2 Physician Oversight Services

Physician oversight is defined as “physician supervision of clients under the care of home health agencies or hospices that require complex or multidisciplinary care modalities.” These modalities involve regular physician status review of related laboratory and other studies, communication with other health professionals involved in patient care, integration of new information into medical treatment plans, and adjustment of medical therapy. Medicaid hospice does not reimburse for physician oversight services.

### 4.3.3.3 Medicaid Services Unrelated to the Terminal Illness

When services are unrelated to the Medicaid Hospice client’s terminal illness, Medicaid (TMHP) pays its providers directly. Providers of services that are unrelated to the terminal illness are required to follow Medicaid prior authorization and claims filing deadlines.

Refer to: “Section 5: Fee-for-Service Prior Authorizations” (Vol. 1, General Information) for more information about prior authorizations for Medicaid hospice clients.

“Section 6: Claims Filing” (Vol. 1, General Information) for more information about filing claims for Medicaid Hospice Clients.

### 4.3.4 Presumptive Eligibility

Presumptive eligibility allows qualified hospitals and entities to determine whether an individual can get short-term Medicaid. Clients who have PE receive immediate, short-term Medicaid eligibility while their regular Medicaid application is processed. PE eligibility categories receive full coverage with the exception of pregnant women who receive ambulatory care services only.

#### 4.3.4.1 PE Medicaid Identification

“PE” on a Your Texas Benefits Medicaid card indicates that a client has presumptive eligibility. PE clients may be identified using TexMedConnect or the Medicaid Client Portal for Providers. An individual who is certified for presumptive eligibility receives the regular Your Texas Benefits Medicaid card.

#### 4.3.4.2 Services

Presumptive eligibility provides full coverage for all PE types of assistance with the exception of pregnant women. Pregnant women only receive ambulatory care services. Labor, delivery, inpatient services, and THSteps medical are not covered during the PE period for pregnant women. If the woman is determined to be eligible for regular Medicaid for the same period of time, regular Medicaid coverage overlays the PE period and provides a full range of services. Client eligibility for PE coverage must be determined by
a qualified hospital or qualified entity. Once eligibility has been determined, services may be obtained from any enrolled Medicaid provider. The claims filing procedures for clients who have PE are the same as those for all other Medicaid clients.

There are five client types of assistance (TA) and one client type program (TP) that provide full Medicaid coverage through presumptive eligibility. Services reimbursed under the presumptive eligibility process are fee-for-service only.

The following are eligible for presumptive eligibility:

- **TA74**—Children under 1 year of age presumptive
- **TA75**—Children 1–5 years of age presumptive
- **TA76**—Children 6–18 years of age presumptive
- **TA83**—Former Foster Care Children presumptive
- **TA86**—Parents and caretaker relatives presumptive
- **TP42**—Pregnant women presumptive

The length of coverage depends on several factors:

- If the individual submits an application for regular Medicaid, the PE Medicaid coverage ends the date the state makes a determination for regular Medicaid.
- If the individual does not submit an application for regular Medicaid, the PE coverage ends the last day of the month following the PE determination.

### 4.3.4.3 Qualified Provider Enrollment

To make PE determinations, the provider must be a qualified hospital or qualified entity. A qualified hospital:

- Is a Medicaid provider.
- Notifies HHSC of its intent to make presumptive eligibility determinations.
- Agrees to make presumptive eligibility determinations according to HHSC policies and procedures.
- Can make presumptive eligibility determinations for pregnant women, children, former foster care children, and parents and other caretakers.
- Helps individuals complete and submit online applications for regular Medicaid.
- Helps individuals understand which documents to send to the state to determine whether they qualify for regular Medicaid.
- Has not been disqualified.

A qualified entity meets the same criteria as a qualified hospital, except that a qualified entity:

- Can be a hospital, clinic, school, or other entity.
- May only make presumptive eligibility determinations for pregnant women.

For more information on how to become a qualified hospital or entity, visit [www.TexasPresumptiveEligibility.com](http://www.TexasPresumptiveEligibility.com).

### 4.3.4.4 Process

Qualified hospital or entity staff determine eligibility for PE coverage based on the client’s statement of residency, income, household composition, citizenship or immigration status, and pregnancy (for Pregnant Women Presumptive). The client is certified for PE if their statement indicates that they meet Medicaid eligibility criteria.
If the client chooses to apply for regular Medicaid, qualified hospital or entity staff assist the client in completing the application. The application is then forwarded to HHSC to determine the client’s eligibility for regular Medicaid.

The period of PE begins on the date the qualified hospital or entity makes the determination and ends when HHSC makes the final Medicaid determination.

### 4.4 CHIP Perinatal Program

The Children’s Health Insurance Program (CHIP) Perinatal Program provides CHIP perinatal benefits for 12 months to the unborn children of non-Medicaid-eligible women. This program allows pregnant women who are ineligible for Medicaid because of income or immigration status to receive prenatal care and provides CHIP benefits to the child upon delivery for the duration of the coverage period. Continuous Medicaid coverage for 12 months is provided from birth to CHIP Perinatal newborns whose mothers received Emergency Medicaid for the labor and delivery. The 12 months of continuous Medicaid coverage for the newborn is available only if the mother received Medicaid for labor and delivery.

#### 4.4.1 Program Benefits

CHIP Perinatal benefits are provided by select CHIP health plans throughout the state. Benefits for the unborn child include:

- Up to 20 prenatal visits:
  - First 28 weeks of pregnancy—one visit every four weeks.
  - From 28 to 36 weeks of pregnancy—one visit every two to three weeks.
  - From 36 weeks to delivery—one visit per week.
  - Additional prenatal visits are allowed if they are medically necessary.
- Pharmacy services, limited laboratory testing, assessments, planning services, education, and counseling.
- Prescription drug coverage based on the current CHIP formulary.
- Hospital facility charges and professional services charges related to the delivery. Preterm labor that does not result in a birth and false labor are not covered benefits.

Program benefits after the child is born include:

- Two postpartum visits for the mother.
- Medicaid benefits for the newborn.

#### 4.4.2 Claims

Providers who serve CHIP Perinatal clients must follow the claims filing guidelines in subsection 6.19.1, “CHIP Perinatal Newborn Transfer Hospital Claims” in “Section 6: Claims Filing” (Vol. 1, General Information).

#### 4.4.3 Client Eligibility Verification

The State Medicaid Manual, Chapter 2, “State Organization,” (Section 2080.18) allows states to contract with outside agents to confirm for providers the eligibility of a Medicaid client. Medicaid providers may contract with these agents for eligibility verification with a cost to the provider. The provider remains responsible for adhering to the claims filing instructions in this manual. The provider, not the agent, is responsible for meeting the 95-day filing deadline and other claims submission criteria.
A number is issued for the baby based on the submission of the Emergency Medical Services Certification Form H3038 or CHIP Perinatal - Emergency Medical Services Certification, Form H3038P for the mother’s labor with delivery.

Establishing Medicaid for the newborn requires the submission of the Emergency Medical Services Certification Form H3038 or CHIP Perinatal - Emergency Medical Services Certification, Form H3038P for the mother’s labor with delivery. If Form H3038 or H3038P is not submitted, Medicaid cannot be established for the newborn from the date of birth for 12 continuous months of Medicaid coverage. Once enrolled, clients are identified as type program (TP) 36 for the mother and TP 45 for the newborn.

Establishing Medicaid (and issuance of a Medicaid number) can take up to 45 days after Form H3038 or H3038P is submitted. Medicaid eligibility for the mother and infant can be verified by:

- Using TexMedConnect.
- Accessing the Medicaid Client Portal for Providers.
- Calling AIS at 1-800-925-9126.

For clients enrolled in the CHIP Program, the CHIP health plan assigns a client ID to be used for billing. Providers should contact the CHIP health plan for billing information.

Newborns whose mother received Medicaid including emergency Medicaid are eligible to receive Medicaid benefits beginning at the date of birth and will not be assigned a client ID from the CHIP health plan.

HHSC requires the expectant mother’s provider to fill out the Emergency Medical Services Certification (Form H3038 or H3038P).

The expectant mother will receive this form from HHSC before her due date, along with a letter reminding her to send information about the birth of her child after delivery. The letter will instruct the expectant mother to take the form to her provider, have the provider fill out the form, then mail the form back to HHSC in a preaddressed, postage-paid envelope. In many cases this activity will occur after delivery when the mother is being discharged from the hospital.

Once HHSC receives the completed Emergency Medical Services Certification (Form H3038 or H3038P), Emergency Medicaid coverage will be added for the mother for the period of time identified by the health care provider. The Emergency Medical Services Certification (Form H3038 or H3038P) is the same form currently required to complete Emergency Medicaid certification.

The CHIP perinatal mother whose income is at or below pregnant women’s Medicaid FPL will not be required to fill out a new application or provide new supporting documentation to apply for Emergency Medicaid. HHSC will determine the woman’s eligibility for Emergency Medicaid by using income and other information the mother to-be provided when she was determined to be eligible for CHIP perinatal coverage, as well as information included on the Emergency Medical Services Certification (Form H3038 or H3038P).

If a woman fails to return the completed Emergency Medical Services Certification (Form H3038 or H3038P) within a month after her due date, HHSC will send her another Emergency Medical Services Certification Form H3038 or H3038P with a postage-paid envelope. If the woman fails to submit Emergency Medical Services Certification (Form H3038 or H3038P), and the hospital cannot locate a Type Program 36 for her in the TMHP online provider lookup tool, then the hospital can bill her for facility fees incurred during her stay.
4.4.3.1 Confirming Receipt of Form H3038 or H3038P

Providers who would like to confirm receipt of form H3038 or H3038P can contact MAXIMUS at 1-877-KIDS-NOW (1-877-543-7669), prompt #6 (for reporting changes) after 48 hours from fax submission. If the submission is by regular mail, providers should allow five business days before contacting MAXIMUS. When calling this number, providers should be prepared to provide the following information:

- National Provider Identifier (NPI)
- Provider name
- Name of person calling
- CHIP perinatal case number (Without the case number, MAXIMUS cannot provide confirmation of receipt. Confirmation of receipt cannot be provided based on client name or address.)

Each form H3038 or H3038P should be faxed one at a time, rather than in a batch. It is important that the form be filled out completely and accurately. If the form is not filled out accurately, it will delay processing and MAXIMUS may not be able to confirm receipt after 48 hours from fax submission.

4.4.3.2 Eligibility Verification for Clients Without a Medicaid ID

Providers should first attempt to verify if a Medicaid number has been issued by calling TMHP at 1-800-925-9126 and using the prompt for AIS or speaking to a representative. Providers can verify eligibility using TexMedConnect or the Medicaid Client Portal for Providers. If a provider is unable to locate a Medicaid number for the mother or infant 45 days after form H3038 or H3038P was faxed, the provider can contact the HHSC Central Processing Center (CPC) in one of the following ways:

- By email at CPC@hhsc.state.tx.us
- By telephone at 1-866-291-1258

CPC needs the following information to respond to requests or inquiries. Providers should submit the information only once. All submissions must be sent in a secure manner. If there are multiple inquiries that are over 45 days, providers can submit them together.

Required information includes the following:

- CHIP perinatal case number
- Mother’s name as it appears on her CHIP Perinatal card
- Dates of service
- Date Form H3038 or H3038P was faxed to MAXIMUS
- Baby’s first and last name
- Baby’s date of birth
- Name and telephone number of the person completing the request

CPC will research inquiries and respond to the provider within 10 business days. This time frame is an approximation and may only apply if all information, including complete contact information, is provided and fewer than 25 names were submitted.

4.4.3.3 Mother’s eligibility

For mothers who currently receive CHIP perinatal, have an income at or below the pregnant women’s Medicaid FPL, and receive emergency Medicaid coverage, providers can check eligibility by:

- Accessing the Medicaid Client Portal for Providers.
- Calling AIS at 1-800-925-9126.
4.4.3.4 **Newborn’s eligibility**

For CHIP Perinatal newborns who have a family income at or below pregnant women’s Medicaid limits FPL, providers can obtain eligibility information and the newborn’s PCN by performing an eligibility verification using the following sources:

- Accessing the Medicaid Client Portal for Providers.
- Calling AIS at 1-800-925-9126.

TMHP cannot provide CHIP Perinatal Program eligibility information for the newborn or mother, regardless of the client’s income level. For CHIP Perinatal Program eligibility information, contact the CHIP health plan.

A report of birth remains an important step to ensure timely Medicaid eligibility for the newborn. A birth must be reported to the state through the typical birth registry process (e.g., use of Texas Electronic Registration system [TER]). In TER, the screen containing the Medicaid/CHIP number should continue to be populated with the mother’s alpha-numeric CHIP Perinatal Program number (e.g., J12345678). In addition, a mother can report the birth by calling 1-877-KIDS-NOW (1-877-543-7669).

4.4.4 **Submission of Birth Information to Texas Vital Statistics Unit**

Hospital providers must submit birth registry information to the DSHS Vital Statistics Unit in a timely manner. Once received by the Vital Statistics Unit, birth information is transmitted to the state’s eligibility systems, so a PCN (Medicaid number) can be issued for newborns whose mothers were at or below the pregnant women’s FPL. Hospitals should use the CHIP Perinatal health plan ID to enter the mother’s CHIP perinatal coverage ID number in the Medicaid/CHIP number field on the Texas Electronic Registration (TER) screen. This number will appear as an alpha-numeric combination, starting with a letter followed by eight digits. For example, G12345678.

For more information, go to the HHSC website at [https://dshs.texas.gov/vs/](https://dshs.texas.gov/vs/), or call Texas Vital Statistics at 1-800-452-9115.

4.5 **Medically Needy Program (MNP)**

The MNP with spend down is limited to children 18 years of age and younger and pregnant women.

The MNP provides Medicaid benefits to children (18 years of age and younger) and pregnant women whose income exceeds the eligibility limits under one of the Medical Assistance Only (MAO) programs but is not enough to meet their medical expenses. Coverage is available for services within the amount, duration, and scope of Texas Medicaid. Individuals are considered adults beginning the month following their 19th birthday.

Medicaid benefits, including family planning and THSteps preventive services through the MNP, are available to:

- Pregnant women.
- Children 18 years of age and younger.

MNP provides access to Medicaid benefits. Applications are made through HHSC. HHSC determines eligibility for the appropriate Medicaid program.

If spend down is applicable, HHSC issues a Medical Bills Transmittal (Form H1120) to the MNP applicant that indicates the spend down amount, months of potential coverage (limited to the month of application and any of the three months before the application month that the applicant has unpaid medical bills), and HHSC contact information.
The applicant is responsible for paying the spend down portion of the medical bills. The TMHP Medically Needy Clearinghouse (MNC) determines which bills may be applied to the applicant’s spend down according to state and federal guidelines. No Medicaid coverage may be granted until the spend down is met.

Newborns of mothers who met spend down and received Medically Needy with Spend Down (TP 56) or Medically Needy with Spend Down-Emergency (TP 32) to cover the newborns birth, are eligible for coverage from the date of birth until the month the child turns one. Hospitals and other providers that complete newborn reporting forms should continue to follow the procedures in subsection 3.2.4, “Newborn Care” in the Inpatient and Outpatient Hospital Services Handbook (Vol. 2, Provider Handbooks) for these newborns.

4.5.1 Spend Down Processing

Applicants are instructed to submit their medical bills or completed claim forms for application toward their spend down to TMHP MNC along with the Medical Bills Transmittal/Insurance Information Form H1120. Charges from the bills or completed claim forms are applied in date of service order to the spend down amount, which is met when the accumulated charges equal the spend down amount.

Providers can assist medically needy clients with their applications by giving them current, itemized statements or completed claim forms to submit to MNC. MNC holds manually completed claim forms used to meet spend down for ten calendar days preceding the completion of the spend down case, then forwards them to claims processing. The prohibition against billing clients does not apply until Medicaid coverage is provided.

Current itemized statements or completed claim forms must include the following:

- Statement date
- Provider name
- Client name
- Date of service
- All services provided and charges
- Current amount due
- Any insurance or client payments with date of payment (the date and amount of any insurance or payments)

**Important:** Amounts used for spend down are deducted from the total billed amount by the provider. Using older bills may provide earlier eligibility for the client.

Bills for past accounts must be current, itemized statements (dated within the last 60 days) that are from the provider and that verify the outstanding status of the account and the current balance due. Accounts that have had payments made by an insurance carrier, including Medicare, must be accompanied by the carrier’s EOB or Remittance Advice and show the specific services covered and amounts paid.

Unpaid bills incurred before the month of potential eligibility (the month with spend down) may be used to meet spend down. Itemized statements must be dated within 60 days of the date they are received at TMHP MNC.

The unpaid balance on currently due accounts may be applied toward the spend down regardless of the date of service. All bills or completed claim forms must be itemized showing the provider’s name, client’s name, dates of service, statement date, services provided, charge for each service, total charges, amounts and dates of payments, and total due.
Clients have 30 days to submit their bills or completed claim forms. Thirty-day extensions are available to the client as necessary to gather all needed information. The provider can assist by furnishing the additional information to the applicant.

All communication about submission of billing information is carried out between MNC and the applicant; however, providers can assist clients by:

- Providing clients with current itemized statements or completed claim forms.
- Encouraging clients to submit all of their medical bills or completed claim forms incurred from all providers at the same time.
- Submitting manual claim forms directly to MNC or to applicants for the MNP, that can be used to meet spend down.

Bills or claim forms submitted to MNC are for application toward the spend down only. Submitting a bill or claim forms for spend down is not a claim for reimbursement. No claims reimbursement is made from such submittals unless the claim form is complete. The provider must file a Medicaid claim after eligibility has been established to have reimbursement considered by Texas Medicaid. If the provider assisted the client with submission of a claim form, the MNC retains all claim forms for ten calendar days preceding the completion of the spend down case. The MNC then forwards all claim forms directly to claims processing to have reimbursement considered by Texas Medicaid.

MNC informs the applicant and HHSC when the spend down is met. HHSC certifies the applicant for Medicaid and sends the Medicaid Identification form to the applicant when Medicaid eligibility is established. The TMHP MNC mails notification letters to providers when clients have met spend down and TMHP has not yet received any claim for the client’s bills. The notification letter states that an invoice was submitted for the spend down and that the provider should submit claims for any bills that fall within the indicated spend down month. Clients are encouraged to inform medical providers of their Medicaid eligibility and make arrangements to pay the charges used to meet the spend down amount. When notified of Medicaid eligibility, the provider asks if the client has retroactive eligibility for previous periods. All bills submitted to MNC are returned to the client, except for claim forms. An automated letter specific to the client’s spend down case is attached, indicating which:

- Bills and charges were used to meet the spend down.
- Bills and charges the client is responsible for paying in part or totally.
- Bills the provider may submit to Medicaid for reimbursement consideration.
- Claims have been received and forwarded to TMHP claims processing.

Providers may inquire about status, months of potential eligibility, Medicaid or case number, and general case information by calling the TMHP Contact Center at 1-800-925-9126.

Medically needy applicants who have a case pending or have not met their spend down are considered private-pay clients and may receive bills and billing information from providers. No claims are filed to Medicaid. A claim that is inadvertently filed is denied because of client ineligibility.

### 4.5.2 Closing an MNP Case

Medically needy cases are closed by MNC for the following reasons:

- Bills were not received within the designated time frame (usually 30 days from the date on which the case is established by the HHSC worker).
- The client failed to provide requested additional case/billing information within 30 days of the MNC request date.
- Insufficient charges were submitted to meet spend down, and the client did not respond to a request for additional charges to be submitted within 30 days of the notification letter.
Charges submitted after the spend down has been met will not reopen the case automatically. The client must call the Client Hotline at 1-800-335-8957.

**Note:** For information regarding the Medically Needy Program for CSHCN Services Program clients refer to the CSHCN Services Program Provider Manual.

### 4.6 Medicaid Buy-in for Children (MBIC) Program

The MBIC program is mandated by S.B. 187, 81st Legislature, Regular Session, 2009, to provide acute care Medicaid coverage for children who are 18 years of age and younger and have disabilities. This program creates a state option for children who are ineligible for Supplemental Security Income (SSI) for reasons other than disability.

Children with disabilities must meet the following requirements to be eligible for MBIC:

- Be 18 years of age or younger.
- Have a family income that is no more than 300 percent of FPL before allowable deductions.
- Meet citizenship, immigration, and residency requirements.
- Be unmarried.
- Not reside in a public institution.

**Exception:** Clients who are enrolled in the MBIC program before they enter a nursing facility or Intermediate Care Facility for Individuals with Intellectual Disabilities (ICF-IID) will continue to receive MBIC benefits until eligibility for the appropriate institutional Medicaid program is determined.

MBIC clients will be enrolled as Medicaid fee-for-service. MBIC clients have access to the same benefits as Medicaid clients who have disabilities. Claims and prior authorization requests for MBIC clients may be submitted according to current guidelines for Medicaid fee-for-service as indicated in this manual.

MBIC benefits are available to enrolled clients through the end of the month that contains their nineteenth birthday. Clients whose birthday falls on the last day of February of a leap year (e.g., February 29, 2004) will be eligible for benefits through the end of March following their nineteenth year.

### 4.7 Healthy Texas Women (HTW) Program

The goal of the HTW program is to expand access to women’s health and family planning services to reduce unintended pregnancies, positively affect the outcome of future pregnancies, and positively impact the health and wellbeing of women and their families in the eligible population.

HTW provides family planning services, related preventive health services that are beneficial to reproductive health and other preventive health services that positively affect maternal health and future pregnancies for women who:

- Are 15 through 44 years of age
  
  **Note:** Women who are 15 through 17 years of age must have a parent or legal guardian apply on their behalf.
- Are a United States citizen or eligible immigrant
- Are a resident of Texas
- Do not currently receive benefits through a Medicaid program (including Medicaid for Pregnant Women), Children’s Health Insurance Program (CHIP), or Medicare Part A or B.
- Have a household income at or below 200 percent of the federal poverty level
• Are not pregnant
• Do not have other insurance that covers the services HTW provides

**Exception:** A client who has other private health insurance may be eligible to receive HTW services if a spouse, parent, or other person would cause physical, emotional, or other harm to the client because the client filed a claim on the health insurance.

**Refer to:** Subsection 2.1, “Guidelines for HTW Providers” in the Healthy Texas Women Program Handbook (Vol. 2, Provider Handbooks).

### 4.8 Medicaid for Breast and Cervical Cancer (MBCC)

The MBCC program provides full Medicaid benefits to women who meet the program’s eligibility requirements. The goal of the program is to improve timely access to breast and cervical cancer treatment for uninsured women who are screened and identified by the National Breast and Cervical Cancer Early Detection Program (NBCCEDP) of the Centers for Disease Control and Prevention (CDC). Additionally, the Texas State Legislature provided funding in 2007 to expand the pool of providers who provide screening and diagnostic services to women so that any provider can diagnose a woman for breast or cervical cancer and she may become eligible for Medicaid through MBCC.

DSHS receives funds from the CDC and awards the funds to providers across the state to perform breast and cervical cancer screenings and diagnostic services under the Breast and Cervical Cancer Services (BCCS) program.

#### 4.8.1 Initial MBCC Program Enrollment

The woman must be diagnosed and in need of treatment for one of the following biopsy-confirmed breast or cervical cancer diagnoses:

- Grade 3 cervical intraepithelial neoplasia (CIN III)
- Severe cervical dysplasia
- Cervical carcinoma in situ
- Primary cervical cancer
- Ductal carcinoma in situ
- Primary breast cancer

A woman may also be eligible for MBCC if she has a diagnosis of metastatic or recurrent breast or cervical cancer and a need for treatment.

Eligibility is determined by a BCCS contractor, DSHS, and HHSC as follows:

- The BCCS contractor screens the client for eligibility if she has a qualifying diagnosis and, if applicable, helps the woman to complete Form H1034, Medicaid for Breast and Cervical Cancer application. The BCCS contractor reviews and collects all of the required eligibility documentation. The woman cannot apply for MBCC at a local HHSC eligibility office.
- DSHS verifies the client’s qualifying diagnosis and submits Form 1034 to HHSC eligibility staff.
- HHSC Centralized Benefits Services staff makes the final Medicaid eligibility determination.

**Refer to:** The Breast and Cervical Cancer Treatment Information page of the DSHS website for more information about the enrollment process.
4.8.2 MBCC Program Eligibility

To be eligible for MBCC, a woman must be:

- 64 years of age or younger and have been screened for breast or cervical cancer and found to need treatment for either breast or cervical cancer.
- A U.S. citizen or eligible immigrant.
- Uninsured or otherwise not eligible for Medicaid.
- A resident of Texas.

A woman who is eligible to receive Texas Medicaid under MBCC receives full Medicaid benefits beginning the day after she received a qualifying diagnosis and for the duration of her cancer treatment. Services are not limited to the treatment of breast and cervical cancer.

4.8.3 Continued MBCC Program Eligibility

After a woman is enrolled in the MBCC program, eligibility may continue if she meets one of the following criteria:

- She is being treated for active disease as defined above,
- She has completed active treatment while in MBCC and is currently receiving hormonal treatment,
- She has completed active treatment while in MBCC and is currently receiving active disease surveillance for TNRBC.

A woman may continue to receive Medicaid benefits as long as she meets the eligibility criteria and provides proof that she is receiving active treatment for breast or cervical cancer. Women who are no longer in MBCC may reapply if they are diagnosed with a new breast or cervical cancer or a metastatic or recurrent breast or cervical cancer.

Note: Active disease surveillance (for the purposes of determining eligibility for MBCC) is periodically monitoring disease progression in order to quickly treat cancerous and precancerous conditions that arise from the presence of a previously diagnosed TNRBC.

If the client’s cancer is in remission and the physician determines that the client requires only routine health screening for a breast or cervical condition (e.g. annual breast examinations, mammograms, and Pap tests as recommended by the American Cancer Society and the U.S. Preventative Services Task Force), the client is not considered to be receiving treatment; and MBCC coverage will not be renewed. A client who is subsequently diagnosed with a new, metastatic, or recurrent breast or cervical cancer may reapply for MBCC benefits.

4.9 Medicare and Medicaid Dual Eligibility

Medicaid clients who are also eligible for Medicare Part A (inpatient coverage), Part B (medical coverage), or Part C (noncontracted Medicare Advantage Plans [MAPs]), may be covered by Texas Medicaid as follows:

- QMB clients are eligible for coinsurance and deductible payments according to the current payment guidelines.
- MQMB clients are eligible for coinsurance and deductible payments according to the current payment guidelines, and receive Medicaid benefits for services that are not a benefit of Medicare or exceed Medicare benefit limitations.

Medicare Part A and Part C (Noncontracted MAPs Only)

For QMB and MQMB clients who are eligible for Medicare Part A, including clients enrolled in MAPs, claims may be reimbursed to providers for the client’s Medicare coinsurance and deductible up to the Medicaid allowed amount for the service less the amount paid by Medicare.
For Medicare Part C, the coinsurance and deductible payment guidelines apply for noncontracted MAPs only.

**Medicare Part B**

For QMB and MQMB clients who are eligible for Medicare Part B, Texas Medicaid reimburses the lesser of the following to providers:

- The coinsurance and deductible payment.
- The amount remaining after the Medicare payment amount is subtracted from the allowed Medicaid fee or encounter rate for the service (If this amount is less than the deductible, then the full deductible is reimbursed instead.)

If the Medicare payment is equal to, or exceeds the Medicaid allowed amount or encounter payment for the service, Texas Medicaid does not make a payment for coinsurance.

**Note:** If the Medicare payment is equal to or exceeds the Medicaid allowed amount or encounter payment for the service, no additional payment is made for coinsurance and deductible.

QMB clients are not eligible for Medicaid coverage for benefits that are not covered by Medicare, and QMB clients are not eligible for THSteps or CCP Medicaid benefits.

QMB and MQMB coverage guidelines do not impact clients who are living in nursing facilities and who receive a vendor rate for client care through HHSC.

Claims for Medicare copayments can also be submitted to TMHP.

**Refer to:** Subsection 2.7.4, “Exceptions” in “Section 2: Texas Medicaid Fee-for-Service Reimbursement” (Vol. 1, General Information) for information about exceptions for Medicare Part B and Part C (noncontracted MAPs).


for more information about filing claims for MQMBs and QMBs.

**4.9.1 QMB/MQMB Identification**

The term “QMB” or “MQMB” on TexMedConnect indicates the client is a Qualified Medicare Beneficiary or a Medicaid Qualified Medicare Beneficiary. The Medicare Catastrophic Coverage Act of 1988 requires Medicare premiums, deductibles, and coinsurance payments to be paid for individuals determined to be QMBs or MQMBs who are enrolled in Medicare Part A and meet certain eligibility criteria (see 1 TAC §§358.201 and 358.202).

**Refer to:** Your Texas Benefits Medicaid card (English and Spanish) on the TMHP website at www.tmhp.com.

**4.9.2 Medicare Part B Crossovers**

The following qualify as Medicare Part B crossover claims: QMB, MQMB, and client TPs 13 or 14, with base plan 10, and category R.

If the provider has not accepted Medicare assignment, the provider may receive payment of the Medicare deductible or coinsurance according to current guidelines on behalf of the QMB, MQMB, client TPs 13 or 14, base plan 10, and category R client. If the provider has collected money from the client and also received reimbursement from TMHP, the provider is required to refund the client’s money.

The Social Security Act requires that Medicaid payment for physician services under Medicare Part B be made on an assignment-related basis.
If Medicaid does not reimburse all or a part of the deductible or coinsurance, the provider is not allowed to bill the client.

Refer to: Subsection 2.7, “Medicare Crossover Claim Reimbursement” in “Section 2: Texas Medicaid Fee-for-Service Reimbursement” (Vol. 1, General Information).

4.9.3 Clients Without QMB or MQMB Status

Medicare is primary to Medicaid, and providers must bill Medicare first for their claims. Medicaid’s responsibility for coinsurance and deductibles is determined in accordance with the Medicaid benefits and limitations including the 30-day spell of illness. TMHP denies claims if the client’s coverage reflects Medicare Part A coverage and Medicare has not been billed first.

Providers must check the client’s Medicare card for Part A coverage before billing Texas Medicaid.

Refer to: Subsection 2.7, “Medicare Crossover Claim Reimbursement” in “Section 2: Texas Medicaid Fee-for-Service Reimbursement” (Vol. 1, General Information).

4.9.4 Medicare Part C

Providers can verify a client’s Medicare Part C eligibility by:

- Checking TMHP EDI.
- Accessing the Medicaid Client Portal for Providers.
- Using TexMedConnect.

In response to an eligibility inquiry, providers receive the client’s Medicare Part C eligibility effective date, end date, and add date.

HHSC contracts with some Medicare Advantage Plans (MAPs) and offers a per-client-per-month payment. The payment to the MAP includes all costs associated with the Medicaid cost sharing for dual-eligible clients. MAPs that contract with HHSC will reimburse providers directly for the cost sharing obligations that are attributable to dual-eligible clients enrolled in the MAP. These payments are included in the capitated rate paid to the HMO and must not be billed to TMHP or a Medicaid client.

TMHP now processes certain claims for clients enrolled in a Medicare Advantage Plan (Part C).

Refer to: Subsection 6.12, “Filing Medicare Primary Claims” in “Section 6: Claims Filing” (Vol. 1, General Information).

A list of MAPs that have contracted with HHSC is available in the “EDI” section of the TMHP website at www.tmhp.com. The list will be updated as additional plans initiate contracts.

4.9.5 Medicare Part D

Providers can verify a client’s Medicare Part D eligibility by accessing the Medicaid Client Portal for Providers.

4.10 Health Insurance Premium Payment (HIPP) Program

The HIPP Program reimburses for the cost of medical insurance premiums. A Medicaid client is eligible for the HIPP Program when Medicaid finds it more cost effective to reimburse a Medicaid client’s group health insurance premiums than to reimburse his or her medical bills directly through Medicaid.

By ensuring access to employer sponsored health insurance, individuals who are eligible for the HIPP Program may receive services that are not normally covered through Medicaid. Also, members of the family who are not eligible for Medicaid may be eligible for the HIPP Program.
Providers can benefit from this program by helping the uninsured population, saving money for the state of Texas, and receiving a higher payment from the group health insurance carrier. Providers can increase HIPP Program enrollment by displaying brochures to educate their Medicaid clients about the program.

For more information, call the TMHP-HIPP Program at 1-800-440-0493 or visit https://hhs.texas.gov/services/financial/health-insurance-premium-payment-hipp-program.

### 4.11 Long-Term Care Providers

A nursing facility, home health services provider, or any other similar long-term care services provider that is Medicare-certified must:

- Seek reimbursement from Medicare before billing Texas Medicaid for services provided to an individual who is eligible to receive similar services under the Medicare program.
- Appeal Medicare claim denials for payment, as directed by the department.

A nursing facility, home health services provider, or any other similar long-term care services provider that is Medicare-certified is not required to seek reimbursement from Medicare before billing Texas Medicaid for a person who is Medicare-eligible and has been determined to not be homebound.

### 4.12 State Supported Living Centers

Inpatient hospital care for individuals who are eligible for Supplemental Security Income (SSI) Medicaid and reside in a State Supported Living Center (SSLC) must be billed to TMHP. Medicaid providers who render off-campus acute care services to Medicaid-eligible SSLC residents are also required to submit claims directly to Medicaid. This is applicable only to residents of the SSLCs operated by HHSC.

Claims and prior authorization requests for acute care services that are rendered to these clients must be submitted directly to Medicaid.

Providers may contact HHSC for assistance or information about billing procedures for state school services.

### 4.13 Forms

The following linked forms can also be found on the [Forms](#) page of the Provider section of the TMHP website at [www.tmhp.com](http://www.tmhp.com):

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<tr>
<td>Other Insurance Form</td>
</tr>
<tr>
<td>Authorization for Use and Release of Health Information</td>
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SECTION 5: FEE-FOR-SERVICE PRIOR AUTHORIZATIONS

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5.1 **General Information About Prior Authorization**

Some fee-for-service Medicaid services require prior authorization as a condition for reimbursement. Information about whether a service requires prior authorization, as well as prior authorization criteria, guidelines, and timelines for the service, is contained in the handbook within Volume 2 that contains the service.

Prior authorization is not a guarantee of payment. Even if a procedure has been prior authorized, reimbursement can be affected for a variety of reasons, e.g., the client is ineligible on the date of service (DOS) or the claim is incomplete. Providers must verify client eligibility status before providing services.

In most instances prior authorization must be approved before the service is provided. Prior Authorization for urgent and emergency services that are provided after business hours, on a weekend, or on a holiday may be requested on the next business day. TMHP considers providers’ business hours as Monday through Friday, from 8 a.m. to 5 p.m., Central Time. Prior authorization requests that do not meet these deadlines may be denied.

To avoid unnecessary denials, the request for prior authorization must contain correct and complete information, including documentation of medical necessity. The documentation of medical necessity must be maintained in the client’s medical record. The requesting provider may be asked for additional information to clarify or complete a request for prior authorization.

Prior authorization requests may be submitted to the TMHP Prior Authorization Department by mail, fax, or the electronic portal. Prescribing or ordering providers, dispensing providers, clients’ responsible adults, and clients may sign prior authorization forms and supporting documentation using electronic or wet signatures.

**Refer to:** Subsection 6.1.4, “Claims Filing Deadlines” in “Section 6: Claims Filing” (Vol. 1, General Information) for the TMHP-approved holidays.

**Note:** Authorization requests for services administered by a client’s managed care organization (MCO) or dental plan must be submitted to the client’s MCO or dental plan according to the guidelines that are specific to the plan under which the client is covered.

**Refer to:** The Medicaid Managed Care Handbook (Vol. 2, Provider Handbooks) for additional information about managed care prior authorizations.

5.1.1 **Prior Authorization Requests for Clients with Retroactive Eligibility**

Retroactive eligibility occurs when the effective date of a client’s Medicaid coverage is before the date the client’s Medicaid eligibility is added to TMHP’s eligibility file, which is called the “add date.”

For clients with retroactive eligibility, prior authorization requests must be submitted after the client’s add date and before a claim is submitted to TMHP.

When an authorization request is submitted for a client who has received retroactive Texas Medicaid eligibility, providers should notify TMHP to avoid potential delays. Providers can notify TMHP of the retroactive client eligibility in one of the following ways:

- Add a comment in the additional comments field for authorization requests that are submitted online on the TMHP website at www.tmhp.com or on the eviCore website at www.evicore.com (for radiological imaging authorizations only).

- Add a comment on the cover sheet or the authorization request form for authorizations that are faxed to TMHP or eviCore (for radiological imaging authorizations only).

If the authorization request is made by telephone, the caller can indicate to the representative at TMHP or eviCore (for radiological imaging authorizations only) that the client has retroactive Texas Medicaid eligibility.
For services provided to fee-for-service Medicaid clients during the client’s retroactive eligibility period, i.e., the period from the effective date to the add date, prior authorization must be obtained within 95 days from the client’s add date and before a claim for those services is submitted to TMHP. For services provided on or after the client’s add date, the provider must obtain prior authorization within 3 business days of the date of service.

The provider is responsible for verifying the client’s eligibility. Providers are strongly encouraged to verify the client’s eligibility frequently. Providers can verify eligibility by:

- Accessing the Medicaid Client Portal for Providers.
- Using TexMedConnect.
- Calling the Automated Inquiry System (AIS) at 1-800-925-9126.

If services are discontinued before the client’s add date, the provider must still obtain prior authorization within 95 days of the add date to be able to submit claims.

Refer to: “Section 4: Client Eligibility” (Vol. 1, General Information).

5.1.2 Prior Authorization Requests for Newly Enrolled Providers

TMHP cannot issue a prior authorization before Medicaid enrollment is complete. Upon notice of Medicaid enrollment, by way of issuance of a provider identifier, the provider must contact the appropriate TMHP Authorization Department to request prior approval before providing services that require prior authorization. Regular prior authorization procedures are followed after the TMHP Prior Authorization Department has been contacted.

Retroactive authorizations are not issued unless the regular authorization procedures for the requested services allow for authorizations to be obtained after services are provided. Providers should refer to specific handbook sections for details about authorization requirements, claims filing, and timeframe guidelines for authorization request submissions. Retroactive authorizations may be granted according to the timeframe guidelines for the specific service requested, and do not exceed those timeframes.

Note: All claims must adhere to the claims filing deadlines as outlined in this manual. Retroactive authorizations cannot exceed the claims filing deadline, and are not issued if the date of services is more than 95 days from the date the new provider identifier is issued as identified by the add date.

Refer to: “Section 1: Provider Enrollment and Responsibilities” (Vol. 1, General Information).

5.1.3 Prior Authorization for Services Rendered Out-of-State

Texas Medicaid covers medical assistance services that are provided to eligible Texas Medicaid clients while they are in a state other than Texas; however, clients are not covered if they leave Texas to receive out-of-state medical care that can be received in Texas. Services that are provided outside of the state are covered by Texas Medicaid to the same extent that medical assistance is furnished and covered in Texas when the service meets one or more requirements of Texas Administrative Code (TAC) Title 1 §352.17.

Note: Border state providers (providers that render services within 50 miles of the Texas border) are considered in-state providers for Texas Medicaid.

Services that are rendered outside of the state must be prior authorized by Texas Medicaid, and TMHP must receive claims from out-of-state providers within 365 days of the date of service. Out-of-state providers that seek reimbursement for services that are rendered outside of the state must submit a Texas Medicaid Provider Enrollment application and be approved for enrollment in Texas Medicaid.

Transplant services that are provided out-of-state but available in Texas will not be reimbursed by Texas Medicaid. When requesting an out-of-state prior authorization for a pre-transplant evaluation, the provider must submit a copy of the transplant evaluation performed by a Texas facility to support the need for an out-of-state pre-transplant evaluation.
Medical assistance and transplant services that are provided to eligible Texas Medicaid clients must meet the criteria included in subsection 1.10, "Enrollment Criteria for Out-of-State Providers" in "Section 1: Provider Enrollment and Responsibilities" (Vol. 1, General Information). If services are rendered to eligible Texas Medicaid clients that do not meet the criteria, the services are not a benefit of Texas Medicaid and will not be considered for reimbursement.

**Refer to:** Subsection 1.10, “Enrollment Criteria for Out-of-State Providers” in “Section 1: Provider Enrollment and Responsibilities” (Vol. 1, General Information).

Subsection 2.6, "Out-of-State Medicaid Providers" in “Section 2: Texas Medicaid Fee-for-Service Reimbursement” (Vol. 1, General Information).

### 5.1.4 Prior Authorization Requests for Clients with Private Insurance

If a client’s primary coverage is private insurance and Medicaid is secondary but prior authorization is required for Medicaid reimbursement, providers must follow the guidelines and requirements listed in the handbook for that service.

### 5.1.5 Prior Authorization Requests for Clients with Medicare/Medicaid

If a client’s primary coverage is Medicare, providers must always confirm with Medicare whether a service is a Medicare benefit for the client.

If a service that requires prior authorization from Medicaid is a Medicare benefit and Medicare approves the service, prior authorization from TMHP is not required for reimbursement of the coinsurance or deductible. If Medicare denies the service, then prior authorization is required. TMHP must receive a prior authorization request within 30 days of the date of Medicare’s final disposition. The Medicare Remittance Advice and Notification (MRAN) that contains Medicare’s final disposition must accompany the prior authorization request.

If a service requires prior authorization through Medicaid and the service is not a benefit of Medicare, providers may request prior authorization from TMHP before receiving the denial from Medicare.

**Note:** Refer to the appropriate handbooks in this manual for additional prior authorization guidelines for clients with dual eligibility.

### 5.1.6 Prior Authorizations for Personal Care Services (PCS)

Before sending a prior authorization request for personal care services to TMHP, the Texas Department of State Health Services (DSHS) will fax the communication tool to the provider. The provider must verify that the information listed on the tool is accurate. If any information on the communication tool is inaccurate, the provider must call the DSHS case manager listed on the tool within three business days of receipt to explain the inaccuracy. The DSHS case manager will correct the communication tool and will fax the updated tool to the provider. The provider must review the updated communication tool and call the DSHS case manager if any inaccuracies remain.

If the provider does not contact the DSHS case manager within three business days of receipt of the communication tool, the case manager will send a prior authorization request to TMHP to have the authorization issued with the information provided on the communication tool.

**Important:** If a provider fails to notify the DSHS case manager of inaccurate information within three business days of receipt of the communication tool, HHSC will not consider making changes to authorizations for past dates of service.
It is the PCS provider’s responsibility to know the prior authorization period for each client who has an open authorization and to ensure that, before the authorization expires, a DSHS case manager has conducted a reassessment and extended the authorization through TMHP. If a provider has not received an updated provider notification letter from TMHP within 30 days of the authorization’s expiration date, the provider should do one of the following:

- Call the TMHP PCS Prior Authorization Inquiry Line at 1-888-648-1517 and ask whether an authorization is in process.
- Call the TMHP PCS Client Line at 1-888-276-0702, Option 2, and ask for a referral to have DSHS conduct a reassessment.
- Call the DSHS regional office, and notify the DSHS case manager that a new authorization has not been received.

Clients can experience a gap in service if an authorization is not updated before it expires. Providers will not be reimbursed for services provided after an authorization has expired and before a new authorization has been issued.

Providers must retain current client information on file.

### 5.1.6.1 Authorizations for Multiple PCS Clients Within the Same Household

DSHS case managers synchronize PCS authorizations within households that have multiple clients who are receiving PCS.

Synchronization of authorizations within households are made as PCS reassessments come due. When clients are due for reassessment, the DSHS case manager assess all eligible clients in the home and submit authorizations for all eligible clients in the household for the same 52-week authorization period. Some authorizations within a household may be shortened or closed and then reinstated to be in alignment with other clients in the same household. DSHS case managers communicate with the provider about the actions that are being taken using the existing Communication Tool.

**Note:** There should be no lapse in services to the client.

### 5.1.6.2 Verifying the Texas Provider Identifier (TPI) on PCS Authorizations

When an authorization notification letter is received by a PCS provider, the provider should verify that the correct TPI was used on the prior authorization for the PCS client. Providers must verify that the TPI on the prior authorization is correct for the location at which the client is receiving services.

Providers who provide services through the Agency option or the Consumer Directed Services (CDS) option must ensure that the TPI on the prior authorization is accurate for the option the client is using. If a provider discovers that the TPI used on the prior authorization is incorrect, the provider should contact the DSHS case manager and ask for the correct TPI to be submitted to TMHP.

### 5.1.7 Prior Authorization for Outpatient Self-Administered Prescription Drugs

**Refer to:** Subsection 1.1, “About the Vendor Drug Program” in the *Outpatient Drug Services Handbook (Vol. 2, Provider Handbooks)* for information about this program.

### 5.1.8 Prior Authorization for Nonemergency Ambulance Transport

According to 1 TAC §354.1111, nonemergency transport is defined as ambulance transport provided for a Medicaid client to or from a scheduled medical appointment, to or from a licensed facility for treatment, or to the client’s home after discharge from a hospital when the client has a medical condition such that the use of an ambulance is the only appropriate means of transportation (i.e., alternate means of transportation are medically contraindicated).

**Refer to:** The *Ambulance Services Handbook (Vol. 2, Provider Handbooks)* for more information about ambulance services.
According to Human Resource Code (HRC) §32.024 (t), a Medicaid-enrolled physician, nursing facility, health-care provider, or other responsible party is required to obtain authorization before an ambulance is used to transport a client in circumstances not involving an emergency.

HRC states that a provider of nonemergency ambulance transport is entitled to payment from the nursing facility, health-care provider, or other responsible party that requested the service if payment under the Medical Assistance Program is denied because of lack of prior authorization and the ambulance provider submits a copy of the claim for which payment was denied.

Refer to: Subsection 5.1.8.1, “Appealing Non-Emergent Ambulance Claims Denied for Missing Prior Authorization Number” in this section for more information about appeals.

The Medical Transportation Program Handbook (Vol. 2, Provider Handbooks) for more information about the Medical Transportation Program.

TMHP responds to nonemergency transport prior authorization requests within 2 business days of receipt of requests for 60 days or less. Providers should submit all requests for a prior authorization number (PAN) in sufficient time to allow TMHP to issue the PAN before the date of the intended transport.

If the client’s medical condition is not appropriate for transport by ambulance, nonemergency ambulance services are not a benefit. Prior authorization is a condition for reimbursement but is not a guarantee of payment. The client and provider must meet all of the Medicaid requirements, such as client eligibility and claim filing deadlines.

Medicaid providers who participate in one of the Medicaid Managed Care health maintenance organization (HMO) plans must follow the HMO’s prior authorization requirements.

The TMHP Ambulance Unit reviews the prior authorization request to determine whether the client’s medical condition is appropriate for transport by ambulance. Incomplete information may cause the request to be suspended for additional medical information or be denied.

The following information helps TMHP determine the appropriateness of the transport:

- An explanation of the client’s physical condition that establishes the medical necessity for transport. The explanation must clearly state the client’s condition requiring transport by ambulance.
- The necessary equipment, treatment, or personnel to be used during the transport.
- The origination and destination points of the client’s transport.

Prior authorization is required when an extra attendant is needed for any nonemergency transport.

When a client’s condition changes, such as a need for oxygen or additional monitoring during transport, the prior authorization request must be updated.


5.1.8.1 Appealing Non-Emergent Ambulance Claims Denied for Missing Prior Authorization Number

Medicaid fee-for-service ambulance providers can appeal claims that have been denied for a missing prior authorization number that was not provided by the requesting physician, facility, or hospital for nonemergent transport. The ambulance provider must include proper documentation with the appeal.

An ambulance provider is required to maintain documentation that represents the client’s medical condition and other clinical information to substantiate medical necessity, the level of service, and the mode of transportation requested. This supporting documentation is limited to documents developed or maintained by the ambulance provider.
The Appeals Process

The ambulance provider must submit the appeal as soon as it is realized that the requesting provider did not obtain the prior authorization. The appeal must be submitted according to all filing deadlines.

Refer to: “Section 7: Appeals” (Vol. 1, General Information) for information about filing deadlines.

The following documentation must be included with the appeal to support the need for transport:

- Documentation representing the client’s medical condition.
- Other clinical information to support medical necessity.
- Level of service given.
- Mode of transportation requested.
- A copy of the run sheet, which must be signed by the emergency medical technician (EMT) in attendance, and which must also include the name and telephone number of the employee requesting the service for the physician, facility, or hospital.

5.1.8.2 Prior Authorization Types, Definitions

One-Time, Nonrepeating

One-time, nonrepeating requests are reserved for those clients who require a one-time transport. The request must be signed and dated by a physician, physician assistant (PA), nurse practitioner (NP), clinical nurse specialist (CNS), registered nurse (RN), or discharge planner with knowledge of the client’s condition. Without a signature and date, the form is considered incomplete.

Recurring

Recurring requests, up to 60 days, are reserved for those clients whose transportation needs are not anticipated to last longer than 60 days. The request must be signed and dated by a physician, PA, NP, or CNS. Without a signature and date, the form is considered incomplete. The request must include the approximate number of visits needed for the repetitive transport (e.g., dialysis, radiation therapy).


5.1.8.3 Nonemergency Prior Authorization Process

To obtain prior authorization, providers must submit a completed Nonemergency Ambulance Prior Authorization Request Texas Medicaid and CSHCN Services Program form by fax to the TMHP Ambulance Unit at 512-514-4205. Prior authorization can also be requested through the TMHP website at www.tmhp.com.

The Nonemergency Ambulance Prior Authorization Form must not be modified. If the form is altered in any way, the request may be denied. The form must be filled out by the facility or the physician’s staff that is most familiar with the client’s condition. For nonemergency ambulance transportation services rendered to a client, ambulance providers may coordinate the nonemergency ambulance prior authorization request between the requesting provider, which may include a physician, nursing facility, healthcare provider, or other responsible party. Ambulance providers may assist in providing necessary information such as their National Provider Identifier (NPI) number, fax number, and business address to the requesting provider. However, the Non-emergency Ambulance Prior Authorization Request form must be signed, dated, and submitted by the Medicaid-enrolled requesting provider, not the ambulance provider.

Medicaid providers may request prior authorization using one of the following methods:

- The client’s physician, nursing facility, Intermediate Care Facility for Individuals with Intellectual Disabilities (ICF-IID), health-care provider, or other responsible party completes the online prior authorization request on the TMHP website at www.tmhp.com.

- Hospitals may call TMHP at 1-800-540-0694 to request prior authorization Monday through Friday, 7 a.m. to 7 p.m., Central Time. A request may be submitted up to 60 days before the date on which the nonemergency transport will occur.

A request for a one-day transport may be submitted on the next business day following the transport in some circumstances; however, every attempt should be made to obtain prior authorization before the transport takes place. Authorization requests for one day transports submitted beyond the next business day will be denied.

A request for a recurring transport must be submitted before the client is transported by ambulance.

After a prior authorization request has been approved, if the client’s condition deteriorates or the need for equipment changes so that additional procedure codes must be submitted for the transport, the requesting provider must submit a new Nonemergency Ambulance Prior Authorization Request form.

Clients who require a hospital-to-hospital or hospital-to-outpatient medical facility transport are issued a PAN for that transport only.

Refer to: Subsection 4.2.1, “Prior Authorization Requirements” in the Inpatient and Outpatient Hospital Services Handbook (Vol. 2, Provider Handbooks) for more information on nonemergency prior authorization for hospitals.

TMHP reviews all of the documentation it receives. An online prior authorization request submitted through the TMHP website is responded to with an online approval or denial. Alternately, a letter of approval or denial is faxed to the requesting provider. The client is notified by mail if the authorization request is denied or downgraded. Reasons for denial include documentation that does not meet the criteria of a medical condition that is appropriate for transport by ambulance, or the client is not Medicaid-eligible for the dates of services requested. Clients may appeal prior authorization request denials by contacting TMHP Client Notification at 1-800-414-3406. Providers may not appeal prior authorization request denials.

The requesting provider must contact the transporting ambulance provider with the PAN and the dates of service that were approved.

Refer to: Subsection 5.5.1, “Prior Authorization Requests Through the TMHP Website” in this section for additional information, including mandatory documentation requirements and retention.

Providers are not required to fax medical documentation to TMHP; however, in certain circumstances, TMHP may request that the hospital fax the supporting documentation. Incomplete online or faxed request forms are not considered a valid authorization request and are denied.

A nonemergency transport will be denied when a claim is submitted with a Nonemergency Ambulance Prior Authorization Request Texas Medicaid and CSHCN Services Program form that is completed and signed after the service is rendered. In addition, a Nonemergency Ambulance Prior Authorization Request Texas Medicaid and CSHCN Services Program form that is completed and signed after the service is rendered will not be accepted on appeal of the denial.

The hospital must maintain documentation of medical necessity, including a copy of the authorization from TMHP in the client’s medical record for any item or service that requires prior authorization. The services provided must be clearly documented in the medical record with all pertinent information regarding the client’s condition to substantiate the need and medical necessity for the services.
5.1.8.4 Nonemergency Ambulance Exception Request

Clients whose physician has documented a debilitating condition and who require recurring trips that will extend longer than 60 days may qualify for an exception to the 60-day prior authorization request.

To request an exception, providers must submit all of the following documentation:

- A completed Nonemergency Ambulance Exception form that is signed and dated by a physician. Without a physician’s signature and date, the form is considered incomplete.

- Medical records that support the client’s debilitating condition, which may include, but is not limited to:
  - Discharge information.
  - Diagnostic images (e.g., MRI, CT, X-rays)
  - Care plan.

  **Note:** Documentation submitted with statements similar to “client has a debilitating condition” are insufficient.

5.1.8.5 Documentation of Medical Necessity and Run Sheets

5.1.8.5.1 Documentation of Medical Necessity

Retrospective review may be performed to ensure documentation supports the medical necessity of the transport.

Documentation to support medical necessity must include one of the following:

- The client is bed-confined before, during and after the trip and alternate means of transport is medically contraindicated and would endanger the client’s health (i.e., injury, surgery, or the use of respiratory equipment). The functional, physical, and mental limitations that have rendered the client bed-confined must be documented.

  **Note:** Bed-confined is defined as a client who is unable to stand, ambulate, and sit in a chair or wheelchair.

- The client’s medical or mental health condition is such that alternate means of the transport is medically contraindicated and would endanger the client’s health (e.g., injury, surgery, or the use of respiratory equipment).

- The client is a direct threat to himself or herself or others, which requires the use of restraints (chemical or physical) or trained medical personnel during transport for client and staff safety (e.g., suicidal).

When physical restraints are needed, documentation must include, but is not limited to, the following:

- Type of restraint
- Time frame of the use of the restraint
- Client’s condition

  **Note:** The standard straps used in an ambulance transport are not considered a restraint.

5.1.8.5.2 Run Sheets

The run sheet is used as a medical record for ambulance services and may serve as a legal document to verify the care that was provided, if necessary. The ambulance provider does not have to submit the run sheet with the claim.
The ambulance provider must have documentation to support the claim. Without documentation that would establish the medical necessity of a nonemergency ambulance transport, the transport may not be covered by Texas Medicaid.

The ambulance provider may decline the transport if the client’s medical or mental health condition does not meet the medical necessity requirements.

It is the responsibility of the ambulance provider to maintain (and furnish to Texas Medicaid upon request) concise and accurate documentation. The run sheet must include the client’s physical assessment that explains why the client requires ambulance transportation and cannot be safely transported by an alternate means of transport.

Coverage will not be allowed if the trip record does not contain a sufficient description of the client’s condition at the time of the transfer for Texas Medicaid to reasonably determine that other means of transportation are contraindicated. Coverage will not be allowed if the description of the client’s condition is limited to statements or opinions such as the following:

- “Patient is nonambulatory.”
- “Patient moved by drawsheet.”
- “Patient could only be moved by stretcher.”
- “Patient is bed-confined.”
- “Patient is unable to sit, stand or walk.”

The run sheet should “paint a picture” of the client’s condition and must be consistent with documentation found in other supporting medical record documentation (including the nonemergency prior authorization request.)

5.1.8.6 Nonemergency Prior Authorization and Retroactive Eligibility

Retroactive eligibility occurs when the effective date of a client’s Medicaid coverage is before the eligibility “add date,” which is the date the client’s Medicaid eligibility is added to TMHP’s eligibility file.

For clients with retroactive eligibility, prior authorization requests must be submitted after the client’s add date and before a claim is submitted to TMHP.

For services that are provided to fee-for-service Medicaid clients during a client’s retroactive eligibility period (i.e., the period from the effective date to the add date), providers must obtain prior authorization within 95 days of the client’s add date and before submitting a claim for those services to TMHP.

For services provided on or after the client’s add date, the provider must obtain prior authorization within three business days of the date of service.

The provider is responsible for verifying the client’s eligibility. Providers are strongly encouraged to verify the client’s eligibility frequently. Providers can verify eligibility by:

- Checking the TMHP electronic data interchange (EDI).
- Accessing the Texas Medicaid Client Portal for Providers.
- Using TexMedConnect.
- Calling the Automated Inquiry System (AIS) at 1-800-925-9126.

If services are discontinued before the client’s add date, the provider must still obtain prior authorization within 95 days of the add date to be able to submit claims.

If a client’s Medicaid eligibility is pending, a PAN must be requested before a nonemergency transport. Initially this request will be denied for Medicaid eligibility. When Medicaid eligibility is established, the requestor has 95 days from the date on which the eligibility was added to TMHP’s files to contact the TMHP Ambulance Unit and request that authorization be considered.
To inquire about Medicaid eligibility, providers can contact AIS at 1-800-925-9126.

5.1.9 Nonemergency Transport Authorization for Medicare and Medicaid Clients

Providers should simultaneously request prior authorization for the nonemergency transport from TMHP for an Medicaid Qualified Medicare Beneficiary (MQMB) client in the event the service requested is denied by Medicare as a non-covered service.

*Note:* Qualified Medicare Beneficiary (QMB) clients are not eligible for Medicaid benefits. Providers can contact Medicare for the Medicare prior authorization guidelines.

5.2 Authorization Requirements for Unlisted Procedure Codes

Providers have the option to obtain prior authorization before rendering the service if all of the required information is available. When requesting a fee-for-service prior authorization for an unlisted procedure code, providers must submit the following information with the prior authorization request:

- Client’s diagnosis.
- Medical records that show the prior treatment for this diagnosis and the medical necessity of the requested procedure.
- A clear, concise description of the procedure to be performed.
- Reason for recommending this particular procedure
- A procedure code that is comparable to the procedure being requested.
- Documentation that this procedure is not investigational or experimental.
- Place of service in which the procedure is to be performed.
- The physician’s intended fee for this procedure including the manufacturer’s suggested retail price (MSRP) or other payment documentation.

If any of this information is unavailable at the time the prior authorization is requested, the request will be returned as incomplete; however, this is not a denial of reimbursement. If the required information becomes available before the service is performed, the prior authorization request can be resubmitted, or the required medical necessity and payment documentation can be submitted with the claim after the service is provided to be considered for reimbursement.

The prior authorization number must appear on the claim when it is submitted to TMHP. Claims submitted without the appropriate prior authorization will be denied.

5.3 Benefit Code

A benefit code is an additional data element that identifies a state program.

Providers that participate in the following programs must use the associated benefit code when they submit prior authorizations:

<table>
<thead>
<tr>
<th>Program</th>
<th>Benefit Code</th>
</tr>
</thead>
<tbody>
<tr>
<td>Comprehensive Care Program (CCP)</td>
<td>CCP</td>
</tr>
<tr>
<td>Texas Health Steps (THSteps) Medical</td>
<td>EP1</td>
</tr>
<tr>
<td>THSteps Dental</td>
<td>DE1</td>
</tr>
<tr>
<td>Family Planning Agencies*</td>
<td>FP3</td>
</tr>
<tr>
<td>Hearing Aid Dispensers</td>
<td>HA1</td>
</tr>
</tbody>
</table>

*Agencies only—Benefit codes should not be used for individual family planning providers.
5.4 Submitting Prior Authorization Forms

A physician or health care provider may submit a medical prior authorization recertification request up to 60 days before the expiration of the current authorization of services. Once submitted, the recertification request will be reviewed and a determination will be issued before the existing authorization ends.

For some services, a prior authorization recertification for a course of treatment cannot be issued 60 days before the end of the current authorization period. For example, physical therapy, occupational therapy, speech therapy, and private duty nursing prior authorizations cannot be recertified 60 days before the end of the current authorization period because of requirements for timely documentation. This limitation applies to both fee-for-service and managed care clients.

When a recertification request cannot be reviewed 60 days before the current prior authorization ends, the provider will receive a response from TMHP explaining that the request has been submitted too early, and the provider will need to resubmit at a later time. For managed care clients, providers should contact the client’s specific MCO for details.

Providers must complete all essential fields on prior authorization forms submitted to TMHP to initiate the prior authorization process.

If any essential field on a prior authorization request has missing, incorrect, or illegible information, TMHP returns the original request to the provider with the following message:

TMHP Prior Authorization could not process this request because the request form submitted has missing, incorrect, or illegible information in one or more essential fields. Please resubmit the request with all essential fields completed with accurate information for processing by TMHP within 14 business days of the request receipt date.

TMHP uses the date that the complete and accurate request form is received to determine the start date for services. Previous submission dates of incomplete forms returned are not considered when determining the start date of service.

Providers that need to update information on a prior authorization request form must strike through the incorrect information with a single line. The original content must remain legible, and the change must be initialed and dated by the original signatory or ordering physician when applicable. Changes that have been made using correction fluid (e.g., Wite-Out) will not be accepted.

Providers must respond to an incomplete prior authorization request within 14 business days of the request receipt date. Incomplete prior authorization requests are requests that are received by TMHP with missing, incomplete, or illegible information.

Prior to denying an incomplete request, TMHP’s Prior Authorization department will attempt to get the correct information from the requesting provider. The Prior Authorization department will make a minimum of three attempts to contact the requesting provider before sending a letter to the client about the status of the request and the need for additional information.

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**Program** | **Benefit Code**
---|---
Maternity | MA1
County Indigent Health Care Program | CA1
Early Childhood Intervention (ECI) providers | ECI
Tuberculosis (TB) Clinics | TB1
Intellectual or Developmental Disability (IDD) case management providers | MH2

*Agencies only—Benefit codes should not be used for individual family planning providers.*
If the information that is necessary to make a prior authorization determination is not received within 14 business days of the request receipt date, the request will be denied as “incomplete.” To ensure timely processing, providers should respond to requests for missing or incomplete information as quickly as possible.

For fee-for-service (FFS) Medicaid requests that require a physician review before a final determination can be made, TMHP’s Physician Reviewer will complete the review within three business days of receipt of the completed prior authorization request. An additional three business days will be allowed for requests that require a peer-to-peer review with the client’s prescribing physician.

For Children with Special Health Care Needs (CSHCN) Services Program requests that do not appear to meet CSHCN medical policy, the TMHP prior authorization nurse will refer those requests to the CSHCN Services Program for review and determination. The CSHCN Services Program will complete the review within three business days of receipt of the completed prior authorization request.

**Note:** Providers may resubmit a new, complete request after receiving a denial for an incomplete request; however, the timeliness submission requirements will apply.

Essential fields contain information needed to process a prior authorization request and include the following:

- Client name
- Client Medicaid number (patient control number [PCN])
- Client date of birth
- Provider name
- TPI
- National Provider Identifier (NPI)
- Quantity of service units requested based on the CPT or HCPCS code requested

### 5.4.1 Recreating TMHP Prior Authorization and Authorization Forms to Fill Out Electronically

Providers are allowed to recreate a Texas Medicaid or CSHCN Services Program prior authorization or authorization form in order to transfer it to an electronic format only if there are no alterations in the form’s content or placement of information (field location).

To ensure that the most current version of the form is utilized, all forms submitted to the prior authorization department for processing must include the form number, effective date and revision date, if applicable, as it appears on the original TMHP form that is published on the TMHP Forms web page.

**Important:** Prior authorization and authorization forms recreated and filled out electronically on the provider’s computer must continue to be printed and faxed or submitted by mail. To submit prior authorization and authorization requests electronically, providers must use the prior authorization tool available on the TMHP website.

### 5.5 Prior Authorization Submission Methods

Prior authorization requests can be submitted by fax, mail, telephone, and online through the TMHP website at [www.tmhp.com](http://www.tmhp.com). The methods to use to request the prior authorization depends on the service being requested.
5.5.1 Prior Authorization Requests Through the TMHP Website

Online prior authorization requests for some services in the following areas can be submitted through the TMHP website at www.tmhp.com:

- Home Health
- Home Telemonitoring
- Comprehensive Care Inpatient Psychiatric (CCIP)
- CCP
- Ambulance
- Substance Use Disorder (SUD) services

The benefits of submitting prior authorization requests through the TMHP website include:

- Online editing to ensure that the required information is being submitted correctly.
- The prior authorization number is issued within seconds of submission and confirms that the prior authorization request was accepted. Before providing services, providers must confirm that the prior authorization was approved.
- Notification of approvals and denials are available more quickly.
- Extension requests and status checks can be performed online for prior authorization requests that were submitted online.

Providers can access online prior authorization requests from the Prior Authorization page located under the Topics tab on the TMHP website at www.tmhp.com.

Instructions for submitting prior authorization requests on the TMHP website are located in the Help section at the bottom of the Prior Authorization page.

Prior authorizations that are submitted online will be processed using the same guidelines as prior authorizations submitted by other methods.

Before providers can submit online prior authorization requests, providers must register on the TMHP website and assign an administrator for each Texas Provider Identifier (TPI) and National Provider Identifier (NPI), if one is not already assigned. Users who are configured with administrator rights automatically have permission to submit prior authorization requests.

The TPI administrator can assign submission privileges for nonadministrator accounts. Billing services and clearinghouses must obtain access to protected health-care information through the appropriate administrator of each TPI/NPI provider number for which they are contracted to provide services.

5.5.1.1 Duplicate Validation Check

Prior authorizations submitted to TMHP electronically on the TMHP website are subject to a duplicate validation check. Prior authorizations that are duplicates of a previously submitted authorization return an immediate response with an associated duplicate submission error message.

The error message instructs providers to either update or remove details of the prior authorization submission.

If there is a modifier needed for authorization or claims processing, providers should indicate the primary modifier in the field labeled “Modifier 1.”

The appropriate modifier that identifies the type of therapy being requested needs to be used when providers submit requests for physical, occupational, and speech therapy.
5.5.1.2  Document Requirements and Retention

If information provided in the online request is insufficient to support medical necessity, TMHP Prior Authorization staff may ask the provider to submit additional paper documentation to support the medical necessity for the service being requested.

Submission of prior authorization requests on the secure pages of the TMHP website does not replace adherence to and completion of the paper forms/documentation requirements outlined in this manual and other publications.

Documentation requirements include, but are not limited to, the following:

- Documentation that supports the medical necessity for the service requested.
- Completion and retention in the client’s medical record of all required prior authorization forms.
- Adherence to signature and date requirements for prior authorization forms and other required forms that are kept in the client record, including the following:
  - All prior authorization forms completed and signed before the online prior authorization request is made
  - Original handwritten or electronic signatures (Stamped signatures and images of wet signatures are not accepted by Texas Medicaid.)
- A printed copy of the Online Request Form, which must be retained in the client’s medical record

Any provider, client, or client’s responsible adult who is required to sign a prior authorization form or any supporting documentation may do so using a wet or electronic signature. Any electronic signature technologies that are used must comply with all federal and state statutes and administrative rules.

Any required documentation that is missing from the client’s medical record subjects the associated payments for services to be recouped.

5.5.1.2.1  Acknowledgement Statement

Before submitting each prior authorization request, providers (and submitters on behalf of providers) must affirm that they have read, understood, and agree to the certification and terms and conditions of the prior authorization request.

Providers and submitters are separately held accountable for their declarations after they have acknowledged their agreement and consent by checking the “We Agree” checkbox after reviewing the certification statement and terms and conditions.

5.5.1.2.2  Certification Statement

“The Provider and Authorization Request Submitter certify that the information supplied on the prior authorization form and any attachments or accompanying information constitute true, correct, and complete information. The Provider and Authorization Request Submitter understand that payment of claims related to this prior authorization will be from federal and state funds, and that falsifying entries, concealment of a material fact, or pertinent omissions may constitute fraud and may be prosecuted under applicable federal and/or state law. Fraud is a felony, which can result in fines or imprisonment.

“By checking ‘We Agree’ you agree and consent to the Certification above and to the TMHP ‘Terms and Conditions.’”

5.5.1.2.3  Terms and Conditions

“I hereby agree to keep such records as are necessary to disclose fully the extent of services provided to individuals under the state’s Title XIX plan and to furnish information regarding any payments claimed for providing such services as the State Agency or U.S. Dept. of Health and Human Services may request. I further agree to accept, as payment in full, the amount paid by Medicaid for those claims submitted for payment under that program, with the exception of authorized deductible, coinsurance, copayment or
similar cost-sharing charge. I certify that the services listed above are/were medically indicated and necessary to the health of this patient and were personally furnished by me or my employee under my personal direction.

“Notice: This is to certify that the foregoing information is true, accurate and complete. I understand that payment and satisfaction of this claim, based on information provided on the Prior Authorization form, will be from federal and state funds, and that any false claims, statements or documents, or concealment of a material fact, may be prosecuted under applicable federal or state law.”

Omission of information or failure to provide true and accurate information or notice of changes to the information previously provided may result in termination of the provider’s Medicaid enrollment and/or personal exclusion from Texas Medicaid.

5.5.2 Prior Authorization Requests to TMHP by Fax, Telephone, or Mail

When submitting prior authorization requests through fax or mail, providers must submit the requests on the approved form. If necessary, providers may submit attachments with the form. Providers must follow the guidelines and requirements listed in the handbook for the service. Providers can refer to the provider handbooks for the guidelines and requirements listed for a specific service.

Prior authorization requests must be signed and dated by a physician or dentist who is familiar with the client’s medical condition before the request is submitted to TMHP. When allowed, prior authorizations must be signed and dated by an advanced practice registered nurse (APRN) or PA who is familiar with the client’s medical condition before the request is submitted to TMHP. Prior authorization requests for services that may be signed by a licensed health-care provider other than a physician, dentist, or when allowed by an APRN and PA, do not require handwritten signatures and dates. Electronic signatures from an RN or therapist are acceptable when submitting therapy requests for CCP.

All signatures must be electronic, digital or handwritten. An electronic or digital signature must be derived using software that creates a digital signature logo with a system-generated date and time stamp or includes the logo of the digital software used. Photocopy or ink stamp of a handwritten signature or a typed signature without a digital stamp are not permitted. TMHP will not authorize any dates of services on the request earlier than the date of the provider’s signature. The prior authorization request that contains the original signature must be kept in the client’s medical record for future access and possible retrospective review. These documentation requirements also apply to telephone authorizations. To avoid delays, providers are encouraged to have all clinical documentation at the time of the initial telephone authorization request.

**Note:** Obstetric (OB) services providers are no longer allowed to initiate prior authorization requests or ultrasound extension requests over the telephone.

To initiate a new prior authorization request or to request an extension of an existing prior authorization, OB services providers are required to submit the request online using the TMHP secure provider portal or on paper by faxing or mailing the applicable prior authorization form to TMHP.

Providers can continue to use the Prior Authorization telephone line to inquire about the status of the prior authorization requests that have been submitted to TMHP online, by fax, or mail.

5.5.2.1 Prior Authorization Calls

Initial prior authorization calls are handled by the TMHP Contact Center. If the contact center representative determines that a prior authorization inquiry can be better handled by the TMHP Prior Authorization department, the TMHP Contact Center representative will transfer the call to a Prior Authorization Clinician.

**Note:** This does not impact the timeframes for the resolution of more complex issues.
5.5.2.2 TMHP Prior Authorization Requests by Fax

Providers who fax new prior authorization requests, resubmitted requests, or additional information to complete a request must include:

- A working fax number on the prior authorization form, so that they can receive faxed responses and correspondence from TMHP.
- The last four digits of the client’s Medicaid identification number on the fax coversheet.

*Note:* Prior authorization cover sheets must not contain any protected health information (PHI) per HIPAA. The faxed cover sheet is not meant to replace the appropriate prior authorization form. Providers cannot include information on a cover sheet that is needed to complete the review of a request.

If a provider is faxing prior authorization requests for more than one client, each client request must be faxed individually with a separate cover sheet. Requests received with multiple clients will be returned to the provider for resubmission to ensure Health Insurance Portability and Accountability Act (HIPAA) compliance.

<table>
<thead>
<tr>
<th>Contact</th>
<th>Fax Number</th>
</tr>
</thead>
<tbody>
<tr>
<td>Ambulance Authorization Fax (includes out-of-state transfers)</td>
<td>1-512-514-4205</td>
</tr>
<tr>
<td>Home Health Services Fax</td>
<td>1-512-514-4209</td>
</tr>
<tr>
<td>CCP Fax</td>
<td>1-512-514-4212</td>
</tr>
<tr>
<td>CCIP Fax</td>
<td>1-512-514-4211</td>
</tr>
<tr>
<td>Outpatient Psychiatric Fax</td>
<td>1-512-514-4213</td>
</tr>
<tr>
<td>TMHP Special Medical Prior Authorization (SMPA) Fax (including transplants)</td>
<td>1-512-514-4213</td>
</tr>
</tbody>
</table>

5.5.2.3 TMHP Prior Authorization Requests by Telephone

<table>
<thead>
<tr>
<th>Contact</th>
<th>Telephone Number</th>
</tr>
</thead>
<tbody>
<tr>
<td>Ambulance Authorization (including out-of-state transfers)</td>
<td>1-800-540-0694</td>
</tr>
</tbody>
</table>

5.5.2.4 TMHP Prior Authorization Requests by Mail

<table>
<thead>
<tr>
<th>Contact</th>
<th>Address</th>
</tr>
</thead>
</table>
| Ambulance (includes out-of-state transfers) | Texas Medicaid & Healthcare Partnership  
Ambulance Prior Authorizations  
PO Box 200735  
Austin, TX 78720-0735 |
| CCP                                  | Texas Medicaid & Healthcare Partnership  
Comprehensive Care Program (CCP) Prior Authorization  
PO Box 200735  
Austin, TX 78720-0735 |
| Dental                               | Texas Medicaid & Healthcare Partnership  
Dental Prior Authorization  
PO Box 204206  
Austin, TX 78720-4206 |
| Home Health Services                 | Texas Medicaid & Healthcare Partnership  
Home Health Services Prior Authorization  
PO Box 202977  
Austin, TX 78720-2977 |
5.5.3 **Home Health Services Prior Authorizations**

Home health services providers cannot initiate new prior authorization requests or request extensions over the telephone. The following home health services are affected:

- Skilled nursing and home health aide visits
- Physical and occupational therapy
- Durable medical equipment (DME)
- Expendable medical supplies

Home health services providers must initiate new prior authorization requests or request extension of existing prior authorizations online using the TMHP secure provider portal or on paper by faxing or mailing TMHP the appropriate paper prior authorization form.

Home health services providers can use telephone number 1-800-925-8957 for follow-up and status inquiry of prior authorizations.

5.5.4 **Radiology Prior Authorizations Through eviCore**

eviCore, Inc., performs radiology prior authorization services on behalf of TMHP.

*Refer to:* Subsection 3.2.6, “Authorization Requirements for CT, CTA, MRI, fMRI, MRA, PET, and Cardiac Nuclear Imaging Services” in the *Radiology and Laboratory Services Handbook* *(Vol. 2, Provider Handbooks)* to determine which radiology services require a prior authorization through eviCore.

5.5.4.1 **Online Prior Authorizations Through eviCore**

Radiology prior authorization requests may be submitted through the eviCore website at [www.evicore.com/provider](http://www.evicore.com/provider).

5.5.4.2 **Prior Authorizations to eviCore by Fax, Telephone, or Mail**

Providers can submit radiology prior authorization requests to eviCore by fax, telephone, or mail at:

- **Telephone:** 1-800-572-2116
- **Fax:** 1-800-572-2119
- **Mail:** Texas Medicaid & Healthcare Partnership
  400 Buckwalter Place Blvd
  Bluffton, SC 29910

5.5.4.3 **Retroactive Authorization Requests**

Retroactive authorization requests for outpatient diagnostic computed tomography (CT), magnetic resonance (MR), positron emission tomography (PET) and cardiac nuclear imaging services for Texas Medicaid fee-for-service clients must be submitted online to eviCore. The retroactive authorizations requests must be submitted to eviCore no later than 14 calendar days after the day on which the study was completed, regardless of the method of submission. If the retroactive authorization request is submitted after the allotted time, the authorization request will not be processed.
5.6 Verifying Prior Authorization Status

Prior authorizations are processed based on the date the request is received. Requests with all required information can take up to three business days after the date of receipt for TMHP to complete the authorization process.

Providers can check the status of prior authorizations requested online through the TMHP website at www.tmhp.com.

Providers may also check status of prior authorizations that are issued by TMHP by using the following numbers.

<table>
<thead>
<tr>
<th>Contact</th>
<th>Telephone Number</th>
</tr>
</thead>
<tbody>
<tr>
<td>Personal Care Services (PCS) Prior Authorization Inquiry Line</td>
<td>1-888-648-1517</td>
</tr>
<tr>
<td>CCP and Home Health Status Line</td>
<td>1-800-846-7470</td>
</tr>
<tr>
<td>All other authorization requests</td>
<td>1-800-925-9126</td>
</tr>
</tbody>
</table>

To check the status of radiology prior authorization requests that are submitted to eviCore, providers should contact eviCore directly at www.evicore.com or 1-800-572-2116.

5.7 Prior Authorization Notifications

TMHP sends a notification to the provider when the prior authorization is approved, denied, or modified. If TMHP receives prior authorization requests with incomplete or insufficient information, TMHP will ask the requesting provider to furnish the additional documentation needed before TMHP can make a decision on the request. If the requesting provider does not respond to the request for additional information, the prior authorization request will be denied. It is the requesting provider’s responsibility to contact the appropriate provider, when necessary, to obtain the additional documentation.

5.8 Prior Authorization Denials Appeals Process

Prior authorizations that are denied by TMHP can be resubmitted to the TMHP Prior Authorization Department with new or additional information for reconsideration.

If the request is denied a second time, or if the provider has no new or additional information, the provider may file an Administrative Appeal to HHSC. Providers must include a copy of the denial letter.

It is strongly recommended that providers maintain a list that details the prior authorizations, including:

- Client’s name
- Client’s Medicaid number
- Date of service
- Provider Identifier
- Items submitted

This information will be required if a provider needs to file an administrative review.

5.9 Closing a Prior Authorization

When a client decides to change providers or elects to discontinue prior-authorized services before the authorization ends, that prior authorization is updated to reflect the early closure date and the reason for closure.
If a client with an active prior authorization changes providers, TMHP must receive a change of provider letter with the request for a new prior authorization in accordance with submission guidelines for the service. The client must sign and date the letter, which must include the name of the previous provider, the current provider, and the effective date for the change.

The client is responsible for notifying the previous provider that the client is discontinuing services and the effective date of the change. TMHP also notifies the previous provider by mail when a prior authorization has been closed early. The letter includes the beginning date of service, the revised ending date of the authorization, and the reason for the early closure.

If a provider submits a Change of Provider letter in the middle of an existing authorization period, the current authorization will be end-dated and the original provider will be notified. TMHP will send the new provider an authorization that begins on the next business day after the end date and lasts through the remainder of the authorization period.

### 5.10 Submitting Claims for Services That Require Prior Authorization

Claims submitted for services that require prior authorization must indicate the authorization number, provider identifier, procedure codes, dates of service, required modifiers, number of units, and the amount for manually priced procedure codes as detailed on the authorization letter. If the prior authorization letter shows itemized details and the provider rendered all services listed, the details on the claim must match the details on the prior authorization letter.

**Important:** Claims processing and payment may be delayed if the detailed information on the authorization letter and the claim details do not match exactly.

Claims for prior authorized services must contain only one prior authorization number per claim. Prior authorization numbers must be indicated on the applicable electronic fields or in the following blocks for paper claim forms:

<table>
<thead>
<tr>
<th>Paper Claim Form</th>
<th>Block for Prior Authorization Number</th>
</tr>
</thead>
<tbody>
<tr>
<td>CMS-1500 (professional) claim form</td>
<td>Block 23</td>
</tr>
<tr>
<td>UB-04 CMS-1450 (institutional) claim form</td>
<td>Block 63</td>
</tr>
<tr>
<td>American Dental Association (ADA) claim form</td>
<td>Block 2</td>
</tr>
<tr>
<td>2017 claim form</td>
<td>Block 30</td>
</tr>
</tbody>
</table>

**Refer to:** Subsection 6.1.3, “TMHP Paper Claims Submission” in “Section 6: Claims Filing” (Vol. 1, General Information).

### 5.10.1 Authorization and Manually Priced Claims

If prior authorization has been obtained for services that use manually priced procedure codes, providers must submit claims for them using the MSRP that was submitted with the authorization request and the following information that is listed on the authorization letter:

- Authorization number
- Provider identifier
- Procedure codes
- Dates of service
- Types of service
- Required modifiers
If the authorization letter shows itemized details, the claim must include all rendered services as they are itemized on the authorization letter and the MSRP rate for each of those services. The procedure codes and MSRP rates that are detailed on the claim must match the procedure codes that are detailed in the authorization letter and the MSRP rates that were submitted with the authorization request. Claims processing and payment may be delayed if there is not an exact match between the detailed information on the authorization letter, the approved authorization, and the information that was submitted on the claim.

**Important:** For appropriate processing and payment, the Pay Price that is indicated on the authorization letter should not be billed on the claim.

Prior authorization is a condition of reimbursement; it is not a guarantee of payment.

### 5.11 Guidelines for Procedures Awaiting Rate Hearing

For procedure codes that require prior authorization but are awaiting a rate hearing, providers must follow the established prior authorization process as defined in the applicable provider handbook. Providers must obtain a timely prior authorization for services provided. Providers must not wait until the rate hearing process for the procedure codes is completed to request prior authorization. In this situation, retroactive prior authorization requests are not granted; the requests are denied as late submissions. Providers are also responsible for meeting the initial 95-day filing deadline and for ensuring that the prior authorization number is on the claim the first time it is submitted to TMHP for consideration of reimbursement.

Claims for procedure codes awaiting a rate hearing are denied. TMHP automatically reprocesses affected claims; providers are not required to appeal the claims unless they are denied for additional reasons after the claims reprocessing is complete. If the required prior authorization number is not on the claim at the time of reprocessing, the claim is denied for lack of prior authorization.
SECTION 6: CLAIMS FILING

TEXAS MEDICAID PROVIDER PROCEDURES MANUAL: VOL. 1

MAY 2021
SECTION 6: CLAIMS FILING

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6.1 Claims Information

Providers that render services to Texas Medicaid fee-for-service and managed care clients must file the assigned claims. Texas Medicaid does not make payments to clients. Federal regulations prohibit providers from charging clients a fee for completing or filing Medicaid claim forms. Providers are not allowed to charge TMHP for filing claims. The cost of claims filing is part of the usual and customary rate for doing business. Providers cannot bill Texas Medicaid or Medicaid clients for missed appointments or failure to keep an appointment. Only claims for services rendered are considered for payment. Medicaid providers are also required to complete and sign authorized medical transportation forms (e.g., Form 3103, Individual Driver Registrant (IDR) Service Record, or Form 3111, Verification of Travel to Healthcare Services by Mass Transit) or provide an equivalent (e.g., provider statement on official letterhead) to attest that services were provided to a client on a specific date. The client presents these forms to the provider.

Providers are not allowed to bill clients or Texas Medicaid for completing these forms.

6.1.1 TMHP Processing Procedures

TMHP processes claims for services rendered to Texas Medicaid fee-for-service clients and carve-out services rendered to Medicaid managed care clients.

**Note:** Claims for services rendered to a Medicaid managed care client must be submitted to the managed care organization (MCO) or dental plan that administers the client’s managed care benefits. Only claims for those services that are carved-out of managed care can be submitted to TMHP.

**Refer to:** The Medicaid Managed Care Handbook (Vol. 2, Provider Handbooks) for more information about carve-out services.

Medicaid claims are subject to the following procedures:

- TMHP verifies all required information is present.
- Claims filed under the same provider identifier and program and ready for disposition at the end of each week are paid to the provider with an explanation of each payment or denial. The explanation is called the Remittance and Status (R&S) Report, which may be received as a downloadable portable document format (PDF) version or on paper. A Health Insurance Portability and Accountability Act (HIPAA)-compliant 835 transaction file is also available for those providers who wish to import claim dispositions into a financial system.

An R&S Report is generated for providers that have weekly claim or financial activity with or without payment. The report identifies pending, paid, denied, and adjusted claims. If no claim activity or outstanding account receivables exist during the time period, an R&S Report is not generated for the week.

- For services that are billed on a claim and have any benefit limitations for providers, the date of service determines which provider’s claims are paid, denied, or recouped. Claims that have been submitted and paid may be recouped if a new claim with an earlier date of service is submitted, depending on the benefit limitations for the services rendered.

Services that have been authorized for an extension of the benefit limitation will not be recouped. Providers can submit an appeal with medical documentation if the claim has been denied.
6.1.1.1 Fiscal Agent

TMHP acts as the state’s Medicaid fiscal agent. A fiscal agent arrangement is one of two methods allowed under federal law and is used by all other states that contract with outside entities for Medicaid claims payment. Under the fiscal agent arrangement, TMHP is responsible for paying claims, and the state is responsible for covering the cost of claims.

Note: The fiscal agent arrangement does not affect Long Term Care (LTC) and Health and Human Services Commission (HHSC) Family Planning providers.

Provider Designations

The fiscal agent arrangement requires that providers be designated as either public or nonpublic. By definition, public providers are those that are owned or operated by a city, state, county, or other government agency or instrumentality, according to the Code of Federal Regulations. In addition, any provider or agency that performs intergovernmental transfers to the state would be considered a public provider. This includes those agencies that can certify and provide state matching funds, (i.e., other state agencies). New providers self-designate (public or private) on the provider enrollment application.

The fiscal agent:

- Rejects all claims not payable under Texas Medicaid rules and regulations.
- Suspends payments to providers according to procedures approved by HHSC.
- Notifies providers of reduction in claim amount or rejection of claim and the reason for doing so.
- Collects payments made in error, affects a current record credit to the department, and provides the department with required data relating to such error corrections.
- Prepares checks or drafts to providers, except for cases in which the department agrees that a basis exists for further review, suspension, or other irregularity within a period not to exceed 30 days of receipt and determination of proper evidence establishing the validity of claims, invoices, and statements.
- Makes provisions for payments to providers who have furnished eligible client benefits.
- Withholds payment of claim when the eligible client has another source of payment.
- Employs and assigns a physician, or physicians, and other professionals as necessary, to establish suitable standards for the audit of claims for services delivered and payment to eligible providers.
- Requires eligible providers to submit information on claim forms.

6.1.1.2 Payment Error Rate Measurement (PERM)

The Improper Payments Information Act (IPIA) of 2002 directs federal agency heads, in accordance with the Office of Management and Budget (OMB) guidance, to annually review agency programs that are susceptible to significant erroneous payments and to report the improper payment estimates to the U.S. Congress.

Every three years the CMS will assess Texas Medicaid using the PERM process to measure improper payments in Texas Medicaid and the Children’s Health Insurance Program (CHIP).

CMS uses PERM to measure the accuracy of Medicaid and CHIP payments made by states for services rendered to clients. Under the PERM program, CMS will use three national contractors to measure improper payments in Medicaid and CHIP:

- The statistical contractor will provide support to the program by identifying the claims to be reviewed and by calculating each state’s error rate.
- The data documentation contractor will collect medical policies from the State and medical records from providers.
• The review contractor will perform medical and data processing reviews of the selected claims in order to identify any improper payments.

Providers are required to provide medical record documentation to support the medical reviews that the federal review contractor will conduct for Texas Medicaid fee-for-service and CHIP claims.

Note: The federal review contractor will also conduct reviews for Primary Care Case Management (PCCM) claims that were submitted to TMHP with dates of service on or before February 29, 2012.

Past studies have shown that the largest cause of error in medical reviews is lack of documentation or insufficient documentation. It is important that information be sent in a timely and complete manner, since a provider’s failure to timely submit complete records in support of the claims filed can result in a higher payment error rate for Texas, which in turn can negatively impact the amount of federal funding received by Texas for Medicaid and CHIP.

Providers must submit the requested medical records to the data documentation contractor and HHSC within 60 calendar days of the receipt of the written notice of request. If providers have not responded within 15 days, the data documentation contractor and possibly state officials will initiate reminder calls and letters to providers. The data documentation contractor and possibly state officials will also initiate reminder calls and letters to providers after 35 days. If providers have not responded in 60 days, the data documentation contractor will submit a letter to the provider and the state PERM director indicating a “no documentation error.” After the provider’s submittal of requested information, the data documentation contractor may request additional information to determine proper payment. In this instance, the provider is given 15 days to provide additional documentation.

If medical records are not received within 60 calendar days, the data documentation contractor will identify the claim as a PERM error and classify all dollars associated with the claim as an overpayment. Providers will be required to reimburse the overpayment in accordance with state and federal requirements.

A provider’s failure to maintain complete and correct documentation in support of claims filed or failure to provide such documentation upon request can result in the provider being sanctioned under Title 1, Texas Administrative Code (TAC) Part 15, Chapter 371. Sanction actions may include, but are not limited to, a finding of overpayment for the claims that are not sufficiently supported by the required documentation. Sanctions may include, but are not limited to, a finding of overpayment for the claims that are not sufficiently supported by the required documentation.

6.1.2 Claims Filing Instructions

This manual references paper claims when explaining filing instructions. HHSC and TMHP encourage providers to submit claims electronically. TMHP offers specifications for electronic claim formats. These specifications are available from the TMHP website and include a cross-reference of the paper claim filing requirements to the electronic format.

Providers can participate in the most efficient and effective method of submitting claims to TMHP by submitting claims through the TMHP Electronic Data Interchange (EDI) claims processing system using TexMedConnect or a third party vendor. The proceeding claim filing instructions in this manual apply to paper and electronic submitters. Although the examples of claims filing instructions refer to their inclusion on the paper claim form, claim data requirements apply to all claim submissions, regardless of the media. Claims must contain the provider’s complete name, address, and provider identifier to avoid unnecessary delays in processing and payment.

Refer to: “Section 3: TMHP Electronic Data Interchange (EDI)” (Vol. 1, General Information) for information on accessing the TMHP website.
6.1.2.1 Wrong Surgery Notification

Providers are required to notify TMHP when a wrong surgery or other invasive procedure is performed on a Texas Medicaid client. Notification is mandated by Senate Bill (SB) 203, Section 3, Regular Session, 81st Texas Legislature, which covers preventable adverse events (PAE) and reimbursement for services associated with PAE.

Professional, inpatient, and outpatient hospital claims that are submitted for the wrong surgery or invasive procedure will be denied. Any corresponding procedures that are rendered to the same client, on the same dates of service (for professional and outpatient hospital claims), or the same date of surgery (for inpatient hospital claims) will be denied. Claims that have already been reimbursed will be recouped.

The law requires providers that are submitting claims for services rendered to Texas Medicaid clients to indicate whether any of the following situations apply to the claim:

- The incorrect operation or invasive procedure was performed on the correct client.
- The operation or invasive procedure was performed on the incorrect client.
- The incorrect operation or invasive procedure was performed on the incorrect body part.

Providers must notify Texas Medicaid of a wrong surgery or invasive procedure by submitting one of the following nonspecific injury, poisoning and other consequences of external causes diagnosis codes or modifiers with the procedure code for the rendered service:

<table>
<thead>
<tr>
<th>Code</th>
<th>Description</th>
<th>Type of Claim</th>
</tr>
</thead>
<tbody>
<tr>
<td>Injury, Poisoning and Other Consequences of External causes Diagnosis Codes</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Y6551</td>
<td>Performance of wrong procedure (operation) on correct patient</td>
<td>Inpatient hospital</td>
</tr>
<tr>
<td>Y6552</td>
<td>Performance of procedure (operation) on patient not scheduled for surgery</td>
<td></td>
</tr>
<tr>
<td>Y6553</td>
<td>Performance of correct procedure (operation) on wrong side or body part</td>
<td></td>
</tr>
</tbody>
</table>

Modifiers

<table>
<thead>
<tr>
<th>Code</th>
<th>Description</th>
<th>Type of Claim</th>
</tr>
</thead>
<tbody>
<tr>
<td>PA</td>
<td>Surgical or other invasive procedure on wrong body part</td>
<td>Professional or outpatient hospital</td>
</tr>
<tr>
<td>PB</td>
<td>Surgical or other invasive procedure on wrong patient</td>
<td></td>
</tr>
<tr>
<td>PC</td>
<td>Wrong surgery or other invasive procedure on patient</td>
<td></td>
</tr>
</tbody>
</table>

Professional or outpatient hospital claims must include a valid diagnosis with up to seven-digit specificity, the procedure code that identifies the service rendered, and the PA, PB, or PC modifier that describes the type of “wrong surgery” performed.

Inpatient hospital claims must be submitted with type of bill (TOB) 110 as an inpatient hospital-nonpayment claim when a “wrong surgery” is reported. If other services or procedures that are unrelated to the “wrong surgery” are provided during the same stay as the “wrong surgery,” the inpatient hospital must submit a claim for the “wrong surgery” and a separate claim or claims for the unrelated services rendered during the same stay as the “wrong surgery.”

The “wrong surgery” claim must include TOB 110, the appropriate diagnosis code, the surgical procedure code for the surgical service rendered, and the date of surgery. The “wrong surgery” claim will be denied.

The unrelated services rendered during the same stay as the “wrong surgery” must include TOB 111, 112, 113, 114, or 115 on a claim separate from the “wrong surgery” claim. The unrelated services that are benefits of Texas Medicaid may be reimbursed by Texas Medicaid.
A claim that is denied for wrong surgery will have one of the following EOB codes:

<table>
<thead>
<tr>
<th>EOB Code</th>
<th>Message</th>
</tr>
</thead>
<tbody>
<tr>
<td>01167</td>
<td>Claim detail denied due to wrong surgery performed on client</td>
</tr>
<tr>
<td>01168</td>
<td>Claim denied due to wrong surgery performed on client</td>
</tr>
<tr>
<td>01185</td>
<td>Claim denied due to wrong surgery claim found in history for the same PCN and DOS</td>
</tr>
<tr>
<td>01186</td>
<td>Claim detail denied due to wrong surgery claim found in history for the same PCN and DOS</td>
</tr>
</tbody>
</table>

PCN = Patient Control Number (also known as the client’s Medicaid number) DOS = Date of service

### 6.1.2.2 Maximum Number of Units allowed per Claim Detail

The total number of units per claim detail can not exceed 9,999. Providers who submit a claim with more than 9,999 units must bill 9,999 units on the first detail of the claim and any additional units on separate details.

### 6.1.2.3 Tips on Expediting Paper Claims

Use the following guidelines to enhance the accuracy and timeliness of paper claims processing.

**General requirements**

- Use original claim forms. Do not use copies of claim forms.
- Detach claims at perforated lines before mailing.
- Use 10 x 13 inch envelopes to mail claims. Do not fold claim forms, appeals, or correspondence.
- Do not use labels, stickers, or stamps on the claim form.
- Do not send duplicate copies of information.
- Use 8 ½ x 11 inch paper. Do not use paper smaller or larger than 8 ½ x 11 inches.
- Do not mail claims with correspondence for other departments.

**Data Fields**

- Print claim data within defined boxes on the claim form.
- Use black ink, but not a black marker. Do not use red ink or highlighters.
- Use all capital letters.
- Print using 10-pitch (12-point) Courier font. Do not use fonts smaller or larger than 12 points. Do not use proportional fonts, such as Arial or Times Roman.
- Use a laser printer for best results. Do not use a dot matrix printer, if possible.
- Do not use dashes or slashes in date fields.

**Attachments**

- Use paper clips on claims or appeals if they include attachments. Do not use glue, tape, or staples.
- Place the claim form on top when sending new claims, followed by any medical records or other attachments.
- Number the pages when sending attachments or multiple claims for the same client (e.g., 1 of 2, 2 of 2).
- Do not total the billed amount on each claim form when submitting multi-page claims for the same client.
• Use the CMS-approved Medicare Remittance Advice Notice (MRAN) printed from Medicare Remit Easy Print (MREP) (professional services) or PC-Print (institutional services) when sending a Remittance Advice from Medicare or the paper MRAN received from Medicare or a Medicare intermediary. You may also download the TMHP Standardized Medicare Advantage Plan (MAP) Remittance Advice Notice Template from the TMHP website at www.tmhp.com.

• Submit claim forms with MRANs and R&S Reports.

6.1.3 TMHP Paper Claims Submission

All paper claims must be submitted with a Texas Provider Identifier (TPI) and National Provider Identifier (NPI) for the billing and performing providers. All other provider fields on the claim forms require an NPI only. If an NPI and TPI are not included in the billing and performing provider fields, or if an NPI is not included on all other provider identifier fields, the claim will be denied.

6.1.4 Claims Filing Deadlines

For claims payment to be considered, providers must adhere to the time limits described in this section. Claims received after the following claims filing deadlines are not payable because Texas Medicaid does not provide coverage for late claims.

Except: Unless otherwise stated, claims must be received by TMHP within 95 days of each DOS. Appeals must be received by TMHP within 120 days of the disposition date on the R&S Report on which the claim appears. A 95-day or 120-day appeal filing deadline that falls on a weekend or a holiday is extended to the next business day following the weekend or holiday.

Only the following holidays extend the deadlines in 2020 and 2021:

<table>
<thead>
<tr>
<th>Date</th>
<th>Holiday</th>
</tr>
</thead>
<tbody>
<tr>
<td>January 1, 2021</td>
<td>New Year’s Day</td>
</tr>
<tr>
<td>January 18, 2021</td>
<td>Martin Luther King, Jr. Day</td>
</tr>
<tr>
<td>February 15, 2021</td>
<td>Presidents Day</td>
</tr>
<tr>
<td>May 31, 2021</td>
<td>Memorial Day</td>
</tr>
<tr>
<td>July 4, 2021</td>
<td>Independence Day</td>
</tr>
<tr>
<td>September 6, 2021</td>
<td>Labor Day</td>
</tr>
<tr>
<td>October 11, 2021*</td>
<td>Columbus Day</td>
</tr>
<tr>
<td>November 11, 2021</td>
<td>Veterans Day</td>
</tr>
<tr>
<td>November 25, 2021</td>
<td>Thanksgiving Day</td>
</tr>
<tr>
<td>November 26, 2021</td>
<td>Day after Thanksgiving</td>
</tr>
<tr>
<td>December 24, 2021</td>
<td>Christmas Eve Day</td>
</tr>
<tr>
<td>December 25, 2021</td>
<td>Christmas Day</td>
</tr>
</tbody>
</table>

*Federal holiday, but not a state holiday. The claims filing deadline will be extended for providers because the Post Office will not be operating on this day.

The following are time limits for submitting claims:

• Inpatient claims that are filed by the hospital must be received by TMHP within 95 days of the discharge date or last DOS on the claim.

• Hospitals that are reimbursed according to diagnosis-related group (DRG) payment methodology may submit an interim claim because the client has been in the facility 30 consecutive days or longer. A total stay claim is needed after discharge to ensure accurate calculation for potential outlier payments for clients who are 20 years of age and younger.
• Hospitals that are reimbursed according to Tax Equity and Fiscal Responsibility Act (TEFRA) of 1982 methodology may submit interim claims before discharge and must submit an interim claim if the client remains in the hospital past the hospital’s fiscal year end.

• When medical services are rendered to a Medicaid client in Texas, TMHP must receive claims within 95 days of the DOS on the claim.

• Re-enrolling providers who are assigned their previous TPI must submit claims so that they are received by TMHP within 95 days of the date of service.

• Providers that are enrolling in Texas Medicaid for the first time or are making a change that requires the issuance of a new TPI can submit claims within 95 days from the date their TPI is issued as long as claims are submitted within 365 days of the date of service.

• Providers who are revalidating an existing enrollment can continue to file claims while they are completing the revalidation process. TMHP must receive claims within 95 days of the date of service.

• TMHP must receive claims from out-of-state providers within 365 days from the DOS. The DOS is the date the service is provided or performed.

• TMHP must receive claims on behalf of an individual who has applied for Medicaid coverage but has not been assigned a Medicaid number on the DOS within 95 days from the date the eligibility was added to the TMHP eligibility file (add date) and within 365 days of the date of service or from the discharge date for inpatient claims.

• Providers should verify eligibility and add date by contacting TMHP (Automated Inquiry System [AIS], TMHP EDI’s electronic eligibility verification, or TMHP Contact Center) when the number is received. Not all applicants become eligible clients. Providers that submit claims electronically within the 365-day federal filing deadline for services rendered to individuals who do not currently have a Texas Medicaid identification number will receive an electronic rejection. Providers can use the TMHP rejection report as proof of meeting the 365-day federal filing deadline and submit an administrative appeal.

Important: Providers should keep documentation of all Texas Medicaid client eligibility verification. Documentation of client eligibility is required for the appeal process.

• If a client becomes retroactively eligible or loses Medicaid eligibility and is later determined to be eligible, the 95-day filing deadline begins on the date that the eligibility start date was added to TMHP files (the add date). However, the 365-day federal filing deadline must still be met.

• When a service is a benefit of Medicare and Medicaid, and the client is covered by both programs, the claim must be filed with Medicare first. TMHP must receive Medicaid claims within 95 days of the date of Medicare disposition.

Providers must submit a paper MRAN received from Medicare or a Medicare intermediary, the computer-generated MRANs from the CMS-approved software application MREP for professional services or PC-Print for institutional services, or the TMHP Standardized Medicare Advantage Plan (MAP) Remittance Advice Notice Template with a completed claim form to TMHP.

• When a client is eligible for Medicare Part B only, the inpatient hospital claim for services covered as Medicaid only is sent directly to TMHP and is subject to the 95-day filing deadline (from date of discharge).

Note: It is strongly recommended that providers who submit paper claims keep a copy of the documentation they send. It is also recommended that paper claims be sent by certified mail with a return receipt requested. This documentation, along with a detailed listing of the claims enclosed, provides proof that the claims were received by TMHP, which is particularly important if it is necessary to prove that the 95-day claims filing deadline has been met.
TMHP will accept certification receipts as proof of the 95-day or 120-filing deadline. If a certified receipt is provided as proof, the certified receipt number must be indicated on the detailed listing along with the Medicaid number, billed amount, DOS, and a signed claim copy. The provider needs to keep such proof of multiple claims submissions if the provider identifier is pending.

- If the provider is attempting to obtain prior authorization for services performed or will be performed, TMHP must receive the claim according to the usual 95-day filing deadline.

- The provider bills TMHP directly within 95 days from the DOS. However, if a non-third party resource (TPR) is billed first, TMHP must receive the claim within 95 days of the claim disposition by the other entity.

**Note:** The provider submits a copy of the disposition with the claim. A non-TPR is secondary to Texas Medicaid and may only pay benefits after Texas Medicaid.

**Refer to:** Subsection 4.12, “Third Party Liability (TPL)” in Section 4, “Client Eligibility” (Vol. 1, General Information) for examples of non-TPRs.

- When a service is billed to another insurance resource, the filing deadline is 95 days from the date of disposition by the other resource.

- When a service is billed to a third party and no response has been received, Medicaid providers must allow 110 days to elapse before submitting a claim to TMHP. However, the 365-day federal filing deadline requirement must still be met.

- A Compass21 (C21) process allows an HHSC Family Planning claim to be paid by Title XIX (Medicaid) if the client is eligible for Title XIX when those services are provided and billed under the HHSC Family Planning Program. In this instance, the Medicaid 95-day filing deadline is in effect and must be met or the claim will be denied.

- For claims re-submitted to TMHP with additional detail changes (i.e., quantity billed), the additional details are subject to the 95-day filing deadline.

**Note:** In accordance with federal regulations, all claims must be initially filed with TMHP within 365 days of the DOS, regardless of provider enrollment status or retroactive eligibility.

**Refer to:** Subsection 6.1.2, “Claims Filing Instructions” in this section.

Subsection 1.1, “Provider Enrollment” in “Section 1: Provider Enrollment and Responsibilities” (Vol. 1, General Information) for information on the provider enrollment process.

Subsection 7.1, “Appeal Methods” in “Section 7: Appeals” (Vol. 1, General Information) for information on the process for submitting appeals.

Subsection 6.1.4.3, “Exceptions to the 95-Day Filing Deadline” in this section.

Subsection A.12.3, “Automated Inquiry System (AIS)” in “Appendix A: State, Federal, and TMHP Contact Information” (Vol. 1, General Information) to learn how to retrieve client eligibility information by telephone.

**Refer to:** “Section 8: Third Party Liability (TPL)” (Vol. 1, General Information).

Subsection 4.1.10, “Eligibility Verification” in “Section 4: Client Eligibility” (Vol. 1, General Information).

Subsection 6.11.6, “Provider Inquiries—Status of Claims” in this section.

**6.1.4.1 Claims for Clients with Retroactive Eligibility**

Claims for clients who receive retroactive eligibility must be submitted within 95 days of the date that the client's eligibility was added to the TMHP eligibility file (add date) and within 365 days of the DOS.
Title 42 of the Code of Federal Regulations (42 CFR), at 447.45 (d) (1), states “The Medicaid agency must require providers to submit all claims no later than 12 months from the date of service.” The 12-month filing deadline applies to all claims. Claims not submitted within 365 days (12 months) from the date of service cannot be considered for payment.

Retroactive eligibility does not constitute an exception to the federal filing deadline. Even if the patient’s Medicaid eligibility determination is delayed, the provider must still submit the claim within 365 days of the date of service. A claim that is not submitted within 365 days of the date of service will not be considered for payment.

If a client is not yet eligible for Medicaid, providers must submit the claim using either 999999999 or 000000000 as the recipient identification number. Although TMHP will deny the claim, providers should retain the denial or electronic rejection report for proof of timely filing, especially if the eligibility determination occurs more than 365 days after the date of service. Claims denied for recipient ineligibility may be resubmitted when the patient becomes eligible for the retroactive date(s) of service. Texas Medicaid may then consider the claim for payment because the initial claim was submitted within the 365-day federal filing deadline and the denial was not the result of an error by the provider.

If the 365-day federal filing deadline requirement has passed, providers must submit the following to TMHP within 95 days from the add date:

- A completed claim form.
- One of the following dated within 365 days from the date of service:
  - A page from an R&S Report documenting a denial of the claim.
  - An electronic rejection report of the claim that includes the Medicaid recipient’s name and date of service.

Providers that have submitted their claims electronically can provide proof of timely filing by submitting a copy of an electronic claims report that includes the following information:

- Client name or Medicaid identification number (PCN)
- DOS
- Total charges
- Batch identification number (Batch ID) (in correct format)

Note: Only reports that were accepted or rejected by TMHP will be honored. The claim filed (client name or PCN, DOS and total charges) should match the information on the batch report.

6.1.4.2 Claims for Newly Enrolled Providers

Claims submitted by newly enrolled providers must be received within 95 days of the date that the new provider identifier is issued, and within 365 days of the date of service. Providers with a pending application should submit any claims that are nearing the 365-day deadline from the date of service. Claims will be rejected by TMHP until a provider identifier is issued. Providers can use the TMHP rejection report as proof of meeting the 365-day deadline and submit an appeal.

Refer to: Subsection 1.1.10.11, “Copy of License, Temporary License, or Certification” in “Section 1: Provider Enrollment and Responsibilities” (Vol. 1, General Information).

All claims for services rendered to Texas Medicaid clients who do not have Medicare benefits are subject to a filing deadline from the date of service of:

- 95 days for in-state providers.
- 365 days for out-of-state providers.
TMHP cannot issue a prior authorization before Medicaid enrollment is complete. Upon notice of Medicaid enrollment, by way of issuance of a provider identifier, the provider must contact the appropriate TMHP Authorization Department before providing services that require a prior authorization number to Medicaid clients. Regular prior authorization procedures are followed after the TMHP Prior Authorization Department has been contacted.

Retroactive authorizations will not be issued unless the regular authorization procedures for the requested services allow for authorizations to be obtained after services are provided. For these services, providers have 95 days from the add date of the client’s retroactive eligibility in TMHP’s system to obtain authorization for services that have already been performed. Providers should refer to the specific manual section for details on authorization requirements, claims filing, and timeframe guidelines for authorization request submissions.

Providers who have not been assigned a provider identifier and have general claim submission questions may refer to this section for assistance with claim submission. If additional general information is needed, providers may call the TMHP Contact Center at 1-800-925-9126 to obtain information. Due to HIPAA privacy guidelines, specific client and claim information cannot be provided.

Providers who have already been assigned a provider identifier and have questions about submitting claims may call the same number and select the option to speak with a TMHP Contact Center representative.

6.1.4.3 Exceptions to the 95-Day Filing Deadline
TMHP is not responsible for appeals about exceptions to the 95-day filing deadline. These appeals must be submitted to the HHSC Claims Administrator Operations Management. TAC allows HHSC to consider exceptions to the 95-day filing deadline under special circumstances.

6.1.4.4 Appeal Time Limits
All appeals of denied claims and requests for adjustments on paid claims must be received by TMHP within 120 days from the date of disposition, the date of the R&S Report on which that claim appears. If the 120-day appeal deadline falls on a weekend or holiday, the deadline will be extended to the next business day.

Refer to: Subsection 6.1.2, “Claims Filing Instructions” in this section.

Hospitals appealing final technical denials, admission denials, DRG changes, continued-stay denials, or cost/day outlier denials refer to “Section 7: Appeals” (Vol. 1, General Information) for complete appeal information.

6.1.4.5 Claims with Incomplete Information and Zero Paid Claims
Claims listed on the R&S Report with $0 allowed and $0 paid may be resubmitted as electronic appeals. Previously, these claims were only accepted as paper claims and were not accepted as electronic appeals. Appeals may be submitted through a third party biller or through TexMedConnect.

Zero-paid claims that are still within the 95-day filing deadline should be submitted as new day claims, which are processed faster than appeals. Electronic appeal for these claims must be submitted within the 120-day appeal deadline. Electronic claims can be resubmitted past the 95-day deadline as new day claims if the following fields have not changed:

- Provider identifiers
- Client Medicaid number
- Dates of service
- Total billed amount
Claims that are past the 95-day filing deadline and require changes to the fields listed above must be appealed on paper, with a copy of the R&S report. All other appeal guidelines remain unchanged.

**Important:** Initial zero-paid claims and appeal submissions must meet the 95-day deadline and 120-day appeal deadline outlined in subsection 6.1.4, “Claims Filing Deadlines” in this section.

### 6.1.4.6 Claims Filing Reminders

After filing a claim to TMHP, providers should review the weekly R&S Report. If within 30 days the claim does not appear in the Claims In Process section, or if it does not appear as a paid, denied, or incomplete claim, the provider should resubmit it to TMHP within 95 days of the DOS.

The provider should allow TMHP 45 days to receive a Medicare-paid claim automatically transmitted for payment of deductible or coinsurance.

Electronic billers should notify TMHP about missing claims when:

- An accepted claim does not appear on the R&S Report within ten workdays of the file submittal.
- A claim or file does not appear on a TMHP Electronic Claims Submission Report within ten days of the file submission.

Certain claims, including those that were submitted for newborn services or that might be covered under Medicare, are suspended for review so that other state agencies can verify information. This review may take longer than 60 days.

These suspended claims will appear on the provider’s R&S Report under “The following claims are being processed” with a message indicating that the client’s eligibility is being investigated. Providers must wait until the claim is finalized and appears under “Paid or Denied” or “Adjustment to Claims” on the R&S Report before appealing the claim. If the claim does not appear on the R&S Report, providers must resubmit the claim to TMHP to ensure compliance with filing and appeal deadlines.

### 6.1.5 HHSC Payment Deadline

Payment deadline rules, as defined by HHSC, affect all providers with the exception of LTC and the HHSC Family Planning Program. The HHSC payment deadline rules for the fiscal agent arrangement ensure that state and federal financial requirements are met.

TMHP is required to finalize and pay claims within 24 months of:

- Each date of service on a claim.
- Discharge date for inpatient claims.

Texas Medicaid and Children with Special Health Care Needs (CSHCN) Service Program payments, excluding crossovers, cannot be made after 24 months. Claims and appeals that are submitted after the designated payment deadlines are denied.

**Note:** Providers may appeal HHSC Office of Inspector General (OIG) initiated claims adjustments (recoupments) after the 24-month deadline but must do so within 120 days from the date of the recoupment. Refer to subsection 7.1.5, “Paper Appeals” in “Section 7: Appeals” (Vol. 1, General Information) for instructions. All appeals of OIG recoupments must be submitted by paper, no electronic or telephone appeals will be accepted.

### 6.1.6 Filing Deadline Calendars

The most current filing deadline calendars are available on the TMHP website at [www.tmhp.com](http://www.tmhp.com):

- Filing Deadline Calendar for 2020
- Filing Deadline Calendar for 2021
6.2 TMHP Electronic Claims Submission

TMHP uses the HIPAA-compliant American National Standards Institute (ANSI) ASC X12 5010 file format through secure socket layer (SSL) and virtual private networking (VPN) connections for maximum security.

Claims may be submitted electronically to TMHP through TexMedConnect on the TMHP website at www.tmhp.com or through billing agents who interface directly with the TMHP EDI Gateway.

Providers must retain all claim and file transmission records. They may be required to submit them for pending research on missing claims or appeals.

Refer to: “Section 3: TMHP Electronic Data Interchange (EDI)” (Vol. 1, General Information).

6.2.1 Benefit and Taxonomy Codes

Providers must submit the Benefit Code field (when applicable), Address field, and Taxonomy Code Field and all other required fields. These fields must be completed before submitting electronic claims.

Taxonomy codes do not affect pricing or the level of pricing, but rather are used to crosswalk the NPI to a TPI. It is critical that the taxonomy code selected as the primary or secondary taxonomy code during a providers enrollment with TMHP is included on all electronic transactions.

Billing providers that are not associated with a group are required to submit a taxonomy code on all electronic claims. Claims submitted without a taxonomy code may be rejected.

Medicare does not require a taxonomy code for Part B claims. Therefore, some claims submitted to TMHP from Medicare for payment of deductible or coinsurance may not include the taxonomy code needed for accurate processing by TMHP.

6.2.2 Electronic Claim Acceptance

Providers should verify that their electronic claims were accepted by Texas Medicaid for payment consideration by referring to their Claim Response report, which is in the 27S batch response file (e.g., file name E085LDS1.27S). Providers should also check their Accepted and Rejected reports in the rej and acc batch response files (e.g., E085LDS1.REJ and E085LDS1.ACC) for additional information. Only claims that have been accepted on the Claim Response report (27S file) will be considered for payment and made available for claim status inquiry. Claims that are rejected must be corrected and resubmitted for payment consideration.

Refer to: Subsection 3.2, “Electronic Billing” in “Section 3: TMHP Electronic Data Interchange (EDI)” (Vol. 1, General Information), visit www.tmhp.com, or call the EDI Help Desk at 1-888-863-3638 for more information about electronic claims submissions.

6.2.3 Electronic Rejections

The most common reasons for electronic professional claim rejections are:

- **Client information does not match.** Client information does not match the PCN on the TMHP eligibility file. The name, date of birth, sex, and nine-digit Medicaid identification number must be an exact match with the client’s identification number on TMHP’s eligibility record. If using TexMedConnect, send an interactive eligibility request to obtain an exact match with TMHP’s record. If not using TexMedConnect, verify through the TMHP website or call AIS at 1-800-925-9126 to verify client information. A lack of complete client eligibility information causes a rejection and possibly delayed payment. To prevent delays when submitting claims electronically:
  - Always include the first and last name of the client on the claim in the appropriate fields.
  - Always enter the client’s complete, valid nine-digit Medicaid number. Valid Medicaid numbers begin with 1, 2, 3, 4, 5, 6 or 7. CSHCN Services Program client numbers begin with a 9.
  - When submitting claims for newborns, use the guidelines in the following section.
• **Referring/Ordering Physician field blank or invalid.** The referring physician’s NPI must be present when billing for consultations, laboratory, or radiology. Consult the software vendor for this field’s location on the electronic claims entry form.

• **Performing Physician ID field blank or invalid.** When the billing provider identifier is a group practice, the performing provider identifier for the physician who performed the service must be entered. Consult the software vendor for this field’s location on the electronic claim form.

• **Facility Provider field blank or invalid.** When place of service (POS) is anywhere other than home or office, the facility’s provider identifier must be present. If the provider identifier is not known, enter the name and address of the facility. Consult the software vendor for this field’s location on the electronic claims entry form.

• **Invalid Type of Service or Invalid Type of Service/Procedure code combination.** In certain cases some procedure codes will require a modifier to denote the procedure’s type of service (TOS).

  **Note:** The C21 claims processing system can accept only 40 characters (including spaces) in the Comments section of electronic submissions for ambulance and dental claims. If providers include more than 40 characters in that field, C21 will accept only the first 40 characters; the other characters will not be imported into C21. Providers must ensure that all of the information that is required for the claim to process appropriately is included in the first 40 characters.

  **Refer to:** Subsection 6.2.5, “Modifier Requirements for TOS Assignment” in this section for TMHP EDI modifier information.

**6.2.3.1 Newborn Claim Hints**

The following are to be used for newborns:

• If the mother’s name is “Jane Jones,” use “Boy Jane Jones” for a male child and “Girl Jane Jones” for a female child.

• Enter “Boy Jane” or “Girl Jane” in first name field and “Jones” in last name field. Always use “boy” or “girl” first and then the mother’s full name. An exact match must be submitted for the claim to process.

• Do not use “NBM” for newborn male or “NBF” for newborn female.

The following are the most common reasons for electronic hospital UB-04 CMS-1450 claim rejections:

• **Admit hour outside allowable range** (such as 24 hours).

• **Billed amount blank.**

• **Health coverage ID blank or invalid.** This number must be the valid nine-digit Medicaid client number. Incorrect data includes: a number less than nine digits; PENDING; 999999999; and Unknown.

• **Referring physician information on outpatient claim is blank** when laboratory/radiology services are ordered or a surgical procedure is performed. The referring physician’s NPI is required in Fields 78–79. Consult the software vendor for the location of this field on the electronic claims entry form.
6.2.4  TMHP EDI Batch Numbers, Julian Dates
All electronic transactions are assigned an eight-character Batch ID immediately upon receipt by the TMHP EDI Gateway. The batch ID format allows electronic submitters to determine the exact day and year that a batch was received. The batch ID format is JJJYSSSS, where each character is defined as follows:

- **JJJ – Julian date.** The three J characters represent the Julian date that the file was received by the TMHP EDI Gateway. The first character (J) is displayed as a letter, where I = 0, J = 1, K = 2, and L = 3. The last two characters (JJ) are displayed as numbers. All three characters (JJJ) together represent the Julian date. For example, a Julian date of 143 would be J43.
- **Y – Year.** The Y character represents the last digit of the calendar year when the TMHP EDI Gateway receives the file. For example, a “2” in this position indicates the year 2012.
- **SSSS = The unique 4-character sequence number assigned by EDI to the batch filed.**

6.2.5  Modifier Requirements for TOS Assignment
Modifiers for TOS assignment are not required for Texas Health Steps (THSteps) Dental claims (claim type 021) and Inpatient Hospital claims (claim type 040). Additionally, procedures submitted by specific provider types such as genetics, eyeglass, and THSteps medical checkup are assigned the appropriate TOS based on the provider type or specific procedure code, and will not require modifiers.

Most procedure codes do not require a modifier for TOS assignment, but modifiers are required for some services submitted on professional claims (claim type 020) and outpatient hospital claims (claim type 023). Services that require a modifier for TOS assignment are listed in the following sections.

6.2.5.1  Assistant Surgery
For assistant surgical procedures, use one of the following modifiers: 80, 81, 82, and AS. Using these modifiers results in TOS 8 being assigned to the procedure.

6.2.5.2  Anesthesia
For anesthesia procedures, use one of the following modifiers: AA, AD, QK, QS, QX, QY, and QZ. Using these modifiers results in TOS 7 being assigned to the procedure.

6.2.5.3  Interpretations
For interpretations or professional components of laboratory, radiology, or radiation therapy procedures, use modifier 26. Using modifier 26 results in TOS I being assigned to the procedure.

*Note:* Procedure codes that only have a TOS I are not required to use modifier 26.

6.2.5.4  Technical Components
For technical components of laboratory, radiology, or radiation therapy procedures, use modifier TC. Using this modifier results in TOS T being assigned to the procedure.

*Exception:* Outpatient hospitals do not include the TC modifier when they provide technical components of lab and radiology services. These services automatically have TOS 4 or 5 assigned and are subject to the facility’s interim reimbursement rate or the clinical lab rate.
6.3 Coding

Electronic billers must code all claims. TMHP encourages all providers to code their paper claims. Claims are processed fast and accurately if providers furnish appropriate information. By coding claims, providers ensure precise and concise representation of the services provided and are assured reimbursement based on the correct code. If providers code claims, a narrative description is not required and does not need to be included unless the code is a not an otherwise classified code.

Important: Claims for anesthesia must have the CPT anesthesia procedure code narrative descriptions or CPT surgical codes; if these codes are not included, the claim will be denied.

The carrier for the Texas Medicare Program has coding manuals available for physicians and suppliers with codes not available in CPT. To order a CPT Coding Manual, write to the following address:

American Medical Association
Book and Pamphlet Fulfillment
PO Box 2964
Milwaukee, WI 53201

6.3.1 Diagnosis Coding

Texas Medicaid requires providers to provide International Classification of Diseases, Tenth Revision, Clinical Modification (ICD-10-CM) diagnosis codes on their claims. The only diagnosis coding structure accepted by Texas Medicaid is the ICD-10-CM. Diagnosis codes must be to the highest level of specificity available. In most cases a written description of the diagnosis is not required.

All diagnosis codes that are submitted on a claim must be appropriate for the age of the client as identified in the ICD-10-CM description of the diagnosis code. Claims that are denied because one or more of the diagnosis codes submitted on the claim are not appropriate for the age of the client may be appealed with the correct diagnosis code or documentation of medical necessity to justify the use of the diagnosis code.

Diagnosis codes in the following categories are not valid as primary or referenced diagnosis:

- Nonspecific injury, poisoning and other consequences of external causes
- Diagnosis in the International Classification of Diseases for Oncology, 3rd Edition (ICD-O-3)
- Factors influencing health status and contact with health services, unless otherwise directed in this manual.
- External causes of morbidity

6.3.1.1 Place of Service (POS) Coding

The POS identifies where services are performed. Indicate the POS by using the appropriate code for each service identified on the claim.

Important: Attention ambulance providers: POS 41 and 42 are accepted by Texas Medicaid for ambulance claims processing. The two-digit origin and destination codes are still required for claims processing.

Use the following codes for POS identification where services are performed:

<table>
<thead>
<tr>
<th>POS</th>
<th>2-Digit Numeric Codes (Electronic Billers)</th>
<th>1-Digit Numeric Codes (Paper Billers)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Office</td>
<td>02, 11, 15, 17, 20, 49, 50, 60, 65, 71, 72</td>
<td>1</td>
</tr>
<tr>
<td>Home</td>
<td>12</td>
<td>2</td>
</tr>
</tbody>
</table>
6.3.2 Type of Service (TOS)

The TOS identifies the specific field or specialty of services provided.

To determine the TOS payable for each procedure code, providers may refer to the online fee lookup (OFL) or the static fee schedules, both are available on the TMHP website at www.tmhp.com.

Refer to: Subsection 6.2.5, “Modifier Requirements for TOS Assignment” in this section for TMHP EDI modifier information.

6.3.2.1 TOS Table

Important: TOS codes are not used for claim submissions, but they do appear on Re&S Reports.

<table>
<thead>
<tr>
<th>TOS</th>
<th>Description</th>
<th>2-Digit Numeric Codes (Electronic Billers)</th>
<th>1-Digit Numeric Codes (Paper Billers)</th>
</tr>
</thead>
<tbody>
<tr>
<td>0</td>
<td>Blood</td>
<td></td>
<td></td>
</tr>
<tr>
<td>1</td>
<td>Medical Services</td>
<td>21, 51, 52, 61</td>
<td>3</td>
</tr>
<tr>
<td>2</td>
<td>Surgery</td>
<td>19, 22, 23, 24, 55, 56, 57, 62</td>
<td>5</td>
</tr>
<tr>
<td>3</td>
<td>Consultations</td>
<td>25</td>
<td>7</td>
</tr>
<tr>
<td>4</td>
<td>Radiology (total component)</td>
<td>01, 03, 04, 05, 06, 07, 08, 16, 18, 26, 34, 41, 42, 53, 99</td>
<td>9</td>
</tr>
<tr>
<td>5</td>
<td>Laboratory (total component)</td>
<td></td>
<td></td>
</tr>
<tr>
<td>6</td>
<td>Radiation Therapy (total component)</td>
<td>13, 31, 32, 54</td>
<td>4</td>
</tr>
<tr>
<td>7</td>
<td>Anesthesia</td>
<td>14, 33</td>
<td>8</td>
</tr>
<tr>
<td>8</td>
<td>Assistant surgery</td>
<td>81</td>
<td>6</td>
</tr>
<tr>
<td>9</td>
<td>Other medical items or services</td>
<td></td>
<td></td>
</tr>
<tr>
<td>C</td>
<td>Home health services</td>
<td></td>
<td></td>
</tr>
<tr>
<td>D</td>
<td>TB clinic</td>
<td></td>
<td></td>
</tr>
<tr>
<td>E</td>
<td>Eyeglasses</td>
<td></td>
<td></td>
</tr>
</tbody>
</table>
6.3.3 Procedure Coding

Texas Medicaid uses the Healthcare Common Procedure Coding System (HCPCS). HCPCS provides health-care providers and third-party payers a common coding structure that uses codes designed around a five-character numeric or alphanumeric base. The procedure codes are updated annually and quarterly.

HCPCS consists of two levels of codes:

  - Numeric, five digits
  - Makes up 80 percent of HCPCS
  - Maintained by AMA, which updates it annually
  - Updates by the AMA are coordinated with CMS before modifications are distributed to third-party payers
  - Anesthesia codes from CPT

- **Level II**—HCPCS
  - Approved and released by CMS
  - Codes for both physician and non-physician services not contained in CPT (for example, ambulance, DME, prosthetics, and some medical codes)
  - Maintained and updated by the CMS Maintenance Task Force
  - Alphanumeric, a single alpha character (A through V) followed by four digits
  - The single alpha character represents one of the following:

<table>
<thead>
<tr>
<th>Alpha</th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td>A</td>
<td>Supplies, ambulance, administrative, miscellaneous</td>
</tr>
<tr>
<td>B</td>
<td>Enteral and parenteral therapy</td>
</tr>
<tr>
<td>E</td>
<td>DME and oxygen</td>
</tr>
<tr>
<td>G</td>
<td>Procedures/professional (temporary)</td>
</tr>
<tr>
<td>H</td>
<td>Rehab and behavioral health services</td>
</tr>
<tr>
<td>J</td>
<td>Drugs (administered other than orally)</td>
</tr>
<tr>
<td>K</td>
<td>Durable Medical Equipment Regional Carriers (DMERC)</td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th>TOS</th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td>F</td>
<td>Ambulatory surgical center (ASC)/hospital-based ambulatory surgical center (HASC)</td>
</tr>
<tr>
<td>G</td>
<td>Genetics</td>
</tr>
<tr>
<td>I</td>
<td>Professional component for radiology, laboratory, or radiation therapy</td>
</tr>
<tr>
<td>J</td>
<td>DME purchase new</td>
</tr>
<tr>
<td>L</td>
<td>DME rental</td>
</tr>
<tr>
<td>R</td>
<td>Hearing aid</td>
</tr>
<tr>
<td>S</td>
<td>THSteps medical</td>
</tr>
<tr>
<td>T</td>
<td>Technical component for radiology, laboratory, or radiation therapy</td>
</tr>
<tr>
<td>W</td>
<td>THSteps dental</td>
</tr>
</tbody>
</table>
6.3.3.1 **HCPCS Updates**

TMHP updates HCPCS codes on both an annual and quarterly basis. Major updates are made annually and minor updates are made quarterly.

Most of the procedure codes that do not replace a discontinued procedure code must go through the rate hearing process. HHSC conducts public rate hearings to provide an opportunity for the provider community to comment on the Medicaid proposed payment rate, as required by Chapter 32 of the Human Resources Code, §32.0282, and Title 1 of the Texas Administrative Code, §355.201.

6.3.3.1.1 Annual HCPCS

Annual HCPCS updates apply additions, changes, and deletions that include the program and coding changes related to the annual HCPCS, Current Dental Terminology (CDT), and CPT updates. These updates ensure that the coding structure is up-to-date by using the latest edition of the CPT and the nationally established HCPCS codes that are released by CMS.

6.3.3.1.2 Quarterly HCPCS

Quarterly HCPCS updates apply HCPCS additions, changes, and deletions that are released by CMS.

6.3.3.1.3 Rate Hearings for New HCPCS Codes

HHSC holds rate hearings for new HCPCS codes on a regular basis. Rate hearings are announced on the HHSC website at [https://hhs.texas.gov/services/health/medicaid-chip/provider-information/texas-medicaid-chip-rate-analysis](https://hhs.texas.gov/services/health/medicaid-chip/provider-information/texas-medicaid-chip-rate-analysis).

Claims for services that are provided before the rates are adopted through the rate hearing process are denied as pending a rate hearing (EOB 02008) until the applicable reimbursement rate is adopted. The client cannot be billed for these services.

Providers are responsible for meeting the initial 95-day filing deadline. Providers must submit the procedure codes that are most appropriate for the services provided, even if the procedure codes have not yet completed the rate hearing process and are denied by Texas Medicaid as pending a rate hearing.

Once the reimbursement rates are established in the rate hearing and applied, TMHP automatically reprocesses affected claims. Providers are not required to appeal the claims unless they are denied for other reasons after the claims reprocessing is complete.

**Refer to:** Subsection 5.11, “Guidelines for Procedures Awaiting Rate Hearing” in “Section 5: Fee-for-Service Prior Authorizations” (Vol. 1, General Information) for more information about the authorization guidelines for procedure codes that are awaiting a rate hearing.

6.3.4 **National Drug Code (NDC)**

The NDC is an 11-digit number on the package or container from which the medication is administered. All Texas Medicaid fee-for-service and Family Planning providers must submit an NDC for professional or outpatient claims submitted with physician-administered prescription drug procedure.

<table>
<thead>
<tr>
<th>Alpha</th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td>L</td>
<td>Orthotic and prosthetic procedures</td>
</tr>
<tr>
<td>M</td>
<td>Medical</td>
</tr>
<tr>
<td>P</td>
<td>Laboratory</td>
</tr>
<tr>
<td>Q</td>
<td>Temporary procedures</td>
</tr>
<tr>
<td>R</td>
<td>Radiology</td>
</tr>
<tr>
<td>S</td>
<td>Private payer</td>
</tr>
<tr>
<td>T</td>
<td>State Medicaid agency</td>
</tr>
<tr>
<td>V</td>
<td>Vision and hearing services</td>
</tr>
</tbody>
</table>
N4 must be entered before the NDC on claims.

National Drug Unit of Measure: The submitted unit of measure should reflect the volume measurement administered. Refer to the NDC Package Measure column on the Texas NDC-to-HCPCS Crosswalk.

The valid units of measurement codes are:
- F2—International unit
- GR—Gram
- ME—Milligram
- ML—Milliliter
- UN—Unit

*Note: Unit quantities are required.*

### 6.3.4.1 Paper Claim Submissions

<table>
<thead>
<tr>
<th>Block No.</th>
<th>Description</th>
<th>Guidelines</th>
</tr>
</thead>
</table>
| 43        | Revenue codes and description        | This block should include the following elements in the following order:                                                                                                                   • NDC qualifier of N4 (e.g., N4)   • The 11-digit NDC number on the package or vial from which the medication was administered. Do not enter hyphens or spaces within this number (e.g., 00409231231). • The unit of measurement code. There are 5 allowed values: F2, GR, ML, UN, or ME (e.g., GR). • The unit quantity with a floating decimal for fractional units (limited to 3 digits, e.g., 0.025).  

*Example: N400409231231GR0.025*

<table>
<thead>
<tr>
<th>Block No.</th>
<th>Description</th>
<th>Guidelines</th>
</tr>
</thead>
</table>
| 24A       | Dates of service                     | In the shaded area, enter the:                                                                                                                   • NDC qualifier of N4 (e.g., N4)   • The 11-digit NDC number on the package or vial from which the medication was administered. Do not enter hyphens or spaces within this number (e.g., 00409231231).  

*Example: N400409231231*

<table>
<thead>
<tr>
<th>Block No.</th>
<th>Description</th>
<th>Guidelines</th>
</tr>
</thead>
<tbody>
<tr>
<td>24D</td>
<td>Procedures, services, or supplies</td>
<td>In the shaded area, enter the NDC quantity of units administered (up to 12 digits, including the decimal point.). A decimal point must be used for fractions of a unit (e.g., 0.025).</td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th>Block No.</th>
<th>Description</th>
<th>Guidelines</th>
</tr>
</thead>
<tbody>
<tr>
<td>24G</td>
<td>Days or units</td>
<td>In the shaded area, enter the NDC unit of measurement code. There are 5 allowed values: F2, GR, ML, UN, or ME.</td>
</tr>
</tbody>
</table>
2017 Claim Form

<table>
<thead>
<tr>
<th>Block No.</th>
<th>Description</th>
<th>Guidelines</th>
</tr>
</thead>
<tbody>
<tr>
<td>32A</td>
<td>Dates of service</td>
<td>In the shaded area, enter the:</td>
</tr>
<tr>
<td></td>
<td></td>
<td>• NDC qualifier of N4 (e.g., N4)</td>
</tr>
<tr>
<td></td>
<td></td>
<td>• The 11-digit NDC number on the package or vial from which the medication was administered. Do not enter hyphens or spaces within this number (e.g., 00409231231).</td>
</tr>
<tr>
<td></td>
<td></td>
<td><strong>Example:</strong> N400409231231</td>
</tr>
<tr>
<td>32D</td>
<td>Procedures, services, or supplies Current Procedural Terminology (CPT)/ HCPCS</td>
<td>In the shaded area, enter the NDC quantity of units administered (up to 12 digits, including the decimal point). A decimal point must be used for fractions of a unit (e.g., 0.025).</td>
</tr>
<tr>
<td>32F</td>
<td>Days or units</td>
<td>In the shaded area, enter the NDC unit of measurement code. There are 5 allowed values: F2, GR, ML, UN, or ME.</td>
</tr>
</tbody>
</table>

### National Drug Unit

Claims will be edited for the value submitted in the NDC quantity field. In order to convert the HCPCS units submitted into the NDC quantity; use the Texas NDC-to-HCPCS Crosswalk to review the “HCPCS Description” and the “NDC Label” description to identify the quantity.

The Texas NDC-to-HCPCS Crosswalk identifies relationships between HCPCS codes and National Drug Codes (NDC). The Texas file is published at least quarterly. The Texas NDC-to-HCPCS Crosswalk can be found at [https://www.txvendordrug.com/formulary/clinician-administered-drugs](https://www.txvendordrug.com/formulary/clinician-administered-drugs). Clinician-administered drugs that do not have an appropriate NDC to HCPCS combination for the procedure code that is submitted are not payable.

#### 6.3.4.2 NDC Requirements for Dual Eligible Clients

The 11-digit NDC, NDC quantity, and NDC Unit of measure information is required on all professional and outpatient clinician-administered drug claims for dual-eligible clients. These drug claims are submitted to Medicare, which will cross over to Medicaid for consideration of coinsurance and deductible liabilities.

*Important:* Claims which cross over without this required information may be denied due to missing, incomplete, or invalid NDC information. This information applies to all Medicaid providers who serve Medicare-Medicaid dual-eligible clients.

Providers may refer to subsection 6.3.4, “National Drug Code (NDC)” in this section for more information on NDC requirements. The [Texas NDC-to-HCPCS Crosswalk](https://www.txvendordrug.com/formulary/clinician-administered-drugs) identifies relationships between HCPCS codes.

#### 6.3.4.3 * Drug Rebate Program

Texas Medicaid will reimburse providers only for clinician-administered drugs and biologicals whose manufacturers participate in the Centers for Medicare & Medicaid Services (CMS) Drug Rebate Program and that show as active on the CMS list for the date of service the drug is administered.

CMS maintains a list of participating manufacturers and their rebate-eligible drug products, which is updated quarterly on the [CMS website](https://www.cms.gov). TMHP will republish this list quarterly in a more accessible format.
When providers submit claims for clinician-administered drug procedure codes, they must include the National Drug Code (NDC) of the administered drug as indicated on the drug packaging. While 340B purchased claims are not eligible for drug rebates, NDCs are required to receive federal funding to pay the claim.

TMHP will deny claims for drug procedure codes under the following circumstances:

- The NDC submitted with the drug procedure code is not on the CMS drug rebate list that was current on the date of service.
- The NDC submitted with the drug procedure code has been terminated.
- The drug procedure code is submitted with a missing or invalid NDC.

To avoid claim denials, providers must speak with the pharmacy or wholesaler with whom they work to ensure the product purchased is on the current CMS list of participating manufacturers and their drugs.

**Note:** Texas Medicaid managed care organizations (MCOs) have their own policies and procedures regarding clinician-administered drugs. Providers must contact the client’s MCO for benefit and limitation information.

Providers can find a complete, downloadable list of procedure codes and the corresponding descriptions on the Vendor Drug Program website at [www.txvendordrug.com](http://www.txvendordrug.com).

Vitamins and minerals procedure codes will be listed on a separate tab of the supplemental file.

### 6.3.5 Modifiers

Modifiers describe and qualify the services provided by Texas Medicaid. A modifier is placed after the five-digit procedure code. Up to two modifiers may apply per service. Examples of frequently used modifiers are listed in the following table. Refer to the service-specific sections for additional modifier requirements.

<table>
<thead>
<tr>
<th>Modifier</th>
<th>Special Instructions/Notes (if applicable)</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>340B Drug Rebate Program</strong></td>
<td></td>
</tr>
<tr>
<td>U8</td>
<td>All eligible organizations and covered entities that are enrolled in the federal 340B Drug Pricing Program to purchase 340B discounted drugs must use modifier U8 when submitting claims for 340B clinician-administered drugs. Non-compliance with this new requirement to use modifier U8 on all claims submitted for 340B clinician-administered drugs may jeopardize a covered entity’s 340B status with the U.S. Department of Health and Human Services Health Resources and Services Administration (HRSA). Providers can refer to the HRSA website at <a href="http://www.hrsa.gov/opa/index.html">www.hrsa.gov/opa/index.html</a> for more information about the 340B Drug Pricing Program.</td>
</tr>
<tr>
<td><strong>Ambulance</strong></td>
<td></td>
</tr>
<tr>
<td>ET</td>
<td>Use for all emergency transport services.</td>
</tr>
<tr>
<td>GY</td>
<td>Use to indicate that no medical necessity existed for a transport.</td>
</tr>
<tr>
<td><strong>Surgeons</strong></td>
<td></td>
</tr>
<tr>
<td>53</td>
<td>Use for physician reporting of a discontinued procedure. For outpatient/ASC reporting of a discontinued procedure, see modifier 73 and 74.</td>
</tr>
<tr>
<td>54+</td>
<td>Surgeon who performs the surgical procedure only must bill the surgical code with modifier 54 and is reimbursed 70% of the global fee.</td>
</tr>
<tr>
<td>55+</td>
<td>Provider who performs the postoperative care only must bill the surgical code with modifier 55 and is reimbursed 20% of the global fee.</td>
</tr>
</tbody>
</table>

* Modifier is required for accurate claims processing.
* Description is defined by the state.
<table>
<thead>
<tr>
<th>Modifier</th>
<th>Special Instructions/Notes (if applicable)</th>
</tr>
</thead>
<tbody>
<tr>
<td>56+</td>
<td>Providers who perform the preoperative care only must bill the surgical code with modifier 56 and is reimbursed 10 percent of the global fee.</td>
</tr>
<tr>
<td>58+</td>
<td>Staged or related procedure or services by the same physician during the postoperative period.</td>
</tr>
<tr>
<td>62+</td>
<td>Cosurgery. Two surgeons perform the specific procedure(s).</td>
</tr>
<tr>
<td>76+</td>
<td>Use modifier 76 or 77 for transplant procedures if it is a second transplant of the same organ.</td>
</tr>
<tr>
<td>77+</td>
<td>Use modifier 76 or 77 for transplant procedures if it is a second transplant of the same organ.</td>
</tr>
<tr>
<td>78+</td>
<td>Return to the operating room for a related procedure during the postoperative period.</td>
</tr>
<tr>
<td>79+</td>
<td>Unrelated procedure or service by the same physician during the postoperative period.</td>
</tr>
<tr>
<td></td>
<td><strong>Assistant Surgeons</strong></td>
</tr>
</tbody>
</table>
| 80 and KX+ | Use modifier 80 and KX together to indicate an assistant surgeon in a teaching facility:  
  • In a case involving exceptional medical circumstances such as emergency or life-threatening situations requiring immediate attention.  
  • When the primary surgeon has a policy of never, without exception, involving a resident in the preoperative, operative, or postoperative care of one of his or her patients.  
  • In a case involving a complex surgical procedure that qualifies for more than one physician. |
| AS       | Use when the physician assistant is not enrolled as an individual provider and provides assistance at surgery. |
|          | **Sterilizations** |
| PM       | Use to indicate post-menopausal. |
| PS       | Use to indicate previously sterilized. |
|          | **Excision of Lesions/Masses** |
| KX+      | Use modifier KX if the excision/destruction is due to one of the following signs or symptoms: inflamed, infected, bleeding, irritated, growing, limiting motion or function. Use of this modifier is subject to retrospective review. |
|          | **Injections** |
| AT       | Use to indicate acute conditions. |
| JA       | Administered intravenously. |
| JB       | Administered subcutaneously. |

+ Modifier is required for accurate claims processing.  
* Description is defined by the state.
<table>
<thead>
<tr>
<th>Modifier</th>
<th>Special Instructions/Notes (if applicable)</th>
</tr>
</thead>
</table>
| KX+      | Use modifier KX to indicate the injection was due to:  
  - Oral route contraindicated or an acceptable oral equivalent is not available.  
  - Injectable medication is the accepted treatment of choice. Oral medication regimens have proven ineffective or are not available.  
  - Patient has a temperature over 102 degrees (documented on the claim) and a high level of antibiotic is needed quickly.  
  - Injection is medically necessary into joints, bursae, tendon sheaths, or trigger points to treat an acute condition or the acute flare up of a chronic condition. |

**Visits**

<table>
<thead>
<tr>
<th>Code</th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td>76+</td>
<td>Use to indicate the repeated non-clinical procedure.</td>
</tr>
<tr>
<td>FP+</td>
<td>Use to indicate that the service was part of an annual family planning examination.</td>
</tr>
<tr>
<td>TH+</td>
<td>Use with external causes of injury and poisoning (E Codes) procedures and morphology of neoplasms (M Codes) procedures to specify antepartum or postpartum care.</td>
</tr>
<tr>
<td>25</td>
<td>Use to describe circumstances in which an office visit was provided at the same time as other separately identifiable services. Refer to the CMS NCCI website for additional information.</td>
</tr>
</tbody>
</table>

**Anesthesia**

One of the following modifier combinations must be used by anesthesiologists directing non-CRNA qualified professionals.

<table>
<thead>
<tr>
<th>AA and U1</th>
<th>Use to indicate that the anesthesia services were performed personally by the anesthesiologist.</th>
</tr>
</thead>
<tbody>
<tr>
<td>AD and U1 (Emergency circumstances only)</td>
<td>Use when directing five or more concurrent procedures provided by non-CRNA qualified professionals. Used in emergency circumstances only and limited to 6 units (90 minutes) per case for each occurrence requiring five or more concurrent procedures.</td>
</tr>
<tr>
<td>QK and U1</td>
<td>Use when directing two, three, or four concurrent procedures provided by non-CRNA qualified professionals.</td>
</tr>
<tr>
<td>QY and U1</td>
<td>Use when directing one procedure provided by a non-CRNA qualified professional.</td>
</tr>
</tbody>
</table>

One of the following modifier combinations must be used by anesthesiologists directing CRNAs.

<table>
<thead>
<tr>
<th>AD and U2 (Emergency circumstances only)</th>
<th>Use when directing five or more concurrent procedures involving CRNA (s). Used in emergency circumstances only and limited to 6 units (90 minutes) per case for each occurrence requiring five or more concurrent procedures.</th>
</tr>
</thead>
<tbody>
<tr>
<td>QK and U2</td>
<td>Use when directing two, three, or four concurrent procedures involving CRNAs.</td>
</tr>
<tr>
<td>QY and U2</td>
<td>Use when directing one procedure by a CRNA.</td>
</tr>
</tbody>
</table>

One of the following modifier combinations must be used by CRNAs.

| QX and U2 | Use to indicate the anesthesia was medically directed by the anesthesiologist. |
| QZ and U1 | Use to indicate the anesthesia was directed by the surgeon. |

**DME**

For DME, use one of the following modifiers:

<table>
<thead>
<tr>
<th>Code</th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td>NU</td>
<td>For DME purchase new</td>
</tr>
<tr>
<td>RR</td>
<td>For DME rental- monthly</td>
</tr>
</tbody>
</table>

* Modifier is required for accurate claims processing.  
  * Description is defined by the state.
<table>
<thead>
<tr>
<th>Modifier</th>
<th>Special Instructions/Notes (if applicable)</th>
</tr>
</thead>
<tbody>
<tr>
<td>UE</td>
<td>For DME other-purchase-used</td>
</tr>
</tbody>
</table>

**FQHC and RHC**

Services provided by a health-care professional require one of the following modifiers:

<table>
<thead>
<tr>
<th>Modifier</th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td>AH</td>
<td>Use to indicate that the services were performed by a clinical psychologist.</td>
</tr>
<tr>
<td>AJ</td>
<td>Use to indicate that the services were performed by a clinical social worker.</td>
</tr>
<tr>
<td>AM</td>
<td>Use to indicate that the services were performed by a physician or team member service (includes clinical psychiatrist).</td>
</tr>
<tr>
<td>SA</td>
<td>Use to indicate that the services were performed by an advanced practice registered nurse (APRN) or CNM rendering services in collaboration with a physician.</td>
</tr>
<tr>
<td>TD</td>
<td>For home services performed by a RN and provided in areas with a shortage of home health agencies.</td>
</tr>
<tr>
<td>TE</td>
<td>For home services performed by an LVN and provided in areas with a shortage of home health agencies.</td>
</tr>
<tr>
<td>TS</td>
<td>Use to indicate a case management follow-up service.</td>
</tr>
<tr>
<td>U1</td>
<td>Licensed professional counselor</td>
</tr>
<tr>
<td>U2</td>
<td>Licensed marriage and family therapist</td>
</tr>
<tr>
<td>U7*</td>
<td>Physician assistant services for other than assistant at surgery</td>
</tr>
</tbody>
</table>

The following modifiers may be used in addition to the modifier identifying the health-care professional that rendered the service:

<table>
<thead>
<tr>
<th>Modifier</th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td>EP</td>
<td>Use to indicate THSteps services (FQHC only).</td>
</tr>
<tr>
<td>FP</td>
<td>Use to indicate that the service was part of an annual family planning examination.</td>
</tr>
<tr>
<td>TH</td>
<td>Use to indicate the encounter is for antepartum care or postpartum care.</td>
</tr>
<tr>
<td>U5*</td>
<td>State-defined modifier for use with case management services.</td>
</tr>
</tbody>
</table>

**Abortion**

<table>
<thead>
<tr>
<th>Modifier</th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td>G7</td>
<td>Use by performing physicians, facilities, anesthesiologists, and CRNAs (with appropriate procedure code) when requesting reimbursement for abortion procedures that are within the scope of the rules and regulations of Texas Medicaid.</td>
</tr>
</tbody>
</table>

**Vision**

<table>
<thead>
<tr>
<th>Modifier</th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td>RB</td>
<td>Use modifier RB to indicate replacement of prosthetic or nonprosthetic eyeglasses or contact lenses.</td>
</tr>
<tr>
<td>VP+</td>
<td>Use when billing prosthetic eyeglasses or contact lenses with a diagnosis of aphakia.</td>
</tr>
</tbody>
</table>

**Laboratory/Radiology**

<table>
<thead>
<tr>
<th>Modifier</th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td>26+</td>
<td>Use for laboratory interpretations and radiological procedures.</td>
</tr>
<tr>
<td>59-</td>
<td>Code to indicate the procedure or service was independent from other services performed on the same day. Refer to the CMS NCCI website for additional information.</td>
</tr>
<tr>
<td>91+</td>
<td>Use for repeat laboratory clinical test.</td>
</tr>
<tr>
<td>76</td>
<td>Use for repeat laboratory nonclinical test.</td>
</tr>
<tr>
<td>SU+</td>
<td>Indicates necessary equipment is in physician’s office for RAST/MAST testing or Pap smears.</td>
</tr>
<tr>
<td>TC+</td>
<td>The modifier TC is used for technical radiological procedures.</td>
</tr>
<tr>
<td>Q4+</td>
<td>Use for lab/radiology/ultrasound interps by other than the attending physician.</td>
</tr>
</tbody>
</table>

* Modifier is required for accurate claims processing.  
* Description is defined by the state.
<table>
<thead>
<tr>
<th>Modifier</th>
<th>Special Instructions/Notes (if applicable)</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>Therapy</strong></td>
<td></td>
</tr>
<tr>
<td>AT+</td>
<td>Must be used to indicate the necessity of an acute condition for occupational therapy (OT), physical therapy (PT), osteopathic manipulation treatment (OMT), or chiropractic services.</td>
</tr>
<tr>
<td>GN</td>
<td>Use to indicate outpatient speech language pathology.</td>
</tr>
<tr>
<td>GO</td>
<td>Use to indicate outpatient occupational therapy.</td>
</tr>
<tr>
<td>GP</td>
<td>Use to indicate outpatient PT.</td>
</tr>
<tr>
<td>U4*</td>
<td>Reassessment</td>
</tr>
<tr>
<td><strong>THSteps Medical</strong></td>
<td></td>
</tr>
<tr>
<td>AM</td>
<td>Physician, team member service</td>
</tr>
<tr>
<td>EP</td>
<td>FQHCs must use modifier EP for services provided under THSteps.</td>
</tr>
<tr>
<td>SA</td>
<td>Nurse practitioner rendering service in collaboration with a physician</td>
</tr>
<tr>
<td>U5*</td>
<td>Intermediate oral examination with dental varnish</td>
</tr>
<tr>
<td>U7*</td>
<td>Physician assistant services for other than assistant at surgery</td>
</tr>
<tr>
<td>TD</td>
<td>Registered nurse</td>
</tr>
<tr>
<td><strong>THSteps Exceptions to Periodicity</strong></td>
<td></td>
</tr>
<tr>
<td>SC</td>
<td>Medically necessary service or supply</td>
</tr>
<tr>
<td>23</td>
<td>Unusual Anesthesia: Occasionally, a procedure, which usually requires either no anesthesia or local anesthesia, because of unusual circumstances must be done under general anesthesia. This circumstance may be reported by adding the modifier 23 to the procedure code of the basic service or by use of the separate five-digit modifier code 09923</td>
</tr>
<tr>
<td>32</td>
<td>Mandated Services: Services related to mandated consultation or related services (e.g., peer review organization [PRO], third party payer, governmental, legislative or regulatory requirement) may be identified by adding the modifier 32 to the basic procedure or the service may be reported by use of the five digit modifier 09932</td>
</tr>
<tr>
<td><strong>Physicians</strong></td>
<td></td>
</tr>
<tr>
<td>Q5</td>
<td>Informal reciprocal arrangement (period not to exceed 14 continuous days)</td>
</tr>
<tr>
<td>Q6</td>
<td>Locum tenens or temporary arrangement (up to 90 days)</td>
</tr>
<tr>
<td><strong>Radiology Services</strong></td>
<td></td>
</tr>
<tr>
<td>U6</td>
<td>CT, CTA, MRI, MRA, Cardiac Nuclear Imaging, and PET Scan studies provided in the emergency department. Obstetric ultrasounds provided in the emergency department or during a hospital observation stay.</td>
</tr>
<tr>
<td><strong>Durable Medical Equipment</strong></td>
<td></td>
</tr>
<tr>
<td>NU</td>
<td>Use to indicate purchased equipment.</td>
</tr>
<tr>
<td>RR</td>
<td>Use to indicate leased equipment.</td>
</tr>
<tr>
<td><strong>Telemedicine/Telehealth</strong></td>
<td></td>
</tr>
<tr>
<td>95</td>
<td>Use with appropriate evaluation and management codes.</td>
</tr>
</tbody>
</table>

* Modifier is required for accurate claims processing.  
* Description is defined by the state.
The following modifiers may appear on R&S Reports (they are not entered by the provider):

- **PT.** The DRG payment was calculated on a per diem basis for an inpatient stay because of patient transfer.
- **PS.** The DRG payment was calculated on a per diem basis because the patient exhausted the 30-day inpatient benefit limitation during the stay.
- **PE.** The DRG payment was calculated on a per diem basis because the patient was ineligible for Medicaid during part of the stay. Also used to adjudicate claims with adjustments to outlier payments.

### 6.3.6 Benefit Code

A benefit code is an additional data element used to identify state programs.

Providers that participate in the following programs must use the associated benefit code when submitting claims and authorizations:

<table>
<thead>
<tr>
<th>Program</th>
<th>Benefit Code</th>
</tr>
</thead>
<tbody>
<tr>
<td>Comprehensive Care Program (CCP)</td>
<td>CCP</td>
</tr>
<tr>
<td>THSteps Medical</td>
<td>EP1</td>
</tr>
<tr>
<td>THSteps Dental</td>
<td>DE1</td>
</tr>
<tr>
<td>Family Planning Agencies*</td>
<td>FP3</td>
</tr>
<tr>
<td>Hearing Aid Dispensers</td>
<td>HA1</td>
</tr>
<tr>
<td>Maternity</td>
<td>MA1</td>
</tr>
<tr>
<td>County Indigent Health Care Program</td>
<td>CA1</td>
</tr>
<tr>
<td>Early Childhood Intervention (ECI) Providers</td>
<td>EC1</td>
</tr>
<tr>
<td>Tuberculosis (TB) Clinics</td>
<td>TB1</td>
</tr>
<tr>
<td>IDD case management providers</td>
<td>MH2</td>
</tr>
</tbody>
</table>

*Agencies only—Benefit codes should not be used for individual family planning providers.

### 6.4 Claims Filing Instructions

This section contains instructions for completion of Medicaid-required claim forms. When filing a claim, providers should review the instructions carefully and complete all requested information. A correctly completed claim form is processed faster.

This section provides a sample claim form and its corresponding instruction table for each acceptable Texas Medicaid claim form.

All providers, except those on prepayment review, should submit paper claims to TMHP to the following address:

Texas Medicaid & Healthcare Partnership
Claims
PO Box 200555
Austin, TX 78720-0555
Providers on prepayment review must submit all paper claims and supporting medical record documentation to the following address:

Texas Medicaid & Healthcare Partnership
Attention: Prepayment Review MC–A11 SURS
PO Box 203638
Austin, TX 78720-3638

6.4.1 National Correct Coding Initiative (NCCI) Guidelines

The Patient Protection and Affordable Care Act (PPACA) mandates that all claims that are submitted to TMHP be filed in accordance with the NCCI guidelines, including claims for services that have been prior authorized or authorized with medical necessity documentation.

The following NCCI MUE limitations have been deactivated as approved by CMS:

<table>
<thead>
<tr>
<th>Procedure Codes</th>
<th>Description</th>
<th>Deactivated Limitation (per date of service)</th>
<th>Approved Limitations</th>
</tr>
</thead>
<tbody>
<tr>
<td>A4281, A4282, A4284, A4286</td>
<td>Breast pump replacement parts</td>
<td>1</td>
<td>2 of each part per rolling year</td>
</tr>
<tr>
<td>H0005</td>
<td>Group therapy for substance use disorder treatment</td>
<td>1</td>
<td>135 units per calendar year</td>
</tr>
</tbody>
</table>

The CMS NCCI and MUE guidelines can be found on the CMS website at [www.cms.gov](http://www.cms.gov).

The NCCI guidelines consist of HCPCS or CPT procedure code pairs that must not be reported together and MUEs that determine whether procedure codes are submitted in quantities that are unlikely to be correct.

The NCCI and MUE spreadsheets are published and updated by CMS and are available on the CMS Medicaid NCCI Coding web page under “NCCI and MUE Edits” as follows:

- **NCCI edit spreadsheets.** The website contains the Medicaid NCCI edit spreadsheet for hospital services and the Medicaid NCCI edit spreadsheet for practitioner services. The spreadsheets list the procedure code pairs that will not be reimbursed separately if they are billed by the same provider with the same date of service. Column 1 procedure codes may be reimbursed and Column 2 procedure codes will be denied. The spreadsheets also contain a column that indicates whether or not a modifier is allowed for services that may be reimbursed separately.

- **MUE edit spreadsheets.** The website contains the Medicaid MUE edit spreadsheets for hospital services, practitioner services, and supplier services. The spreadsheets list procedure codes and the number of units that may be reimbursed for each procedure code. Units that are submitted beyond these limitations will be denied.

**Note:** Providers are required to comply with NCCI and MUE guidelines as well as the guidelines that are published in the Texas Medicaid Provider Procedures Manual, all currently published website articles, fee schedules, and all other application information published on the TMHP website at [www.tmhp.com](http://www.tmhp.com). In instances when Texas Medicaid medical policy is more restrictive than NCCI or MUE guidance, Texas Medicaid medical policy prevails.

HHSC continue to implement and enforce correct coding initiatives. Providers may see additional claim denials related to NCCI and MUE edits including those services that were prior authorized or authorized with medical necessity documentation.
If a rendered service does not comply with a guideline as defined by NCCI, medical necessity documentation may be submitted with the claim for the service to be considered for reimbursement; however, medical necessity documentation does not guarantee payment for the service.

**Important:** Prior authorization and authorization based on documentation of medical necessity is a condition for reimbursement; it is not a guarantee of payment.

Claims that were submitted with dates of service from October 1, 2010, through June 30, 2013, will not be reprocessed in accordance with the NCCI guidelines; however, any claims with dates of service on or after October 1, 2010, that are appealed or reprocessed for reasons other than NCCI auditing will be subject to NCCI auditing guidelines.

### 6.4.1.1 NCCI Processing Categories

The following coding rule categories are applied to claims that are submitted with dates of service on or after October 1, 2010:

<table>
<thead>
<tr>
<th>Coding Rule Category</th>
<th>Description</th>
</tr>
</thead>
</table>
| Maximum units        | CMS has assigned to all procedure codes a maximum number of units that may be submitted for a client per day, regardless of the provider. The maximum number of units for each procedure code is based on the following criteria:  
  - Procedure code description  
  - Anatomical site  
  - CMS sources  
  - Clinical guidelines  
**Important:** If the maximum number of units has been exceeded on a particular line item, the line item will be denied. The line item will not be cut back to the allowable quantity. The line item may be appealed with the appropriate quantity for consideration. |
| NCCI                 | NCCI is a collection of bundling edits created and sponsored by CMS that are separated into two major categories: Column I and Column II procedure code edits (previously referred to as “Comprehensive” and “Component”) and Mutually Exclusive procedure code edits. NCCI edits are applied to services that are performed by the same provider on the same date of service only and do not apply to services that are performed within the global surgical period. Each NCCI code pair edit is associated with a policy as defined in the National Correct Coding Initiative Policy Manual. Effective dates apply to code pairs in NCCI and represent the date when CMS added the code pair combination to the NCCI edits. Code combinations are processed based on this effective date. Termination dates also apply to code pairs in NCCI. This date represents the date when CMS removed the code pair combination from the NCCI edits. Code combinations are refreshed quarterly. |

### 6.4.1.2 CPT and HCPCS Claims Auditing Guidelines

Claims with dates of service on or after October 1, 2010, must be filed in accordance with Current Procedural Terminology (CPT) and Healthcare Common Procedure Coding System (HCPCS) guidelines as defined in the American Medical Association (AMA) and Centers for Medicare & Medicaid Services (CMS) coding manuals. Claims that are not filed in accordance with CPT and HCPCS guidelines may be denied, including claims for services that were prior authorized or authorized based on documentation of medical necessity.
If a rendered service does not comply with CPT or HCPCS guidelines, medical necessity documentation may be submitted with the claim for the service to be considered for reimbursement; however, medical necessity documentation does not guarantee payment for the service.

**Important:** Prior authorization and authorization based on documentation of medical necessity is a condition for reimbursement; it is not a guarantee of payment.

The following coding rule categories apply to claims submissions:

<table>
<thead>
<tr>
<th>Coding Rule Category</th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td>Add-on codes</td>
<td>Certain services are commonly carried out in addition to the rendering of the primary procedure and are associated with the primary procedures. These additional or supplemental procedures are referred to as “add-on” procedures. Add-on codes are identified in the CPT Manual with a plus mark (“+”) symbol and are also listed in Appendix D of the CPT Manual. Add-on codes are always performed in addition to a primary procedure, and should never be reported as a stand-alone service. When an add-on code is submitted and the primary procedure has not been identified on either the same or different claim, then the add-on code will be denied as an inappropriately-coded procedure. If the primary procedure is denied for any reason, then the add-on code will be denied also.</td>
</tr>
<tr>
<td>Deleted HCPCS codes</td>
<td>Procedure codes undergo revision by the AMA and CMS on a regular basis. Revisions typically include adding new procedure codes, deleting procedure codes, and redefining the description of existing procedure codes. These revisions are normally made on an annual basis by the governing entities with occasional quarterly updates. Claims that are received with deleted procedure codes will be validated against the date of service. If the procedure code is valid for the date of service, the claim will continue processing. If the procedure code is invalid for the date of service, the invalid procedure code will be denied.</td>
</tr>
<tr>
<td>Diagnosis validity</td>
<td>ICD-10-CM diagnosis codes undergo revision by the Centers for Disease Control and Prevention (CDC) and CMS on a regular basis. Revisions typically include adding new diagnosis codes, deleting diagnosis codes, and redefining the description of existing diagnosis codes. These revisions are normally made on an annual basis. Claims that are received with invalid diagnosis codes will be validated against the date of service. If the diagnosis code is valid for the date of service, the claim will continue processing. If the diagnosis code is invalid for the date of service, the procedure that is referenced to the invalid diagnosis code will be denied.</td>
</tr>
<tr>
<td>Diagnosis-age</td>
<td>Certain diagnosis codes are age-specific. If a diagnosis code that is billed does not match the age of the client on that date of service, all services associated with that diagnosis code will be denied.</td>
</tr>
<tr>
<td>Diagnosis-gender</td>
<td>Certain diagnosis codes are gender-specific. If the diagnosis code that is billed does not match the gender of the client, all services associated with that diagnosis code will be denied.</td>
</tr>
<tr>
<td>Duplicate claim</td>
<td>A duplicate claim is defined as a claim or procedure code detail that exactly matches a claim or procedure code detail that has been reimbursed to the same provider for the same client. Duplicate claims or details include the same date of service, procedure code, modifier, and number of units. Duplicate claims or procedure code details will be denied.</td>
</tr>
</tbody>
</table>

**Note:** Modifiers may be used to identify separate services.
6.4.2 Claim Form Requirements

6.4.2.1 Provider Signature on Claims

Every paper CMS-1500, American Dental Association (ADA) Dental Claim Form, and 2017 Claim Form must be submitted with the provider’s or an authorized representative’s handwritten signature (or signature stamp) in the appropriate block of the claim form. Signatory supervision of the authorized
representative is required. Providers delegating signature authority to a member of the office staff or to a billing service remain responsible for the accuracy of all information on a claim submitted for payment. Initials are only acceptable for first and middle names. The last name must be spelled out. An acceptable example is J.A. Smith for John Adam Smith. An unacceptable example is J.A.S. for John Adam Smith. Typewritten names must be accompanied by a handwritten signature; in other words, a typewritten name with signed initials is not acceptable. The signature must be contained within the appropriate block of the claim form. Claims prepared by computer billing services or office-based computers may have “Signature on File” printed in the signature block, but it must be in the same font that is used in the rest of the form. For claims prepared by a billing service, the billing service must retain a letter on file from the provider authorizing the service.

*Printing the provider’s name instead of “Signature on File” is unacceptable.* Because space is limited in the signature block, providers should not type their names in the block. Claims not meeting these specifications appear in the “Paid or Denied Claims” sections of the R&S Reports.


6.4.2.2 Group Providers

Providers billing as a group must give the performing provider identifier on their claims as well as the group provider identifier. This requirement excludes THSteps medical providers.

6.4.2.3 Supervising Physician Provider Number Required on Some Claims

The supervising physician provider number is required on claims for services that are ordered or referred by one provider at the direction of or under the supervision of another provider, and the referral or order is based on the supervised provider’s evaluation of the client.

If a referral or order for services to a Texas Medicaid client is based on a client evaluation that was performed by the supervised provider, the billing provider’s claim must include the names and NPIs of both the ordering provider and the supervising provider. The billing provider must obtain all of the required information from the ordering or referring provider before submitting the claim to TMHP.

Providers who submit TexMedConnect electronic claims for professional, ambulance, or vision services can provide the claim information in the designated field for the supervising provider of the referring or ordering provider.

Providers can refer to TexMedConnect instructions on the TMHP website at www.tmhp.com for details about the “Referring/Other Supervising Provider” field for professional, ambulance, and vision electronic claims.

Note: Pharmacy claims are currently excluded from this requirement.

6.4.2.4 Ordering or Referring Provider NPI

All Texas Medicaid claims for services that require a physician order or referral must include the ordering or referring provider’s NPI:

- If the ordering or referring provider is enrolled in Texas Medicaid as a billing or performing provider, the billing or performing provider NPI must be used on the claim as the ordering or referring provider.

- If the ordering or referring provider is not currently enrolled in Texas Medicaid as a billing or performing provider, the provider must enroll to receive an ordering or referring-only TPI. After the ordering or referring provider is enrolled, the ordering or referring provider’s NPI must be used on the claim as the ordering or referring provider.

Important: The billing provider is responsible for confirming that the ordering or referring provider is enrolled as an ordering or referring-only provider.
Claims that are submitted without the ordering or referring provider’s NPI and claims submitted with an NPI for a provider who is not enrolled in Texas Medicaid may be subject to retrospective review and denial for a missing or invalid NPI.

*Note:* Providers who enroll in Texas Medicaid as ordering- and referring-only providers receive a TPI that can be used for orders and referrals for Texas Medicaid clients and CSHCN Services Program clients.

6.4.2.5 Attending Provider NPI Requirements

The attending provider is the individual who would normally be expected to certify and re-certify the medical necessity of the number of services rendered or who has primary responsibility for the patient’s medical care and treatment.

*Note:* Outpatient claim providers may be instructed to submit the ordering provider name and NPI number in the attending provider field.

6.4.2.6 Prior Authorization Numbers on Claims

Claims filed to TMHP must contain only one prior authorization number per claim. Prior authorization numbers must be indicated on the appropriate electronic field or on the paper claim forms in the indicated block:

- CMS-1500—Block 23
- UB-04 CMS-1450—Block 63
- ADA—Block 2
- Family Planning—Block 30

6.4.2.7 Newborn Clients Without Medicaid Numbers

If a Medicaid eligible newborn has not been assigned a Medicaid number on the DOS, the provider must wait until a Medicaid client number is assigned to file the claim. The provider writes the number instead of “Pending.” The 95-day filing period begins on the “add date,” which is the date the eligibility is received and added to the TMHP eligibility file. Providers verify eligibility and add date through TexMedConnect or by calling AIS or the TMHP Contact Center at 1-800-925-9126 after the number is received.

Providers must check Medicaid eligibility regularly to file claims within the required 95-day filing deadline.

*Refer to:* “Section 4: Client Eligibility” (*Vol. 1, General Information*).

6.4.2.8 Multipage Claim Forms

6.4.2.8.1 Professional Claims

The approved electronic claims format is designed to list 50 line items. The total number of details allowed for a professional claim by the TMHP claims processing system (C21) is 28. If the services provided exceed 28 line items on an approved electronic claims format or 28 line items on paper claims, the provider must submit another claim for the additional line items.

The CMS-1500 paper claim form is designed to list six line items in Block 24. If more than six line items are billed on a paper claim, a provider may attach additional forms (pages) totaling no more than 28 line items. The first page of a multipage claim must contain all the required billing information. On subsequent pages of the multipage claim, the provider should identify the client’s name, diagnosis,
information required for services in Block 24, and the page number of the attachment (for example, page 2 of 3) in the top right-hand corner of the form and indicate “continued” in Block 28. The combined total charges for all pages should be listed on the last page in Block 28.

Note: Providers who submit professional claims for inpatient services are required to include only the facility’s NPI on the CMS-1500 paper claim form or electronic equivalent. The CMS-1500 paper claim form and electronic equivalents do not have a field for the facility’s TPI.

6.4.2.8.2 Institutional Claims

The total number of details allowed for an institutional claim by the TMHP claims processing system (C21) is 28. C21 merges like revenue codes together for inpatient claims to reduce the lines to 28 or less. If the C21 merge function is unable to reduce the lines to 28 or less, the claim will be denied, and the provider will need to reduce the number of details and resubmit the claim.

An EDI approved electronic format of the UB-04 CMS-1450 is designed to list 71 lines. C21 merges like revenue codes together to reduce the lines to 28 or less.

Providers submitting electronic claims using TexMedConnect may not submit more than 28 lines. If the services exceed the 28 lines, the provider may submit another claim for the additional lines or merge codes.

The paper UB-04 CMS-1450 is designed to list 23 lines in Block 43. If services exceed the 23-line limitation, the provider may attach additional pages. The first page of a multipage claim must contain all required billing information. On subsequent pages, the provider identifies the client’s name, diagnosis, all information required in Block 43, and the page number of the attachment (e.g., page 2 of 3) in the top right-hand corner of the form and indicate “continued” on Line 23 of Block 47. The combined total charges for all pages should be listed on the last page on Line 23 of Block 47.

When splitting a claim, all pages must contain the required information. Usually, there are logical breaks to a claim. For example, the provider may submit the surgery charges in one claim and the subsequent recovery days in the next claim.

TEFRA hospitals are required to submit all charges.

6.4.2.8.3 Inpatient Hospital Claims

Medicaid present-on-admission (POA) reporting is required for all inpatient hospital claims that are paid under prospective payment basis methodology. No hospitals are exempt from this POA requirement.

Medicare crossover hospital claims must also comply with the Medicaid requirement to include the POA values. Claims submitted without the POA indicators are denied. POA values are:

<table>
<thead>
<tr>
<th>POA Value</th>
<th>Description</th>
<th>Payment</th>
</tr>
</thead>
<tbody>
<tr>
<td>Y</td>
<td>Diagnosis was present at the time of admission.</td>
<td>Payment will be made by Texas Medicaid when a hospital acquired condition (HAC) is present.</td>
</tr>
<tr>
<td>N</td>
<td>Diagnosis was not present at the time of admission.</td>
<td>No payment will be made by Texas Medicaid when an HAC is present.</td>
</tr>
<tr>
<td>U</td>
<td>Documentation was insufficient.</td>
<td>No payment will be made by Texas Medicaid when an HAC is present.</td>
</tr>
<tr>
<td>W</td>
<td>Clinically undetermined.</td>
<td>Payment will be made by Texas Medicaid when an HAC is present.</td>
</tr>
<tr>
<td>Blank</td>
<td>Exempt from POA Reporting</td>
<td>Exempt from POA Reporting</td>
</tr>
</tbody>
</table>

Note: Texas Medicaid follows Medicare guidelines for payments referenced in the above table.
Depending on the POA indicator value, the DRG may be recalculated, which could result in a lower payment to the hospital facility provider. If the number of days on an authorization is higher than the number of days allowed as a result of a POA DRG recalculation, the lesser of the number of days is reimbursed.

Refer to: Federal Register, Vol. 76, No. 108 (for CMS).

### 6.4.2.9 Attachments to Claims

To expedite claims processing, providers must supply all information on the claim form itself and limit attachments to those required by TMHP or necessary to supply information to properly adjudicate the claim. The following claim form attachments are required when appropriate:

- All claims for services associated with an elective sterilization must have a valid Sterilization Consent Form attached or on file at TMHP.
- Nonemergency ambulance transfers must have documentation of medical necessity including out-of-locality transfers.
- For fee-for-service clients, providers filing to TMHP for Medicaid payment of Medicare coinsurance and deductible according to current payment guidelines must attach the paper MRAN received from Medicare or a Medicare intermediary or the computer generated MRANs from the CMS-approved software applications MREP for professional services or PC-Print for institutional services. Providers that submit paper crossover claims must submit only one of the approved MRAN formats.
- For MAP clients, providers filing to TMHP for Medicaid payment of Medicare coinsurance and deductible according to current payment guidelines must submit with the paper claim the TMHP Standardized Medicare Advantage Plan (MAP) Remittance Advice Notice Template with the MAP EOB. If the template and MAP EOB contain conflicting information, the claim will not be processed and will be returned to the provider.
- Medically necessary abortions performed (on the basis of a physician’s professional judgement, the life of the mother is endangered if the fetus were carried to term), or abortions provided for pregnancy related to rape or incest must have a signed and dated physician certification statement. Elective abortions are not benefits of Texas Medicaid.
- Hysterectomies must have a Hysterectomy Acknowledgment Statement attached or on file at TMHP.

Refer to: Texas Medicaid - Title XIX Acknowledgment of Hysterectomy Information on the TMHP website at www.tmhp.com.

- Claims for services that were paid by an MCO and then recouped must contain the recoupment EOB from the MCO for consideration of payment. The claims must meet the 95-day deadline from the recoupment disposition date.

Note: Letter requests for refunds will not be accepted. A recoupment EOB with a disposition date is required.

### 6.4.2.10 Clients with a Designated or Primary Care Provider

Claims for clients with a primary care provider or designated provider (i.e., Texas Medicaid fee-for-service clients enrolled as Limited Program clients) must indicate the primary care provider or designated provider identifiers in the billing or performing provider fields.

When clients receive services from a different provider, such as a specialist, the primary care provider or designated provider’s information must be included in the referring provider fields on the claim.
### 6.5 CMS-1500 Paper Claim Filing Instructions

The following providers bill for services using the ANSI ASC X12 837P 5010 electronic specifications or the CMS-1500 paper claim form:

<table>
<thead>
<tr>
<th>Providers</th>
</tr>
</thead>
<tbody>
<tr>
<td>Ambulance</td>
</tr>
<tr>
<td>ASC (freestanding)</td>
</tr>
<tr>
<td>Case Management for Blind and Visually Impaired Children (BVIC), Case Management for Early Childhood Intervention (ECI), and Case Management for Children and Pregnant Women</td>
</tr>
<tr>
<td>Certified nurse-midwife (CNM)</td>
</tr>
<tr>
<td>Certified registered nurse anesthetist (CRNA)</td>
</tr>
<tr>
<td>Certified respiratory care practitioner (CRCP)</td>
</tr>
<tr>
<td>Chemical dependency treatment facilities</td>
</tr>
<tr>
<td>Chiropractor</td>
</tr>
<tr>
<td>Clinical nurse specialist (CNS)</td>
</tr>
<tr>
<td>Dentist (doctor of dentistry practicing as a limited physician)</td>
</tr>
<tr>
<td>DME</td>
</tr>
<tr>
<td>Family planning agency that does not also receive funds from the HHSC Family Planning Program</td>
</tr>
<tr>
<td>FQHC</td>
</tr>
<tr>
<td>Genetic service agency</td>
</tr>
<tr>
<td>Hearing aid</td>
</tr>
<tr>
<td>IDD case management</td>
</tr>
<tr>
<td>In-home total parenteral nutrition (TPN) supplier</td>
</tr>
<tr>
<td>Laboratory</td>
</tr>
<tr>
<td>Licensed dietitian (CCP only)</td>
</tr>
<tr>
<td>Licensed clinical social worker (LCSW)</td>
</tr>
<tr>
<td>Licensed professional counselor (LPC)</td>
</tr>
<tr>
<td>Maternity service clinic (MSC)</td>
</tr>
<tr>
<td>Mental health (MH) targeted case management</td>
</tr>
<tr>
<td>Mental health (MH) rehabilitative services</td>
</tr>
<tr>
<td>Nurse practitioner (NP)</td>
</tr>
<tr>
<td>Occupational therapist (CCP only)</td>
</tr>
<tr>
<td>Optician/optometrist/ophthalmologist</td>
</tr>
<tr>
<td>Orthotic and prosthetic supplier (CCP only)</td>
</tr>
<tr>
<td>Pharmacy</td>
</tr>
<tr>
<td>Physical therapist</td>
</tr>
<tr>
<td>Physician (group and individual)</td>
</tr>
<tr>
<td>Physician assistant (PA)</td>
</tr>
<tr>
<td>Podiatrist</td>
</tr>
<tr>
<td>Private duty nurse (PDN) (CCP only)</td>
</tr>
<tr>
<td>Psychologist</td>
</tr>
<tr>
<td>Radiology</td>
</tr>
</tbody>
</table>


Providers

| Rural Health Clinics rendering services to THSteps clients |
| School Health and Related Services (SHARS) |
| Speech language pathologist (CCP only) |
| THSteps medical |
| Tuberculosis clinic |

Providers obtain copies of the CMS-1500 paper claim form from a vendor of their choice; TMHP does not supply them.

### 6.5.1 CMS-1500 Electronic Billing

Electronic billers must submit CMS-1500 paper claim forms with TexMedConnect or approved vendor software that uses the ANSI ASC X12 837P 5010 format. Specifications are available to providers developing in-house systems, software developers, and vendors on the TMHP website at [www.tmhp.com/topics/edi](http://www.tmhp.com/topics/edi). Because each software developer is different, location of fields may vary. Contact the software developer or vendor for this information. Direct questions and development requirements to the TMHP EDI Help Desk at 1-888-863-3638.

**Refer to:** Subsection 3.2, “Electronic Billing” in “Section 3: TMHP Electronic Data Interchange (EDI)” (Vol. 1, General Information) for information about electronic billing.

### 6.5.2 CMS-1500 Claim Form (Paper) Billing

Claims must contain the billing provider’s complete name, address, and a provider identifier. Claims without a provider name, address, and provider identifier cannot be processed. Each claim form must have the appropriate signatory evidence in the signature certification block.

**Refer to:** The Professional Paper Claim Form (CMS-1500) page of the CMS website at [www.cms.gov](http://www.cms.gov) for more information about the CMS-1500 paper claim form. Providers can purchase CMS-1500 paper claim forms from the vendor of their choice. TMHP does not supply the forms. Providers can find examples of completed claim forms on the Claim Form Examples page of the TMHP website at [www.tmhp.com](http://www.tmhp.com).

**Important:** When completing a CMS-1500 paper claim form, all required information must be included on the claim in the appropriate block. Information is not keyed from attachments. Superbills or itemized statements are not accepted as claim supplements.

### 6.5.3 CMS-1500 Provider Definitions

The following definitions apply to the provider terms used on the CMS-1500 paper claim form:

**Referring Provider**

The referring provider is the individual who directed the patient for care to the provider that rendered the services being submitted on the claim form.

Examples include, but are not limited to the following:

- A primary care provider referring to a specialist
- An orthodontist referring to an oral and maxillofacial surgeon
- A physician referring to a physical therapist
- A provider referring to a home health agency
Ordering Provider
The ordering provider is the individual who requested the services or items listed in Block D of the CMS-1500 paper claim form.
Examples include, but are not limited to, a provider ordering diagnostic tests, medical equipment, or supplies.

Rendering Provider
The rendering provider is the individual who provided the care to the client. In the case where a substitute provider was used, that individual is considered the rendering provider.
An individual such as a lab technician or radiology technician who performs services in a support role is not considered a rendering provider.

Supervising Provider
The supervising provider is the individual who provided oversight of the rendering provider and the services listed on the CMS-1500 paper claim form.
An example would be the supervision of a resident physician.

Purchased Service Provider
A purchased service provider is an individual or entity that performs a service on a contractual or reassignment basis.
Examples of services include the following:
- Processing a laboratory specimen
- Grinding eyeglass lenses to the specifications of the referring provider
- Performing diagnostic testing services (excluding clinical laboratory testing) subject to Medicare’s antimarkup rule

In the case where a substitute provider is used, that individual is not considered a purchased service provider.

6.5.4 CMS-1500 Instruction Table
The instructions describe what information must be entered in each of the block numbers of the CMS-1500 paper claim form. Block numbers not referenced in the table may be left blank. They are not required for claim processing by TMHP.

<table>
<thead>
<tr>
<th>Block No.</th>
<th>Description</th>
<th>Guidelines</th>
</tr>
</thead>
<tbody>
<tr>
<td>1a</td>
<td>Insured’s ID No. (for program checked above, include all letters)</td>
<td>Enter the client’s nine-digit patient number from the Medicaid identification form. For other property &amp; casualty claims: Enter the Federal Tax ID or SSN of the insured person or entity.</td>
</tr>
<tr>
<td>2</td>
<td>Patient’s name</td>
<td>Enter the client’s last name, first name, and middle initial as printed on the Medicaid identification form. If the insured uses a last name suffix (e.g., Jr, Sr) enter it after the last name and before the first name.</td>
</tr>
<tr>
<td>3</td>
<td>Patient’s date of birth Patient’s sex</td>
<td>Enter numerically the month, day, and year (MM/DD/YYYY) the client was born. Indicate the client’s gender by checking the appropriate box. Only one box can be marked.</td>
</tr>
<tr>
<td>5</td>
<td>Patient’s address</td>
<td>Enter the client’s complete address as described (street, city, state, and ZIP code).</td>
</tr>
<tr>
<td>Block No.</td>
<td>Description</td>
<td>Guidelines</td>
</tr>
<tr>
<td>-----------</td>
<td>-------------</td>
<td>-------------</td>
</tr>
</tbody>
</table>
| 9         | Other insured’s name | For special situations, use this space to provide additional information such as:  
If the client is deceased, enter “DOD” in block 9 and the time of death in 9a if the services were rendered on the date of death. Enter the date of death in block 9b. |
| 10a       | Is patient’s condition related to: | Check the appropriate box. If other insurance is available, enter appropriate information in blocks 11, 11a, and 11b. |
| 10b       | a. Employment (current or previous)? | |
| 10c       | b. Auto accident? | |
|           | c. Other accident? | |
| 11        | Other health insurance coverage | • If another insurance resource has made payment or denied a claim, enter the name of the insurance company. The other insurance EOB or denial letter must be attached to the claim form.  
• If the client is enrolled in Medicare attach a copy of the MRAN to the claim form.  
• For Workers Compensation and other property and casualty claims: (Required if known) Enter Workers’ Compensation or property and casualty claim number assigned by the payer. |
| 11a       | | |
| 11b       | | |
| 11c       | Insurance plan or program name | Enter the benefit code, if applicable, for the billing or performing provider. |
| 12        | Patient’s or authorized person’s signature | Enter “Signature on File,” “SOF,” or legal signature. When legal signature is entered, enter the date signed in eight digit format (MMDDYYYY).  
TMHP will process the claim without the signature of the patient. |
| 14        | Date of current | Enter the first date (MM/DD/YYYY) of the present illness or injury. For pregnancy enter the date of the last menstrual period.  
If the client has chronic renal disease, enter the date of onset of dialysis treatments.  
Indicate the date of treatments for PT and OT. |
<table>
<thead>
<tr>
<th>Block No.</th>
<th>Description</th>
<th>Guidelines</th>
</tr>
</thead>
</table>
| 17       | Name of referring physician or other source          | Enter the name (First Name, Middle Initial, Last Name) and credentials of the professional who referred, ordered, or supervised the service(s) or supplies on the claim. If multiple providers are involved, enter one provider using the following priority order:  
1. Referring Provider  
2. Ordering Provider  
3. Supervising Provider  
Do not use periods or commas within the name. A hyphen can be used for hyphenated names. Enter the applicable qualifier to identify which provider is being reported.  
DN = Referring Provider  
DK = Ordering Provider  
DQ = Supervising Provider  
Supervising Physician for Referring Physicians:  
If there is a Supervising Physician for the referring or ordering provider that is listed in Block 17, the name and NPI of the supervising provider must go in Block 19. |
| 17b      | NPI                                                   | Enter the NPI number of the referring, ordering, or supervising provider.                                                                                                                                 |
| 19       | Additional claim information                         | **Ambulance transfers of multiple clients**  
If the claim is part of a multiple transfer, indicate the other client’s complete name and Medicaid number.  
**Ambulance Hospital-to-Hospital Transfers**  
Indicate the services required from the second facility and unavailable at the first facility  
**Supervising Physician for Referring Physicians:**  
If there is a Supervising Physician for the referring or ordering provider that is listed in Block 17, the name and NPI of the supervising provider must go in Block 19. |
| 20       | Outside lab                                          | Check the appropriate box. The information may be requested for retrospective review.  
If “yes,” enter the provider identifier of the facility that performed the service in block 32. |
| 21       | Diagnosis or nature of illness or injury             | Enter the applicable ICD indicator to identify which version of ICD codes is being reported.  
9 = ICD-9-CM  
0 = ICD-10-CM  
Enter the patient’s diagnosis and/or condition codes. List no more than 12 diagnosis codes.  
Relate lines A-L to the lines of service in 24E by the letter of the line. Use the highest level of specificity.  
Do not provide narrative description in this field. |
<table>
<thead>
<tr>
<th>Block No.</th>
<th>Description</th>
<th>Guidelines</th>
</tr>
</thead>
<tbody>
<tr>
<td>23</td>
<td>Prior authorization number</td>
<td>Enter the PAN issued by TMHP. For Workers Compensation and other property and casualty claims, this is required when prior authorization, referral, concurrent review, or voluntary certification was received.</td>
</tr>
</tbody>
</table>
| 24       | (Various)                                        | General notes for blocks 24a through 24j:  
  • Unless otherwise specified, all required information should be entered in the unshaded portion.  
  • If more than six line items are billed for the entire claim, a provider must attach additional claim forms with no more than 28-line items for the entire claim.  
  • For multi-page claim forms, indicate the page number of the attachment (for example, page 2 of 3) in the top right-hand corner of the claim form. |
| 24a      | Date(s) of service                               | Enter the date of service for each procedure provided in a MM/DD/YYYY format. If more than one date of service is for a single procedure, each date must be given on a separate line.  
  **NDC**  
  In the shaded area, enter the:  
  • NDC qualifier of N4 (e.g., N4).  
  • The 11-digit NDC number on the package or vial from which the medication was administered. Do not enter hyphens or spaces within this number (e.g., 00409231231).  
  **Example:** N400409231231  
  **Refer to:** Subsection 6.3.4, “National Drug Code (NDC)” in this section. |
| 24b      | Place of service                                 | Select the appropriate POS code for each service from the table under subsection 6.3.1.1, “Place of Service (POS) Coding” in this section.                                                               |
| 24c      | EMG (THSteps medical checkup condition indicator) | Enter the appropriate condition indicator for THSteps medical checkups.  
  **Refer to:** Subsection 5.3.6, “THSteps Medical Checkups” in the Children’s Services Handbook (Vol. 2, Provider Handbooks).                                                                           |
| 24d      | Fully describe procedures, medical services, or supplies furnished for each date given | Enter the appropriate procedure codes and modifier for all services billed. If a procedure code is not available, enter a concise description.  
  **NDC**  
  In the shaded area, enter a 1- through 12-digit NDC quantity of unit. A decimal point must be used for fractions of a unit (e.g., 0.025).  
  **Refer to:** Subsection 6.3.4, “National Drug Code (NDC)” in this section. |
<table>
<thead>
<tr>
<th>Block No.</th>
<th>Description</th>
<th>Guidelines</th>
</tr>
</thead>
<tbody>
<tr>
<td>24e</td>
<td>Diagnosis pointer</td>
<td>In 24 E, enter the diagnosis code reference letter (pointer) as shown in Form Field 21 to relate the date of service and the procedures performed to the primary diagnosis. When multiple services are performed, the primary reference number for each service should be listed first, other applicable services should follow. The reference letter(s) should be A-L or multiple letters as applicable. Diagnosis codes must be entered in Form Field 21 only. Do not enter diagnosis codes in Form Field 24E.</td>
</tr>
<tr>
<td>24f</td>
<td>Charges</td>
<td>Indicate the usual and customary charges for each service listed. Charges must not be higher than fees charged to private-pay clients.</td>
</tr>
<tr>
<td>24g</td>
<td>Days or units</td>
<td>If multiple services are performed on the same day, enter the number of services performed (such as the quantity billed). Note: The maximum number of units per detail is 9,999.</td>
</tr>
<tr>
<td>24j</td>
<td>Rendering provider ID # (performing)</td>
<td>Enter the provider identifier of the individual rendering services unless otherwise indicated in the provider specific section of this manual. Enter the TPI in the shaded area of the field. Entered the NPI in the unshaded area of the field.</td>
</tr>
<tr>
<td>26</td>
<td>Patient’s account number</td>
<td>Optional: Enter the client identification number if it is different than the subscriber/insured’s identification number. Used by provider’s office to identify internal client account number.</td>
</tr>
<tr>
<td>27</td>
<td>Accept assignment</td>
<td>Required All providers of Texas Medicaid must accept assignment to receive payment by checking Yes.</td>
</tr>
<tr>
<td>28</td>
<td>Total charge</td>
<td>Enter the total charges. For multi-page claims enter “continue” on initial and subsequent claim forms. Indicate the total of all charges on the last claim. Note: Indicate the page number of the attachment (for example, page 2 of 3) in the top right-hand corner of the form.</td>
</tr>
<tr>
<td>29</td>
<td>Amount paid</td>
<td>Enter any amount paid by an insurance company or other sources known at the time of submission of the claim. Identify the source of each payment and date in block 11. If the client makes a payment, the reason for the payment must be indicated in block 11.</td>
</tr>
<tr>
<td>30</td>
<td>Balance due</td>
<td>If appropriate, subtract block 29 from block 28 and enter the balance.</td>
</tr>
</tbody>
</table>
6.6 UB-04 CMS-1450 Paper Claim Filing Instructions

The following provider types may bill electronically or use the UB-04 CMS-1450 paper claim form when requesting payment:

<table>
<thead>
<tr>
<th>Provider Types</th>
<th>Guidelines</th>
</tr>
</thead>
<tbody>
<tr>
<td>ASCs (hospital-based)</td>
<td></td>
</tr>
<tr>
<td>Comprehensive outpatient rehabilitation facilities (CORFs) (CCP only)</td>
<td></td>
</tr>
<tr>
<td>FQHCs</td>
<td></td>
</tr>
<tr>
<td>Note: Must use CMS-1500 when billing THSteps.</td>
<td></td>
</tr>
<tr>
<td>Home health agencies</td>
<td></td>
</tr>
<tr>
<td>Hospitals</td>
<td></td>
</tr>
<tr>
<td>• Inpatient (acute care, rehabilitation, military, and psychiatric hospitals)</td>
<td></td>
</tr>
<tr>
<td>• Outpatient</td>
<td></td>
</tr>
<tr>
<td>Indian Health</td>
<td></td>
</tr>
<tr>
<td>Renal dialysis center</td>
<td></td>
</tr>
<tr>
<td>Personal Care Services (PCS)</td>
<td></td>
</tr>
<tr>
<td>RHHCs (freestanding and hospital-based)</td>
<td></td>
</tr>
<tr>
<td>Note: Must use CMS-1500 when billing THSteps.</td>
<td></td>
</tr>
</tbody>
</table>

If a service is rendered in the facility setting but the facility’s medical record does not clearly support the information submitted on the facility claim, the facility may request additional information from the physician before submitting the claim to ensure the facility medical record supports the filed claim.

**Note:** In the case of an audit, facility providers will not be allowed to submit an addendum to the original medical records for finalized claims.
6.6.1 UB-04 CMS-1450 Electronic Billing

Electronic billers must submit UB-04 CMS-1450 claims with TexMedConnect or approved vendor software that uses the ANSI ASC X12 837I 5010 format. Specifications are available to providers developing in-house systems and software developers and vendors. Because each software package is different, field locations may vary. Contact the software developer or vendor for this information. Direct questions and development requirements to the TMHP EDI Help Desk at 1-888-863-3638.

Refer to: Subsection 3.2, “Electronic Billing” in “Section 3: TMHP Electronic Data Interchange (EDI)” (Vol. 1, General Information) for more information about electronic billing.

Note: The maximum number of electronic claim details that will be accepted electronically is 71. Only 28 details will be processed.

6.6.2 UB-04 CMS-1450 Claim Form (Paper) Billing

Providers obtain the UB-04 CMS-1450 paper claim forms from a vendor of their choice.

Note: To avoid claim denial, only the provider’s NPI should be placed in form locators 76-79 of the UB-04 CMS-1450 paper claim form or in the referring provider field on the electronic claim unless the client is a limited client.

Completed UB-04 CMS-1450 claims must contain the billing provider’s full name, address, and provider identifier (TPI/NPI in the appropriate fields). Claims without a provider name, address, and provider identifier in the appropriate fields cannot be processed.

Refer to: The Institutional paper claim form (CMS-1450) CMS website at www.cms.gov for more information about the CMS-1450 paper claim form. Providers can purchase CMS-1450 paper claim forms from the vendor of their choice. TMHP does not supply the forms. Providers can find examples of completed claim forms on the Claim Form Examples page of the TMHP website at www.tmhp.com.

subsection 6.6.3, “UB-04 CMS-1450 Instruction Table” in this section.

6.6.3 UB-04 CMS-1450 Instruction Table

The instructions describe what information must be entered in each of the block numbers of the UB-04 CMS-1450 paper claim form. Block numbers not referenced in the table may be left blank. They are not required for claim processing by TMHP.

<table>
<thead>
<tr>
<th>Block No.</th>
<th>Description</th>
<th>Guidelines</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>Unlabeled</td>
<td>Enter the hospital name, street, city, state, ZIP+4 Code, and telephone number.</td>
</tr>
<tr>
<td>3a</td>
<td>Patient control number</td>
<td><strong>Optional:</strong> Any alphanumeric character (limit 16) entered in this block is referenced on the R&amp;S Report.</td>
</tr>
<tr>
<td>3b</td>
<td>Medical record number</td>
<td>Enter the patient’s medical record number (limited to ten digits) assigned by the hospital.</td>
</tr>
<tr>
<td>4</td>
<td>Type of bill (TOB)</td>
<td>Enter a TOB code. First Digit—Type of Facility: 1 Hospital, 2 Skilled nursing, 3 Home health agency, 7 Clinic (rural health clinic [RHC], federally qualified health center [FQHC], and renal dialysis center [RDC]), 8 Special facility</td>
</tr>
<tr>
<td>Block No.</td>
<td>Description</td>
<td>Guidelines</td>
</tr>
<tr>
<td>-----------</td>
<td>----------------------</td>
<td>---------------------------------------------------------------------------------------------------------------------------------------------</td>
</tr>
</tbody>
</table>
|           | Second Digit—Bill Classification (except clinics and special facilities):                                                                 | 1 Inpatient (including Medicare Part A)  
2 Inpatient (Medicare Part B only)  
3 Outpatient  
4 Other (for hospital-referenced diagnostic services, for example, laboratories and X-rays)  
7 Intermediate care  

Second Digit—Bill Classification (clinics only):                                                                 | 1 Rural health  
2 Hospital-based or independent renal dialysis center  
3 Free standing  
5 CORFs  

Third Digit—Frequency:                                                                 | 0 Nonpayment/zero claim  
1 Admit through discharge  
2 Interim-first claim  
3 Interim-continuing claim  
4 Interim-last claim  
5 Late charges-only claim  
6 Adjustment of prior claim  
7 Replacement of prior claim  

6 | Statement covers period | Enter the beginning and ending dates of service billed.                                                                                       |
| 8a | Patient identifier     | Optional: Enter the patient identification number if it is different than the subscriber/insured’s identification number.  
Used by providers office to identify internal patient account number.       |

8b | Patient name           | Enter the patient’s last name, first name, and middle initial as printed on the Medicaid identification form.                          |

9a–9b | Patient address       | Starting in 9a, enter the patient’s complete address as described (street, city, state, and ZIP+4 Code).                  |

10 | Birthdate             | Enter the patient’s date of birth (MM/DD/YYYY).                                                                                         |

11 | Sex                   | Indicate the patient’s gender by entering an “M” or “F.”                                                                               |

12 | Admission date        | Enter the numerical date (MM/DD/YYYY) of admission for inpatient claims; date of service (DOS) for outpatient claims; or start of care (SOC) for home health claims.  
Providers that receive a transfer patient from another hospital must enter the actual dates the patient was admitted into each facility. |

13 | Admission hour        | Use military time (00 to 23) for the time of admission for inpatient claims or time of treatment for outpatient claims.               |

14 | Priority (Type) of Admission or Visit | Providers can refer to the National Uniform Billing Code website at www.nubc.org for the current list of Priority (Type) of Admission or Visit codes. |

15 | Point of Origin for Admission or Visit | Providers can refer to the National Uniform Billing Code website at www.nubc.org for the current list of Point of Origin for Admission or Visit codes. |

16 | Discharge hour        | For inpatient claims, enter the hour of discharge or death. Use military time (00 to 23) to express the hour of discharge. If this is an interim bill (patient status of “30”), leave the block blank. |
<table>
<thead>
<tr>
<th>Block No.</th>
<th>Description</th>
<th>Guidelines</th>
</tr>
</thead>
<tbody>
<tr>
<td>17</td>
<td>Patient Discharge Status</td>
<td>Providers can refer to the National Uniform Billing Code website at <a href="http://www.nubc.org">www.nubc.org</a> for the current list of Patient Discharge Status Codes.</td>
</tr>
<tr>
<td>18–28</td>
<td>Condition codes</td>
<td>Enter the two-digit condition code “05” to indicate that a legal claim was filed for recovery of funds potentially due to a patient.</td>
</tr>
<tr>
<td>29</td>
<td>ACDT state</td>
<td>Optional: Accident state.</td>
</tr>
<tr>
<td>31–34</td>
<td>Occurrence codes and dates</td>
<td>Providers can refer to the National Uniform Billing Code website at <a href="http://www.nubc.org">www.nubc.org</a> for the current list of Occurrence Codes.</td>
</tr>
<tr>
<td>35–36</td>
<td>Occurrence span codes and dates</td>
<td>For inpatient claims, enter code “71” if this hospital admission is a readmission within seven days of a previous stay. Enter the dates of the previous stay.</td>
</tr>
<tr>
<td>39–41</td>
<td>Value codes</td>
<td>Accident hour—For inpatient claims, if the patient was admitted as the result of an accident, enter value code 45 with the time of the accident using military time (00 to 23). Use code 99 if the time is unknown. For inpatient claims, enter value code 80 and the total days represented on this claim that are to be covered. Usually, this is the difference between the admission and discharge dates. In all circumstances, the number in this block is equal to the number of covered accommodation days listed in Block 46. For inpatient claims, enter value code 81 and the total days represented on this claim that are not covered. The sum of Blocks 39–41 must equal the total days billed as reflected in Block 6.</td>
</tr>
</tbody>
</table>
| 42–43    | Revenue codes and description  | For inpatient hospital services, enter the description and revenue code for the total charges and each accommodation and ancillary provided. List accommodations in the order of occurrence. List ancillaries in ascending order. The space to the right of the dotted line is used for the accommodation rate. **NDC** This block should include the following elements in the following order:  
  - NDC qualifier of N4 (e.g., N4)  
  - The 11-digit NDC number on the package or vial from which the medication was administered. Do not enter hyphens or spaces within this number (e.g., 00409231231).  
  - The unit of measurement code. There are 5 allowed values: F2, GR, ML, UN, or ME (e.g., GR).  
  - The unit quantity with a floating decimal for fractional units (limited to 3 digits, e.g., 0.025).  
  **Example:** N400409231231GR0.025  
  **Referto:** Subsection 6.3.4, “National Drug Code (NDC)” in this section. |
<table>
<thead>
<tr>
<th>Block No.</th>
<th>Description</th>
<th>Guidelines</th>
</tr>
</thead>
</table>
| 44       | HCPCS/rates | **Inpatient:**  
Enter the accommodation rate per day.  
Match the appropriate diagnoses listed in Blocks 67A through 67Q corresponding to each procedure. If a procedure corresponds to more than one diagnosis, enter the primary diagnosis.  
Each service and supply must be itemized on the claim form.  

**Home Health Services**  
Outpatient claims must have the appropriate revenue code and, if appropriate, the corresponding HCPCS code or narrative description.  

**Refer to:** Subsection 4.5.5, “Outpatient Hospital Revenue Codes” in the *Inpatient and Outpatient Hospital Services Handbook* (Vol. 2, Provider Handbooks) for additional information on which revenue codes require HCPCS codes.  

**Outpatient:**  
Outpatient claims must have the appropriate Healthcare Common Procedure Coding System (HCPCS) code.  

**Refer to:** Subsection 4.5.5, “Outpatient Hospital Revenue Codes” in the *Inpatient and Outpatient Hospital Services Handbook* (Vol. 2, Provider Handbooks) for additional information on which revenue codes require HCPCS codes.  

Each service, except for medical/surgical and intravenous (IV) supplies and medication, must be itemized on the claim form or an attached statement.  

**Note:** The UB-04 CMS-1450 paper claim form is limited to 28 items per inpatient and outpatient claim.  
If necessary, combine IV supplies and central supplies on the charge detail and consider them to be single items with the appropriate quantities and total charges by dates of service. Multiple dates of service may not be combined on outpatient claims.  

| 45       | Service date | Enter the numerical date of service that corresponds to each procedure for outpatient claims. Multiple dates of service may not be combined on outpatient claims. |
| 45 (line 23) | Creation date | Enter the date the bill was submitted. |
| 46       | Serv. units  | Provide units of service, if applicable.  
For inpatient services, enter the number of days for each accommodation listed. If applicable, enter the number of pints of blood.  
When billing for observation room services, the units indicated in this block should always represent hours spent in observation. |
| 47       | Total charges | Enter the total charges for each service provided.  

**Note:** For multi-page claims enter “continue” on initial and subsequent claim forms. Indicate the total of all charges on the last claim and the page number of the attachment (for example, page 2 of 3) in the top right-hand corner of the form.
<table>
<thead>
<tr>
<th>Block No.</th>
<th>Description</th>
<th>Guidelines</th>
</tr>
</thead>
<tbody>
<tr>
<td>48</td>
<td>Noncovered charges</td>
<td>If any of the total charges are noncovered, enter this amount.</td>
</tr>
<tr>
<td>50</td>
<td>Payer Name</td>
<td>Enter the health plan name.</td>
</tr>
<tr>
<td>51</td>
<td>Health Plan ID</td>
<td>Enter the health plan identification number.</td>
</tr>
</tbody>
</table>
| 54        | Prior payments | Enter amounts paid by any TPR, and complete Blocks 32, 61, 62, and 80 as required:  
  - Block 32 - Occurrence code and date.  
  - Block 61 - Insured group name  
  - Block 62 - Insurance group number  
  - Block 80 - Remarks. This section is used for requesting the 110-day rule for a third party insurance. |
| 56        | NPI | Enter the NPI of the billing provider. |
| 57        | Other identification (ID) number | Enter the TPI number (non-NPI number) of the billing provider. |
| 58        | Insured’s name | If other health insurance is involved, enter the insured’s name. |
| 60        | Medicaid identification number | Enter the patient’s nine-digit Medicaid identification number. |
| 61        | Insured group name | Enter the name and address of the other health insurance. |
| 62        | Insurance group number | Enter the policy number or group number of the other health insurance. |
| 63        | Treatment authorization code | Enter the prior authorization number if one was issued. |
| 65        | Employer name | Enter the name of the patient’s employer if health care might be provided. |
| 66        | Diagnosis/Procedure Code Qualifier | Enter the applicable ICD indicator to identify which version of ICD codes is being reported.  
  9 = ICD-9-CM  
  0 = ICD-10-CM |
| 67        | Principal diagnosis (DX) code and present on admission (POA) indicator | Enter the ICD-10-CM diagnosis code in the unshaded area for the principal diagnosis to the highest level of specificity available.  
**Required:** POA Indicator—Enter the applicable POA indicator in the shaded area for inpatient claims.  
**Refer to:** Subsection 6.4.2.8.3, “Inpatient Hospital Claims” in this section for POA values.
<table>
<thead>
<tr>
<th>Block No.</th>
<th>Description</th>
<th>Guidelines</th>
</tr>
</thead>
<tbody>
<tr>
<td>67A-67Q</td>
<td>Secondary DX codes and POA indicator</td>
<td>Enter the ICD-10-CM diagnosis code in the unshaded area to the highest level of specificity available for each additional diagnosis. Enter one diagnosis per block, using Blocks A through J only. A diagnosis is not required for clinical laboratory services provided to nonpatients (TOB “141”). <strong>Exception:</strong> A diagnosis is required when billing for estrogen receptor assays, plasmapheresis, and cancer antigen CA 125, immunofluorescent studies, surgical pathology, and alphafetoprotein. <strong>Note:</strong> ICD-10-CM diagnosis codes entered in 67K–67Q are not required for systematic claims processing. <strong>Required:</strong> POA indicator—Enter the applicable POA indicator in the shaded area for inpatient claims. <strong>Refer to:</strong> Subsection 6.4.2.8.3, “Inpatient Hospital Claims” in this section for POA values.</td>
</tr>
<tr>
<td>69</td>
<td>Admit DX code</td>
<td>Enter the ICD-10-CM diagnosis code indicating the cause of admission or include a narrative <strong>Note:</strong> The admitting diagnosis is only for inpatient claims.</td>
</tr>
<tr>
<td>70a-70c</td>
<td>Patient’s reason DX</td>
<td><strong>Optional:</strong> New block indicating the patient’s reason for visit on unscheduled outpatient claims.</td>
</tr>
<tr>
<td>71</td>
<td>Prospective Payment System (PPS) code</td>
<td><strong>Optional:</strong> The PPS code is assigned to the claim to identify the DRG based on the grouper software called for under contract with the primary payer.</td>
</tr>
<tr>
<td>72a-72c</td>
<td>External cause of injury (ECI) and POA indication</td>
<td><strong>Optional:</strong> Enter the ICD-10-CM diagnosis code in the unshaded area to the highest level of specificity available for each additional diagnosis. <strong>Required:</strong> POA indicator—Enter the applicable POA indicator in the shaded area for inpatient claims. <strong>Refer to:</strong> Subsection 6.4.2.8.3, “Inpatient Hospital Claims” in this section for POA values.</td>
</tr>
<tr>
<td>74</td>
<td>Principal procedure code and date</td>
<td>Enter the ICD-10-CM procedure code for each surgical procedure and the date (MM/DD/YYYY) each was performed.</td>
</tr>
<tr>
<td>74a-74e</td>
<td>Other procedure codes and dates</td>
<td>Enter the ICD-10-CM procedure code for each surgical procedure and the date (MM/DD/YYYY) each was performed.</td>
</tr>
<tr>
<td>76</td>
<td>Attending provider</td>
<td>Enter the attending provider name and NPI. Outpatient claims require an attending provider. Inpatient claims, services that require an attending provider are defined as those listed in the ICD-10-CM coding manual volume 3, which includes surgical, diagnostic, or medical procedures.</td>
</tr>
<tr>
<td>77</td>
<td>Operating</td>
<td>Enter operating provider’s name (last name and first name) and NPI number of the operating provider.</td>
</tr>
</tbody>
</table>
### 6.6.4 Filing Tips for Outpatient Claims

The following are outpatient claim filing tips:

- Use HCPCS codes in Block 44 when available and give a narrative description in Block 43 for all services and supplies provided.

**Important:** Services and supplies that exceed the 28 items per claim limitation must be submitted on an additional UB-04 CMS-1450 paper claim form and will be assigned a different claim number by TMHP.

<table>
<thead>
<tr>
<th>Block No.</th>
<th>Description</th>
<th>Guidelines</th>
</tr>
</thead>
<tbody>
<tr>
<td>78-79</td>
<td>Other</td>
<td>Other provider’s name (last name and first name) and NPI. NPI number of the referring and prescribing provider. Other operating physician—An individual performing a secondary surgical procedure or assisting the operating physician. Required when another operating physician is involved. Rendering provider—The health-care professional who performed, delivered, or completed a particular medical service or nonsurgical procedure. <strong>Important:</strong> Qualifier 82 is required to identify the rendering provider for acute care inpatient and outpatient institutional services. <strong>Note:</strong> If the referring physician is a resident, Blocks 76 through 79 must identify the physician who is supervising the resident.</td>
</tr>
</tbody>
</table>
| 80        | Remarks     | This block is used to explain special situations such as the following:  
- The home health agency must document in writing the number of Medicare visits used in the nursing plan of care and also in this block.  
- If a patient stays beyond dismissal time, indicate the medical reason if additional charge is made.  
- If billing for a private room, the medical necessity must be indicated, signed, and dated by the physician.  
- If services are the result of an accident, the cause and location of the accident must be entered in this block. The time must be entered in Block 39.  
- If laboratory work is sent out, the name and address or the provider identifier of the facility where the work was forwarded must be entered in this block.  
- If the services resulted from a family planning provider’s referral, write “family planning referral.”  
- If services were provided at another facility, indicate the name and address of the facility where the services were rendered.  
- Request for 110-day rule for a third party insurance. |
| 81A-81D   | Code code (CC) | **Optional:** Area to capture additional information necessary to adjudicate the claims. required when, in the judgment of the provider, the information is needed to substantiate the medical treatment and is not support elsewhere on the claim data set. |
• Combine central supplies and bill as one item. IV supplies may be combined and billed as one item. Include appropriate quantities and total charges for each combined procedure code used. Using combination procedure codes conserves space on the claim form.

• The 28-item limitation per claim: a UB-04 CMS-1450 paper claim form submitted with 28 or fewer items is given an internal control number (ICN) by TMHP. Multipage claim forms are processed as one claim for that client if all pages contain 28 or fewer items.

• Itemized Statements: Itemized statements are not used for assignment of procedure codes. HCPCS codes or narrative descriptions of procedures must be reflected on the face of the UB-04 CMS-1450 paper claim form. Attachments will only be used for clarification purposes.

Refer to: Subsection 6.3.3, “Procedure Coding” in this section.

6.7 American Dental Association (ADA) Dental Claim Filing Instructions

Providers billing for dental services and Intermediate Care Facility for Individuals with Intellectual Disabilities (ICF-IID) dental services may bill electronically or use the ADA claim form.

Note: TMHP is responsible for reimbursing all THSteps dental services provided by dentists.

6.7.1 ADA Dental Claim Electronic Billing

Electronic billers must submit THSteps dental claims using TexMedConnect or an approved vendor software that uses the ANSI ASC X12 837D 5010 format. Specifications are available to providers developing in-house systems and software developers and vendors. Because each software package is different, block locations may vary. Contact the software developer or vendor for this information. Direct questions and development requirements to the TMHP EDI Help Desk at 1-888-863-3638.

Note: Dental providers who submit American National Standards Institute, Accredited Standards Committee X12 (ANSI ASC X12N) 837D transactions through the TMHP Electronic Data Interchange (EDI) are required to include the header date of service (HDOS) to comply with International Classification of Diseases, Tenth Revision, Clinical Modification (ICD-10-CM) claims processing guidelines.

Refer to: “Section 3: TMHP Electronic Data Interchange (EDI)” (Vol. 1, General Information) for more information about electronic filing.

6.7.2 ADA Dental Claim Form (Paper) Billing

All participating THSteps claim providers are required to submit a ADA Dental claim form for paper claim submissions to Texas Medicaid. These forms may be obtained by contacting the ADA at 1-800-947-4746.

Claims must contain the billing provider’s complete name, address and a provider identifier. Claims without a provider name, address, and provider identifier cannot be processed.

6.7.3 ADA Dental Claim Form

Samples of the ADA Dental Claim form can be found on the ADA website at www.ada.org.

6.7.4 ADA Dental Claim Form Instruction Table

The following table is an itemized description of the questions appearing on the form. Thoroughly complete the ADA Dental claim form according to the instructions in the table to facilitate prompt and accurate reimbursement and reduce follow-up inquiries.
<table>
<thead>
<tr>
<th>ADA Block No.</th>
<th>ADA Description</th>
<th>Instructions</th>
</tr>
</thead>
</table>
| 1            | Type of Transaction                                  | For Texas Medicaid, check the Statement of Actual Services Box. The other two boxes are not applicable. Do not use the ADA Dental Claim Form as a Texas Medicaid Prior Authorization form.  
**Refer to:** THSteps Dental Mandatory Prior Authorization Request Form on the TMHP website at www.tmhp.com. |
| 2            | Predetermination/ Preauthorization Number             | Enter prior authorization number if assigned by Medicaid.                                                                                                                                                                                                                                                                                      |
| 3            | Company/Plan Name, Address, City, State, ZIP Code    | Enter TMHP and the address.  
**Refer to:** Subsection A.11, “Written Communication With TMHP” in “Appendix A: State, Federal, and TMHP Contact Information” (Vol. 1, General Information).                                                                                     |
| 4            | Other Dental or Medical Coverage                     | Check applicable box. If both “Dental” and “Medical” are marked, complete blocks 5–11 for dental only.                                                                                                                                                                                                                                       |
| 5-11         | Other Coverage Information                           | General notes:  
- Enter the information for non-Medicaid insurance coverage.  
- Enter the information for the policyholder or subscriber, not necessarily the patient. May be a parent or legal guardian of the patient receiving treatment.                                                                                           |
<p>| 5            | Name of Policyholder/ Subscriber in # 4              | Enter the policyholder/subscriber name.                                                                                                                                                                                                                                                                                                      |
| 6            | Date of Birth (MM/DD/CCYY)                           | Enter policyholder/subscriber eight-digit date of birth (MM/DD/YYYY).                                                                                                                                                                                                                                                                       |
| 7            | Gender                                               | Check the appropriate box for the policyholder/subscriber gender                                                                                                                                                                                                                                                                            |
| 8            | Policyholder/Subscriber ID                           | Enter policyholder/subscriber identifier.                                                                                                                                                                                                                                                                                                   |
| 9            | Plan/Group Number                                    | Enter policyholder/subscriber plan/group number.                                                                                                                                                                                                                                                                                           |
| 10           | Patient’s Relationship to Person Named in # 5        | Enter the patient’s relationship to policyholder/subscriber.                                                                                                                                                                                                                                                                               |
| 11           | Other Insurance Company/Dental Benefit Plan Name, Address, City, State, ZIP Code | Enter the contact information for the insurance company providing the non-Medicaid coverage.                                                                                                                                                                                                                                              |
| 12           | Policyholder/Subscriber Name (Last, First, Middle Initial, Suffix), Address, City, State, ZIP Code | Enter the Medicaid patient’s last name, first name, and middle initial as printed on the Medicaid identification form.                                                                                                                                                                                                                   |
| 13           | Date of Birth (MM/DD/CCYY)                           | Enter the Medicaid patient’s date of birth (MM/DD/YYYY).                                                                                                                                                                                                                                                                                     |</p>
<table>
<thead>
<tr>
<th>ADA Block No.</th>
<th>ADA Description</th>
<th>Instructions</th>
</tr>
</thead>
<tbody>
<tr>
<td>14</td>
<td>Gender</td>
<td>Check the appropriate box for the Medicaid patient’s gender.</td>
</tr>
<tr>
<td>15</td>
<td>Policyholder/Subscriber ID</td>
<td>Enter nine-digit patient number from the Medicaid identification form.</td>
</tr>
<tr>
<td>16</td>
<td>Plan/Group/Number</td>
<td>Enter the billing or performing provider’s benefit code, if applicable.</td>
</tr>
<tr>
<td>17</td>
<td>Employer Name</td>
<td>Not applicable to Texas Medicaid.</td>
</tr>
<tr>
<td>18</td>
<td>Relationship to Policyholder/Subscriber in #12 Above</td>
<td>Not applicable to Texas Medicaid.</td>
</tr>
<tr>
<td>19</td>
<td>Reserved for Local Use</td>
<td>Leave blank and skip to Item 20. (Field was previously used to report “Student Status”) Include the appropriate modifier.</td>
</tr>
<tr>
<td>20</td>
<td>Name (Last, First, Middle Initial, Suffix), Address, City, State, ZIP Code</td>
<td>Not applicable to Texas Medicaid.</td>
</tr>
<tr>
<td>21</td>
<td>Date of Birth (MM/DD/CCYY)</td>
<td>Not applicable to Texas Medicaid.</td>
</tr>
<tr>
<td>22</td>
<td>Gender</td>
<td>Not applicable to Texas Medicaid.</td>
</tr>
<tr>
<td>23</td>
<td>Patient ID/Account # (Assigned by Dentist)</td>
<td><strong>Optional:</strong> Enter the patient identification number if it is different than the subscriber/insured’s identification number. Used by dental office to identify internal patient account number.</td>
</tr>
<tr>
<td>24</td>
<td>Procedure Date (MM/DD/CCYY)</td>
<td>Enter the eight-digit date of service (MM/DD/YYYY).</td>
</tr>
<tr>
<td>25</td>
<td>Area of Oral Cavity</td>
<td>Not applicable to Texas Medicaid.</td>
</tr>
<tr>
<td>26</td>
<td>Tooth System</td>
<td>Not applicable to Texas Medicaid.</td>
</tr>
</tbody>
</table>
| 27           | Tooth Number(s) or Letter(s) | Enter the Tooth ID as required for procedure code.  
**Refer to:** Subsection 4.2.13, “Tooth Identification (TID) and Surface Identification (SID) Systems” in the *Children’s Services Handbook* (Vol. 2, Provider Handbooks). |
| 28           | Tooth Surface   | Enter Surface ID as required for procedure code.  
**Refer to:** Subsection 4.2.13, “Tooth Identification (TID) and Surface Identification (SID) Systems” in the *Children’s Services Handbook* (Vol. 2, Provider Handbooks). |
| 29           | Procedure Code  | Use an appropriate Current Dental Terminology (CDT) procedure code. |
| 29a          | Diagnosis Code Pointer | Enter the letter(s) from Box 34 that identified the diagnosis code(s) applicable to the dental procedure. List the primary diagnosis pointer first. |
| 29b          | Procedure Quantity | Enter the number of times (01-99) the procedure identified in Item 29 is delivered to the patient on the date of service shown in item 24. The default value is “01”. |
### ADA Block No. ADA Description Instructions

<table>
<thead>
<tr>
<th>ADA Block No.</th>
<th>ADA Description</th>
<th>Instructions</th>
</tr>
</thead>
<tbody>
<tr>
<td>30</td>
<td>Description</td>
<td>Provide a brief description of the service provided (e.g., abbreviation of the procedure code’s nomenclature).</td>
</tr>
<tr>
<td>31</td>
<td>Fee</td>
<td>Enter usual and customary charges for each service listed. Charges must not be higher than the fees charged to private pay clients.</td>
</tr>
<tr>
<td>31a</td>
<td>Other Fee(s)</td>
<td>When other changes applicable to dental services provided must be reported, enter the amount here. Charges may include state tax and other charges imposed by regulatory bodies. Identify the source of each payment date in Block 11. If the client makes a payment, the reason for the payment must be identified in Block 11.</td>
</tr>
</tbody>
</table>
| 32            | Total Fee             | Enter the sum of all fees in Block 31. For multi-page claims, enter “continue” on initial and subsequent claim forms. Indicate the total of all charges on the last claim.  
**Note:** Indicate the page number of the attachment (for example, page 2 of 3) in the top right-hand corner of the form. |
| 33            | Missing Teeth Information | Mark an “X” on each missing tooth. For identifying missing permanent dentition only. Report missing teeth when pertinent to periodontal, prosthodontic (fixed and removable), or implant services procedures on a particular claim. |
| 34            | Diagnosis Code List Qualifier | Enter “AB= ICD-10” to identify the diagnosis code source.  |
| 34a           | Diagnosis Codes(s)    | Enter up to four applicable diagnosis codes after each letter (A-D). The primary diagnosis code is entered adjacent to the letter “A”. |
| 35            | Remarks               | Use this space for:  
- Explanation of exception to periodicity.  
- The facility name and address and NPI if the place of treatment indicated in Block 38 is not the provider’s office.  
- Explanation of emergency if indicated in Block 45.  
- To provide more information such as reports for local orthodontia codes, 999 codes, multiple supernumerary teeth, or remarks. |
<p>| 36            | Patient/Guardian signature | Not applicable to Texas Medicaid.                                             |
| 37            | Subscriber signature  | Not applicable to Texas Medicaid.                                             |</p>
<table>
<thead>
<tr>
<th>ADA Block No.</th>
<th>ADA Description</th>
<th>Instructions</th>
</tr>
</thead>
</table>
| 38           | Place of Treatment               | Enter the 2-digit place of service (POS) code for professional claims, which is a Health Insurance Portability and Accountability Act (HIPAA) standard. Frequently used POS codes include the following:  
  • 11 = Office  
  • 12 = Home  
  • 21 = Inpatient hospital  
  • 22 = Outpatient hospital  
  • 31 = Skilled nursing facility  
  • 32 = Nursing facility |
| 39           | Enclosures                       | Enter a “Y” or “N” to indicate whether or not there are enclosures of any type included with the claim submission (e.g., radiographs, oral images, models).                                                       |
| 40           | Is Treatment for Orthodontics?   | Check Yes or No as appropriate.                                                                                                                                                                             |
| 41           | Date Appliance Placed            | Not applicable to Texas Medicaid.                                                                                                                                                                           |
| 42           | Months of Treatment Remaining    | Not applicable to Texas Medicaid.                                                                                                                                                                           |
| 43           | Replacement of Prosthesis?       | Not applicable to Texas Medicaid.                                                                                                                                                                           |
| 44           | Date Prior Placement             | Not applicable to Texas Medicaid.                                                                                                                                                                           |
| 45           | Treatment Resulting from (Check applicable box) | Providers are required to check the Other Accident box for emergency claim reimbursement. If the Other Accident box is checked, information about the emergency must be provided in Block 35. |
| 46           | Date of Accident (MM/DD/CCYY)    | Not applicable to Texas Medicaid.                                                                                                                                                                           |
| 47           | Auto Accident State              | Not applicable to Texas Medicaid.                                                                                                                                                                           |
| 48           | Name, Address, City, State, ZIP Code | Enter the name and address of the billing group or individual provider. Do not enter the name and address of a provider employed within a group.                                                           |
| 49           | NPI                              | Enter the billing provider’s NPI for a group or an individual. Do not enter the NPI for a provider employed within a group.                                                                              |
| 50           | License Number                   | Not applicable to Texas Medicaid.                                                                                                                                                                           |
| 51           | Social Security Number (SSN) or Tax Identification Number (TIN) | Not applicable to Texas Medicaid.                                                                                                                                                                           |
| 52           | Telephone Number                 | Enter the area code and number for the billing group or individual. Do not enter the telephone number of a provider employed within a group.                                                              |
| 52A          | Additional Provider ID           | Enter the nine-digit TPI assigned to the billing dentist or dental entity. Do not enter the TPI for a provider employed within a group.                                                                      |
6.8 Family Planning Claim Filing Instructions

The following providers bill for services using the ANSI ASC X12 837P 5010 electronic specifications or the CMS-1500 paper claim form:

| Providers |
|-----------------|-----------------|-----------------|
| Clinical nurse specialist (CNS) |
| Family Planning title agencies contracted with HHSC |
| Federally Qualified Health Center (FQHC) |
| Nurse practitioner (NP) |
| Physician |
| Physician assistant (PA) |

6.8.1 Family Planning Electronic Billing

Electronic billers must submit family planning claims with TexMedConnect or approved vendor software that uses the ANSI ASC X12 837P 5010 format. Specifications are available to providers developing in-house systems, software developers, and vendors on the TMHP website at [www.tmhp.com/topics/edi](http://www.tmhp.com/topics/edi). Because each software developer is different, location of fields may vary. Contact the software developer or vendor for this information. Direct questions and development requirements to the TMHP EDI Help Desk at 1-888-863-3638.


6.9 Family Planning Claim Form (Paper Billing)

Claims must contain the billing providers complete name, address, and a provider identifier. Claims without a provider name, address, and provider identifier cannot be processed.
### 6.9.1 2017 Claim Form
A copy of a blank 2017 Claim Form and copies of example completed claim forms are available on the Claim Form Examples page of the TMHP website at www.tmhp.com.

### 6.9.2 2017 Claim Form Instruction Table
The instructions describe what information must be entered in each of the block numbers of the 2017 Claim Form.

<table>
<thead>
<tr>
<th>Block No.</th>
<th>Description</th>
<th>Guidelines</th>
<th>Required (Paper)</th>
</tr>
</thead>
</table>
| 1         | Program     | Check the box for the specific program to which these services are billed:  
- Family Planning Program: XIX (Check this box for Title XIX family planning services and for Healthy Texas Women (HTW) program services)  
- DSHS Family Planning Program (DFPP)                                                                                                                                                                                                                                                 | XIX, DFPP       |
| 2a        | Billing provider TPI | Enter the billing provider’s nine-digit TPI.                                                                                                                                                                                                                                         | All             |
| 2b        | Billing provider NPI | Enter the billing provider’s NPI.                                                                                                                                                                                                                                               | All             |
| 3         | Provider name | Enter the provider’s name as enrolled with TMHP.                                                                                                                                                                                                                                   | All             |
| 4         | Eligibility date (DFPP) | Enter the date (MM/DD/CCYY) this client was designated eligible for DFPP services.  
For DFPP, the eligibility date can be found on the following forms:  
- INDIVIDUAL Eligibility Form (EF05-14215)  
- HOUSEHOLD Eligibility Form (EF05-14214)  
- HOUSEHOLD Eligibility Worksheet (EF05-13227)  
- An approved DSHS substitute                                                                                                                                                                                                                                                        | DFPP            |
| 5         | DSHS Client no. (Medicaid PCN if XIX) | If previous DFPP, claims or encounters have been submitted to TMHP, enter the client’s nine-digit DSHS client number, which begins with “F.”  
If the client has Title XIX Medicaid, enter the client’s nine-digit client number from the Medicaid Identification form.  
If this is a new client, without Medicaid, leave this block blank and TMHP will assign a DSHS client number for the client.                                                                                       | XIX             |
<p>| 6         | Patient’s name (last name, first name, middle initial) | Enter the client’s last name, first name, and middle initial as printed on the Medicaid Identification Form, if Title XIX, or as printed in the provider’s records, if DFPP.                                                                                                                                 | All             |
| 7         | Address (street, city, state) | Enter the client’s complete home address as described by the client (street, city, and state). This reflects the location where the client lives.                                                                                                                                                                                                 | All             |
| 7a        | ZIP Code    | Enter the client’s ZIP Code.                                                                                                                                                                                                                                                     | All             |
| 8         | County of residence | Enter the county code that corresponds to the client’s address. Please use the HHSC county codes.                                                                                                                                                                                     | All             |</p>
<table>
<thead>
<tr>
<th>Block No.</th>
<th>Description</th>
<th>Guidelines</th>
<th>Required (Paper)</th>
</tr>
</thead>
<tbody>
<tr>
<td>9</td>
<td>Date of birth</td>
<td>Enter numerically the month, day, and year (MM/DD/CCYY) the client was born.</td>
<td>All</td>
</tr>
<tr>
<td>10</td>
<td>Sex</td>
<td>Indicate the client’s sex by checking the appropriate box.</td>
<td>All</td>
</tr>
<tr>
<td>11</td>
<td>Patient status</td>
<td>Indicate if this is the client’s first visit to this provider (new patient) or if this client has been to this provider previously (established patient). If the provider’s records have been purged and the client appears to be new to the provider, check “New Patient.”</td>
<td>All</td>
</tr>
<tr>
<td>12</td>
<td>Patient’s Social Security number</td>
<td>Enter the client’s nine-digit Social Security number (SSN). If the client does not have a SSN, or refuses to provide the number, enter 000-00-0001.</td>
<td>All</td>
</tr>
<tr>
<td>13</td>
<td>Race (code #)</td>
<td>Indicate the client’s race by entering the appropriate race code number in the box.</td>
<td>All</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Aggregate categories used here are consistent with reporting requirements of the Office of Management and Budget Statistical Direction.</td>
<td></td>
</tr>
<tr>
<td></td>
<td></td>
<td>Race is independent of ethnicity and all clients should be self-categorized as White, Black or African American, American Indian or Native Alaskan, Asian, Native Hawaiian or other Pacific Islander, or Unknown or Not Reported. An “Hispanic” client must also have a race category selected.</td>
<td></td>
</tr>
<tr>
<td>13a</td>
<td>Ethnicity</td>
<td>Indicate whether the client is of Hispanic descent by entering the appropriate code number in the box.</td>
<td>All</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Ethnicity is independent of race and all clients should be counted as either Hispanic or non-Hispanic. The Office of Management and Budget defines Hispanic as “a person of Mexican, Puerto Rican, Cuban, Central, or South American culture or origin, regardless of race.”</td>
<td></td>
</tr>
<tr>
<td>14</td>
<td>Marital status</td>
<td>Indicate the client’s marital status by entering the appropriate marital code number in the box.</td>
<td>All</td>
</tr>
<tr>
<td>15</td>
<td>Family income (all)</td>
<td>DFPP: Use the gross monthly income calculated and reported on the INDIVIDUAL Eligibility Form (EF05-14215), the HOUSEHOLD Eligibility Form (EF05-14214), or the HOUSEHOLD Eligibility Worksheet (EF05-13227). Title XIX: Enter the gross monthly income reported by the client. Be sure to include all sources of income. If income is received in a lump sum, or if it is for a period of time greater than a month (e.g., for seasonal employment), divide the total income by the number of months included in the payment period. If income is paid weekly, multiply weekly income by 4.33. If paid every two weeks, multiply amount by 2.165. If paid twice a month, multiply by 2. Enter $1.00 for clients not wishing to reveal income information.</td>
<td>All</td>
</tr>
<tr>
<td>Block No.</td>
<td>Description</td>
<td>Guidelines</td>
<td>Required (Paper)</td>
</tr>
<tr>
<td>----------</td>
<td>-------------------------------------------</td>
<td>---------------------------------------------------------------------------</td>
<td>------------------</td>
</tr>
<tr>
<td>15a</td>
<td>Family size</td>
<td>DFPP: Use the family size reported on the eligibility assessment tool.</td>
<td>All</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Title XIX providers: Enter the number of family members supported by the income listed in Box 15. Must be at least “one.”</td>
<td></td>
</tr>
<tr>
<td>16</td>
<td>Number times pregnant</td>
<td>Enter the number of times this client has been pregnant. If male, enter zero.</td>
<td>XIX</td>
</tr>
<tr>
<td>17</td>
<td>Number live births</td>
<td>Enter the number of live births for this client. If male, enter zero.</td>
<td>XIX</td>
</tr>
<tr>
<td>18</td>
<td>Number living children</td>
<td>Enter the number of living children this client has. This also must be completed for male clients.</td>
<td>XIX</td>
</tr>
<tr>
<td>19</td>
<td>Primary birth control method before initial visit</td>
<td>Enter the appropriate code letter (a through r) in the box.</td>
<td>XIX</td>
</tr>
<tr>
<td>20</td>
<td>Primary birth control method at end of this visit</td>
<td>Enter the appropriate code letter (a through r) in the box.</td>
<td>XIX</td>
</tr>
<tr>
<td>21</td>
<td>If no method used at end of this visit, give reason (required only if #20=r)</td>
<td>If the primary birth control method at the end of the visit was “no method” (r), you must complete this box with an appropriate code letter from this block (a through g).</td>
<td>XIX (only if #20=r)</td>
</tr>
<tr>
<td>22</td>
<td>Is there other insurance available?</td>
<td>Check the appropriate box.</td>
<td>Optional</td>
</tr>
<tr>
<td>23</td>
<td>Other insurance name and address</td>
<td>Enter the name and address of the health insurance carrier.</td>
<td>Optional</td>
</tr>
<tr>
<td>24a</td>
<td>Insured’s policy/group no.</td>
<td>Enter the insurance policy number or group number.</td>
<td>Optional</td>
</tr>
<tr>
<td>24b</td>
<td>Benefit code</td>
<td>Benefit code, if applicable for the billing or performing provider.</td>
<td>Optional</td>
</tr>
<tr>
<td>25</td>
<td>Other insurance paid amount</td>
<td>Enter the amount paid by the other insurance company. If payment was denied, enter “Denied” in this block.</td>
<td>Optional</td>
</tr>
<tr>
<td>25a</td>
<td>Date of notification</td>
<td>Enter the date of the other insurance payment or denial in this block. This must be in the format of MM/DD/CCYY.</td>
<td>Optional</td>
</tr>
<tr>
<td>26</td>
<td>Name of referring provider</td>
<td>If a non-family planning service is being billed, and the service requires a referring provider, enter the provider’s name.</td>
<td>XIX (if available)</td>
</tr>
<tr>
<td>27b</td>
<td>Referring NPI</td>
<td>If a non-family planning service is being billed and the service requires a referring provider identifier, enter the referring provider’s NPI.</td>
<td>XIX</td>
</tr>
<tr>
<td>Block No.</td>
<td>Description</td>
<td>Guidelines</td>
<td>Required (Paper)</td>
</tr>
<tr>
<td>-----------</td>
<td>-------------</td>
<td>------------</td>
<td>-----------------</td>
</tr>
<tr>
<td>28</td>
<td>Level of practitioner</td>
<td>Enter the level of practitioner that performed the service. Primary care or generalist physicians and specialists are correctly classified as “Physicians.” Certified nurse-midwives, nurse practitioners, clinical nurse specialists, and physician assistants providing encounters are correctly categorized as “Midlevel.” Encounters provided by a registered nurse or a licensed vocational nurse would be categorized as “Nurse.” Encounters provided by staff not included in the preceding classifications would be correctly categorized as “Other.” If a client has encounters with staff members of different categories during one visit, select the highest category of staff with whom the client interacted. Optional for agencies not receiving any DFPP funding.</td>
<td>DFPP</td>
</tr>
<tr>
<td>29</td>
<td>Diagnosis code (Relate Items A-L to service line 32E)</td>
<td>Enter the applicable ICD indicator to identify which version of ICD codes is being reported. 9 = ICD-9-CM 0 = ICD-10-CM Enter the patient’s diagnosis and/or condition codes. List no more than 12 diagnosis codes. Relate lines A-L to the lines of service in 24E by the letter of the line. Use the highest level of specificity. Do not provide narrative description in this field.</td>
<td>All</td>
</tr>
<tr>
<td>30</td>
<td>Authorization number</td>
<td>Enter the authorization number for the client, if appropriate.</td>
<td>Optional</td>
</tr>
<tr>
<td>31</td>
<td>Date of occurrence</td>
<td>Use this section when billing for complications related to sterilizations, contraceptive implants, or intrauterine devices (IUDs). This block should contain the date (MM/DD/CCYY) of the original sterilization, implant, or IUD procedure associated with the complications currently being billed.</td>
<td>All, if billing complications</td>
</tr>
<tr>
<td>32A</td>
<td>Dates of service</td>
<td>Enter the dates of service (DOS) for each procedure provided in a MM/DD/CCYY format. If more than one DOS is for a single procedure, each date must be given (such as 3/16, 17, 18/2010). Electronic Billers Medicaid does not accept multiple (to-from) dates on a single-line detail. Bill only one date per line. NDC In the shaded area, enter the: • NDC qualifier of N4 (e.g., N4) • The 11-digit NDC number on the package or vial from which the medication was administered. Do not enter hyphens or spaces within this number (e.g., 00409231231). Example: N400409231231</td>
<td>All</td>
</tr>
<tr>
<td>Block No.</td>
<td>Description</td>
<td>Guidelines</td>
<td>Required (Paper)</td>
</tr>
<tr>
<td>----------</td>
<td>-------------</td>
<td>------------</td>
<td>-----------------</td>
</tr>
<tr>
<td>32B</td>
<td>Place of service</td>
<td>Enter the appropriate POS code for each service from the POS table in the Texas Medicaid Provider Procedures Manual. If the client is registered at a hospital, the POS must indicate inpatient or outpatient status at the time of service.</td>
<td>All</td>
</tr>
<tr>
<td>32C</td>
<td>Reserved for local use</td>
<td>Leave this block blank. <strong>Note:</strong> TOS codes are no longer required for claims submission.</td>
<td>Optional</td>
</tr>
<tr>
<td>32D</td>
<td>Procedures, services, or supplies CPT/HCPCS modifier</td>
<td>Enter the appropriate CPT or HCPCS procedure codes for all procedures/services billed. <strong>NDC</strong> In the shaded area, enter the NDC quantity of units administered (up to 12 digits, including the decimal point.). A decimal point must be used for fractions of a unit.</td>
<td>All</td>
</tr>
<tr>
<td>32E</td>
<td>Dx. ref. (29)</td>
<td>Enter the diagnosis line item reference (A-L) for each service or procedure as it relates to each ICD diagnosis code identified in Block 29. When multiple services are performed, the primary reference number for each service should be listed first, other applicable services should follow. The reference letter(s) should be A-L or multiple letters as applicable. Diagnosis codes must be entered in Form Field 29 only. Do not enter diagnosis codes in Form Field 32E.</td>
<td>All</td>
</tr>
<tr>
<td>32F</td>
<td>Units or days (quantity)</td>
<td>If multiple services are performed on the same day, enter the number of services performed (such as the quantity billed). <strong>NDC</strong> In the shaded area, enter the NDC unit of measurement code. There are 5 allowed values: F2, GR, ML, UN or ME.</td>
<td>All</td>
</tr>
<tr>
<td>32G</td>
<td>$ Charges</td>
<td>Indicate the charges for each service listed (quantity multiplied by reimbursement rate). Charges must not be higher than fees charged to private-pay clients.</td>
<td>All</td>
</tr>
<tr>
<td>32H (a)</td>
<td>Performing provider number (XIX only)-TPI</td>
<td>Members of a group practice (except pathology and renal dialysis groups) must identify the nine-digit TPI of the provider within the group who performed the service. <strong>Note:</strong> To avoid unnecessary denials, DFPP providers should include the performing provider’s TPI on the claim. Although not required for DFPP claims, if a claim or encounter that was submitted through DFPP is later determined eligible to be paid under Title XIX, the claim will be denied if the performing provider information is missing.</td>
<td>XIX</td>
</tr>
<tr>
<td>Block No.</td>
<td>Description</td>
<td>Guidelines</td>
<td>Required (Paper)</td>
</tr>
<tr>
<td>----------</td>
<td>--------------------------------------------------</td>
<td>----------------------------------------------------------------------------------------------</td>
<td>------------------</td>
</tr>
</tbody>
</table>
| 32H (b)  | Performing provider number (XIX only)-NPI          | Optional: Members of a group practice (except pathology and renal dialysis groups) must identify NPI of the provider within the group who performed the service.  
**Note:** To avoid unnecessary denials, DFPP providers should include the performing provider’s TPI on the claim. Although not required for DFPP claims, if a claim or encounter that was submitted through DFPP is later determined eligible to be paid under Title XIX, the claim will be denied if the performing provider information is missing. | XIX              |
| 33       | Federal tax ID number/EIN (optional)              | Enter the federal TIN (Employer Identification Number [EIN]) that is associated with the provider identifier enrolled with TMHP.  
**Note:** To avoid unnecessary denials, PHC and EPHC providers should include the federal tax ID on the claim. Although not required for PHC and EPHC claims, if a claim or encounter that was submitted through PHC or EPHC is later determined eligible to be paid under Title XIX, the claim will be denied if the tax ID information is missing. | XIX, DFPP       |
| 34       | Patient’s account number (optional)               | Enter the client’s account number that is used in the provider’s office for its payment records. | Optional         |
| 35       | Patient copay assessed (DFPP)                     | If the client was assessed a copayment (DFPP), enter the dollar amount assessed.  
If no copay was assessed, enter $0.00. Copay cannot be assessed for Title XIX clients.  
Copayment must not exceed $30.00 for DFPP patients. | DFPP             |
| 36       | Total charges                                     | Enter the total of separate charges for each page of the claim. Enter the total of all pages on last claim if filing a multipage claim. | All              |
| 37       | Signature of physician or supplier               | The physician/supplier or an authorized representative must sign and date the claim. Billing services may print “Signature on file” in place of the provider’s signature if the billing service obtains and retains on file a letter signed and dated by the provider authorizing this practice.  
When providers enroll to be an electronic biller, the “Signature on file” requirement is satisfied during the enrollment process. | All              |
6.10 Vision Claim Form

All vision services must be billed on a CMS-1500 paper claim form or the appropriate electronic formats. Vision claims submitted on other forms are denied with EOB 01145, “Claim form not allowed for this program.”

For eyewear claims beyond program benefits, (e.g., replacing lost or destroyed eye wear), providers must have the patient sign the “Patient Certification Form” and retain in their records. Do not submit form to TMHP.

Refer to: Vision Care Eyeglass Patient (Medicaid Client) Certification Form on the TMHP website at www.tmhp.com.

The following table shows the blocks required for vision claims on a CMS-1500 paper claim form.

<table>
<thead>
<tr>
<th>Block No.</th>
<th>Description</th>
<th>Guidelines</th>
<th>Required (Paper)</th>
</tr>
</thead>
<tbody>
<tr>
<td>38</td>
<td>Name and address of facility where services were rendered (if other than home or office)</td>
<td>If the services were provided in a place other than the client’s home or the provider’s facility, enter name, address, and ZIP Code, of the facility (such as the hospital or birthing center) where the service was provided. Independently practicing health-care professionals must enter the name and number of the school district/cooperative where the child is enrolled (SHARS). For laboratory specimens sent to an outside laboratory for additional testing, the complete name and address of the outside laboratory should be entered. The laboratory should bill Texas Medicaid for the services performed.</td>
<td>XIX</td>
</tr>
<tr>
<td>38a</td>
<td>NPI</td>
<td>Enter the NPI of the provider where services were rendered (if other than home or office).</td>
<td>XIX</td>
</tr>
<tr>
<td>39</td>
<td>Physician’s, supplier’s billing name, address, ZIP Code, and telephone number</td>
<td>Enter the billing provider name, street, city, state, ZIP Code, and telephone number.</td>
<td>Optional</td>
</tr>
</tbody>
</table>
6.11 Remittance and Status (R&S) Report

The R&S Report provides information on pending, paid, denied, and adjusted claims. TMHP provides weekly R&S Reports to give providers detailed information about the status of claims submitted to TMHP. The R&S Report also identifies accounts receivables established as a result of inappropriate payment. These receivables are recouped from claim submissions. All claims for the same provider identifier and program processed for payment are paid at the end of the week, either by a single check or

<table>
<thead>
<tr>
<th>Block No.</th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td>10</td>
<td>Was condition related to:</td>
</tr>
<tr>
<td></td>
<td>a. Patient’s employment</td>
</tr>
<tr>
<td></td>
<td>b. Auto accident</td>
</tr>
<tr>
<td></td>
<td>c. Other accident</td>
</tr>
<tr>
<td>11</td>
<td>Medicare number</td>
</tr>
<tr>
<td>12</td>
<td>Patient’s or authorized person’s signature</td>
</tr>
<tr>
<td>13*</td>
<td>Insured or authorized person’s signature</td>
</tr>
<tr>
<td>17</td>
<td>Name of referring physician or other source</td>
</tr>
<tr>
<td>17b NPI</td>
<td>Name, provider identifiers, and address of prescribing medical doctor or doctor of optometry</td>
</tr>
<tr>
<td>21</td>
<td>Diagnosis or nature of illness or injury</td>
</tr>
<tr>
<td>24A</td>
<td>DOS</td>
</tr>
<tr>
<td>24B</td>
<td>POS</td>
</tr>
<tr>
<td>24D</td>
<td>Describe procedures, medical services, or supplies furnished for each date given</td>
</tr>
<tr>
<td>24D, Line “5” for new prescription</td>
<td>Prescription/description of lenses and frames</td>
</tr>
<tr>
<td>24D, Line “6” for old prescription</td>
<td></td>
</tr>
<tr>
<td>24E</td>
<td>Diagnosis pointer</td>
</tr>
<tr>
<td>24F</td>
<td>Charges</td>
</tr>
<tr>
<td>26*</td>
<td>The account number for the patient that is used in the provider’s office for its billing records.</td>
</tr>
<tr>
<td>27</td>
<td>Check “YES” or “NO”</td>
</tr>
<tr>
<td>28</td>
<td>Total charges</td>
</tr>
<tr>
<td>29</td>
<td>Amount paid by other insurance</td>
</tr>
<tr>
<td>31</td>
<td>Signature of physician or supplier</td>
</tr>
<tr>
<td>32</td>
<td>Name and address of facility where services were rendered if other than home or office</td>
</tr>
<tr>
<td>33</td>
<td>Telephone number</td>
</tr>
<tr>
<td>33</td>
<td>Physician’s or supplier’s name, address, city, state, and ZIP code</td>
</tr>
<tr>
<td>No longer used</td>
<td>Referral from screening program (THSteps)</td>
</tr>
</tbody>
</table>
or with Electronic Funds Transfer (EFT). If no claim activity or outstanding account receivables exist during the cycle week, the provider does not receive an R&S Report. Providers are responsible for reconciling their records to the R&S to determine payments and denials received.

**Note:** Providers receive a single R&S Report that details Texas Medicaid activities and provides individual program summaries. Combined provider payments are made based on the provider’s settings for Texas Medicaid fee-for-service.

Providers must retain copies of all R&S Reports for a minimum of five years. Providers must not use R&S Report originals for appeal purposes, but must submit copies of the R&S Reports with appeal documentation.

**Refer to:** “Section 3: TMHP Electronic Data Interchange (EDI)” (Vol. 1, General Information) for information on electronic claims submissions.

### 6.11.1 R&S Report Delivery Options

TMHP offers two options for the delivery of the R&S Report:

- A PDF version that is available on the TMHP website through the secure provider portal.
- An Electronic Remittance and Status (ER&S) Report that is available through EDI.

The PDF version of the R&S Report is available through TexMedConnect, and can be downloaded by registered users of the TMHP website at [www.tmhp.com](http://www.tmhp.com). The report is available each Monday morning, immediately following the weekly claims cycle. Payments associated with the R&S Report are released the next Friday following the weekly claims cycle. Newly-enrolled providers are initially set up to receive the PDF version of the R&S Report.

The EDI delivery method is also available. Using HIPAA-compliant EDI standards, the (ER&S 835 file) can be downloaded through the TMHP EDI Gateway using third party software. The ER&S Report is available on Thursday the week the provider payments are released.

**Note:** In rare instances, payments and R&S delivery may be delayed due to a system outage or holiday.

In addition to the PDF R&S Report, an optional R&S Report delivery method is also available. Using HIPAA-compliant EDI standards, the ER&S Report can be downloaded through the TMHP EDI Gateway using TexMedConnect or third party software. The ER&S Report is also available each Monday after the completion of the claims processing cycle.

**Refer to:** “Section 3: TMHP Electronic Data Interchange (EDI)” (Vol. 1, General Information) for more information about EDI formats and enrollment for the ER&S Report.

### 6.11.2 Banner Pages

Banner pages serve two purposes:

- They identify the provider’s name and address.
- They are used to inform providers of new policies and procedures.

The title pages include the following information:

- TMHP address for submitting paper appeals
- Provider’s name, address, and telephone number
- Unique R&S Report number specific to each report
- Provider identifier (TPI, NPI, and atypical provider identifier [API])
- Report sequence number (indicates the week number of the year)
- Date of the week being reported on the R&S Report
• Tax Identification Number
• Page number (R&S Report begins with page 1)
• AIS telephone number
• Taxonomy code

### 6.11.3 R&S Report Field Explanation

- **Patient name.** Lists the client’s last name and first name, as indicated on the eligibility file.
- **Claim number.** The 24-digit Medicaid ICN for a specific claim. The format for the TMHP claim number is expanded to PPP/CCC/MMM/CCYY/JJJ/BBBBB/SSS.

<table>
<thead>
<tr>
<th>Acronym</th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td>PPP</td>
<td>Program</td>
</tr>
<tr>
<td>CCC</td>
<td>Claim type</td>
</tr>
<tr>
<td>MMM</td>
<td>Media source (region)</td>
</tr>
<tr>
<td>CCYY</td>
<td>Year in which the claim was received</td>
</tr>
<tr>
<td>JJJ</td>
<td>Julian date on which the claim was received</td>
</tr>
<tr>
<td>BBBBB</td>
<td>TMHP internal batch number</td>
</tr>
<tr>
<td>SSS</td>
<td>TMHP internal claim sequence within the batch</td>
</tr>
</tbody>
</table>

#### Program Type

<table>
<thead>
<tr>
<th>PPP</th>
<th>Program</th>
</tr>
</thead>
<tbody>
<tr>
<td>001</td>
<td>Long Term Care</td>
</tr>
<tr>
<td>100</td>
<td>Medicaid</td>
</tr>
<tr>
<td>200</td>
<td>Managed Care (for carve-out services administered by TMHP and PCCM claims with dates of service before March 1, 2012)</td>
</tr>
<tr>
<td>300</td>
<td>Family Planning (DSHS Family Planning Program)</td>
</tr>
<tr>
<td>400</td>
<td>CSHCN Services Program</td>
</tr>
<tr>
<td>999</td>
<td>Default/summary for all media regions</td>
</tr>
</tbody>
</table>

#### Claim Type

<table>
<thead>
<tr>
<th>Claim Type</th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td>020</td>
<td>Physician/supplier (Medicaid only) (genetics agencies, THSteps [medical only], FQHC, optometrist, optician)</td>
</tr>
<tr>
<td>021</td>
<td>THSteps (dental)</td>
</tr>
<tr>
<td>023</td>
<td>Outpatient hospital, home health, RHC, FQHC</td>
</tr>
<tr>
<td>030</td>
<td>Physician crossovers</td>
</tr>
<tr>
<td>031</td>
<td>Hospital outpatient crossovers, home health crossovers, RHC crossovers</td>
</tr>
<tr>
<td>040</td>
<td>Inpatient hospital</td>
</tr>
<tr>
<td>050</td>
<td>Inpatient crossover</td>
</tr>
<tr>
<td>056</td>
<td>DSHS Family Planning Program</td>
</tr>
<tr>
<td>058</td>
<td>Family Planning Title XIX</td>
</tr>
</tbody>
</table>
Media Source (MMM)

<table>
<thead>
<tr>
<th>Region</th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td>010</td>
<td>Paper</td>
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<tr>
<td>011</td>
<td>Paper adjustment</td>
</tr>
<tr>
<td>030</td>
<td>Electronic (including TexMedConnect)</td>
</tr>
<tr>
<td>031</td>
<td>Electronic adjustment (including TexMedConnect)</td>
</tr>
<tr>
<td>041</td>
<td>AIS adjustment</td>
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<tr>
<td>051</td>
<td>Mass adjustment</td>
</tr>
<tr>
<td>061</td>
<td>Crossover adjustment</td>
</tr>
<tr>
<td>071</td>
<td>Retroactive eligibility adjustment</td>
</tr>
<tr>
<td>080</td>
<td>State Action Request</td>
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<tr>
<td>081</td>
<td>State Action Request adjustment</td>
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<td>090</td>
<td>Telephone</td>
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<tr>
<td>100</td>
<td>Fax</td>
</tr>
<tr>
<td>110</td>
<td>Mail</td>
</tr>
<tr>
<td>120</td>
<td>Encounter</td>
</tr>
<tr>
<td>121</td>
<td>Encounter Adjustment</td>
</tr>
</tbody>
</table>

- Medicaid #. The client’s Medicaid number.
- Patient Account #. If a patient account number is used on the provider’s claim, it appears here.
- Medical Record #. If a medical record number is used on the provider’s claim, it appears here.
- Medicare #. If the claim is a result of an automatic crossover from Medicare, the last ten digits of the Medicare claim number appears directly under the TMHP claim number.
- Diagnosis. Primary diagnosis listed on the provider’s claim.
- Service Dates. Format MMDDYYYY (month, day, year) in “From” and “To” dates of service.
- TOS/Proc. Indicates by code the specific service provided to the client. The one-digit TOS appears first followed by a HCPCS procedure code. A three-digit code represents a hospital accommodation or ancillary revenue code. For claims paid under prospective payment methodology, it is the code of the DRG.
- Billed Quantity. Indicates the quantity billed per claim detail.
- Billed Charge. Indicates the charge billed per claim detail.
- Allowed Quantity. Indicates the quantity TMHP has allowed per claim detail.
- Allowed Charge. Indicates the charges TMHP has allowed per claim detail. For inpatient hospital claims, the allowed amount for the DRG appears.
- POS Column. The R&S Report includes the POS to the left of the Paid Amount. A one-digit numeric code identifying the POS is indicated in this column. Refer to subsection 6.3.1.1, “Place of Service (POS) Coding” in this section for the appropriate cross-reference among the two-digit numeric POS codes (Medicare), and one-digit numeric code on the R&S Report. Providers using electronic claims submission should continue using the same POS codes.
- Paid Amt. The final amount allowed for payment per claim detail. The total paid amount for the claim appears on the claim total line.
• **EOB Codes and Explanation of Pending Status (EOPS) Codes.** These codes explain the payment or denial of the provider’s claim. The EOB codes are printed next to or directly below the claim. The EOPS codes appear only in “The Following Claims Are Being Processed” section of the R&S Report. The codes explain the status of pending claims and are not an actual denial or final disposition. An explanation of all EOB and EOPS codes appearing on the R&S Report are printed in the Appendix at the end of the R&S Report. Up to five EOB codes are displayed.

• **Total TEFRA Billed and Allowed Charges.** Indicates claim details that have been denied or reduced.

• **Benefit.** Indicates the three digit benefit code associated with the claim.

• **Modifier.** Modifiers have been developed to describe and qualify services provided. For THSteps dental services two modifiers are printed. The first modifier is the TID and the second is the SID.

### 6.11.4 R&S Report Section Explanation

#### 6.11.4.1 Claims – Paid or Denied

The heading “Claims – Paid or Denied Claims” is centered on the top of each page in this section. Claims in this section finalized the week before the preparation of the R&S Report. The claims are sorted by claim status, claim type, and by order of client names. The reported status of each claim will not change unless further action is initiated by the provider, HHSC, or TMHP.

The following information is provided on a separate line for all inpatient hospital claims processed according to prospective payment methodology:

- **Age.** Client’s age according to TMHP records
- **Sex.** Client’s sex according to TMHP records: M = Male, F = Female, U = Unknown
- **Pat-Stat.** Indicates the client’s status at the time of discharge or the last DOS on the claim (refer to instructions for UB-04 CMS-1450 paper claim form, Block 17)
- **Proc.** ICD-10-PCS code indicates the primary surgical procedure used in determining the DRG

**Important:** Only paper claims appear in this section of the R&S Report. Claims filed electronically without required information are rejected. Users are required to retrieve the response file to determine reasons for rejections.

TMHP cannot process incomplete claims. Incomplete claims may be submitted as original claims only if the resubmission is received by TMHP within the original filing deadline.

**Refer to:** Subsection 6.1, “Claims Information” in this section for a description of different claim types.

#### 6.11.4.2 Adjustments to Claims

*Adjustments – Paid or Denied* is centered at the top of each page in this section. Adjustments are sorted by claim type and then patient name and Medicaid number. Media types 011, 021, 031, 041, 051, 061, 071, and 081 appear in this section. An adjustment prints in the same format as a paid or denied claim.

The adjusted claim is listed first on the R&S Report. EOB 00123, “This is an adjustment to previous claim XXXXXXXXXXXXXXXXXXXXXXXX which appears on R&S Report dated XX/XX/XX” follows this claim. Immediately below is the claim as originally processed. An accounts receivable is created for the original claim total as noted by EOB 00601, “A receivable has been established in the amount of the original payment: $XXX,XXX,XXX.XX. Future payments will be reduced or withheld until such amount is paid in full.” prints below the claim indicating the amount to be recouped. This amount appears under the heading, “Financial Transactions Accounts Receivable.” EOB 06065, “Account Receivable is due to the adjusted claim listed. For details, refer to your R&S Report for the date listed within the original date field.”
Claims adjusted as a result of a rate change will be listed on the R&S Report with EOB 01154 “This adjustment is a result of a rate change.”

**Refer to:** Subsection 6.2.5, “Modifier Requirements for TOS Assignment” in this section for a list of the most commonly used modifiers.

### 6.11.4.3 Financial Transactions

All claim refunds, reissues, voids/stops, recoupments, backup withholdings, levies, and payouts appear in this section of the R&S Report. The Financial Transactions section does not use the R&S Report form headings. Additional subheadings are printed to identify the financial transactions. The following descriptions are types of financial items.

#### 6.11.4.3.1 Accounts Receivable

This label identifies money subtracted from the provider’s current payment owed to TMHP. Specific claim data are not given on the R&S Report unless the accounts receivable control number is provided which should be referenced when corresponding with TMHP. Accounts receivable appear on the R&S Report in the following format:

- **Control Number.** A number to reference when corresponding with TMHP.
- **Recoupment Rate.** The percentage of the provider’s payment that is withheld each week unless the provider elects to have a specific amount withheld each week.
- **Maximum Periodic Recoupment Amount.** The amount to be withheld each week. This area is blank if the provider elects to have a percentage withheld each week.
- **Original Date.** The date the financial transaction was processed originally.
- **Original Amount.** The total amount owed TMHP.
- **Prior Date.** The date the last transaction on the accounts receivable occurred.
- **Medical Record Number.** A number assigned by the provider, if available. This area is blank for purged claims.
- **Prior Balance.** The amount owed from a previous R&S Report.
- **Applied Amount.** The amount subtracted from the current R&S Report.
- **Balance.** Indicates the total outstanding accounts receivable (AR) balance that remains due to TMHP.
- **FYE.** The fiscal year end (FYE) for cost reports.
- **EOB.** The EOB code that corresponds to the reason code for the accounts receivable.
- **Patient Name.** The name of the patient on the claim, if the accounts receivable are claim-specific.
- **Claim Number.** The ICN of the original claim, if the accounts receivable are claim-specific.
- **Backup Withholding Penalty Information.** A penalty assessed by the Internal Revenue Service (IRS) for noncompliance due to a B-Notice. Although the current payment amount is lowered by the amount of the backup withholding, the provider’s 1099 earnings are not lowered.
- **Control Number.** TMHP control number to reference when corresponding with TMHP.
- **Original Date.** The date the backup withholding was set up originally.
- **Withheld Amount.** Amount withheld (31 percent) of the provider’s checkwrite.
6.11.4.3.2 IRS Levies
The payments withheld from a provider’s checkwrite as a result of a notice from the IRS of a levy against the provider appear in the “IRS Levy Information” section of the R&S Report. Payments are withheld until the levy is satisfied or released. Although the current payment amount is lowered by the amount of the levy payment, the provider’s 1099 earnings are not lowered. IRS levies are reported in the following format:

- **Control Number.** TMHP control number to reference when corresponding with TMHP.
- **Maximum Recoupment Rate.** The percentage of the provider’s payment that is withheld each week, unless the provider elects to have a specific amount withheld each week.
- **Maximum Recoupment Amount.** The amount to be withheld periodically.
- **Original Date.** The date the levy was set up originally.
- **Original Amount.** The total amount owed to the IRS.
- **Prior Balance.** The amount owed from a previous R&S Report.
- **Prior Date.** The date the last transaction on the levy occurred.
- **Current Amount.** The amount subtracted from the current R&S Report and paid to the IRS.
- **Remaining Balance.** The amount still owed on the levy. (This amount becomes the “previous balance” on the next R&S Report.)

6.11.4.3.3 Refunds
Refunds are identified by EOB 00124, “Thank you for your refund; your 1099 liability has been credited.” This statement is verification that dollars refunded to TMHP for incorrect payments have been received and posted. The provider’s check number and the date of the check are printed on the R&S Report. Claim refunds appear on the R&S Report in the following format:

- **Claim Specific:**
  - **ICN.** The claim number of the claim to which the refund was applied this cycle.
  - **Patient Name.** The first name, middle initial, and last name of the patient on the applicable claim.
  - **Medicaid Number.** The patient’s Medicaid or CSHCN Services Program number.
  - **Date of Service.** The format MMDDCCYY (month, day, and year) in “From” DOS.
  - **Total Billed.** The total amount billed for the claim being refunded.
  - **Amount Applied This Cycle.** The refund amount applied to the claim.
  - **EOB.** Corresponds to the reason code assigned.

- **Nonclaim Specific:**
  - **Control Number.** A control number to reference when corresponding with TMHP.
  - **FYE.** The fiscal year for which this refund is applicable.
  - **EOB.** Corresponds to the reason code assigned.
6.11.4.3.4 Payouts

Payouts are dollars TMHP owes to the provider. TMHP processes two types of payouts: system payouts that increase the weekly check amount and manual payouts that result in a separate check being sent to the provider. Specific claim data are not given on the R&S Report for payouts. A control number is given, which should be referenced when corresponding with TMHP. System and manual payouts appear on the R&S Report in the following format:

- **Payout Control Number.** A control number to reference when corresponding with TMHP.
- **Payout Amount.** The amount of the payout.
- **FYE.** The fiscal year for which the payout is applicable.
- **EOB.** Corresponds to the reason code assigned.
- **Patient Name.** Name of the patient (if available).
- **PCN.** Medicaid number of the patient (if available).
- **DOS.** Date of service (if available).

6.11.4.3.5 Reissues

The provider’s 1099 earnings are not affected by reissues. A messages states, “Your payment has been increased by the amount indicated below”:

- **Check Number.** The number of the original check.
- **Check Amount.** The amount of the original check.
- **R&S Number.** The number of the original R&S Report.
- **R&S Date.** The date of the original R&S Report.

6.11.4.3.6 Voids and Stops

The provider’s 1099 earnings are credited by the amount of the voided/stopped payment.

- **Check Number.** The number of the voided/stopped payment.
- **Check Amount.** The amount of the voided/stopped payment.
- **R&S Number.** The number of the voided/stopped payment.
- **R&S Date.** The date of the voided/stopped payment.

6.11.4.4 Claims Payment Summary

This section summarizes all payments, adjustments, and financial transactions listed on the R&S Report. The section has two categories: one for amounts “Affecting Payment This Cycle” and one for “Amount Affecting 1099 Earnings.”

If the provider is receiving a check on this particular R&S Report, the following information is given: “Payment summary for check XXXXXXXXX in the amount of XXX,XXX,XXX.XX.” If the payment is EFT: “Payment summary for direct deposit by EFT XXXXXXXXX in the amount of XXX,XXX,XXX.XX.” The check number also is printed on the check that accompanies the R&S Report.

**Headings for the Payment Summary for “Affecting Payment This Cycle” and “Amount Affecting 1099 Earnings”**

- **Claims Paid.** Indicates the number of claims processed for the week and the year-to-date total.
- **System Payouts.** The total amount of system payouts made to the provider by TMHP.
- **Manual Payouts (Remitted by separate check or EFT).** The total amount of manual payouts made to the provider by TMHP.
• **Amount Paid to IRS for Levies.** The amount remitted to IRS and withheld from the provider’s payment due to an IRS levy.

• **Amount Paid to IRS for Backup Withholding.** The amount paid to the IRS for backup withholding.

• **Accounts Receivable Recoupments.** The total amount withheld from the provider’s payment due to accounts receivable.

• **Miscellaneous Levies.** The amount withheld from the provider’s payment and remitted to HHSC for a SHARS Admin Fee levy.

• **Amounts Stopped/Voided.** The total amount of the payment that was voided or stopped with no reissuance of payment.

• **System Reissues.** The amount of the reissued payment.

• **Claim Related Refunds.** The total amount of claim-related refunds applied during the weekly cycle.

• **Nonclaim Related Refunds.** The total amount of nonclaim-related refunds applied during the weekly cycle.

• **Approved to Pay/Deny Amount.** The total amount of claim payments that were approved to pay/deny within the week. (This column will not be used at this time.)

• **Pending Claims.** The total amount billed for claims in process as of the cutoff date for the report.

### 6.11.4.5 The Following Claims are Being Processed

In the “Following Claims are Being Processed” section, the R&S Report may list up to five EOPS codes per claim. The claims listed in this section are in process and cannot be appealed for any reason until they appear in either the “Claims Paid or Denied,” or “Adjustments Paid and Denied” sections of the R&S Report. TMHP is listing the pending status of these claims for informational purposes only. The pending messages should not be interpreted as a final claim disposition. Weekly, all claims and appeals on claims TMHP has “in process” from the provider are listed on the R&S Report. The Following Claims are Being Processed claim prints in the same format as a paid or denied claim.

### 6.11.4.6 Explanation of Benefit Codes Messages

This section lists the descriptions of all EOBs that appeared on the R&S Report. EOBs appear in numerical order.

EDI ANSI X12 5010 835 files display the appropriate Claims Adjustment Reason Code (CARC), Claims Adjustment Group Code (CAGC), and Remittance Advice Remarks Code (RARC) explanation codes that are associated with EOB denials.

The 835 file includes the CARC, CAGC, and RARC explanation codes that are associated with the highest priority detail EOB to provide a clearer explanation for the denial.

### 6.11.4.7 Explanation of Pending Status Codes Appendix

This section lists the description of all EOPS codes that appeared on the R&S Report. EOPS appear in numerical order.

EOB and EOPS codes may appear on the same pending claim because some details may have already finalized while others may have questions and are pending.

### 6.11.5 R&S Report Examples

Examples of R&S Reports are available on the TMHP website at [www.tmhp.com](http://www.tmhp.com).

<table>
<thead>
<tr>
<th>R&amp;S Report Examples</th>
</tr>
</thead>
<tbody>
<tr>
<td>Accounts Receivables R&amp;S Report</td>
</tr>
<tr>
<td>(For purposes of example, accounts receivables, void, and stop pay appear together.)</td>
</tr>
</tbody>
</table>
6.11.6 Provider Inquiries—Status of Claims

TMHP provides several effective mechanisms for researching the status of a claim. Weekly, TMHP provides the R&S Report reflecting all claims with a paid, denied, or pending status. Providers verify claim status using the provider’s log of pending claims.

Electronic billers allow ten business days for a claim to appear on their R&S Reports. If the claim does not appear on an R&S Report as paid, pending, or denied, a transmission failure, file rejection, or claims rejection may exist. Providers check records for transmission reports correspondence from the TMHP EDI Help Desk.

The provider allows at least 30 days for a Medicaid paper claim to appear on an R&S Report after the claim has been submitted to TMHP. If a claim has not been received by TMHP and must be submitted a second time, the second claim must also meet the 95-day filing deadline.

The provider allows TMHP 45 days to receive a Medicare-paid claim automatically transmitted for payment of coinsurance or deductible according to current payment guidelines. Claims that fail to cross over from Medicare may be filed to TMHP by submitting a paper MRAN received from Medicare or a Medicare intermediary, the computer generated MRANs from the CMS-approved software applications MREP for professional services or PC-Print for institutional services or, for MAP clients, TMHP Standardized Medicare Advantage Plan (MAP) Remittance Advice Notice Template with the completed claim form.

If the claim does not appear on an R&S Report as paid, pending, or denied, providers can use any of the following procedures to inquire about the status of the claim:

- The provider can use the claim status inquiry function of TexMedConnect on the TMHP website at www.tmhp.com.
- The provider can call AIS at 1-800-925-9126 to determine if the claim is pending, paid, denied, or if TMHP has no record of the claim.
- If any of the three options above indicates that TMHP has no record of the claim, the provider can call the TMHP Contact Center at 1-800-925-9126 and speak to a TMHP contact center representative.
• If the TMHP Contact Center has no record of a claim that was submitted within the original filing deadline, the provider can submit a copy of the original claim to TMHP for processing. Electronic billers may refile the claim electronically. For claims submitted by a hospital for inpatient services, the filing deadline is 95 days from the discharge date or the last DOS on the claim. For all other types of providers, the filing deadline is 95 days from each DOS on the claim.

• If the 95-day filing deadline has passed and the claim is still within 120 days of the date of the rejection report or the R&S Report, the provider can submit a signed copy of the claim and all of the documentation that supports the original claim submission, including any electronic rejection reports, to:

Texas Medicaid & Healthcare Partnership  
Inquiry Control Unit  
12357-A Riata Trace Parkway, Suite 100  
Austin, TX 78727

Providers must retain copies of all R&S Reports for a minimum of five years. Providers must not send original R&S Reports back with appeals. Providers must submit one copy of the R&S Report to TMHP per appeal.


6.12 Filing Medicare Primary Claims

When a service is a benefit of both Medicare and Medicaid, the claim must be filed to Medicare first. Providers should not file a claim with Medicaid until Medicare has dispositioned the claim unless the service is a Medicaid-only service.

All Medicare providers and suppliers who offer services and supplies to Qualified Medicaid Beneficiaries (QMB) or Medicaid Qualified Medicare Beneficiaries (MQMB) must not bill dual eligible clients for Medicare cost-sharing. This includes deductible, coinsurance, and copayments for any Medicaid covered items and services.

Medicaid claims for Qualified Medicare Beneficiary (QMB) and Medicaid Qualified Medicare Beneficiary (MQMB) clients can be filed to Medicaid for consideration of coinsurance and deductible payment as follows:

- Medicare primary claims filed to Medicare Administrative Contractors (MACs) may be transferred electronically to TMHP through a Benefit Coordination and Recovery Center (BCRC).
- Providers can submit crossover claims directly to TMHP using a paper claim form only for the specific circumstances indicated in the following section.

Note: These guidelines do not apply to services that are rendered to clients who are living in a nursing facility.

Refer to: Subsection 2.7, “Medicare Crossover Claim Reimbursement” in “Section 2: Texas Medicaid Fee-for-Service Reimbursement” (Vol. 1, General Information) for information about reimbursement for QMBs and MQMBs.

Subsection 4.9, “Medicare and Medicaid Dual Eligibility” in “Section 4: Client Eligibility” (Vol. 1, General Information) for information about MQMBs and QMBs eligibility.

6.12.1 Electronic Crossover Claims

Medicare primary claims filed to MACs may be transferred electronically to TMHP through a BCRC for claims that are processed as assigned. Providers should contact their MAC for more information.
This electronic crossover process allows providers to receive disposition from both carriers while only filing the claim once. Providers must allow 60 days from the date of Medicare’s disposition for a claim to appear on the Medicaid R&S Report.

If all services on the claim are denied by Medicare, the claim is not automatically transferred to TMHP by the MAC through the BCRC. Providers must submit the denied crossover claims to TMHP on paper.

Claims that are submitted to Medicare must include the facility’s NPI. Medicare crossover claims must comply with the Medicaid requirement to include a facility NPI. If a Medicare crossover claim includes a service for which Medicaid requires a facility NPI but the claim does not include the facility’s NPI number, the claim will be denied by Texas Medicaid.

Important: TMHP accepts only electronic crossover claims that are automatically transferred to TMHP by the MAC through the BCRC. TMHP accepts only paper crossover claims from providers and other entities. TMHP does not accept electronic crossover new day claims or appeals from providers and other entities. TMHP accepts only paper appeals.

6.12.1.1 Type of Bills Values for Medicare Crossover Claims

Type of bills (TOB) values in the 12x series may be billed to Medicare for Medicare Inpatient Part B services as appropriate, but TOB values in the 12x series are not valid for Medicaid claims.

Reminder: Texas Medicaid only allows interim billing and late changes to be submitted on inpatient claims.

6.12.1.2 Medicare Copayments

Claims for Medicare copayments can also be submitted to TMHP. TMHP processes and pays Medicare HMO and Medicare PPO copayments for dual-eligible clients according to Medicaid guidelines.

The following procedure codes may be reimbursed for Medicare copayments:

The following Medicaid codes have been created for copayments, which are considered an atypical service:

<table>
<thead>
<tr>
<th>Procedure Code</th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td>CP003</td>
<td>Medicare HMO copayment-professional</td>
</tr>
<tr>
<td>CP004</td>
<td>Medicare PPO copayment-professional</td>
</tr>
<tr>
<td>CP007</td>
<td>Medicare HMO copayment-outpatient</td>
</tr>
<tr>
<td>CP008</td>
<td>Medicare PPO copayment-outpatient</td>
</tr>
</tbody>
</table>

TMHP may reimburse the copayment in addition to a service the HMO or PPO has denied if the client is eligible for Texas Medicaid and the procedure is reimbursed under Medicaid guidelines. Providers are not allowed to hold the client liable for the copayment.

An office or emergency room (ER) visit (the ER physician is paid only when the ER is not staffed by the hospital) is reimbursed a maximum copayment of $10 per visit. The hospital ER visit is reimbursed at a maximum of $50 to the facility. TMHP pays up to four copayments per day, per client. ER visits are limited to one per day, per client, and are considered one of the four copayments allowed per day.

Refer to: Subsection 2.7.4.2, “Nephrology (Hemodialysis, Renal Dialysis) and Renal Dialysis Facility Providers” in “Section 2: Texas Medicaid Fee-for-Service Reimbursement” (Vol. 1, General Information) for information about claims for nephrology (hemodialysis, renal dialysis) and renal dialysis facility providers for Medicare crossover Claims.

Subsection 2.7.4, “Exceptions” in “Section 2: Texas Medicaid Fee-for-Service Reimbursement” (Vol. 1, General Information) for information about exceptions for Medicare Part A, Part B, and Part C (noncontracted MAPs) reimbursement.
6.12.1.3 Requirement for Group Billing Providers – Professional Claims

The performing provider NPI must be included on the professional electronic claim if the billing provider is a group. Claims are processed using the performing provider NPI that is submitted on the Medicare claim.

**Important:** The performing provider who is identified on the claim must be a member of the billing provider’s group. If the performing provider is not a member of the billing provider group, the detail line item will be denied.

A claim is denied if the performing provider NPI is missing, invalid, or is not a member of the billing provider’s group. Denied claims may be appealed on paper with the appropriate performing provider information.

6.12.2 Paper Crossovers Claims

TMHP accepts only paper crossover claims or appeals from providers and other entities.

The following paper crossover claims may be submitted to TMHP:

- For QMB and MQMB clients, if a crossover claim is not transferred to TMHP electronically through the BCRC, the provider can submit a paper claim to TMHP for coinsurance and deductible reimbursement consideration.

- For MQMB clients, if a claim is denied by Medicare because the services are not a benefit of Medicare or because Medicare benefits have been exhausted, the provider can submit a paper claim to TMHP for coinsurance and deductible reimbursement consideration, and reimbursement consideration for the Medicaid-only services that were denied by Medicare. The Medicare EOB that contains the relevant claim denial must be submitted to TMHP with the completed claim from within 95 days from the Medicare disposition date and 365 days from the date of service. The denied services are processed as Medicaid-only services.

Claims that are submitted to Medicare must include the facility’s NPI. Medicare crossover claims must comply with the Medicaid requirement to include a facility NPI. If a Medicare crossover claim includes a service for which Medicaid requires a facility NPI but the claim does not include the facility’s NPI number, the claim will be denied by Texas Medicaid.

**Important:** Claims that are denied by Medicare for administrative reasons must be appealed to Medicare before they are submitted to Texas Medicaid.

The paper submission must include all of the following:

- The Medicare Remittance Advice (RA) or Remittance Notice (RN), using the CMS-approved software MREP, for professional services, or PC-Print or a paper MRAN from Medicare.

- The appropriate, completed paper CMS-1500 or UB-04 CMS-1450 paper claim form.

  **Note:** Although it is not required, it is strongly recommended that providers send claim forms with their Medicare appeals in case one is needed for further processing.

- The appropriate TMHP Standardized Medicare Advantage Plan (MAP) Remittance Advice Notice Template for Medicare Advantage Plan only. The template must be submitted with the claims form and the MAP EOB.

Providers that receive Remittance Advice Notices from a Medicare intermediary may submit these in place of the MRAN to TMHP which must contain the following required information:

- Client name
- Medicare number
- NPI
• Dates of service
• Procedure code (Professional and Outpatient claims)
• Billed amount
• Medicare allowed amount or non-covered amount
• Deductible amount
• Co-insurance amount
• Medicare paid amount
• Medicare ICN
• Quantity billed

6.12.2.1 Deductible or Coinsurance Amount Balancing

The Texas Medicaid claims processing system validates that the total Medicare deductible and coinsurance amounts on the claim header match the sum of the detail Medicare deductible and coinsurance amounts.

For paper crossover claims, providers must submit the same information to Texas Medicaid that was received from Medicare.

Texas Medicaid will reimburse Medicare crossover claims up to the Texas Medicaid allowed amount for Medicaid-covered services. System enhancements have been identified to ensure appropriate age restrictions are enforced applicable to the services rendered.

Example: For a Medicare service provided to an adult client, if that service is only payable to Medicaid for clients who are 20 years of age and younger, the age restriction will be applied and the Medicaid allowed amount will be zero. Since the Medicare payment exceeds the Medicaid allowed amount or encounter payment for the service, Texas Medicaid will not make a payment for coinsurance liabilities.

Because Medicare reimbursed more than Medicaid allowed, the client has no liability for any balance or Medicare coinsurance related to the rendered services.

6.12.2.2 TMHP Standardized Medicare Advantage Plan (MAP) Remittance Advice Notice Template

The TMHP Standardized Medicare Advantage Plan (MAP) Remittance Advice Notice Template must be submitted for paper MAP claims only. The template must be submitted with the claim form and the MAP EOB.

Note: Providers must not submit the template for traditional Medicare crossover claims.

The following guidelines apply for the submission of the TMHP Standardized Medicare Advantage Plan (MAP) Remittance Advice Notice Templates:

• The Medicare ICN must be included on the form. Claims are denied if the Medicare ICN is omitted.

• For the TMHP Crossover Professional Claim Type 30 form, the performing provider NPI and TPI must be submitted on each detail line item. A detail line item is denied if the performing provider NPI or TPI is omitted, if the performing provider NPI is not associated with the TPI according to the performing provider’s enrollment information, or if the performing provider is not a member of the group billing provider.

• For the TMHP Crossover Outpatient Facility Claim Type 31 form, the detail line items are required. Claims are denied if the details are omitted.
The TMHP Standardized Medicare Advantage Plan (MAP) Remittance Advice Notice Template must be submitted with a completed claim form and MAP EOB, must be legible, and must identify only one client per page. Providers must not submit handwritten MAP templates.

Claims that do not meet these standards are not processed and are returned to the provider.

By submitting the TMHP Standardized Medicare Advantage Plan (MAP) Remittance Advice Notice Templates to TMHP, the provider attests that the information included in the template matches the EOB that was received from the MAP. If the information on the template does not exactly match the information on the RA or RN, the claim may be denied.

Refer to: subsection 6.20, “Forms” in this section for the TMHP Standardized Medicare Advantage Plan (MAP) Remittance Advice Notice Templates and instructions.

Subsection 2.7, “Medicare Crossover Claim Reimbursement” in “Section 2: Texas Medicaid Fee-for-Service Reimbursement” (Vol. 1, General Information).

6.12.2.3 Crossover Paper Claims Filing Deadlines

The paper crossover claim with all required, EOBs, templates, and forms must be received by TMHP within 95 days of the Medicare date of disposition and 365 days from the date of service in order to be considered for processing.

6.12.3 Filing Medicare-Adjusted Claims

TMHP accepts crossover appeals only on paper.

Providers may submit Medicare-adjusted claims by submitting the adjusted Medicare RA/RNs (paper or electronic) and the appropriate TMHP Standardized Medicare Advantage Plan (MAP) Remittance Advice Notice Template. The information on the Medicare RA/RN must exactly match the information submitted on the TMHP Standardized Medicare Advantage Plan (MAP) Remittance Advice Notice Template.

Refer to: Subsection 3.7.1, “Medicaid Relationship to Medicare” in the Inpatient and Outpatient Hospital Services Handbook (Vol. 2, Provider Handbooks) for additional information on hospital Medicare claims filing requirements.

Important: TMHP does not accept electronic crossover appeals.

6.13 Medically Needy Claims Filing

TMHP must receive claims for unpaid bills not applied toward spend down within 95 days from the date eligibility was added to the TMHP client eligibility file (add date). These bills must be on the appropriate claim form (for example, CMS-1500 or UB-04 CMS-1450). Providers are allowed to submit completed CMS claim forms directly to the Medically Needy Clearinghouse (MNC) or to applicants for the Medically Needy Program (MNP) to be used to meet spend down. The completed CMS claim forms used to meet spend down are held for ten calendar days by the MNC, then forwarded to TMHP claims processing. Claims for services provided after the spend down is met must be received within 95 days from the date eligibility is added. Inpatient hospital facility claims must be received within 95 days from the date of discharge or last DOS on the claim. This applies when eligibility is not retroactive.

The client’s payment responsibilities are as follows:

- If the entire bill was used to meet spend down, the client is responsible for payment of the entire bill.
- If a portion of one of the bills was used to meet the spend down, the client is responsible for paying the portion applied toward the spend down, unless it exceeds the Medicaid allowable amount.
- The claim must show the total billed amount for the services provided. Charges for ineligible days or spend down amounts should not be deducted or noncovered on the claim.
• A client’s payment toward spend down is not reflected on the claim submitted to TMHP.
• A client is not required to pay the spend down amount before a claim is filed to Medicaid.
• Payments made by the client for services not used in the spend down but were incurred during an eligible period must be reimbursed to the client before the provider files a claim to TMHP.
• Services that require prior authorization and are provided before the client becomes eligible for Medicaid by meeting spend down are not reimbursable by Texas Medicaid.
• If a bill or a completed CMS claim form was not used to meet spend down and the dates of service are within the client’s eligible period, submit the total bill to TMHP.

When eligibility has been established, a TP 55 with spend down client can receive the same care and services available to all other Medicaid clients. If eligibility is established through TP 30 with spend down, the client’s Medicaid eligibility is restricted to coverage for an emergency medical condition only. Emergency medical condition is defined under subsection 4.3.2.2, “Exceptions to Lock-in Status” in “Section 4: Client Eligibility” (Vol. 1, General Information).

6.14 Claims Filing for Consumer-Directed Services (CDS)
Clients who participate in the CDS option for both PCS and a waiver program, through HHSC are required to choose one Financial Management Services Agency (FMSA) to provide services through both programs. FMSAs are permitted to file only the financial management services (FMS) fee, also known as the monthly administrative fee, through one program. The FMSA should file the FMS claim through the program with the highest reimbursement rate. Currently, the waiver programs have a higher reimbursement rate for the FMS fee than the Texas Medicaid PCS benefit, so a FMSA should file claims for the monthly FMS fee through the waiver programs.

The U8 modifier, which is used when submitting claims for the monthly PCS administrative fee, must be prior authorized. The DSHS case managers have two options when sending a prior authorization request for PCS to TMHP:
• If a client is only using the CDS option for Texas Medicaid PCS, a case manager will submit a prior authorization request to TMHP that approves the U8 modifier and either the U7 or UB modifier. In this case, the provider authorization notification letter will include the U8 modifier and the U7 or UB modifier.
• If a client is using the CDS option for both Texas Medicaid PCS and a waiver program, a case manager will submit a prior authorization request to TMHP that approves either the U7 or UB modifier. The U8 modifier will not be prior authorized in this situation.

When a provider authorization notification letter is received by a FMSA, the provider should verify that the correct modifiers have been prior authorized for each PCS client. Providers who think that the approved modifiers are incorrect should contact the DSHS case manager and ask for the correct modifiers to be submitted to TMHP for prior authorization.

6.15 Claims Filing for Home Health Agency Services
Providers must use only type of bill (TOB) 321 in Form Locator (FL) 4 of the UB-04 CMS-1450. Other TOBs are invalid and will result in a claim denial. Home Health Services must be submitted to TMHP in an approved electronic format or on a CMS-1500 or a UB-04 CMS-1450 paper claim form. Submit home health DME and medical supplies to TMHP in an approved electronic format, or on a CMS-1500 or on a UB-04 CMS-1450 paper claim form. Providers may purchase CMS-1500 or UB-04 CMS-1450 paper claim forms from the vendor of their choice. TMHP does not supply them.

When completing a CMS-1500 or a UB-04 CMS-1450 paper claim form, all required information must be included on the claim, as TMHP does not key information from attachments.
Refer to: “Section 3: TMHP Electronic Data Interchange (EDI)” (Vol. 1, General Information) for information about electronic claims submissions.

Subsection 6.5, “CMS-1500 Paper Claim Filing Instructions” in this section for instructions on how to complete paper claims. Blocks that are not referenced are not required for processing by TMHP and may be left blank.

Subsection 6.6, “UB-04 CMS-1450 Paper Claim Filing Instructions” in this section.

Outpatient claims must have the appropriate revenue code and, if appropriate, the corresponding Healthcare Common Procedure Coding System (HCPCS) code or narrative description. The prior authorization number must appear on the CMS-1500 paper claim form in Block 23 and in Block 63 of the UB-04 CMS-1450 paper claim form. The certification dates or the revised request date on the POC must coincide with the DOS on the claim. Prior authorization does not waive the 95-day filing deadline requirement.

6.16 Claims for Medicaid Hospice Clients Not Related to the Terminal Illness

When the services are unrelated to the terminal illness, providers must submit a claim for Medicaid services to TMHP. The claim must include a statement and documentation from the hospice that the services billed are not related to the client’s terminal illness.

If TMHP denies the claim, the following information must be submitted with the providers appeal.

- A copy of the R&S Report, with the client or claim number in question circled
- Clinical records, which may be obtained from the hospice provider
- Supporting documentation giving reasons the services billed are not related to the terminal illness

Refer to: Subsection 4.3.3, “Hospice Program” in “Section 4: Client Eligibility” (Vol. 1, General Information) for more information related to Medicaid hospice client benefits and eligibility.

6.16.1 Medical Services When Client is Discharged From Hospice

Submit claims to TMHP for Medicaid services with a statement that the services billed were provided after the client was discharged from the Hospice Program. The provider must obtain a copy of Form 3071, Medicaid Hospice Cancellation, from the Hospice Program to support the discharge.

If TMHP denies the claim, the provider may appeal the decision with the following information:

- A copy of the R&S Report, with the client or claim number in question circled
- Supporting documentation stating that the client was not in hospice at the time

6.16.2 Claims Address for Medicaid Hospice Clients Not Related to the Terminal Illness

Mail paper claims to the following address:

Texas Medicaid & Healthcare Partnership
PO Box 200105
Austin, TX 78720-0105

Appeal claims by writing to the following address:

Texas Medicaid & Healthcare Partnership
PO Box 200645
Austin, TX 78720-0645
6.16.3  Lab and X-Ray
Submit claims for services unrelated to the terminal illness to TMHP. Submit claims for services related to the terminal illness to the hospice provider.

6.17  Claims for Texas Medicaid and CSHCN Services Program Eligible Clients
The CSHCN Services Program is the payer of last resort when clients have other insurance, including Texas Medicaid and private carriers. The CSHCN Services Program does not supplement a client’s Texas Medicaid benefits; however, services that are not a benefit of Texas Medicaid, such as hospice and medical foods, may be covered by the CSHCN Services Program.

6.17.1  New Claim Submissions
New claims that are submitted for clients who are eligible for both Texas Medicaid and CSHCN Services Program benefits during the same eligibility period will be processed through the appropriate program and may result in a separate claim for each program. The Medicaid claim number and disposition will be listed under the “Claims – Paid or Denied” section of the Medicaid/Managed Care R&S Report. If the claim includes services that are not benefits of Texas Medicaid but are benefits of the CSHCN Services Program, a claim will be created with a unique claim number that will be listed under the “Claims – Paid or Denied” section of the CSHCN Services Program R&S Report.

Note:  If all of the services that are submitted on the claim are Texas Medicaid benefits, a CSHCN Services Program claim will not be created. Only a Texas Medicaid claim will be created, and the claim number will appear on the provider’s Medicaid/Managed Care R&S Report.

6.17.2  CSHCN Services Program Claims Reprocessing for Retroactive Texas Medicaid Eligibility
Claims that have already been paid by the CSHCN Services Program for clients who received retroactive Texas Medicaid eligibility for dates of service covered on the paid claims will be reprocessed to pay under the appropriate program. The reprocessed CSHCN Services Program claim number will appear under the “Adjustments – Paid or Denied” section of the CSHCN Services Program R&S Report. An accounts receivable will be created for services covered by Texas Medicaid that will be reflected on the “Financial Transactions” page under the “Accounts Receivable” section of the CSHCN Services Program R&S Report. The claim will be reprocessed to Texas Medicaid and given a new claim number. The new Texas Medicaid claim number and disposition will appear under the “Claims – Paid or Denied” section of the Medicaid/Managed Care R&S Report.

TMHP will contact providers when it reprocesses claims for services that require a Texas Medicaid prior authorization. Providers will be informed that a Texas Medicaid prior authorization must be submitted within a specified time frame for the claim to be considered for processing through Texas Medicaid.

6.18  Claims for State Supported Living Center Residents (SSLC)
Medicaid providers who render off-campus acute care services to Medicaid-eligible State Supported Living Center (SSLC) residents must submit claims directly to Medicaid. This is applicable only to residents of the SSLCs operated by HHSC.

Claims and prior authorization requests for acute care services rendered to these individuals must be submitted to Medicaid. These requests must be submitted according to guidelines for acute care services as indicated in this manual.

Refer to: “Section 5: Fee-for-Service Prior Authorizations” (Vol. 1, General Information) for more information on prior authorizations.
6.19 Children’s Health Insurance Program (CHIP) Perinatal Claims

Claims for services provided to CHIP Perinatal Program clients are submitted to and considered for reimbursement as follows:

- For women with income at or below 198 percent FPL:
  - Hospital facility charges are paid through Emergency Medicaid and processed by TMHP.
  - Professional service charges are paid through the CHIP Perinatal Program and processed through CHIP.

  **Note:** Delivery-related professional services claims denied by the CHIP Perinatal health plan will be considered for reimbursement through Emergency Medicaid and will require the CHIP Perinatal health plan denial notice. These claims should be submitted through the existing Medicaid appeals process within 95 days from the date of the CHIP Perinatal Health plan denial notice. The provider must provide a copy of the complete explanation of benefits that includes the complete description of the reason for denial.

- For newborns with a family income at or below 198 percent FPL:
  - Hospital facility charges are paid through Medicaid and processed by TMHP.
  - Professional service charges are paid through Medicaid and processed by TMHP.

Inpatient services (limited to labor with delivery) for unborn children and women with income at or below 202 of FPL will be covered under CHIP Perinatal, and these claims will be paid by the CHIP Perinatal health plan.

6.19.1 CHIP Perinatal Newborn Transfer Hospital Claims

TMHP processes CHIP Perinatal newborn transfer hospital claims even if the claim from the initial hospital stay has not been received.

The hospital transfer must have occurred within 24 hours of the discharge date from the initial delivery hospital stay. This change applies only to CHIP Perinatal newborns with a family income at or below 198 percent of the FPL.

Transfer claims must be filed with TMHP on an electronic institutional claim or the UB-04 CMS-1450 paper claim form using admission type 1, 2, 3, or 5 in block 14, source of admission code 4 or 6 in block 15, and the actual date and time the client was admitted in block 12 of the UB-04 CMS-1450 paper claim form.

6.20 Forms

The following linked forms can also be found on the [Forms](www.tmhp.com) page of the Provider section of the TMHP website at www.tmhp.com:

<table>
<thead>
<tr>
<th>Forms</th>
</tr>
</thead>
<tbody>
<tr>
<td>MRAN Form Crossover Claim Type 30 Form and Instructions</td>
</tr>
<tr>
<td>MRAN Form Crossover Claim Type 31 Form and Instructions</td>
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<tr>
<td>MRAN Form Crossover Claim Type 50 Form and Instructions</td>
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<td>Sample Letter XUB Computer Billing Service Inc</td>
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7.1 Appeal Methods

An appeal is a request for reconsideration of a previously dispositioned claim.

Providers may use three methods to appeal Medicaid fee-for-service and carve-out service claims to Texas Medicaid & Healthcare Partnership (TMHP): electronic, Automated Inquiry System (AIS), or paper.

TMHP must receive all appeals of denied claims and requests for adjustments on paid claims within 120 days from the date of disposition of the Remittance and Status (R&S) Report on which that claim appears. If the 120-day appeal deadline falls on a weekend or holiday, the deadline is extended to the next business day.

Standard administrative requests and medical appeals must be sent first to TMHP or the claims processing entity as a first-level appeal. After the provider has exhausted all aspects of the appeals process for the entire claim, the provider may submit a second-level appeal to HHSC.

1) A first-level appeal is a provider’s initial standard administrative or medical appeal of a claim that has been denied or adjusted by TMHP. This appeal is submitted by the provider directly to TMHP for adjudication and must contain all required information to be considered.

2) A second-level appeal is a provider’s final medical or standard administrative appeal to HHSC of a claim that meets all of the following requirements:
   - It has been denied or adjusted by TMHP.
   - It has been appealed as a first-level appeal to TMHP.
   - It has been denied again for the same reason(s) by TMHP.

This appeal is submitted by the provider to HHSC, which may subsequently require TMHP to gather information related to the original claim and the first-level appeal. HHSC is the sole adjudicator of this final appeal.

All providers must submit second-level administrative appeals and exceptions to the 95-day filing deadline appeals to the following address:

Texas Health and Human Services Commission
HHSC Claims Administrator Operations Management
Mail Code 91X
PO BOX 204077
Austin, Texas 78720-4077

TMHP is not responsible for managing appeals resulting from utilization review (UR) decisions by the HHSC Office of Inspector General (OIG) UR Unit. These must be submitted to HHSC Medical and UR Appeals.

**Note:** Appeals for managed care claims must be submitted to the managed care organization (MCO) or dental plan that administers the client’s managed care benefits. The only managed care appeals administered by TMHP are those for carve-out services.

**Refereto:** Subsection 7.3.3, “Utilization Review Appeals” in this section.

The *Medicaid Managed Care Handbook* (Vol. 2, Provider Handbooks) for additional information about managed care appeals.

7.1.1 Electronic Appeal Submission

Electronic appeal submission is a method of submitting appeals using a personal computer. The electronic appeals feature can be accessed by a business organization (e.g., billing agents) interfacing directly with the TMHP Electronic Data Interchange (EDI) Gateway or through TexMedConnect, the free web-based application available from TMHP.
The Health Insurance Portability and Accountability Act (HIPAA) standard American National Standards Institute (ANSI) ASC X12 837 format is accepted by TMHP EDI.

For other information, contact the TMHP EDI Help Desk at 1-888-863-3638.

### 7.1.1.1 Advantages of Electronic Appeal Submission

Using electronic appeal submission provides the following advantages to the users:

- Increased accuracy of appeals filed to potentially improve cash flow.
- Maintained audit trails through print and download capabilities.
- Appeal submission windows can be automatically filled in with electronic R&S (ER&S) Report information, thereby reducing data entry time.

### 7.1.1.2 Disallowed Electronic Appeals

The following claims may not be appealed electronically:

- Claims that require supporting documentation (e.g., operative report, medical records, home health, hearing aid, and dental X-rays).
- Diagnosis-related group (DRG) assignment.
- Medicare crossovers.
- Claims listed as pending or in process with explanation of pending status (EOPS) messages.
- Claims denied as past filing deadline except when retroactive eligibility deadlines apply.
- Claims denied as past the payment deadline.
- Claims with quantity billed changes in the claims details.
- Claims that are the result of a mass adjustment.

**Exception:** Inpatient hospital claims denied for lack of a Hysterectomy Acknowledgment Statement or a Sterilization Consent Form may be appealed electronically if the requested form has been faxed according to the instructions in the Texas Medicaid - Title XIX Acknowledgment of Hysterectomy Information on the TMHP website at www.tmhp.com.

### 7.1.2 Resubmission of TMHP Electronic Data Interchange (EDI) Rejections

TMHP EDI transactions that fail HIPAA edits will be rejected, and the submitter will receive a 277CA claim response file. The 277CA claims response file lists activity by submitter, provider, and payer.

The 277CA claims response file includes member identifier, patient last name and first initial, patient control number (PCN), type of bill or place of service, charge, transaction from and to dates, receipt date, rejection code, and rejection description.

Providers must send the batch ID, PCN, date of service, transaction from and to dates, receipt date, and rejection codes from the 277CA claims response file to TMHP when appealing denied claims.

The batch ID is located in the file name of the returned 277CA claims response, and not within the file. Providers must include the batch ID in all electronic response files submitted to TMHP for appeals to denied claims. Handwritten batch IDs are not acceptable for submission to TMHP. Providers who cannot identify or retrieve the batch ID from the 277CA claims response file name should contact the clearinghouse or vendor to have the filename included in the response document. If not, the provider must request a copy of the response file that contains the filename from the clearinghouse.

Providers who receive a rejection on the 277CA claims response file may resubmit an electronic claim within 95 days of the date of service.
A paper appeal may also be submitted with a copy of the response document within 120 days of the 277CA claims response file rejection to meet the filing deadline. A copy of the electronic response file rejection to include the batch ID must accompany each corrected claim that is submitted on paper.

### 7.1.3 Automated Inquiry System (AIS) Appeals

The following appeals may be submitted using AIS:

- **Client eligibility.** The client’s correct Medicaid number, name, and date of birth are required.
- **Provider information (excluding Medicare crossovers).** The correct provider identifier is required for the billing provider, performing provider, referring provider, and limited provider. The name and address of the provider are required for the facility and outside laboratory.
- **Claim corrections.** Providers may correct the:
  - Patient control number (PCN).
  - Date of birth.
  - Date of onset.
  - X-ray date.
  - Place of service (POS).
  - Quantity billed.
  - Prior authorization number (PAN).
  - Beginning date of service (DOS), as long as the new date is within the filing deadline for the claim.
  - Ending date of service, as long as the new date is within the filing deadline for the claim.

The following appeals may not be appealed through AIS:

- Claims listed on the R&S Report as Incomplete Claims
- Claims listed on the R&S Report with $0 allowed and $0 paid
- Claims requiring supporting documentation (for example, operative report, medical records, home health, hearing aid, and dental X-rays)
- DRG assignment
- Procedure code, modifier, or diagnosis code
- Medicare crossovers
- Claims listed as *pending* or *in process* with EOPS messages
- Claims denied as *past filing deadline* except when retroactive eligibility deadlines apply
- Claims denied as *past the payment deadline*
- Inpatient hospital claims that require supporting documentation
- Third party liability (TPL)/other insurance

Providers may appeal these denials either electronically or on paper.
Refer to: Subsection 7.1.1.2, “Disallowed Electronic Appeals” in this section to determine whether these appeals can be billed electronically. If these appeals cannot be billed electronically, a paper claim must be submitted.

**Exception:** Inpatient hospital claims denied for lack of a Hysterectomy Acknowledgment Statement or a Sterilization Consent Form may be appealed if the requested form has been faxed according to the instructions under subsection 6.13, “Hysterectomy Services” in the Gynecological, Obstetrics, and Family Planning Title XIX Services Handbook (Vol. 2, Provider Handbooks).

### 7.1.4 Automated Inquiry System Automated Appeals Guide

To access the AIS automated appeals guide, providers can call 1-800-925-9126. Providers may submit up to three fields per claim and 15 appeals per call. If during any step invalid information is entered three times, the call transfers to the TMHP Contact Center for assistance.

### 7.1.5 Paper Appeals

Claim appeal requests that cannot be appealed electronically or by using AIS may be appealed on paper. Completed claim forms are not required to be submitted with paper appeals. Providers who submit paper appeals must clearly document on the attached R&S Report the information that is being appealed and identify the claim being appealed.

If a provider determines that a claim cannot be appealed electronically or through AIS, the claim may be appealed on paper by completing the following:

1) Submit a copy of the R&S Report page on which the claim is paid or denied. A copy of other official notification from TMHP may also be submitted.
2) Submit one copy of the R&S Report for each claim appealed.
3) Circle only one claim per R&S Report page.
4) Identify the reason for the appeal.
5) If applicable, indicate the incorrect information and provide the corrected information that should be used to appeal the claim.
6) Attach a copy of any supporting medical documentation that is required or has been requested by TMHP. Supporting documentation must be on a separate page and not copied on the opposite side of the R&S Report.

**Note:** It is strongly recommended that providers submitting paper appeals retain a copy of the documentation being sent. It also is recommended that paper documentation be sent by certified mail with a return receipt requested. This documentation, along with a detailed listing of the claims enclosed, provides proof that the claims were received by TMHP, which is particularly important if it is necessary to prove that the 120-day appeals deadline has been met. If a certified receipt is provided as proof, the certified receipt number must be indicated on the detailed listing along with the Medicaid number, billed amount, date of service (DOS), and a signed claim copy. The provider may need to keep such proof regarding multiple claims submissions if the provider identifier is pending.

Medicare crossovers and inpatient hospital appeals related to medical necessity denials or DRG assignment/adjustment must be submitted on paper with the appropriate documentation.
Submit correspondence, adjustments, and appeals (including routine inpatient hospital claims) to the following address:

Texas Medicaid & Healthcare Partnership
Appeals/Adjustments
PO Box 200645
Austin, TX 78720-0645

Exception: Hospitals appealing HHSC OIG UR Unit final technical denials, admission denials, DRG revisions, continued-stay denials for Tax Equity and Fiscal Responsibility Act (of 1982) (TEFRA) Hospitals, or cost/day outliers must appeal to HHSC at the following address:

Texas Health and Human Services Commission
Medical and UR Appeals, H-230
PO Box 85200
Austin, TX 78708-5200

All other provider fields on the claim forms (referring, facility, admitting, operating, and other) require only an NPI.

Providers that choose to appeal the claim with NPI information must continue submitting both a TPI and an NPI until the claim is finalized.

7.1.5.1 Texas Medicaid Fee-for-Service DRG Adjustment Appeal
Texas Medicaid fee-for-service hospital providers who are appealing a DRG adjustment (higher weight DRG) must provide the original and revised UB-04 CMS-1450 paper claim form, the complete medical record, and a statement defining the reason for the requested change. Hospitals have 120 days from the date of the R&S Report to request an addition of a diagnosis or procedure resulting in a DRG adjustment. Providers appealing a DRG that has not been revised by the OIG Utilization Review Unit should appeal to TMHP.

Refer to: Subsection 7.3.3, “Utilization Review Appeals” in this section.

7.1.5.2 Medical Necessity Denial Appeals
Appeals of denials relating to medical necessity decisions made for all medical services with the exception of HHSC Inpatient UR cases may be submitted for further review if providers find denials are inappropriate. All necessary documentation must accompany the request for review. Incomplete appeals and adjustment requests are denied by TMHP with an explanation of benefits (EOB) code requesting additional information.

TMHP reviews each appeal (DRG adjustment and medical necessity) and forwards written notice of final action in the form of a letter or an adjustment transaction on the R&S Report.

7.1.5.3 Other Insurance Appeals
To appeal a claim denial due to other insurance coverage, the provider must submit complete other insurance information including the disposition date. The disposition date indicates when the other insurance company processed the payment or denial. An appeal submitted without this information will be denied.

If submitting a paper appeal the provider must submit EOBs containing disposition dates. If the disposition date appears only on the first page of an EOB that has multiple pages and the claim that is being appealed is on a subsequent page, the provider must also include the first page of the EOB that shows the disposition date.

7.1.6 Appeals Submitted Incorrectly
If an incomplete appeal is received, it is returned to the sender with further appeal instructions and a request for more information. Documentation (either by letter or facsimile) that does not clearly indicate the reason for submission is returned to the sender for clarification.
If an appeal is received that may be more appropriately addressed in another department, the appeal is forwarded to the appropriate department for research and response.

If the TMHP Medical Director or designee identifies a pattern of ineffective use of the appeals process, the provider may be referred to a provider relations representative for assistance.

### 7.2 Refunds to TMHP

The TMHP Cash Reimbursement Unit is responsible for processing financial adjustments when any of the following occur: overpayment, duplicate payment, payment to incorrect providers, and overlapping payments by Medicaid and a third party resource (TPR).

Providers have the option of refunding payments by issuing a check to TMHP or requesting a recoupment through the paper appeal process. The paper appeal process does not require a provider to issue a check because the refund amount is reduced on the R&S Report. To accurately process claim refunds, the TMHP Cash Reimbursement Unit requests that the refund check be accompanied by Texas Medicaid Refund Information Form, with the following information:

- Refunding provider’s name and provider identifier.
- Client’s name and Medicaid ID number.
- Date of service.
- A copy of the R&S Report showing the claim to which the refund is being applied.
- The specific reason for the refund.
- Name and address of the attorney or casualty insurance company (including the policy and claim number).
- TPR subscriber information.
- Amount of insurance payment.

Refer to: “Section 8: Third Party Liability (TPL)” (Vol. 1, General Information) for additional TPL information.

### 7.3 Appeals to HHSC Texas Medicaid Fee-for-Service

#### 7.3.1 Administrative Claim Appeals

An administrative appeal is a request for review of (not a hearing on) claims that are denied by TMHP or claims processing entity for technical and nonmedical reasons. There are two types of administrative appeals:

- **Exception requests to the 95-day filing deadline or 120-day appeal deadline.** A provider’s formal written request for review of (not a hearing on) a claim that is denied or adjusted by TMHP for failure to meet the 95-day filing deadline or 120-day appeal deadline. Exception requests to the 95-day filing deadline should meet one of the five exceptions in subsection 7.3.1.2, “Exceptions to the 95-Day Filing Deadline” in this section. Exceptions to the 120-day appeal deadline should meet one of the situations in subsection 7.3.1.3, “Exceptions to the 120-day Appeal Deadline” in this section.

- **Standard Administrative Appeal.** A provider’s formal written request for review of (not a hearing on) a claim or prior-authorization that is denied by TMHP for technical or non-medical reasons.

An administrative claims appeal is a request for a review as defined in Title 1 TAC §354.2201(2).

An administrative appeal must be:

- Submitted in writing to HHSC Claims Administrator Operations Management by the provider delivering the service or claiming reimbursement for the service.
• Received by HHSC Claims Administrator Operations Management after the appeals process with TMHP or the claims processing entity has been exhausted, and must contain evidence of appeal dispositions from TMHP or the claims processing entity:
  • All correspondence and documentation from the provider to TMHP or the claims processing entity including copies of supporting documentation submitted during the appeal process.
  • All correspondence from TMHP or the claims processing entity to the provider including TMHP’s final decision letter or such from the claims processing entity.
• Complete and contain all of the information necessary for consideration and determination by HHSC Claims Administrator Operations Management to include the following:
  • A written explanation specifying the reason/request for appealing the claim.
  • Supporting documentation for the request.
  • All R&S Reports identifying the claims/services in question.
  • Identification of the incorrect information and the corrected information that is to be used to appeal the claim.
  • A copy of the original claim, if available. Claim copies are helpful when the appeal involves medical policy or procedure coding issues. Also provide a corrected signed claim.
  • A copy of supporting medical documentation that is necessary or requested by TMHP.
  • Provider’s internal notes and logs or ticket numbers from the TMHP Contact Center when pertinent (cannot be used as proof of timely filing).
  • Memos from HHSC, TMHP, or claims processing entity indicating any problems, policy changes, or claims processing discrepancies that may be relevant to the appeal.
  • Other documents, such as receipts (i.e., certified mail along with a detailed listing of the claims enclosed), in-service notes, minutes from meetings, if relevant to the appeals. Receipts can be helpful when the issue is late filing.
• Received by HHSC Claims Administrator Operations Management within 120 days from the date of disposition by TMHP or the claims processing entity as evidenced by the weekly R&S Report.

Administrative appeals can be submitted electronically through the My Account page on the TMHP portal. Once an appeal has been submitted, providers can login to My Account to view the following information:
• The status of administrative appeals that were submitted electronically or on paper
• Communications from TMHP and Texas HHSC

Providers who have submitted their claims electronically must identify the batch submission ID with the date on the electronic claims report. This report must indicate the TMHP assigned batch ID. In addition, this report must include the individual claim that is being appealed. The claim information on the batch report, including date of service and billed amount, must match the information on the claim that is being appealed. This required information constitutes proof of timely filing.

Note: Only reports accepted or rejected from TMHP or the claims processing entity to the vendor will be honored unless the provider is a direct submitter (TexMedConnect). Office notes indicating claims were submitted on time or personal screen prints of claim submissions are not considered proof of timely filing.

HHSC Claims Administrator Operations Management only reviews appeals that are received within 18 months from the DOS. All claims must be paid within 24 months from the DOS as outlined in 1 TAC §354.1003.
Providers must adhere to all filing and appeal deadlines for an appeal to be reviewed by HHSC Claims Administrator Operations Management. The filing and appeal deadlines are described in 1 TAC §354.1003.

Additional information requested by HHSC Claims Administrator Operations Management must be returned to HHSC Claims Administrator Operations Management within 21 calendar days from the date of the letter from HHSC Claims Administrator Operations Management. If the information is not received within 21 calendar days, the case is closed.

A determination made by HHSC Claims Administrator Operations Management is the final decision for claim appeals. No additional consideration is available. Therefore, ensure that all documents pertinent to the appeal are submitted. New evidence is required for an additional appeal to HHSC Claims Administrator Operations Management.

Mail appeal requests to the following address:

Texas Health and Human Services Commission  
HHSC Claims Administrator Operations Management  
Mail Code-91X  
PO Box 204077  
Austin, Texas 78720-4077

Providers may request the status of an administrative appeal by sending an email to HHSC at MCD_Administrative_Appeals@hhsc.state.tx.us. The email must include the TPI number, client name, Medicaid number, date of service, and, if available, the case review number.

7.3.1.1 Requirements for Exception Requests

HHSC Claims Administrator Operations Management makes the final decision on whether claims fall within one of the exceptions to the 95-day or 120-day filing deadlines.

Providers must submit the following documentation for all exception requests:

- Exception requests must be in writing and mailed directly to HHSC.
- Adequate back-up documentation must accompany the exception request. Failure to provide adequate documentation results in the case being closed. Providers are notified of the reason for denial.
- All claims that are to be considered for an exception must accompany the request. HHSC will consider only the claims that are attached to the request.
- Additional claims cannot be added to an exception request after the exception request has been completed by HHSC. Additional claims must be submitted as a separate request and must include all required documentation. Information from a previous request will not be linked by HHSC to process additional claims.
- All exception requests must include an affidavit or statement from the provider stating the details of the cause for the delay, the exception being requested, and verification that the delay was not caused by neglect, indifference, or lack of diligence of the provider or the provider’s employee or agent. This affidavit or statement must be made by the person with personal knowledge of the facts.
- Multiple requests submitted simultaneously must be sorted by provider identifier first, and then alphabetically by client name. The orderly submission of exception requests facilitates the review process. Exception requests are returned to the provider if not submitted in the required format.

HHSC may request additional information which must be received within 21 calendar days from the date of the letter from HHSC. If the information is not received within 21 calendar days, the case will remain closed.

HHSC notifies providers about the outcome of the case upon completion of an exception request review.
7.3.1.2 Exceptions to the 95-Day Filing Deadline

HHSC Claims Administrator Operations Management is responsible for reviewing requests for exceptions to the 95-day filing deadline for Texas Medicaid fee-for-service. Only providers can submit exception requests. Requests from billing companies, vendors, or clearinghouses are not accepted unless accompanied by a signed authorization from the provider (with each appeal). Without provider authorization, these requests are returned without further action.

HHSC will only consider exceptions to the 95-day filing deadline for claims that are submitted within the 365-day federal filing deadline from the date of service as outlined in 1 TAC §354.1003.

Exceptions to the filing deadline are considered when one of the following situations exists:

- Catastrophic event that substantially interferes with normal business operations of the provider, or damage or destruction of the provider’s business office or records by a natural disaster, including, but not limited to, fire, flood, or earthquake; or damage or destruction of the provider’s business office or records by circumstances that are clearly beyond the control of the provider, including, but not limited to, criminal activity. The damage or destruction of business records or criminal activity exception does not apply to any negligent or intentional act of an employee or agent of the provider because these persons are presumed to be within the control of the provider. The presumption can only be rebutted when the intentional acts of the employee or agent lead to termination of employment and filing of criminal charges against the employee or agent.

  Providers requesting an exception for catastrophic events must include independent evidence of insurable loss; medical, accident, or death records; or police or fire report substantiating the exception of damage, destruction, or criminal activity.

- Delay or error in the eligibility determination of a client, or delay due to erroneous written information from HHSC, its designee, or another state agency.

  Providers requesting an exception for the delay or error in the eligibility determination of a client or delay due to erroneous written information from HHSC, its designee, or another state agency must include the written document from HHSC or its designee that contains the erroneous information or explanation of the delayed information.

- Delay due to electronic claim or system implementation problems experienced by HHSC, its designee, or Texas Medicaid providers.

  Providers requesting an exception for the delay due to electronic claim or system implementation problems experienced by HHSC, its designee, or Texas Medicaid providers must include the written repair statement, invoice, computer or modem generated error report (indicating attempts to transmit the data failed for reasons outside the control of the provider), or the explanation for the system implementation problems.

  The documentation must include a detailed explanation made by the person making the repairs or installing the system, specifically indicating the relationship and impact of the computer problem or system implementation to claims submission, and a detailed statement explaining why alternative billing procedures were not initiated after the delay in repairs or system implementation was known.

  If the provider is requesting an exception based upon an electronic claim or system implementation problem experienced by HHSC or its designee, the provider must submit a written statement outlining the details of the electronic claim or system implementation problems experienced by HHSC or its designee that caused the delay in the submission of claims by the provider, any steps taken to notify the state or its designee of the problem, and a verification that the delay was not caused by the neglect, indifference, or lack of diligence on the part of the provider or its employees or agents.
• Submission of claims occurred within the 365-day federal filing deadline, but the claim was not filed within 95 days from the date of service because the service was determined to be a benefit of the Medicaid program, and an effective date for the new benefit was applied retroactively.

Providers requesting an exception for claims that were submitted within the 365-day federal filing deadline, but were not filed within the 95-days of the date of service because the service was determined to be a benefit of Texas Medicaid and an effective date for the new benefit was applied retroactively, must include a written, detailed explanation of the facts and documentation to demonstrate the 365-day federal filing deadline for the benefit was met.

• Client eligibility is determined retroactively and the provider is not notified of retroactive coverage.

Providers requesting an exception for client eligibility determined retroactively and the provider is not notified of retroactive coverage must include a written, detailed explanation of the facts and activities illustrating the provider’s efforts in requesting eligibility information for the client. The explanation must contain dates, contact information, and any responses from the client.

7.3.1.3 Exceptions to the 120-day Appeal Deadline

HHSC must receive a written exception request within 120 days of TMHP’s final action. HHSC shall consider exceptions to the 120-day appeal deadline for the following listed situations. This is a one-time exception request; therefore, all claims that are to be considered within the request for an exception must accompany the request. Claims submitted after HHSC’s determination has been made for the exception will be denied consideration because they were not included in the original request.

• An exception request must be received by HHSC within 18 months from the date of service to be considered. This requirement will be waived for the exceptions listed in the following bullets (b) and (c), as well as the situation listed under “Exceptions to the 24-month deadline.”

• The following exceptions to the 120-day appeal deadline are considered if the criteria in the previous bullet is met and there is evidence to support one of the following bullets:
  
  (a) Errors made by a third party payor that were outside the control of the provider. The provider must submit a statement outlining the details of the cause for the error, the exception being requested, and verification that the error was not caused by neglect, indifference, or lack of diligence of the provider, the provider’s employee, or agent. This affidavit or statement should be made by the person with personal knowledge of the facts. In lieu of the above affidavit or statement from the provider, the provider may obtain an affidavit or statement from the third party payor including the same information, and provide this to HHSC as part of the request for appeal.

  (b) Errors made by the reimbursement entity that were outside the control of the provider. The provider must submit a statement from the original payor outlining the details of the cause of the error, the exception being requested, and verification that the error was not caused by neglect, indifference, or lack of diligence on the part of the provider, the provider’s employee, or agent. In lieu of the above reimbursement entity’s statement, the provider may submit a statement including the same information and provide this to HHSC as part of the request for appeal.

  (c) Claims were adjudicated, but an error in the claim’s processing was identified after the 120-day appeal deadline. The error is not the fault of the provider. An error occurred in the claims processing system that is identified after the 120-day appeal deadline has passed.

7.3.1.4 Exceptions to the 24-Month Payment Deadline

HHSC shall consider exceptions to the 24-month claims payment deadline for the following listed situations. The final decision about whether a claim falls within one of the following exceptions will be made by HHSC.
• Claims for providers with retroactive adjustments who are reimbursed under a retrospective payment system.
• Claims paid within six months from the Medicare paid date.
• Claims from providers under investigation for fraud or abuse.
• Claims paid at any time in accordance with a court order, to carry out hearing decisions or agency corrective actions taken to resolve a dispute, or to extend the benefits of a hearing decision, corrective action, or court order to others in the same situation as those directly affected by it.

Mail exception requests to HHSC at the following address:

Texas Health and Human Services Commission
HHSC Claims Administrator Operations Management
Mail Code 91X
PO Box 204077
Austin, TX 78720-4077

7.3.2 Medical Necessity Appeals

Medical necessity appeals are defined as disputes regarding medical necessity of services. Providers must appeal to TMHP and exhaust the appeal/grievance process before submitting an appeal to HHSC.

Medical necessity appeals related to UR decisions made by the HHSC OIG UR Unit must be appealed to HHSC not TMHP.

When filing appeals to HHSC, providers must submit copies of all supporting documentation, including information sent to TMHP.

Refer to: Subsection 7.1.5.1, “Texas Medicaid Fee-for-Service DRG Adjustment Appeal” in this section.

7.3.3 Utilization Review Appeals

Hospitals may appeal adverse UR decisions made by the HHSC OIG UR Unit to the HHSC Medical and UR Appeals Unit. The written appeal request, with complete medical record and approved Business Records Affidavit Form must be received by the Medical and UR Appeals Unit within 120 days of the date of the original HHSC OIG UR decision letter. If the appeal request with all required documentation is not received within 120 days, the appeal is not conducted, and the HHSC OIG UR decision is considered final. Any claim the facility may have to the Medicaid funds at issue are barred. Extensions of time are not granted for filing the written appeal request and submitting all of the required documentation. The procedures and specific requirements for appealing these decisions can be found in the sections that follow.

Hospitals may appeal adverse HHSC OIG UR Unit determinations to the following address:

HHSC Medical and UR Appeals
Mail Code H-230
PO Box 85200
Austin, TX 78708

or

Broadmoor Building 902
11501 Burnet Road (Express Mail Only)
Austin, TX 78758
7.3.3.1 Admission Denials, Continued Stay Denials for TEFRA Hospitals, DRG Revisions, and Cost/Day Outlier Denials

If a hospital is dissatisfied with the original retrospective review conducted by the HHSC OIG UR Unit, it may submit a written request for an appeal to the HHSC Medical and Utilization Review Appeals Unit.

The request for an appeal must include a copy of the complete medical record, a letter explaining the reasons why the HHSC OIG UR decision is incorrect, a copy of the HHSC OIG UR decision letter, and an original, properly completed, and notarized affidavit in the format approved by HHSC. The affidavit allows the hospital to certify the record as a business and legal document.

**Refer to:** [Business Records Affidavit Form](https://www.tmhp.com) on the TMHP website at [www.tmhp.com](http://www.tmhp.com).

Complete medical records must be provided to HHSC at no charge. A complete medical record must include, but is not limited to, a discharge summary, history and physical, emergency room record, operative report, pathology report, anesthesia record, consultation reports, physician progress notes, physician orders, laboratory reports, X-ray reports, special diagnostic reports, nurses’ notes, and medication records.

Any additional information requested by the HHSC Medical and UR Appeals Unit must be returned to the HHSC Medical and UR Appeals Unit within 21 calendar days of the request. If the requested documentation is not received within this time frame, the case is closed without an opportunity for further review and the original HHSC OIG UR decision is considered the final decision.

If a hospital is notified that it failed to submit any required documentation with the initial appeal request, the required information must be returned to the HHSC Medical and UR Appeals Unit within 21 calendar days of the date of notification, or within 120 days of the date of the original HHSC OIG UR decision letter, whichever is sooner. If the required documentation is not received within the time frames, the case is closed without an opportunity for further review and the original HHSC OIG UR decision is considered the final decision. Extensions of time are not granted for filing the written appeal request and submitting all required documentation.

The HHSC Medical and UR Appeals Unit is responsible for conducting an independent review in response to a provider’s appeal. The professional staff uses only the documentation submitted in the medical record to determine whether an inpatient admission was appropriate and whether the diagnoses and procedures were correct. The HHSC UR and Medical Appeals physician or designee performs a complete review for the medical necessity of inpatient admission, DRG validation, quality of care, continued stay medical necessity, and ancillary charges (TEFRA cases) using the medical record documentation submitted on appeal. After completion of the review, the physician or designee renders a final decision on the case. The final decision may include determinations regarding multiple aspects of the admission. The hospital is notified in writing of the final decision. Inpatient admission denials cannot be rebilled as outpatient claims except as noted in subsection 4.2.4, “Outpatient Observation Room Services” in the *Inpatient and Outpatient Hospital Services Handbook* (Vol. 2, Provider Handbooks).

The HHSC Medical and Utilization Review Unit recognizes that hospital staff may use guidelines, such as the American Hospital Association’s Coding Clinic, to assist them in identifying diagnoses or procedures for statistical and billing purposes. However, the HHSC Medical and Utilization Review Appeals Unit determines the appropriate diagnoses or procedures for reimbursement purposes using the documentation in the medical record (submitted on appeal) and the following guidelines:

- **Principal diagnosis assignment.** The diagnosis (condition) established after study to be chiefly responsible for causing the admission of the client to the hospital for care. The principal diagnosis must be treated or evaluated during the admission to the hospital.

- **Secondary diagnosis assignment.** Conditions that affect patient care in terms of requiring clinical evaluation, therapeutic treatment, diagnostic procedures, extended length of hospital stay, or increased nursing care or monitoring, or, in the case of a newborn (birth through 28 days of age),
which the physician deems to have clinically significant implications for future health care needs. Normal newborn conditions or routine procedures should not be considered as complications or comorbidities for DRG assignment.

If the principal diagnosis, secondary diagnoses, or procedures are not substantiated in the medical record, not sequenced correctly, or have been omitted, the codes may be changed, added, or deleted by the HHSC Medical and UR Appeals physician or designee. When it is determined the diagnoses or procedures are substantiated and sequenced correctly, a final DRG assignment is made.

If the hospital is displeased with the appeals decision, the attending physician or medical director of the hospital may request an educational conference with the HHSC Medical and UR Appeals physician or designee. The educational conference is held by telephone between the physician or designee and the hospital medical director or attending physician. This is an opportunity for the physicians to discuss the deciding factors in the case and any hospital billing processes that may have affected the adjudication of the case. The educational conference will not alter the previous appeal decision.

### 7.3.3.2 Final Technical Denials

Hospitals may submit a request for a written appeal to HHSC Medical and UR Appeals only if the hospital has evidence that the HHSC OIG UR Unit issued a final technical denial in error, or did not provide proper notification of the preliminary technical denial. The request must include a letter explaining the reasons why the HHSC OIG UR decision is incorrect and a copy of the HHSC OIG UR decision letter.

The written appeal request must be received by HHSC Medical and UR Appeals within 120 days of the date of the original HHSC OIG UR decision letter. If the request is not received within the 120 days, the appeal is not conducted and the HHSC OIG UR decision is considered final. Any claim the facility may have to the Medicaid funds at issue are barred. Extensions of time are not granted for filing the written appeal request.

If the appeal time frame is met, the HHSC Medical and UR Appeals Unit reviews all the documentation and renders a final decision on the case. If it is determined the technical denial was issued correctly by the HHSC OIG UR Unit, HHSC’s decision is upheld. The hospital is notified in writing of the decision. This decision is the final decision.

If it is determined that the final technical denial decision should be overturned, the HHSC Medical and UR Appeals Unit will request a copy of the complete medical record and an original, properly completed, notarized affidavit in the format approved by HHSC. The affidavit allows the hospital to certify the record as a business and legal document. The HHSC Medical and UR Appeals physician or designee performs a complete review for the medical necessity of the admission, DRG validation, quality of care or continued stay, and ancillary charges (for TEFRA Hospitals) using only the medical record documentation. After completion of the review, the physician or designee renders a final decision on the case. The hospital is notified in writing of the final decision.

If the requested documentation is not received within the required 21-day time frame, the case is closed without further opportunity for review and the original HHSC OIG UR decision is considered final.

### 7.3.4 Provider Complaints

TMHP provides for due process for resolving all provider complaints. A complaint is defined as any dissatisfaction expressed by telephone or in writing by the provider, or on behalf of that provider, concerning Texas Medicaid. The definition of complaint does not include a misunderstanding or a problem of misinformation that is resolved promptly by clearing up the misunderstanding or supplying the appropriate information to the provider’s satisfaction. The definition also does not include a provider’s oral or written dissatisfaction with an adverse determination or appeals regarding claim payments and denials.
Procedures governing the provider complaints process are designed to identify and resolve provider complaints in a timely and satisfactory manner. Most complaints are resolved within 30 calendar days. Complaints to TMHP may be submitted using the following methods:

- By using the Email Us button on the TMHP Contact web page
- By telephone at 1-800-925-9126
- In writing to:

  TMHP
  Complaints Resolution Department
  PO Box 204270
  Austin, TX 78720-4270

Questions regarding the complaint process or the status of a complaint should be directed to the TMHP Contact Center at 1-800-925-9126.

7.3.4.1 Provider Complaint Policy

TMHP takes seriously and acts on each provider complaint. Depending on the level and nature of the complaint, TMHP works with the provider to resolve the issue or directs the complaint to the appropriate department.

The Medical Affairs Division handles complaints that relate to utilization of services (including ER use), denial of continued stay, and all clinical and access issues. This includes provider’s appeal of an adverse authorization decision.

If the complaint relates to a medical issue, the Medical Affairs Division staff may assist in resolving the complaint. The provider complaints process applies only to the resolution of disputes within the control of Texas Medicaid, such as administrative or medical issues. The provider complaint process does not apply to allegations of negligence against third parties, including other Texas Medicaid providers. These complaints are referred to HHSC for review and evaluation and are resolved by HHSC.

7.3.4.2 Provider Complaint Process

The TMHP Complaints Resolution Department handles all provider complaints. The processing of a provider’s complaint is described as follows:

- Providers must submit their complaint by telephone or in writing (mail or fax).
- Providers will receive a written acknowledgement letter from TMHP within five business days of receipt of the complaint.
- Referrals to other departments, such as Provider Relations or Medical Affairs, are made when appropriate.
- If the complaint cannot be resolved within 30 calendar days, the provider is notified in writing of the status of the complaint.
- If the Email Us option on the TMHP contact page is used, providers can select to receive communication by text, email, or both.

Providers who believe they did not receive due process regarding their complaint from TMHP may file a complaint with HHSC. Providers are encouraged to utilize the appeals and grievance process with TMHP before filing a complaint with HHSC.
7.3.4.3 Complaints to HHSC—Texas Medicaid Fee-for-Service

Texas Medicaid fee-for-service providers may file complaints to the HHSC Claims Administrator Operations Management if they find they did not receive full due process from TMHP in the management of their appeal. Texas Medicaid fee-for-service providers must exhaust the appeals and grievance process with TMHP before filing a complaint with the HHSC Claims Administrator Operations Management.

Refer to: Subsection 7.3, “Appeals to HHSC Texas Medicaid Fee-for-Service” in this section for information about submission of an appeal to HHSC.

A complaint is defined as any dissatisfaction expressed in writing by the provider, or on behalf of that provider, concerning Texas Medicaid. The term complaint does not include the following:

- A misunderstanding or a problem of misinformation that is resolved promptly by clearing up the misunderstanding or supplying the appropriate information to the provider’s satisfaction.
- A provider’s oral or written dissatisfaction with an adverse determination.

Under the complaint process, the HHSC Claims Administrator Operations Management works with TMHP and providers to verify the validity of the complaint, determine if the established due process was followed in resolving appeals and grievances, and addresses other program and contract issues, as applicable.

Complaints must be in writing and received by the HHSC Claims Administrator Operations Management within 60 calendar days from TMHP’s written notification of the final appeal decision.

When filing a complaint, providers must submit a letter explaining the specific reasons they believe the final appeal decision by TMHP is incorrect and copies of the following documentation:

- All correspondence and documentation from the provider to TMHP, including copies of supporting documentation submitted during the appeal process.
- All correspondence from TMHP to the provider, including TMHP’s final decision letter.
- All R&S Reports of the claims and services in question, if applicable.
- Provider’s original claim or billing record, electronic or manual, if applicable.
- Provider’s internal notes and logs when pertinent.
- Memos from HHSC or TMHP indicating any problems, policy changes, or claims’ processing discrepancies that may be relevant to the complaint.
- Other documents, such as certified mail receipts, original date-stamped envelopes, in-service notes, or minutes from meetings if relevant to the complaint. Receipts can be helpful when the issue is late filing.

Complaint requests may be mailed to the following address:

Texas Health and Human Services Commission
HHSC Claims Administrator Operations Management
Mail Code 91X
PO Box 204077
Austin, TX 78720-4077

7.4 Cost Report Settlement Appeal Process

A provider who is dissatisfied with the determination contained in the Notice of Program Reimbursement (NPR) from TMHP Medicaid Audit may request an appeal as follows:

- The request for appeal must be in writing.
- The request for appeal must be filed within 180 calendar days from the date of receipt of the NPR.
• If the amount in controversy is at least $1,000, the request for the appeal must be filed with TMHP Medicaid Audit.

• If the NPR shows that the provider is indebted to Texas Medicaid, TMHP must take the necessary action to recover the overpayment, including a suspension of interim payments. This process will take place even if an appeal has been requested.

### 7.4.1 Appeals to TMHP Medicaid Audit

A provider’s request to appeal his or her NPR must include the following:

- Identify specific individual items in TMHP Medicaid Audit’s determination with which the provider disagrees.
- Give the reasons the provider believes these are incorrect.
- Identify the amount in controversy for each item and provide a calculation of that amount.

The appeal may include any materials the provider believes will support its position.

TMHP Medicaid Audit completes a desk review of the appeal within six months of the date of receipt of complete documentation supporting the appeal. TMHP does the following:

- Reviews the materials submitted by the provider.
- Informs the provider if it appears that the request for an appeal was not timely or the amount of controversy is not at least $1,000.
- Reviews the record that formed the basis for the determination of the total payment due to the provider.
- Attempts to resolve as many points in controversy as possible with the provider and inform him or her in writing the issues that have been resolved and those that the provider may appeal to HHSC.
- Ensures all available documentation in support of the provider or TMHP Medicaid Audit is part of the record.

To appeal to TMHP Medicaid Audit, send the written notice within 120 days of receipt of the NPR letter to the following address:

Texas Medicaid & Healthcare Partnership  
Medicaid Audit Operations Director  
PO Box 200345  
Austin, TX 78720-0345

### 7.5 Forms

The following linked forms can be found on the Forms page of the Provider section of the TMHP website at [www.tmhp.com](http://www.tmhp.com):

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<thead>
<tr>
<th>Forms</th>
</tr>
</thead>
<tbody>
<tr>
<td>Business Records Affidavit Form</td>
</tr>
<tr>
<td>Texas Medicaid Refund Information Form</td>
</tr>
<tr>
<td>Credit Balance Refund Worksheet</td>
</tr>
</tbody>
</table>
SECTION 8: THIRD PARTY LIABILITY (TPL)

TEXAS MEDICAID PROVIDER PROCEDURES MANUAL: VOL. 1

MAY 2021
SECTION 8: THIRD PARTY LIABILITY (TPL)

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8.1 Third Party Liability (TPL)

Texas Medicaid Third Party Liability program recovers payments from third parties that are responsible for paying towards a medical claim for services rendered to a Texas Medicaid client. A third party resource (TPR) is the entity, individual, or other source (other than Medicaid, Medicaid managed care organizations (MCOs), Medicaid managed care dental plans, the client, non-TPR sources, or Medicare) that is legally responsible for paying the medical claims of Texas Medicaid clients. The Third Party Liability program helps reduce Medicaid costs by shifting claims expenses to third party payers.

As a condition of eligibility, Medicaid clients assign their rights (and the rights of any other eligible individuals on whose behalf he or she has legal authority under state law to assign such rights) to medical support and payment for medical care from any third party to Medicaid.

Federal and state laws require the use of Medicaid funds for the payment of most medical services only after all reasonable measures have been made to use a client’s TPR or other insurance. To the extent allowed by federal law, a health-care service provider must seek reimbursement from available third party insurance that the provider knows about or should know about before billing Texas Medicaid. Medicaid pays only after the third party has met its legal obligation to pay (i.e., Medicaid is the payer of last resort). All claims submitted for Texas Medicaid payment for clients with other insurance coverage must reference the information, regardless of whether a copy of the explanation of benefits (EOB) from the insurance company is submitted with the claim.

Refer to: Subsection 7.2, “Refunds to TMHP” in “Section 7: Appeals” (Vol. 1, General Information) for information regarding refunds to TMHP resulting from other insurance payments and conditions surrounding provider billing of third party insurers.

Eligible clients enrolled in private health maintenance organizations (HMOs) must not be charged the co-payment amount because the provider has accepted Medicaid assignment.

8.2 Verifying a Client’s TPR

Providers can verify a client’s TPR and other insurance information by:

- Using TexMedConnect.
- Accessing the Medicaid Client Portal for Providers.

To ensure receipt of TPR disposition of payment or denial, providers must obtain an assignment of insurance benefits from the client at the time of service. Providers are asked not to provide claim copies or statements to the client.

Providers that are aware that a client has other health insurance that is not indicated on TexMedConnect or the Medicaid Client Portal for Providers must notify TMHP of the details concerning the type of policy and scope of benefits.

Providers can notify TMHP by calling TPR at 1-800-846-7307, sending a fax to 1-512-514-4225, or mailing the Other Insurance Form to the following address:

Texas Medicaid & Healthcare Partnership
Third Party Resources Unit
PO Box 202948
Austin, TX 78720-2948

Any indemnity insurance policy that pays cash to the insured for wages lost or for days of hospitalization rather than for specific medical services is considered a TPR if the policy is assignable to someone else. The Health and Human Services Commission (HHSC) has assignment to any Medicaid applicant’s or client’s right of recovery from a third party health insurer, to the extent of the cost of medical care services paid by Medicaid. Texas Medicaid requires a provider take all reasonable measures to use a client’s TPR before billing Medicaid.
Medicaid-eligible clients may not be held responsible for billed charges that are in excess of the TPR payment for services covered by Texas Medicaid. If the TPR pays less than the Medicaid-allowable amount for covered services, the provider should submit a claim to TMHP for any additional allowable amount.

8.3 TPR Sources
A provider who furnishes services and participates in Texas Medicaid may not refuse to furnish services to an eligible client because of a third party’s potential liability for payment of the services.

TPR includes payments from any of the following sources:
- Other health insurance including assignable indemnity contract
- Health maintenance organization (HMO)
- Public health programs available to clients with Medicaid such as Medicare and Tricare
- Profit and nonprofit health plan
- Employment-related health insurance
- Self-insured plans
- Casualty coverage resulting from an accidental injury such as automobile or property insurance (including no-fault insurance); No-fault automobile insurance such as personal injury protection (PIP) and automobile medical insurance
- Liability insurance
- Life insurance policies, trust funds, cancer policies, or other supplemental policies
- Workers’ Compensation
- Medical support from absent parents
- Court judgments or settlements from a liability insurer
- First party probate-estate recoveries
- Other federal programs unless excluded by statute
- Other liable third parties

Claims for family planning services (including Title XIX and the Department of State Health Services [DSHS] Family Planning Program) must be submitted to TMHP before billing a client’s TPR. Federal regulations protect the client’s confidential choice of birth control and family planning services. Confidentiality is jeopardized when seeking information from TPRs.

School Health and Related Services (SHARS) providers are not required to file claims with private insurance before billing Medicaid.

Early Childhood Intervention (ECI) providers are not required to file claims with private insurance before billing Medicaid for Targeted Case Management services and Specialized Skills Training (SST).

Case Management for Children and Pregnant Women providers are not required to file claims with other health insurance before filing with Medicaid.

THSteps medical and dental providers are not required to bill other insurance before billing Medicaid. If the provider is aware of other insurance, however, the provider must choose whether or not to bill the other insurance. The provider has the following options:
- If the provider chooses to bill the other insurance, the provider must submit the claim to the client’s other insurance before submitting the claim to Medicaid.
• If the provider chooses to bill Medicaid and not the client’s other insurance, the provider is indicating acceptance of the Medicaid payment as payment in full. Medicaid then has the right to recovery from the other insurance. The provider does not have the right to recovery and cannot seek reimbursement from the other insurance after Medicaid has made payment.

• If the provider learns of other insurance coverage after Medicaid has paid, the provider must refund Medicaid before billing the other insurance.

8.3.1 Exceptions

Certain funding sources are an exception to TPR. Adoption agencies or foster parents are not considered a TPR. Medicaid is primary in the STAR Health program (except when court-ordered to provide health insurance). This is an exception to the rule that Medicaid is payer of last resort. Providers must not bill other health insurance unless there is a court order that places this responsibility elsewhere. For Texas Health Steps (THSteps), providers must refer to the MCO, or dental plan that administers the client’s managed care benefits for additional information.

Refer to: Subsection 1.1, “About the Vendor Drug Program” in the Outpatient Drug Services Handbook (Vol. 2, Provider Handbooks) for information about this program.

Non-TPR sources are secondary to Texas Medicaid and may only pay benefits after Texas Medicaid. The following are the most common non-TPR sources:

• Texas Workforce Solutions–Vocational Rehabilitation Services
• Texas Kidney Health Care Program
• Crime Victims’ Compensation Program
• Muscular Dystrophy Association
• Children with Special Health Care Needs (CSHCN) Services Program
• Texas Band of Kickapoo Equity Health Program
• Maternal and Child Health
• DSHS Family Planning Program
• State Legalization Impact Assistance Grant (SLIAG)
• Adoption Agencies
• Home and Community-based Waivers Programs through HHSC
• Blind and Visually Impaired Services
• Healthy Texas Women

If providers have questions about others not listed, they may contact a provider relations representative.

Note: Claims for clients who are seeking disability determination must be submitted to HHSC for consideration of reimbursement. Refer to the HHS website at https://hhs.texas.gov/ for additional information about disability determinations and claims filing.

Denied claims or services that are not a benefit of Medicaid may be submitted to non-TPR sources. If a claim is submitted inadvertently to a non-TPR source listed above before submission to TMHP, the claim may be submitted to TMHP within the filing deadlines.


If a non-TPR source erroneously makes a payment for a dual-eligible client for services also covered by Medicaid, the payment is refunded to the non-TPR source.
8.3.1.1 Adoption Cases

Texas Medicaid, not the adoption agency, should be billed for all medical services that are a benefit of Texas Medicaid. If a claim is inadvertently sent to the adoption agency before it is sent to TMHP, TMHP must receive the claim within 95 days of the date of disposition from the adoption agency denial, payment, requests for refund or recoupment, to be considered for payment.

If the adoption agency inadvertently makes a payment for services that are covered by Medicaid, the provider must refund the payment to the agency.


A copy of the non-TPR disposition must be submitted with the claim and received at TMHP within 95 days of the date of the disposition (denial, payment, request for refund, or recoupment of payment by the non-TPR source).

8.4 Non-TPR Sources

A non-TPR is secondary to Texas Medicaid and may only pay benefits after Texas Medicaid. Non-TPR sources include, but are not limited to, accident-only policies, managed care plans, Indigent Health Care, and Medicaid/Children’s Health Insurance Program (CHIP).

The provider bills TMHP directly within 95 days from the date of service. However, if a non-TPR is billed first, TMHP must receive the claim within 95 days of the claim disposition by the other entity.

Note: The provider must submit a copy of the disposition with the claim.

8.5 TPL for Hospitals

Inpatient and outpatient hospitals and providers enrolled in Texas Medicaid are required to inform TMHP about circumstances that may result in third party liability for health-care claims. After receiving this information, TMHP pursues reimbursement from responsible third parties.

Hospitals and providers must fax the Other Insurance Form for Health Insurance or the Tort Response Form for accidents to 1-512-514-4225 or mail it to the following address:

Texas Medicaid & Healthcare Partnership
TPL Correspondence
Third Party Liability Unit
PO Box 202948
Austin, TX 78720-2948

8.6 TPL for Managed Care

Medicaid MCO and dental plan claims payments for services rendered to Texas Medicaid managed care clients are subject to the federal and state requirement for other insurance payments. As directed by HHSC, Texas Medicaid will recover payments based on claims data processed by the Medicaid MCO or dental plan.

TPR includes payments from any of the following sources:

- Other health insurance, including assignable indemnity contracts
- Commercial MCOs (other insurance available through a source other than Texas Medicaid or Medicare)

A provider who furnishes services and participates in Texas Medicaid may not refuse to furnish services to an eligible client because of a third party’s potential liability for the payment of the services.
8.7 Tort

HHSC contracts with TMHP to administer third party liability cases. To ensure that Texas Medicaid is the payer of last resort, TMHP performs postpayment investigations of potential casualty and liability cases. TMHP also identifies and recovers Medicaid expenditures in casualty cases involving Medicaid clients.

The Human Resources Code, chapter 32, section 32.033 establishes automatic assignment of a Medicaid client’s right of recovery from personal insurance as a condition of Medicaid eligibility.

Investigations are a result of referrals from many sources, including attorneys, insurance companies, health-care providers, Medicaid clients, and state agencies. Referrals should be faxed to 1-512-514-4225 or mailed to:

TMHP TPL/Tort Department
PO Box 202948
Austin, TX, 78720-2948


TMHP releases Medicaid claims information when an HHSC Authorization for Use and Release of Health Information Form is submitted. The form must be signed by the Medicaid client. Referrals are processed within ten business days.


An attorney or other person who represents a Medicaid client in a third party claim or action for damages for personal injuries must send written notice of representation. The written notice must be submitted within 45 days of the date on which the attorney or representative undertakes representation of the Medicaid client, or from the date on which a potential third party is identified. The following information must be included:

- The Medicaid client’s name, address, and identifying information.
- The name and address of any third party or third party health insurer against whom a third party claim is or may be asserted for injuries to the Medicaid client.
- The name and address of any health-care provider that has asserted a claim for payment for medical services provided to the Medicaid client for which a third party may be liable for payment, whether or not the claim was submitted to or paid by TMHP.

If any of the information described above is unknown at the time the initial notice is filed, it should be indicated on the notice and revised if and when the information becomes known.

An authorization to release information about the Medicaid client directly to the attorney or representative may be included as a part of the notice and must be signed by the Medicaid client. The HHSC Authorization for Use and Release of Health Information Form must be used.

HHSC must approve all trusts before any proceeds from a third party are placed into a trust.

Providers may direct third party liability questions to the TMHP TPL/Tort Contact Center at 1-800-846-7307.

8.7.1 Accident-Related Claims

TMHP monitors all accident claims to determine whether another resource may be liable for the medical expenses of clients with Medicaid coverage. Providers are requested to ask clients whether medical services are necessary because of accident-related injuries. If the claim is the result of an accident, providers enter the appropriate code and date in Block 10 of the CMS-1500 paper claim form, and Blocks 31-34 on the UB-04 CMS-1450 paper claim form.
If payment is immediately available from a known third party such as Workers’ Compensation or Personal Injury Protection (PIP) automobile insurance, that responsible party must be billed before Medicaid, and the insurance disposition information must be filed with the Medicaid claim. If the third party payment is substantially delayed because of contested liability or unresolved legal action, a claim may be submitted to TMHP for consideration of payment.

TMHP processes the liability-related claim and pursues reimbursement directly from the potentially liable party on a postpayment basis. Include the following information on these claims:

- Name and address of the liable third party
- Policy and claim number
- Description of the accident including location, date, time, and alleged cause
- Reason for delayed payment by the liable third party

8.7.1.1 Accident Resources, Refunds

Acting on behalf of HHSC, TMHP has specific rights of recovery from any settlement, court judgment, or other resources awarded to a client with Medicaid coverage (Texas Human Resources Code, Chapter 32.033). In most cases, TMHP works directly with the attorneys, courts, and insurance companies to seek reimbursement for Medicaid payments. If a provider receives a portion of a settlement for services also paid by Medicaid, the provider must make a refund to TMHP. Any provider filing a lien for the entire billed amount must contact the TPL/Tort Department at TMHP for Medicaid postpayment activities to be coordinated. A provider may not file a lien for the difference between the billed charges and the Medicaid payment. A lien may be filed for services not covered by Medicaid. A lien is the liability of the client with Medicaid coverage.

Providers should contact the TPL/Tort Department at TMHP after submitting an itemized statement and/or claim copies for any accident-related services billed to Medicaid if they received a request from an attorney, a casualty insurance company, or a client.

The provider must submit the following information to TMHP:

- Client’s name
- Medicaid ID number
- Dates of service involved
- Name and address of the attorney or casualty insurance company (including the policy and claim number)

This information enables TMHP to pursue reimbursement from any settlement. Providers must use the Tort Response Form to report accident information to TMHP. When the form is completed, providers must remit it to the TMHP TPL/Tort Department (the address and fax number are on the form).

Providers may contact the TMHP TPL/Tort Department by calling 1-800-846-7307, Option 3, sending a fax to 1-512-514-4225, or mailing to the following address:

Texas Medicaid & Healthcare Partnership
TPL/Tort Department
PO Box 202948
Austin, TX 78720-2948

8.7.1.2 Workers’ Compensation

Payment of covered services under Workers’ Compensation is considered reimbursement in full. The client must not be billed. Services not covered by Workers’ Compensation must be billed to TMHP.
### 8.7.2 Providers Filing Liens for Third Party Reimbursement

Any provider filing a lien for the entire billed amount must contact the TMHP TPL/Tort Department for Medicaid postpayment activities to be coordinated.

A provider may file a lien for the entire billed amount only after meeting the criteria in Title 1 Texas Administrative Code (TAC) §354.2322. Providers who identify a third party, within 12 months of the date of service, and wish to submit a bill or other written demand for payment or collection of debt to a third party after a claim for payment has been submitted and paid by Medicaid must refund any amounts paid before submitting a bill or other written demand for payment or collection of debt to the third party for payment, and they must comply with the provisions set forth in 1 TAC §354.2322, which states:

Providers may retain a payment from a third party in excess of the amount Medicaid would otherwise have paid only if the following requirements are met:

- The provider submits an informational claim to TMHP within the claims filing deadline. (Refer to subsection 8.7.3, “Informational Claims” in this section.)
- The provider gives notice to the client or the attorney or representative of the client that the provider may not or will not submit a claim for payment to Medicaid and the provider may or will pursue a third party, if one is identified, for payment of the claim. The notice must contain a prominent disclosure that the provider is prohibited from billing the client or a representative of the client for any Medicaid-covered services, regardless of whether there is an eventual recovery or lack of recovery from the third party or Medicaid.
- The provider establishes the right to payment separate of any amounts claimed and established by the client.
- The provider obtains a settlement or award in its own name separate from a settlement obtained by or on behalf of the client or award obtained by or on behalf of the client, or there is an agreement between the client or attorney or representative of the client and the provider, that specifies the amount that will be paid to the provider after a settlement or award is obtained by the client.

### 8.7.3 Informational Claims

If providers determine that a third party may be liable for a Medicaid client’s accident-related claim, they can submit an informational claim to the TMHP Tort Department to indicate that a third party is being pursued for payment. This allows providers to secure the 95-day claims filing deadline in the event that the payment is not received from the third party.

TMHP processes informational claims for all claims administered by TMHP, including fee-for-service claims and carve-out services. TMHP does not process informational claims for managed care claims that are administered by the client’s MCO or dental plan.

### 8.7.4 Submission of Informational Claims

Providers must submit informational claims to TMHP:

- On a CMS-1450 UB-04 or CMS-1500 paper claim form. Informational claims cannot be submitted to TMHP electronically or by fax.
- On an Informational Claims Submission Form. Providers should complete only one form per client, regardless of how many separate informational claims are being submitted with the form.
- By certified mail.
- Within the 95-day claims filing deadline. Informational claims will not be accepted after the 95-day claims filing deadline.

**Refer to:** Informational Claims Submission Form on the TMHP website at [www.tmhp.com](http://www.tmhp.com).
Providers must complete either the Insurance Information field (liable third party) or the Attorney Information field on the Informational Claims Submission Form.

Providers must send the informational claims and the Informational Claims Submission Form by certified mail to TMHP at:

TMHP TPL/Tort Department
PO Box 202948
Austin, TX 78720-2948

TMHP will send providers a letter to confirm that the informational claim was received. The letter will provide the date on which TMHP must receive a request from the provider to convert the informational claim to a claim for payment. If TMHP receives an informational claim that cannot be processed, TMHP will notify the provider of the reason.

Providers can inquire about the status of an informational claim by calling the TMHP TPL/Tort Department at 1-800-846-7307. If a provider has not received confirmation that TMHP has received the informational claim within 30 days, the provider should contact the TMHP TPL/Tort Department at 1-800-846-7307 to validate the status of the request.

### 8.7.5 Informational Claim Converting to Claims for Payment

If providers have submitted an informational claim to TMHP but have not received payment from the liable third party, they must make one of the following determinations and notify TMHP within 18 months of the date of service:

- Providers can continue to pursue a claim for payment against the third party and forego the right to convert an informational claim to a claim for payment by Texas Medicaid.

- Providers can submit a request to convert to the informational claim to a claim for payment consideration from Texas Medicaid.

Providers that decide to convert an informational claim to a claim for payment consideration must submit a request to TMHP. The request must be submitted:

- On provider letterhead.
- With the client’s name and Medicaid ID, the date of service, and total billed amount that was originally submitted on the UB-04 CMS-1450 or CMS-1500 paper claim form.
- By fax 1-512-514-4225 or by mail to:

  TMHP/Tort Department
  PO Box 202948
  Austin, TX 78720-2948

TMHP will not accept any conversion request that is submitted more than 18 months after the date of service, regardless of whether an informational claim was submitted timely to TMHP. All payment deadlines are enforced regardless of whether the provider decides to pursue a third party claim. The conversion of informational claims to actual claims is not a guarantee of payment by TMHP.

### 8.8 Other Insurance Claims Filing

The following information must be provided in the “Other Insurance” field on the paper claim and in the appropriate field of electronic claims. On the CMS-1500 paper claim form, Blocks 9 or 11, and 29 must contain the appropriate information:

- Name of the other insurance resource
- Address of the other insurance resource
- Policy number and group number
• Policyholder
• Effective date if available
• Date of disposition by other insurance resource (used to calculate filing deadline)
• Payment or specific denial information

**Important:** By accepting assignment on a claim for which the client has Medicaid coverage, providers agree to accept payment made by insurance carriers and Texas Medicaid when appropriate as payment in full. The client cannot be held liable for any balance or copays related to Medicaid-covered services.

### 8.8.1 Unbundled Services That Are Prior Authorized and Manually Priced Procedure Codes

Providers that submit prior authorization requests and claims to TMHP must:

• Unbundle any bundled procedure codes that have been submitted to the client’s other insurance company.
• Itemize the rates.

If prior authorization has been obtained for services that use manually priced procedure codes, providers must submit claims for the services using the manufacturer’s suggested retail price (MSRP) that was submitted with the authorization request and the following information that is listed on the authorization letter:

• The authorization number
• The provider identifier
• The procedure codes
• The dates of service
• The types of service
• The required modifiers

If the authorization letter shows itemized details, the claim must include all rendered services as they are itemized on the authorization letter and the MSRP rate for each of those services. The procedure codes and MSRP rates that are detailed on the claim must match the procedure codes that are detailed in the authorization letter and the MSRP rates that were submitted with the authorization request. Claims processing and payment may be delayed if there is not an exact match between the detailed information on the authorization letter, the approved authorization, and the information that was submitted on the claim.

**Important:** For appropriate processing and payment, the Pay Price that is indicated on the authorization letter should not be submitted on the claim.

Prior authorization is a condition of reimbursement; it is not a guarantee of payment.

### 8.8.2 Other Insurance Credits

Providing other insurance payment information, even when no additional payment is expected from TMHP, provides benefit to all parties involved in Texas Medicaid. When a TPR issues a payment or partial payment to a provider, the other insurance credit must be indicated on the claim form submitted to TMHP.

This procedure benefits both providers and TMHP even if the TPR payment exceeds the Medicaid allowed amount. Although additional payment may not be issued by TMHP, informing TMHP of the other insurance credit allows TMHP to track the appropriate use of TPRs. Informing TMHP of a TPR
credit provides hospitals with a more accurate standard dollar amount (SDA) rate setting and assists the program in tracking recoveries and reducing Medicaid medical expenditures by informing TMHP of liable third parties.

Providers must report TPR payments correctly in the appropriate electronic field or the paper claim form block as follows:

<table>
<thead>
<tr>
<th>Claim Form</th>
<th>Reference</th>
</tr>
</thead>
<tbody>
<tr>
<td>CMS-1500</td>
<td>Block 29, CMS-1500 Blank Claim Form</td>
</tr>
<tr>
<td>UB-04 CMS-1450</td>
<td>Block 54, UB-04 CMS-1450 Blank Claim Form</td>
</tr>
<tr>
<td>THSteps Dental</td>
<td>Block 31, ADA Dental Claim Form</td>
</tr>
</tbody>
</table>

Refer to: “Section 6: Claims Filing” (Vol. 1, General Information) for claim filing instructions.

8.8.2.1 Deductibles

TMHP will consider deductibles for reimbursement when the original third party payor applied the payment amount directly to the client’s deductible. The explanation of benefit reflecting the application of the payment by the other insurance (third party payor) and a completed signed claim copy must be submitted to TMHP for consideration.

8.8.2.2 HMO Copayments

TMHP processes and pays HMO copayments for private and Medicare HMOs as well as private and Medicare preferred provider organization (PPO) copayments for clients who are eligible for reimbursement under Medicaid guidelines.

TMHP pays the copayment in addition to the service the HMO or PPO has denied, if the client is eligible for Texas Medicaid and the procedure is reimbursed under Medicaid guidelines. Providers are not allowed to hold the client liable for the copayment.

An office or emergency room (ER) visit (the ER physician is paid only when the ER is not staffed by the hospital) is reimbursed a maximum copayment of $10 per visit. The hospital ER visit is reimbursed at a maximum of $50 to the facility. TMHP pays up to four copayments per day, per client. ER visits are limited to one per day, per client, and are considered one of the four copayments allowed per day.

Important: By accepting assignment on a claim for which the client has Medicaid coverage, providers agree to accept payment made by insurance carriers and Texas Medicaid when appropriate as payment in full. The client cannot be held liable for any balance related to Medicaid-covered services.

The following Medicaid codes have been created for copayments, which are considered an atypical service:

<table>
<thead>
<tr>
<th>POS 1 - Office</th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td>CP001</td>
<td>Private HMO copayment-professional</td>
</tr>
<tr>
<td>CP002</td>
<td>Private PPO copayment-professional</td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th>POS 5 - Outpatient</th>
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<tr>
<td>CP005</td>
<td>Private HMO copayment-outpatient</td>
</tr>
<tr>
<td>CP006</td>
<td>Private PPO copayment-outpatient</td>
</tr>
</tbody>
</table>

Important: Providers must submit a new claim when filing for procedure codes CP001 and CP002. No explanation of benefits (EOB) or any other accompanying documentation is required to be attached to the claim form when filing for these services.
8.8.2.3 PPO Discounts

PPO discounts are not considered a part of other insurance payments. Electronic submitters must supply the PPO discount amount when submitting other insurance information; however, this information is not included in the total other insurance payment during claims processing. Paper submitters are not required to add the PPO discount to the other insurance payment.

8.8.2.4 Verbal Denial

Providers may call the other insurance resource and receive a verbal denial. The other insurance record can either be updated when the provider files the claim or calls the TPL/Tort Customer Service line at 1-800-846-7307. When calling the TPL/Tort Customer Service line and when filing claims to TMHP, the provider must have the following information before any updates are made.

Verbal denial requirements:
- Date of the telephone call to the other insurance resource
- Insurance company’s name and telephone number
- Name of the individual contacted at the insurance company
- Policyholder and group information for the client
- Specific reason for the denial, including the client’s type of coverage to enhance the accuracy of future claims processing (for example, a policy that covers inpatient services or physician services only)

Providers that update a client’s insurance records through the TMHP TPL/Tort Customer Service line must follow the current appeal process once the other insurance information has been updated on the client’s file.

8.8.2.5 110-Day Rule

When a service is billed to a third party and no response has been received, Medicaid providers must allow 110 days to elapse before submitting a claim to TMHP. If a TPR has not responded or delays payment or denial of a provider’s claim for more than 110 days after the date the claim was billed, Medicaid considers the claim for reimbursement. However, the 365-day federal filing deadline requirement must still be met. The following information is required:
- Name and address of the TPR
- Date the TPR was billed
- Statement signed and dated by the provider that no disposition has been received from the TPR within 110 days of the date the claim was billed

When TMHP denies a claim because of the client’s other coverage, information that identifies the other insurance appears on the provider’s Remittance and Status (R&S) Report. The claim is not to be refiled with TMHP until disposition from the TPR has been received or until 110 days have lapsed since the billing of the claim with no disposition from the TPR. A statement from the client or family member which indicates that they no longer have this resource is not sufficient documentation to reprocess the claim.

When a provider is advised by a TPR that benefits have been paid to the client, the information must be included on the claim with the date and amount of payment made to the client if available. If a denial was sent to the client, refer to the verbal denial guidelines above for required information. This enables TMHP to consider the claim for reimbursement.
8.8.2.6 Filing Deadlines
In accordance with federal regulations, all claims must initially be filed with TMHP within 365 days of the date of service (DOS). Claims that involve filing to a TPR have the following deadlines:

- Claims with a valid disposition (payment or denial) must be received by TMHP within 95 days of the date of disposition by the TPR and within 365 days of the DOS. Appealed claims that were originally denied with EOB 00260, which indicates that the provider files with a TPR, must be received within 95 days of the date of disposition by the TPR or within 120 days of the date on which TMHP denied the claim.
- The provider must appeal the claim to TMHP with complete other insurance information, which includes all EOBs and disposition dates. The disposition date is the date on which the other insurance company processed the payment or denial.
- If a provider submits other insurance EOBs without disposition dates, the appeal will be denied. If the other insurance disposition date appears only on the first page of an EOB that has multiple pages and the claim that is being submitted to TMHP is on a subsequent page or pages, the provider must submit the first page that shows the disposition date and all of the pages that show the claim that is being submitted to TMHP.
- If more than 110 days have passed from the date a claim was filed to the TPR without a response, the claim is submitted to TMHP for consideration of payment.

Refer to: Subsection 6.12.2.5, “Filing Deadlines” in “Section 6: Claims Filing” (Vol. 1, General Information) for information about filing deadlines for clients with other insurance.

8.9 Other Insurance Appeals
To appeal a claim denial due to other insurance coverage, the provider must submit complete other insurance information including the disposition date. The disposition date indicates when the other insurance company processed the payment or denial. An appeal submitted without this information will be denied.

If submitting a paper appeal the provider must submit EOBs containing disposition dates. If the disposition date appears only on the first page of an EOB that has multiple pages and the claim that is being appealed is on a subsequent page, the provider must also include the first page of the EOB that shows the disposition date.

Note: Claims denied for TPL/other insurance cannot be appealed through the TMHP Automated Inquiry System (AIS).

8.10 Refunds Resulting from Other Insurance Payments
The TMHP Cash Reimbursement Unit is responsible for processing financial adjustments when overlapping payments by Medicaid and a TPR occur.

Providers can use the Texas Medicaid Refund Information Form to:

- Refund the overpayment by issuing a check to TMHP. Providers must submit the refund check to TMHP along with the Texas Medicaid Refund Information Form and all required information requested on the form.
- Request that the claim be reprocessed and the money recouped. The overpayment will be reduced from next weekly payment made after claims are processed.

Refer to: The Texas Medicaid Refund Information Form, which is available in the Forms section of the TMHP website at www.tmhp.com.

If the amount paid by the other insurance carrier is less than Medicaid’s allowed amount, providers may bill TMHP for the difference. All claims must meet all timely filing deadlines.
Providers are prohibited from receiving payment from Medicaid and billing a TPR without refunding the Medicaid payment.

If within 12 months of the date of service a provider identifies that the client has other insurance and wants to submit a claim for payment to the other insurance company, the provider must refund any amounts previously paid by TMHP before submitting the claim to the other insurance.

If other insurance paid for the services submitted on the claim, the provider must submit the following to TMHP:
- The exact amount paid.
- The insurance company’s name and address.
- The client’s policy number and group number.

Providers are limited to the Medicaid allowed amount for the services. Providers are required to accept the TMHP paid amount as payment in full. If the provider fails to refund a payment to TMHP before submitting a claim to the other insurance, TMHP will recoup the entire other insurance payment.

In accordance with 1 TAC §§354.2321 [g] and 354.2322 [i], providers that do not follow TPR rules “may be referred for investigation and prosecution for violations of state or federal Medicaid or false claims laws.” Providers should refer to the full text of these rules for a full description of payment requirements.

### 8.11 Contact Information

TPL/Tort Telephone and Fax Communication

<table>
<thead>
<tr>
<th>Contact</th>
<th>General Inquiry Telephone number</th>
<th>Fax Number</th>
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</thead>
<tbody>
<tr>
<td>Third Party Resources (TPR)</td>
<td>1-800-846-7307</td>
<td>1-512-514-4225</td>
</tr>
<tr>
<td>Tort</td>
<td>1-800-846-7307</td>
<td>1-512-514-4225</td>
</tr>
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</table>

Written Communication with TMHP

<table>
<thead>
<tr>
<th>Contact</th>
<th>Address</th>
</tr>
</thead>
<tbody>
<tr>
<td>TPL/Tort</td>
<td>Texas Medicaid &amp; Healthcare Partnership Third Party Liability/Tort</td>
</tr>
<tr>
<td></td>
<td>PO Box 202948</td>
</tr>
<tr>
<td></td>
<td>Austin, TX 78720-2948</td>
</tr>
<tr>
<td>HHSC - Third Party Liability (TPL)</td>
<td>Health and Human Services Commission Attn: Third Party Liability</td>
</tr>
<tr>
<td></td>
<td>4900 N. Lamar Blvd.</td>
</tr>
<tr>
<td></td>
<td>Mail Code: 1354</td>
</tr>
<tr>
<td></td>
<td>Austin, TX 78751</td>
</tr>
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</table>

### 8.12 Forms

<table>
<thead>
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<th>Forms</th>
</tr>
</thead>
<tbody>
<tr>
<td>Your Texas Benefits Medicaid card (English and Spanish)</td>
</tr>
<tr>
<td>Informational Claims Submission Form</td>
</tr>
<tr>
<td>Other Insurance Form</td>
</tr>
<tr>
<td>Authorization for Use and Release of Health Information</td>
</tr>
<tr>
<td>Authorization for Use and Release of Health Information (Spanish)</td>
</tr>
<tr>
<td>Tort Response Form</td>
</tr>
<tr>
<td>Forms</td>
</tr>
<tr>
<td>--------------------------------------------</td>
</tr>
<tr>
<td>Texas Medicaid Refund Information Form</td>
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# APPENDIX A: STATE, FEDERAL, AND TMHP CONTACT INFORMATION

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<td>Texas Health and Human Services Commission Blind Children’s Vocational Discovery and Development Program (BCVDDP)</td>
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<td>TMHP Telephone and Fax Communication</td>
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<td>Written Communication With TMHP</td>
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<td>A.12</td>
<td>Other TMHP Information</td>
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<td>TMHP Website</td>
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<td>A.12.4</td>
<td>TMHP Provider Relations</td>
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<tr>
<td>A.12.5</td>
<td>Reporting a Complaint to TMHP</td>
</tr>
<tr>
<td>A.12.6</td>
<td>TMHP Electronic Data Interchange (EDI) Help Desk</td>
</tr>
</tbody>
</table>
A.1 Texas Health and Human Services Commission (HHSC) and Texas Department of State Health Services (DSHS) Office Addresses

Use the following address for general inquiries or for any group that is not listed in the following table:

Texas Health and Human Services Commission
PO Box 13247
Austin, TX 78711-3247

*Note:* Remember to use the four-digit addition to the ZIP Code.

Use the following address for the HHSC Inspector General:

Texas Health and Human Services Commission
Office of Inspector General
PO Box 85200
Austin, TX 78708-5200

*Note:* Remember to use the four-digit addition to the ZIP Code.

For the following groups, use the corresponding address and include the group name on the second line of the address.

<table>
<thead>
<tr>
<th>Address</th>
<th>Group Name</th>
</tr>
</thead>
<tbody>
<tr>
<td>HHSC</td>
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</tr>
<tr>
<td>Medicaid CHIP–H-200</td>
<td></td>
</tr>
<tr>
<td>PO Box 85200</td>
<td></td>
</tr>
<tr>
<td>Austin, TX 78708</td>
<td></td>
</tr>
<tr>
<td>HHSC</td>
<td></td>
</tr>
<tr>
<td>Quality Review/Limited Program—1323</td>
<td></td>
</tr>
<tr>
<td>PO Box 85200</td>
<td></td>
</tr>
<tr>
<td>Austin, TX 78708</td>
<td></td>
</tr>
<tr>
<td>HHSC</td>
<td></td>
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<tr>
<td>Third Party Liability (TPL)</td>
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</tr>
<tr>
<td>PO Box 85200</td>
<td></td>
</tr>
<tr>
<td>Mail Code 1354</td>
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</tr>
<tr>
<td>Austin, TX 78708-5200</td>
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<tr>
<td>HHSC</td>
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<td>Medical &amp; UR Appeals</td>
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</tr>
<tr>
<td>Mail Code H-230</td>
<td></td>
</tr>
<tr>
<td>PO Box 85200</td>
<td></td>
</tr>
<tr>
<td>Austin, TX 78708</td>
<td></td>
</tr>
<tr>
<td>or</td>
<td></td>
</tr>
<tr>
<td>4900 North Lamar (Express Mail Only)</td>
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</tr>
<tr>
<td>Austin, TX 78751</td>
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<tr>
<td>HHSC</td>
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<tr>
<td>Medicaid Vendor Drug H-630</td>
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<tr>
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<tr>
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<td>HHSC</td>
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<tr>
<td>Mail Code 0224</td>
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</tr>
<tr>
<td>Austin, TX 78714-9347</td>
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<tr>
<td>Region</td>
<td>Regional Director</td>
</tr>
<tr>
<td>--------</td>
<td>------------------</td>
</tr>
<tr>
<td>01</td>
<td>Beth Miller</td>
</tr>
<tr>
<td></td>
<td>6302 Iola Avenue</td>
</tr>
<tr>
<td></td>
<td>Lubbock, TX 79424</td>
</tr>
<tr>
<td></td>
<td>Amarillo: 1-806-356-3151</td>
</tr>
<tr>
<td>02/09</td>
<td>Sandra McKinney</td>
</tr>
<tr>
<td>Abilene</td>
<td>4601 South First Street</td>
</tr>
<tr>
<td></td>
<td>Abilene, TX 79604</td>
</tr>
<tr>
<td></td>
<td>PO Box 521</td>
</tr>
<tr>
<td></td>
<td>Abilene, TX 79604</td>
</tr>
<tr>
<td>03</td>
<td>Tracy Hays</td>
</tr>
<tr>
<td>Grand Prairie</td>
<td>801 South State Hwy 161</td>
</tr>
<tr>
<td></td>
<td>Grand Prairie, TX 75051</td>
</tr>
<tr>
<td></td>
<td>PO Box 532089</td>
</tr>
<tr>
<td></td>
<td>Grand Prairie, TX 75053-2089</td>
</tr>
<tr>
<td>04</td>
<td>Fay Booker</td>
</tr>
<tr>
<td>Tyler</td>
<td>302 East Rieck Road</td>
</tr>
<tr>
<td></td>
<td>Tyler, TX 75703</td>
</tr>
<tr>
<td></td>
<td>Toll Free: 211</td>
</tr>
<tr>
<td>05</td>
<td>Stephanie Semien</td>
</tr>
<tr>
<td>Beaumont</td>
<td>350 Pine Street, 9th Floor</td>
</tr>
<tr>
<td></td>
<td>Beaumont, TX 77701</td>
</tr>
<tr>
<td></td>
<td>Toll Free: 211</td>
</tr>
<tr>
<td>06</td>
<td>Gracie Perez</td>
</tr>
<tr>
<td>Houston</td>
<td>5425 Polk Street, Suite 230</td>
</tr>
<tr>
<td></td>
<td>Houston, TX 77023</td>
</tr>
<tr>
<td></td>
<td>PO Box 16017</td>
</tr>
<tr>
<td></td>
<td>Houston, TX 77222-6017</td>
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### A.2 HHSC Access and Eligibility Services (AES)

<table>
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<th>Address</th>
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<tbody>
<tr>
<td>DSHS</td>
<td>Children with Special Health Care Needs (CSHCN) Services Program (Mail Code 1938)</td>
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<tr>
<td>PO Box 149347</td>
<td>Genetic Services (1918)</td>
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<tr>
<td>Austin, TX 78714-9347</td>
<td>Indigent Health Care (Mail Code 2831)</td>
</tr>
<tr>
<td></td>
<td>Texas Health Steps (THSteps) (Mail Code 1938)</td>
</tr>
<tr>
<td></td>
<td>Case Management for Children and Pregnant Women (Mail Code 1938)</td>
</tr>
<tr>
<td>Region</td>
<td>Regional Director</td>
</tr>
<tr>
<td>--------</td>
<td>------------------</td>
</tr>
<tr>
<td>07 Austin</td>
<td>Sandra Dillett</td>
</tr>
<tr>
<td></td>
<td></td>
</tr>
<tr>
<td></td>
<td></td>
</tr>
<tr>
<td>08 San Antonio</td>
<td>Teriann Lyons</td>
</tr>
<tr>
<td></td>
<td></td>
</tr>
<tr>
<td></td>
<td></td>
</tr>
<tr>
<td></td>
<td></td>
</tr>
<tr>
<td>10 El Paso</td>
<td>Kate Hill</td>
</tr>
<tr>
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</tr>
<tr>
<td></td>
<td></td>
</tr>
<tr>
<td>11 Edinburg</td>
<td>Cynthia Pena</td>
</tr>
<tr>
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<tr>
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</tr>
<tr>
<td>ART &amp; Customer Care Center</td>
<td>Debra Gault</td>
</tr>
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<td></td>
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<tr>
<td>MEPD</td>
<td>Rachel Patton</td>
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For additional office information, visit the HHSC website at hhs.texas.gov.

Refer to: subsection A.5, “DSHS Health Service Regions Map” in this section to identify the regional boundaries.

### A.2.1 Telephone Communication with HHSC and DSHS

<table>
<thead>
<tr>
<th>Contact</th>
<th>Telephone Number</th>
</tr>
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<tbody>
<tr>
<td>Assessment Utilization Services (limited program) (Option 4)</td>
<td>1-800-436-6184</td>
</tr>
<tr>
<td>HHSC Hearing Services for Children (HSC) (hearing aid, evaluations)</td>
<td>1-800-925-9126</td>
</tr>
<tr>
<td>DSHS Emergency Medical Services Division</td>
<td>1-512-834-6700</td>
</tr>
<tr>
<td>DSHS ImmTrac2 Help Desk</td>
<td>1-800-348-9158</td>
</tr>
<tr>
<td>DSHS Immunization Branch</td>
<td>1-800-252-9152</td>
</tr>
<tr>
<td>Medical Transportation Program (MTP) Hotline</td>
<td>1-877-633-8747</td>
</tr>
<tr>
<td>THSteps/EPSDT Hotline</td>
<td>1-877-847-8377</td>
</tr>
<tr>
<td>Medicaid Vendor Drug Program Pharmacy Provider Resolution Helpdesk (fee-for-service)</td>
<td>1-800-435-4165</td>
</tr>
</tbody>
</table>

### A.3 Client Telephone Communication with HHSC

Clients can call the client toll-free number at 1-800-252-8263.
### A.4 Federal and State Telephone Numbers

<table>
<thead>
<tr>
<th>Telephone Number</th>
<th>Department/Program</th>
</tr>
</thead>
<tbody>
<tr>
<td>1-800-CDC-INFO</td>
<td>AIDS Hotline (Nationwide, distributed by Centers for Disease Control and Prevention [CDC], Atlanta, Georgia)</td>
</tr>
<tr>
<td>1-800-299-2437</td>
<td>HIV/STD InfoLine</td>
</tr>
<tr>
<td>1-800-255-1090</td>
<td>Texas HIV Medication Program</td>
</tr>
<tr>
<td>1-800-252-5400</td>
<td>Child/Elder Abuse Intake (Department of Family and Protective Services [DFPS])</td>
</tr>
<tr>
<td>1-512-776-7420</td>
<td>Vision and Hearing Screening Program (DSHS)</td>
</tr>
<tr>
<td>1-512-834-6650, Ext. 2601</td>
<td>Clinical Laboratory Improvement Amendments (CLIA) Certification Line</td>
</tr>
<tr>
<td>1-800-458-9858</td>
<td>Client Abuse Hotline for Long Term Care Services and Support—Nursing Facilities (HHSC)</td>
</tr>
<tr>
<td>1-800-252-8263</td>
<td>Client Inquiry Hotline (HHSC) (Medicaid questions from clients with Medicaid only)</td>
</tr>
<tr>
<td>1-512-776-7745</td>
<td>THSteps Program (DSHS)</td>
</tr>
<tr>
<td>1-888-963-7111 ext. 7318 or 1-512-776-7318</td>
<td>THSteps Laboratory Services (DSHS)</td>
</tr>
<tr>
<td>Fax: 1-512-776-7294</td>
<td></td>
</tr>
<tr>
<td>1-888-963-7111 ext. 7661 or 1-512-776-7661</td>
<td>Laboratory Supply Orders (DSHS)</td>
</tr>
<tr>
<td>Fax: 1-512-776-7672</td>
<td></td>
</tr>
<tr>
<td>1-888-963-7111 ext. 7578 or 1-512-776-7578</td>
<td>Report of Laboratory Test Results (DSHS)</td>
</tr>
<tr>
<td>1-512-776-7796</td>
<td>HHSC Family Planning</td>
</tr>
<tr>
<td>1-800-436-6184</td>
<td>Fraud or Abuse of Provider Services (HHSC Office of Inspector General)</td>
</tr>
<tr>
<td>1-800-436-6184</td>
<td>Fraud or Abuse/Long Term Care Services and Support—Nursing Facilities/HHSC</td>
</tr>
<tr>
<td>1-800-436-6184</td>
<td>Fraud or Abuse/Client/HHSC Office of the Inspector General</td>
</tr>
<tr>
<td>1-800-792-1109</td>
<td>Goal-Directed Therapy</td>
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<tr>
<td>1-512-438-3169 or 1-800-252-8010</td>
<td>Hospice Program (HHSC Policy Development division)</td>
</tr>
<tr>
<td>1-800-252-9152</td>
<td>Immunization Branch (DSHS)</td>
</tr>
<tr>
<td>1-800-925-9126</td>
<td>Medically Needy Spend Down Unit</td>
</tr>
<tr>
<td>1-800-MEDICARE or 1-800-633-4227</td>
<td>Medicare/Social Security Administration</td>
</tr>
<tr>
<td>1-800-925-9126</td>
<td>Newborn Screening (DSHS)</td>
</tr>
<tr>
<td>1-800-925-9126</td>
<td>HHSC Hearing Services for Children (HSC)</td>
</tr>
<tr>
<td>1-877-787-8999</td>
<td>HHS Office of the Ombudsman (for information about ECI or to refer a child.)</td>
</tr>
<tr>
<td>1-800-436-6184</td>
<td>Recipient Utilization Control Unit (HHSC) (for limited status review and for referrals from providers for potential client overutilization, etc.)</td>
</tr>
<tr>
<td>1-512-776-7796</td>
<td>Breast and Cervical Cancer Services (DSHS)</td>
</tr>
<tr>
<td>1-713-526-2559</td>
<td>Snellen Letter (Tumbling E Wall Chart)</td>
</tr>
<tr>
<td>1-800-435-4165</td>
<td>Medicaid Vendor Drug Program (HHSC) (fee-for-service) (specifically for pharmacy use)</td>
</tr>
<tr>
<td>1-877-728-3927</td>
<td>Medicaid Vendor Drugs Prior Authorization Center (fee-for-service)</td>
</tr>
<tr>
<td>1-866-993-9972</td>
<td>Healthy Texas Women (HTW) program Eligibility</td>
</tr>
<tr>
<td>1-512-776-7796</td>
<td>Cervical Cancer Screening</td>
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A.5 DSHS Health Service Regions Map

Local and Regional Public Health Coverage

A.6 DSHS Health Service Region Contacts

<table>
<thead>
<tr>
<th>Health Service Region 1 Regional Office (Lubbock)</th>
<th>Health Service Regions 2 &amp; 3 Regional Office (Arlington)</th>
</tr>
</thead>
<tbody>
<tr>
<td>DSHS/PHR 1</td>
<td>DSHS/PHR 2 &amp; 3</td>
</tr>
<tr>
<td>6302 Iola Ave.</td>
<td>1301 S. Bowen, Suite 200</td>
</tr>
<tr>
<td>Lubbock, TX 79424</td>
<td>Arlington, TX 76013</td>
</tr>
<tr>
<td>1-806-744-3577</td>
<td>1-817-264-4500</td>
</tr>
<tr>
<td>Fax: 1-806-783-6435</td>
<td>Fax: 1-817-264-4506</td>
</tr>
<tr>
<td>Regional Medical Director</td>
<td>Regional Medical Director</td>
</tr>
<tr>
<td>Peter W. Pendergrass, MD, MPH</td>
<td>James A. Zoretic, MD</td>
</tr>
<tr>
<td>Deputy Regional Director</td>
<td>Deputy Regional Director (acting)</td>
</tr>
<tr>
<td>Barry Wilson</td>
<td>Earlene Quinn</td>
</tr>
<tr>
<td>Manager of Social Work Services and Case Management</td>
<td>Manager of Social Work Services and Case Management</td>
</tr>
<tr>
<td>Judy Lara, LBSW</td>
<td>Lindsay Matousek, LMSW, CCM</td>
</tr>
<tr>
<td>Communicable Disease Manager</td>
<td>Director of Clinic Operations</td>
</tr>
<tr>
<td>Vacant</td>
<td>Dorothy Kuhlmann, RN</td>
</tr>
<tr>
<td>Immunization Program Manager</td>
<td>Immunization Program Manager</td>
</tr>
<tr>
<td>Keila Johnson</td>
<td>Sonna Sanders</td>
</tr>
</tbody>
</table>
## Health Service Region 1
### Regional Office (Lubbock)
- **Tuberculosis Team Leader**
  - Melanie Lee
- **THSteps Operations Lead**
  - Elizabeth Stanford
  - 6302 Iola Ave
  - Lubbock, TX 79424
  - Mail Code: 1899
  - 1-806-783-6445
  - Fax: 1-806-783-6455

## Health Service Region 2 & 3
### Regional Office (Arlington)
- **Communicable Disease Program Manager**
  - Gary Willett
- **Tuberculosis Team Leader**
  - Jeff Ralston
- **Emergency Preparedness**
  - Bryan Flow, DVM

## Health Service Regions 4 & 5 (North)
### Regional Office (Tyler)
- **Regional Office (Tyler)**
  - DSHS/PHR 4 & 5 North
  - 2521 West Front Street
  - Tyler, TX 75702
  - 1-903-595-3585
  - Fax: 1-903-593-4187
- **Regional Medical Director**
  - Dr. Paul K. McGaha, DO, MPH
- **Deputy Regional Director**
  - Vacant
- **Manager of Social Work Services and Case Management**
  - Caleb Rackley, LMSW
- **Director of Nursing**
  - Barbara Lay, RN, MSN
- **Immunization Program Manager**
  - Toni Wright
- **HIV/STD Program Manager**
  - Charles O’Brien
- **Tuberculosis Program Manager**
  - Teresa Santiago, RN
- **THSteps Operations Lead**
  - Sherrylinn Adams, LBSW
  - 2521 W. Front, Mail Code 1358
  - Tyler, TX 75702
  - 1-903-533-5357
  - Fax: 1-903-595-4706

## Health Service Regions 6 & 5 (South)
### Regional Office (Houston)
- **Regional Office (Houston)**
  - DSHS 6 & 5 South
  - 5425 Polk Avenue, Suite J
  - Houston, TX 77023
  - 1-713-767-3000
  - Fax: 1-713-767-3049
- **Regional Medical Director**
  - Regional Medical Director (Acting)
  - John G. Jordan, M.D., MPH
- **Deputy Regional Director**
  - Greta Etnyre, MS, RD
- **Manager of Social Work Services and Case Management**
  - Vacant
- **Director of Nursing**
  - Melinda Denson, RN, MPH
- **Immunization Program Manager**
  - Angel H. Angco, MBA, RN
- **HIV/STD Program Manager**
  - Linda Hollins
- **Tuberculosis Program Manager**
  - Lewis Gonzalez, MD
- **THSteps Operations Lead**
  - Shannon Jones
  - 5425 Polk Avenue, Suite J, Mail Code 1906
  - Houston, TX 77023-1497
  - 1-713-767-3105
  - Fax: 1-713-767-3125
**APPENDIX A: STATE, FEDERAL, AND TMHP CONTACT INFORMATION MAY 2021**

<table>
<thead>
<tr>
<th><em>Health Service Region 7</em> Regional Office (Temple)</th>
<th>Health Service Region 8 Regional Office (San Antonio)</th>
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<tbody>
<tr>
<td>DSHS/PHR 7</td>
<td>DSHS/PHR 8</td>
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<tr>
<td>2408 S 37th Street</td>
<td>7430 Louis Pasteur Drive</td>
</tr>
<tr>
<td>Temple, TX 76504-7168</td>
<td>San Antonio, TX 78229</td>
</tr>
<tr>
<td>1-254-778-6744</td>
<td>1-210-949-2000</td>
</tr>
<tr>
<td>Fax: 1-254-778-4066</td>
<td>Fax: 1-210-949-2015</td>
</tr>
<tr>
<td>Regional Medical Director</td>
<td>Regional Medical Director</td>
</tr>
<tr>
<td>Lisa Cornelius, MD, MPH</td>
<td>Sandra Guerra-Cantu, MD, MPH</td>
</tr>
<tr>
<td>Deputy Regional Director</td>
<td>Deputy Regional Director</td>
</tr>
<tr>
<td>Jon Huss</td>
<td>Gail Morrow, MPH</td>
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<tr>
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<td>Manager of Social Work Services and Case Management</td>
</tr>
<tr>
<td>Leesa Ferrero, LMSW</td>
<td>Leticia Guerra, LBSW</td>
</tr>
<tr>
<td>Director of Nursing</td>
<td>Director of Nursing</td>
</tr>
<tr>
<td>Pauline Culbert, MSN, RN</td>
<td>Sandra Jones, MSN, CNS</td>
</tr>
<tr>
<td>Immunization Program Manager</td>
<td>Immunization Program Manager</td>
</tr>
<tr>
<td>Diane Romnes</td>
<td>Laurie Henefery</td>
</tr>
<tr>
<td>HIV/STD Program Manager</td>
<td>Communicable Disease Program Manager</td>
</tr>
<tr>
<td>Al Gonzales</td>
<td>Cherise Rohr-Allegrini, PhD</td>
</tr>
<tr>
<td>Tuberculosis Program Manager/Nurse Consultant</td>
<td>HIV/STD Program Manager</td>
</tr>
<tr>
<td>Dana Schoepf, RN</td>
<td>Joanna Nichols, MPH</td>
</tr>
<tr>
<td>THSteps Operations Co-Leads</td>
<td>THSteps Operations Lead</td>
</tr>
<tr>
<td>Kimberly Langley</td>
<td>Velma Stille</td>
</tr>
<tr>
<td>1-254-760-1176</td>
<td>7430 Louis Pasteur Drive, Mail Code 5716</td>
</tr>
<tr>
<td>2408 S. 37th Street, Mail Code 1902</td>
<td>San Antonio, TX 78229</td>
</tr>
<tr>
<td>Temple, TX 76504</td>
<td>1-210-949-2159</td>
</tr>
<tr>
<td>Fax: 1-254-778-5490</td>
<td>Fax: 1-210-949-2041</td>
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<table>
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<tr>
<th>Health Service Regions 9 &amp; 10 Regional Office (El Paso)</th>
<th>Health Service Region 11 Regional Office (Harlingen)</th>
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<tr>
<td>DSHS/PHR 9 &amp; 10</td>
<td>DSHS/PHR 11</td>
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<tr>
<td>401 E. Franklin, Suite 210</td>
<td>601 W. Sesame Drive</td>
</tr>
<tr>
<td>El Paso, TX 79901</td>
<td>Harlingen, TX 78550</td>
</tr>
<tr>
<td>1-915-834-7682</td>
<td>1-956-423-0130</td>
</tr>
<tr>
<td>Fax: 1-915-834-7808</td>
<td>Fax: 1-956-444-3293</td>
</tr>
<tr>
<td>Regional Medical Director</td>
<td>Regional Medical Director</td>
</tr>
<tr>
<td>James A. Zoretic, M.D., M.P.H., Interim Regional Medical Director</td>
<td>Brian Smith, MD, MPH</td>
</tr>
<tr>
<td>Deputy Regional Director</td>
<td>Deputy Regional Director</td>
</tr>
<tr>
<td>Blanca Serrano, MPH, RS</td>
<td>Sylvia Garces-Hobs</td>
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<tr>
<td>Manager of Social Work Services and Case Management</td>
<td>Manager of Social Work Services and Case Management</td>
</tr>
<tr>
<td>Patrice Loge, LMSW</td>
<td>Angelica Martinez, MS, LBSW</td>
</tr>
<tr>
<td>Director of Nursing</td>
<td>Director of Nursing</td>
</tr>
<tr>
<td>Sharon Lindsey, RN</td>
<td>Darlene Farias, RN</td>
</tr>
<tr>
<td>Immunization Program Manager</td>
<td>Immunization Program Manager</td>
</tr>
<tr>
<td>Jose Padilla</td>
<td>Ivette Nunez</td>
</tr>
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</table>
### A.7 State Participating Local Health Departments and Public Health Districts

<table>
<thead>
<tr>
<th>Health Service Regions 9 &amp; 10 Regional Office (El Paso)</th>
<th>Health Service Region 11 Regional Office (Harlingen)</th>
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<tbody>
<tr>
<td>HIV/STD Program Manager</td>
<td>HIV/STD Program Manager</td>
</tr>
<tr>
<td>Oscar Hernandez</td>
<td>Richard Anguiano</td>
</tr>
<tr>
<td>Communicable Disease Manager</td>
<td>Tuberculosis Program Manager</td>
</tr>
<tr>
<td>Gale Morrow, MPH, CHES</td>
<td>Maria San Pedro, MSPHN, RN</td>
</tr>
<tr>
<td>THSteps Operations Lead</td>
<td>THSteps Operations Lead</td>
</tr>
<tr>
<td>Patrice Loge, LMSW</td>
<td>Ray Garza</td>
</tr>
<tr>
<td>401 E. Franklin, Suite 200, Mail Code 1903</td>
<td>601 W. Sesame Drive, Mail Code 1907</td>
</tr>
<tr>
<td>El Paso, TX 79901</td>
<td>Harlingen, TX 78550 bou</td>
</tr>
<tr>
<td>1-915-834-7733</td>
<td>1-956-421-5563</td>
</tr>
<tr>
<td>Fax: 1-915-834-7802</td>
<td>Fax: 1-956-444-3293</td>
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<table>
<thead>
<tr>
<th>State Participating Local Health Departments and Public Health Districts</th>
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<tbody>
<tr>
<td>Abilene Public Health Department</td>
</tr>
<tr>
<td>Region 2/3</td>
</tr>
<tr>
<td>Larry Johnson, Administrator</td>
</tr>
<tr>
<td>PO Box 6489 (79608-6489)</td>
</tr>
<tr>
<td>850 N. 6th Street</td>
</tr>
<tr>
<td>Abilene, TX 79605</td>
</tr>
<tr>
<td>1-325-692-5600</td>
</tr>
<tr>
<td>Fax: 1-325-734-5370</td>
</tr>
<tr>
<td>Hidalgo County Health Department</td>
</tr>
<tr>
<td>Region 11</td>
</tr>
<tr>
<td>Eduardo Olivarez, Administrator</td>
</tr>
<tr>
<td>Omar Garza, MD, Director</td>
</tr>
<tr>
<td>1304 South 25th Street</td>
</tr>
<tr>
<td>Edinburg, TX 78539-7205</td>
</tr>
<tr>
<td>1-956-383-6221</td>
</tr>
<tr>
<td>Fax: 1-956-444-3298</td>
</tr>
<tr>
<td>Amarillo Bi-City-County Health District</td>
</tr>
<tr>
<td>Department of Health</td>
</tr>
<tr>
<td>Roger Smalligan, MD, Health Authority</td>
</tr>
<tr>
<td>Matt Richardson, Director for the City of Amarillo</td>
</tr>
<tr>
<td>Department of Health</td>
</tr>
<tr>
<td>1000 Martin Road</td>
</tr>
<tr>
<td>Amarillo, TX 79107</td>
</tr>
<tr>
<td>1-806-378-6300</td>
</tr>
<tr>
<td>Fax: 1-806-378-6306</td>
</tr>
<tr>
<td>Houston Health &amp; Human Services Department Region 6/5 S</td>
</tr>
<tr>
<td>Stephen L. Williams, MD, MPH, Director</td>
</tr>
<tr>
<td>8000 North Stadium Drive</td>
</tr>
<tr>
<td>Houston, TX 77054</td>
</tr>
<tr>
<td>1-713-794-9311</td>
</tr>
<tr>
<td>Fax: 1-713-798-0862</td>
</tr>
<tr>
<td>Andrews City-County Health Department</td>
</tr>
<tr>
<td>Region 9/10</td>
</tr>
<tr>
<td>Robert Garcia, MD, Director</td>
</tr>
<tr>
<td>208 NW 2nd street</td>
</tr>
<tr>
<td>Andrews, TX 79714</td>
</tr>
<tr>
<td>1-432-524-1434</td>
</tr>
<tr>
<td>Fax: 1-432-524-1461</td>
</tr>
<tr>
<td>Jackson County Health Department</td>
</tr>
<tr>
<td>Region 8</td>
</tr>
<tr>
<td>Bain C. Cate, MD, Director</td>
</tr>
<tr>
<td>411 North Wells, Room 206</td>
</tr>
<tr>
<td>Edna, TX 77957</td>
</tr>
<tr>
<td>1-361-782-5221</td>
</tr>
<tr>
<td>Fax: 1-361-782-7312</td>
</tr>
<tr>
<td>Angelina County &amp; Cities Health District</td>
</tr>
<tr>
<td>Region 4/5N</td>
</tr>
<tr>
<td>Sharon Shaw, Administrator</td>
</tr>
<tr>
<td>John Rudis, MD, Director</td>
</tr>
<tr>
<td>Lufkin, TX 75904</td>
</tr>
<tr>
<td>1-936-632-1139</td>
</tr>
<tr>
<td>Fax: 1-936-632-2640</td>
</tr>
<tr>
<td>Jasper-Newton County Public Health District Region 4/5 N</td>
</tr>
<tr>
<td>Danny Brackin</td>
</tr>
<tr>
<td>139 West Lamar Street</td>
</tr>
<tr>
<td>Jasper, TX 75951</td>
</tr>
<tr>
<td>1-409-384-6829</td>
</tr>
<tr>
<td>Fax: 1-409-384-7861</td>
</tr>
<tr>
<td>Atascosa County Health Department</td>
</tr>
<tr>
<td>Region 8</td>
</tr>
<tr>
<td>Gerald B. Phillips, MD, Director</td>
</tr>
<tr>
<td>1102 Campbell Avenue</td>
</tr>
<tr>
<td>Jourdanton, TX 78026</td>
</tr>
<tr>
<td>1-830-769-3451</td>
</tr>
<tr>
<td>Fax: 1-210-769-2349</td>
</tr>
<tr>
<td>Jefferson County Health Authority</td>
</tr>
<tr>
<td>Cecil A. Walkes, MD</td>
</tr>
<tr>
<td>1295 Pearl Street</td>
</tr>
<tr>
<td>Beaumont, TX 77701</td>
</tr>
<tr>
<td>1-409-835-8530</td>
</tr>
<tr>
<td>Fax: 1-409-839-2353</td>
</tr>
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</table>
## State Participating Local Health Departments and Public Health Districts

<table>
<thead>
<tr>
<th>Region</th>
<th>Department/Agency</th>
<th>State</th>
<th>County</th>
<th>Contact Information</th>
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<tbody>
<tr>
<td>7</td>
<td>Austin Department of Health &amp; Human Services</td>
<td>Liberty County Health Authority</td>
<td>Steven C. Ellerbe, DO</td>
<td>720 Travis, Liberty, TX 77575, 1-936-336-6439, Fax: 1-936-336-6517</td>
</tr>
<tr>
<td></td>
<td>David Lurie, Director</td>
<td></td>
<td></td>
<td>7201 Levander Loop, Bldg. E, Austin, TX 78744, Fax: 1-512-972-5016</td>
</tr>
<tr>
<td>7/6 S</td>
<td>Beaumont City Health Department</td>
<td>Live Oak County Health Department</td>
<td>Alan Crouther, Director</td>
<td>Drawer 670 (78022), Live Oak County Courthouse, George West, TX 78022, 1-361-449-2733, Fax: 1-361-449-1013</td>
</tr>
<tr>
<td></td>
<td>Region 6/5 S</td>
<td></td>
<td></td>
<td>Ingrid West-Holmes, Director</td>
</tr>
<tr>
<td>7</td>
<td>Bell County Public Health District</td>
<td>Lubbock City Health Department</td>
<td>Nancy Haney, Director</td>
<td>PO Box 2548 (79408), 1902 Texas Avenue, Lubbock, TX 79405, 1-806-775-2899, Fax: 1-806-775-3209</td>
</tr>
<tr>
<td></td>
<td>Region 7</td>
<td></td>
<td></td>
<td>Wayne Farrell, Director</td>
</tr>
<tr>
<td>6/5 S</td>
<td>Brazoria County Health Department</td>
<td>Maverick County Health Department</td>
<td>Arturo Batres, MD, Director</td>
<td>432 East Mulberry, Angleton, TX 77515, 1-979-864-1484, Fax: 1-979-756-1456</td>
</tr>
<tr>
<td></td>
<td>Region 2</td>
<td></td>
<td></td>
<td>Leo D. O’Gorman, MD, MPH, Director</td>
</tr>
<tr>
<td>7</td>
<td>Brazos County Health Department</td>
<td>Marshall-Harrison County Health District</td>
<td>Robert Palmer, MD, Director</td>
<td>201 North Texas Avenue, Bryan, TX 77803-5317, 1-979-361-4440, Fax: 1-409-823-6993</td>
</tr>
<tr>
<td></td>
<td>Region 8</td>
<td></td>
<td></td>
<td>Ken Bost, Executive Director</td>
</tr>
<tr>
<td>2/3</td>
<td>Brownwood-Brown County Health Department</td>
<td>Medina County Health Department</td>
<td>John W. Meyer, MD, Director</td>
<td>Russell Skinner, MD, Director</td>
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<tr>
<td></td>
<td>Region 4/5 N</td>
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<td></td>
<td>805 Lindsey Drive, Marshall, TX 75670, 1-903-938-8338, Fax: 1-903-938-8330</td>
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<tr>
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<td>201 North Texas Avenue, Bryan, TX 77803-5317, 1-979-361-4440, Fax: 1-409-823-6993</td>
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<td>8</td>
<td>Calhoun County Health Department</td>
<td>Midland County Health Department</td>
<td>Celestino Garcia, RS, Administrator</td>
<td>117 West Ash, Port Lavaca, TX 77979, 1-361-552-9721, Fax: 1-361-552-9722</td>
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<td>Region 8</td>
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<td>Bain C. Cate, Director</td>
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<tr>
<td>State Participating Local Health Departments and Public Health Districts</td>
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<tr>
<td>Region 11</td>
<td>Region 7</td>
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<tr>
<td>Yvette Salinas, Administrator</td>
<td>Patsy Gaines, Director</td>
<td></td>
<td></td>
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</tr>
<tr>
<td>1390 W. Ex. 83</td>
<td>PO Box 469 (76520)</td>
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<tr>
<td>San Benito, Tx. 78586</td>
<td>209 South Houston Street</td>
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<tr>
<td>1-956-247-3685</td>
<td>Cameron, TX 76520</td>
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<tr>
<td>Fax: 1-956-361-8280</td>
<td>1-254-697-7039</td>
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<td>Cass County Health Department</td>
<td>Montgomery County Health Department</td>
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</tr>
<tr>
<td>R. Bruce LeGrow, MD, Director</td>
<td>Vickie Modeland, Director</td>
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## State Participating Local Health Departments and Public Health Districts

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<tr>
<th>State Participating Health Department and Public Health District</th>
<th>Contact Information</th>
</tr>
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<tbody>
<tr>
<td>Ector County Health Department Region 9/10</td>
<td>Uvalde City-County Health Department Region 8</td>
</tr>
<tr>
<td>Gino Solla, Director</td>
<td>Liz Barrett, RN, Director</td>
</tr>
<tr>
<td>Nathan Galloway, MD, Director</td>
<td>1021 Garnerfield Road</td>
</tr>
<tr>
<td>221 North Texas</td>
<td>Uvalde, TX 78801</td>
</tr>
<tr>
<td>Odessa, TX 79761</td>
<td>1-830-278-1705</td>
</tr>
<tr>
<td>1-432-498-4141</td>
<td>Fax: 1-830-278-1881</td>
</tr>
<tr>
<td>Fax: 1-432-498-4143</td>
<td></td>
</tr>
<tr>
<td>El Paso City-County Health and Environmental District/</td>
<td>Victoria County Health Department Region 8</td>
</tr>
<tr>
<td>Region 9/10</td>
<td>Bain C. Cate, MD, Director</td>
</tr>
<tr>
<td>Jorge Magaña, MD, Director</td>
<td>PO Box 2350 (77902)</td>
</tr>
<tr>
<td>1148 Airway Blvd</td>
<td>2805 N. Navarro</td>
</tr>
<tr>
<td>El Paso, TX 79925-3692</td>
<td>Victoria, TX 77901</td>
</tr>
<tr>
<td>1-915-771-5701</td>
<td>1-361-578-6281</td>
</tr>
<tr>
<td>Fax: 1-915-543-3541</td>
<td>Fax: 1-361-578-7046</td>
</tr>
<tr>
<td>Fort Bend County Health Department Region 6/5 S</td>
<td>Waco-McLennan County Public Health District Region 7</td>
</tr>
<tr>
<td>Jean Galloway, MD, Director</td>
<td>Roger Barker, MBA, Director</td>
</tr>
<tr>
<td>PO Box 668 (77471)</td>
<td>225 West Waco Drive</td>
</tr>
<tr>
<td>4520 Reading Road, Suite A</td>
<td>Waco, TX 76707</td>
</tr>
<tr>
<td>Rosenberg, TX 77471</td>
<td>1-254-750-5450</td>
</tr>
<tr>
<td>1-281-342-6414</td>
<td>Fax: 1-254-750-5452</td>
</tr>
<tr>
<td>Fax: 1-281-342-7371</td>
<td></td>
</tr>
<tr>
<td>Galveston County Health District Region 6/5 S</td>
<td>Wichita Falls-Wichita County Public Health District Region 2/3</td>
</tr>
<tr>
<td>H. Mark Guidry, MD, MPH, Director</td>
<td>Lou Franklin, RN, BSN, Director</td>
</tr>
<tr>
<td>PO Box 939 (77568)</td>
<td>Tom Edmonson, Administrator</td>
</tr>
<tr>
<td>1207 Oak Street</td>
<td>1700 Third Street</td>
</tr>
<tr>
<td>La Marque, TX 77568</td>
<td>Wichita Falls, TX 76301</td>
</tr>
<tr>
<td>1-409-938-2401</td>
<td>1-940-761-7805</td>
</tr>
<tr>
<td>Fax: 1-409-938-2243</td>
<td>Fax: 1-940-767-5242</td>
</tr>
<tr>
<td><a href="mailto:rmorris@gchd.org">rmorris@gchd.org</a></td>
<td></td>
</tr>
<tr>
<td>Grayson County Health Department Region 2/3</td>
<td>Williamson County and Cities Public Health District Region 7</td>
</tr>
<tr>
<td>Wayne Bell</td>
<td>W.S. Riggins, Jr., MD, MPH, Director</td>
</tr>
<tr>
<td>515 North Walnut</td>
<td>PO Box 570 (78627)</td>
</tr>
<tr>
<td>Sherman, TX 75090</td>
<td>100 W. 3rd Street</td>
</tr>
<tr>
<td>1-903-893-0131</td>
<td>Georgetown, TX 78626</td>
</tr>
<tr>
<td>Fax: 1-903-892-3776</td>
<td>1-512-943-3600</td>
</tr>
<tr>
<td>Greenville-Hunt County Health Department Region 2/3</td>
<td>Fax: 1-512-943-1499</td>
</tr>
<tr>
<td>Mark Memahan, DO, Director</td>
<td>Wilson County Health Department Region 8</td>
</tr>
<tr>
<td>Henry Underwood, DO, Director</td>
<td>Edwin Baker, Director</td>
</tr>
<tr>
<td>4815 King Street, Ste. B</td>
<td>PO Box 396 (78114)</td>
</tr>
<tr>
<td>Greenville, TX 75401</td>
<td>Wilson County Courthouse</td>
</tr>
<tr>
<td>1-903-455-4433</td>
<td>Floresville, TX 78114</td>
</tr>
<tr>
<td>Fax: 1-903-455-4956</td>
<td>1-830-393-8503</td>
</tr>
<tr>
<td>Wilson County Health Department Region 8</td>
<td>Fax: 1-830-393-6031</td>
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## State Participating Local Health Departments and Public Health Districts

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<tr>
<td>Region 6/5 S</td>
<td>Region 4/5 N</td>
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<tr>
<td>Mary Pastor, Director</td>
<td>David C. Murley, MD, Director</td>
</tr>
<tr>
<td>PO Box 820 (77625)</td>
<td>Wood County Courthouse</td>
</tr>
<tr>
<td>440 W. Monroe</td>
<td>PO Box 596 (75783)</td>
</tr>
<tr>
<td>Kountze, TX 77625</td>
<td>Quitman, TX 75783</td>
</tr>
<tr>
<td>1-409-246-5188</td>
<td>1-903-763-5406</td>
</tr>
<tr>
<td>Fax: 1-409-246-4373</td>
<td>Fax: 1-903-763-2902</td>
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<th>Harris County Health Department</th>
<th>Zavala County Health Department</th>
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<tr>
<td>Region 6/5 S</td>
<td>Region 8</td>
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<tr>
<td>2223 W. Loop South</td>
<td>Antonio Rivera, MD, Director</td>
</tr>
<tr>
<td>Houston, TX 77027</td>
<td>600 North John F. Kennedy Drive</td>
</tr>
<tr>
<td>1-713-439-6016</td>
<td>Crystal City, TX 78839</td>
</tr>
<tr>
<td>Fax: 1-713-439-6080</td>
<td>1-210-374-3010</td>
</tr>
<tr>
<td><a href="mailto:thyslop@hc.co.harris.tx.us">thyslop@hc.co.harris.tx.us</a></td>
<td>Fax: 1-210-374-3007</td>
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<tr>
<td>Region N</td>
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<tr>
<td>Larry Birdwell, DO, Director</td>
<td></td>
</tr>
<tr>
<td>401-A Broadway Drive</td>
<td></td>
</tr>
<tr>
<td>San Marcos, TX 78666</td>
<td></td>
</tr>
<tr>
<td>1-512-353-4353</td>
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<tr>
<td>Fax: 1-512-396-4656</td>
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## A.8 Texas Health and Human Services Commission Blind Children’s Vocational Discovery and Development Program (BCVDDP)

### HHSC, BCVDDP

<table>
<thead>
<tr>
<th>Abilene</th>
<th>Irving</th>
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<tbody>
<tr>
<td>4601 S. 1st Street, Suite M</td>
<td>440 S. Nursery Rd.</td>
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<tr>
<td>Abilene, TX 79605-1463</td>
<td>MC: 1469</td>
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<tr>
<td>PO BOX 521</td>
<td>Irving, TX 75060</td>
</tr>
<tr>
<td>MC: 6846</td>
<td>1-972-721-6580</td>
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<tr>
<td>Abilene, TX 79604-0521</td>
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<tr>
<td>1-325-795-5845</td>
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<tr>
<th>Amarillo</th>
<th>Laredo</th>
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<tr>
<td>28 Western Plaza Dr.</td>
<td>1500 N. Arkansas</td>
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<tr>
<td>MC: 0052</td>
<td>MC: 2031</td>
</tr>
<tr>
<td>Amarillo, TX 79109</td>
<td>Laredo, TX 78043</td>
</tr>
<tr>
<td>1-806-351-3881</td>
<td>1-956-764-6284</td>
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<table>
<thead>
<tr>
<th>Austin</th>
<th>Lubbock</th>
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<tbody>
<tr>
<td>7701 Metropolis Dr., BLDG 12, Ste. 100</td>
<td>6302 Iola St.</td>
</tr>
<tr>
<td>MC: 0172</td>
<td>MC: 2171</td>
</tr>
<tr>
<td>Austin, TX 78744</td>
<td>Lubbock, TX 79424</td>
</tr>
<tr>
<td>1-512-416-5263</td>
<td>1-806-783-6630</td>
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<th>Beaumont</th>
<th>McAllen</th>
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<tr>
<td>3105 Executive Blvd.</td>
<td>4501 West Business 83</td>
</tr>
<tr>
<td>MC: 0291</td>
<td>MC: 2222</td>
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<tr>
<td>Beaumont, TX 77708</td>
<td>McAllen, TX 78501</td>
</tr>
<tr>
<td>1-409-730-1098</td>
<td>1-956-971-1281</td>
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<table>
<thead>
<tr>
<th>Bryan College Station</th>
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<tr>
<td>3000 East Villa Maria Rd</td>
<td>3016 Kermit Hwy.</td>
</tr>
<tr>
<td>MC: 7331</td>
<td>MC: 2503</td>
</tr>
<tr>
<td>Bryan, TX 77803</td>
<td>Odessa, TX 79764</td>
</tr>
<tr>
<td>1-979-776-7492</td>
<td>1-432-334-5624</td>
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### HHSC, BCVDDP

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<th>PA Request Telephone Number</th>
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<tbody>
<tr>
<td>Corpus Christi</td>
<td></td>
<td>San Angelo 622 South Oakes, Suite D 6979 76903 325-659-7653</td>
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<tr>
<td>Corpus Christi, TX 78415</td>
<td>361-878-7707</td>
<td></td>
<td></td>
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<tr>
<td>Dallas</td>
<td></td>
<td>San Antonio 11307 Roszell 2794 78217 210-619-8147</td>
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<tr>
<td>1545 W. Mockingbird Ln.</td>
<td>214-638-7575</td>
<td></td>
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<tr>
<td>Dallas, TX 75235</td>
<td></td>
<td></td>
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<tr>
<td>El Paso</td>
<td></td>
<td>Tyler 3303 Mineola Highway 3137 75702 903-595-4841</td>
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</tr>
<tr>
<td>401 E. Franklin, Suite #240</td>
<td>361-878-7707</td>
<td></td>
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</tr>
<tr>
<td>El Paso, TX 79901</td>
<td></td>
<td></td>
<td></td>
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<tr>
<td>Fort Worth</td>
<td></td>
<td>Victoria 2306 Leary Ln. 3192 77901 361-734-732</td>
<td></td>
</tr>
<tr>
<td>4733 E. Lancaster Ave.</td>
<td>817-535-3893</td>
<td></td>
<td></td>
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<tr>
<td>Fort Worth, TX 76103</td>
<td></td>
<td></td>
<td></td>
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<tr>
<td>Harlingen</td>
<td></td>
<td>Waco 801 Austin Street, Suite 710 6820 76701 254-750-9623</td>
<td></td>
</tr>
<tr>
<td>3625 W. Business 83</td>
<td>956-365-0129</td>
<td></td>
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<tr>
<td>Harlingen, TX 78552</td>
<td></td>
<td></td>
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<tr>
<td>Houston</td>
<td></td>
<td>Wichita Falls 1328 Oakhurst Dr. 3323 76302 940-720-8446</td>
<td></td>
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<tr>
<td>1459 E. 45th St.</td>
<td>713-696-3669</td>
<td></td>
<td></td>
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<tr>
<td>Houston, TX 77022</td>
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<tr>
<td>Houston-Southeast</td>
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<tr>
<td>10060 Fuqua</td>
<td>713-948-7957</td>
<td></td>
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<tr>
<td>Houston, TX 77089</td>
<td></td>
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### A.9 HHSC Open Records Requests

Providers who would like to make open records requests should visit the [Open Records](#) page of the HHSC website. For general inquiries, such as claim status inquiries, eligibility verification, and prior authorization requests, refer to subsection A.10, “TMHP Telephone and Fax Communication” in this section for contact information.

### A.10 TMHP Telephone and Fax Communication

<table>
<thead>
<tr>
<th>Contact</th>
<th>PA Request Telephone Number</th>
<th>General Inquiry Telephone Number</th>
<th>Fax Number</th>
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<tbody>
<tr>
<td>TMHP Contact Center</td>
<td>N/A</td>
<td>1-800-925-9126 or 1-512-335-5986</td>
<td></td>
</tr>
<tr>
<td>Automated Inquiry System</td>
<td>N/A</td>
<td>1-800-925-9126 or 1-512-335-5986</td>
<td></td>
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<tr>
<td>(AIS)</td>
<td></td>
<td>TMHP Contact Center 514-4214</td>
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<tr>
<td>Provider Enrollment</td>
<td>N/A</td>
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<td></td>
</tr>
<tr>
<td>Telephone appeals</td>
<td>N/A</td>
<td>1-800-745-4452</td>
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# Appendix A: State, Federal, and TMHP Contact Information

## Program and Prior Authorization Information

<table>
<thead>
<tr>
<th>Contact</th>
<th>PA Request Telephone Number</th>
<th>General Inquiry Telephone Number</th>
<th>Fax Number</th>
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<tbody>
<tr>
<td>TMHP Electronic Data Interchange (EDI) Help Desk</td>
<td>N/A</td>
<td>1-888-863-3638</td>
<td>1-512-514-4228</td>
</tr>
<tr>
<td></td>
<td></td>
<td></td>
<td>1-512-514-4230</td>
</tr>
<tr>
<td><strong>Ambulance (Medicaid and CSHCN Services Program)</strong></td>
<td>1-800-540-0694</td>
<td>TMHP Contact Center</td>
<td>1-512-514-4205</td>
</tr>
<tr>
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<td>Prior Authorization Only</td>
</tr>
<tr>
<td><strong>Children with Special Health Care Needs (CSHCN) Services Program</strong></td>
<td>N/A</td>
<td>1-800-568-2413</td>
<td>1-512-514-4222</td>
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<td></td>
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<td>Prior Authorization Only</td>
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<tr>
<td><strong>Comprehensive Care Program (CCP) (for CCP prior authorization status and general CCP and Home Health Services information)</strong></td>
<td>N/A</td>
<td>1-800-846-7470</td>
<td>1-512-514-4212</td>
</tr>
<tr>
<td></td>
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<td>Prior Authorization Only</td>
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<tr>
<td><strong>Comprehensive Care Inpatient Psychiatric (CCIP) Unit</strong></td>
<td>N/A</td>
<td>1-800-213-8877</td>
<td>1-512-514-4211</td>
</tr>
<tr>
<td>Option 1 – CCIP Option 2 – Substance use</td>
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<td></td>
<td>Prior Authorization Only</td>
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<tr>
<td><strong>Home Health Services</strong></td>
<td>N/A</td>
<td>Requests for new prior authorization or extensions cannot be made over the telephone.</td>
<td>1-512-514-4209</td>
</tr>
<tr>
<td></td>
<td></td>
<td>TMHP Contact Center</td>
<td>Prior Authorization Only</td>
</tr>
<tr>
<td><strong>Obstetric ultrasound</strong></td>
<td>1-800-302-6167</td>
<td>TMHP Contact Center</td>
<td>1-512-302-5039</td>
</tr>
<tr>
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<td>Prior Authorization Only</td>
</tr>
<tr>
<td><strong>Special Medical Prior Authorization (SMPA)</strong></td>
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<td>TMHP Contact Center</td>
<td>1-512-514-4213</td>
</tr>
<tr>
<td></td>
<td></td>
<td></td>
<td>Prior Authorization Only</td>
</tr>
<tr>
<td><strong>Texas Health Steps (THSteps) dental inquiries</strong></td>
<td>N/A</td>
<td>1-800-568-2460</td>
<td>N/A</td>
</tr>
<tr>
<td><strong>THSteps medical inquiries</strong></td>
<td>N/A</td>
<td>1-800-757-5691</td>
<td>N/A</td>
</tr>
<tr>
<td><strong>Family Planning (tubal ligation and vasectomy consent forms)</strong></td>
<td>N/A</td>
<td>TMHP Contact Center</td>
<td>1-512-514-4229</td>
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<td><strong>Hysterectomy acknowledgment statements</strong></td>
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<td>TMHP Contact Center</td>
<td>1-512-514-4218</td>
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## Other Program and Reimbursement Information

<table>
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<th>Contact</th>
<th>PA Request Telephone Number</th>
<th>General Inquiry Telephone Number</th>
<th>Fax Number</th>
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<tbody>
<tr>
<td><strong>Health Insurance Premium Payment (HIPP)</strong></td>
<td>N/A</td>
<td>1-800-440-0493</td>
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<tr>
<td><strong>Long Term Care (LTC) operations</strong></td>
<td>N/A</td>
<td>1-800-626-4117</td>
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<tr>
<td><strong>LTC—Nursing Facilities</strong></td>
<td>N/A</td>
<td>1-800-727-5436</td>
<td></td>
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<tr>
<td><strong>Medicaid Audit/Cost Reports</strong></td>
<td>N/A</td>
<td>1-512-506-6117</td>
<td>1-512-506-7811</td>
</tr>
<tr>
<td><strong>Third Party Liability (TPL) (Option 2)</strong></td>
<td>N/A</td>
<td>1-800-846-7307</td>
<td>1-512-514-4225</td>
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<tr>
<td><strong>Tort (Option 3)</strong></td>
<td>N/A</td>
<td>1-800-846-7307</td>
<td>1-512-514-4225</td>
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---

1-800-302-6167 TMHP Contact Center 1-512-514-4209 Prior Authorization Only
A.11 **Written Communication With TMHP**

All CMS-1500 forms (excluding ambulance, radiology/laboratory, immunization services, rural health, and mental health rehabilitation) sent to TMHP for the first time, as well as claims being resubmitted because they were initially denied as incomplete claims, must be sent to the following address:

Texas Medicaid & Healthcare Partnership  
Claims  
PO Box 200555  
Austin, TX 78720-0555

The post office box addresses must be used for the specific items listed in the following table:

<table>
<thead>
<tr>
<th>Correspondence</th>
<th>Address</th>
</tr>
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</table>
| Appeals/adjustments of claims (except zero paid/zero allowed on Remittance & Status [R&S] Reports) | Texas Medicaid & Healthcare Partnership Appeals/Adjustments  
PO Box 200645  
Austin, TX 78720-0645 |
| Electronically rejected claims past the 95-day filing deadline and within 120 days of electronic rejection report | Texas Medicaid & Healthcare Partnership  
Appeals/Adjustments  
PO Box 200645  
Austin, TX 78720-0645 |
| All first-time claims                                                          | Texas Medicaid & Healthcare Partnership Claims  
PO Box 200555  
Austin, TX 78720-0555 |
| Ambulance Authorization (includes out-of-state transfers)                      | Texas Medicaid & Healthcare Partnership Ambulance Prior Authorizations  
P O Box 200735  
Austin, TX 78720-0735 |
| CCP requests (prior authorization and appeals)                                 | Texas Medicaid & Healthcare Partnership Comprehensive Care Program (CCP)  
PO Box 200735  
Austin, TX 78720-0735 |
| CSHCN Services Program claims                                                  | Texas Medicaid & Healthcare Partnership CSHCN Services Program Claims  
PO Box 200855  
Austin, TX 78720-0855 |
| Home Health Services prior authorizations                                     | Texas Medicaid & Healthcare Partnership Home Health Services  
PO Box 202977  
Austin, TX 78720-2977 |
| Medicaid audit correspondence                                                  | Texas Medicaid & Healthcare Partnership Medicaid Audit  
PO Box 200345  
Austin, TX 78720-0345 |
| Medically Needy Clearinghouse (MNC) or Spend Down Unit correspondence          | Texas Medicaid & Healthcare Partnership Medically Needy Clearinghouse  
PO Box 202947  
Austin, TX 78720-2947 |
| Provider Enrollment correspondence                                            | Texas Medicaid & Healthcare Partnership Provider Enrollment  
PO Box 200795  
Austin, TX 78720-0795 |
## A.12 Other TMHP Information

### A.12.1 TMHP Website

The TMHP website at [www.tmhp.com](http://www.tmhp.com) is a valuable resource that provides:

- Provider education information and training.
- Publications, such as bulletins, banner messages, and provider manuals.
- A TMHP News section with announcements of program changes and other important information.
- Real-time and static fee schedules.
- Online provider enrollment.
- Complete instructions for setting up a Provider Administrator account and the use of online claims status inquiries (CSI), eligibility verification, and Electronic Remittance and Status (ER&S) Reports.

Additional advanced features are available for those providers who create an account. All enrolled providers are eligible for this free account. Once an account is activated, providers will have access to:

- Texas Medicaid enrollment information.
- CSIs.
- Eligibility verification.
- ER&S Report download option.
- Claims submission.
- Claims appeals.
- Online provider lookup.
- Online fee lookup (OFL) to obtain real-time fee information for an individual or a range of procedure codes. Benefits and limitations for certain services and history up to 2-years is also available.
- Payment amounts search, view, and print capabilities.
- Notification of an invalid address on file for any Texas Provider Identifier (TPI) associated with a provider’s National Provider Identifier (NPI).

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<table>
<thead>
<tr>
<th>Correspondence</th>
<th>Address</th>
</tr>
</thead>
<tbody>
<tr>
<td>Other provider correspondence</td>
<td>Texas Medicaid &amp; Healthcare Partnership Provider Relations</td>
</tr>
<tr>
<td></td>
<td>PO Box 202978</td>
</tr>
<tr>
<td></td>
<td>Austin, TX 78720-0978</td>
</tr>
<tr>
<td>Send all other written communication to TMHP</td>
<td>Texas Medicaid &amp; Healthcare Partnership (Department)</td>
</tr>
<tr>
<td></td>
<td>12357-B Riata Trace Parkway, Suite 100</td>
</tr>
<tr>
<td></td>
<td>Austin, TX 78727</td>
</tr>
<tr>
<td>TMHP Fee-for-Service and ICF-IID Dental prior</td>
<td>Texas Medicaid &amp; Healthcare Partnership Fee-for-Service and ICF-IID Dental</td>
</tr>
<tr>
<td>authorization requests</td>
<td>PO Box 204206</td>
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<tr>
<td></td>
<td>Austin, TX 78720-4206</td>
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<td>TPL/Tort correspondence</td>
<td>Texas Medicaid &amp; Healthcare Partnership Third Party Liability/Tort</td>
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<td>PO Box 202948</td>
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<tr>
<td></td>
<td>Austin, TX 78720-2948</td>
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</tbody>
</table>
• Notification of pending payments because of inaccurate or incomplete provider information.
• Manage hospital admission and discharge information on clients residing in an institution for mental diseases (IMD)

**Important:** Natural disasters, such as floods or hurricanes, can impact the delivery of health care to Texas Medicaid clients. When disaster strikes, providers should monitor the TMHP website for special instructions.

New services are always being added to the website. Please visit [www.tmhp.com](http://www.tmhp.com) for the latest information on TMHP online services.

### A.12.2 TMHP Contact Center

The TMHP Contact Center is available from 7 a.m. to 7 p.m., Central Time, Monday through Friday.

The TMHP Contact Center assists with questions such as:
- Provider enrollment procedures
- Claims filing procedures
- Policy information

The TMHP Contact Center is available to assist providers and clients. Please review the telephone and fax communication guides in this section for a list of contact telephone and fax numbers.

Provider calls, including those that were previously made to the Provider Relations territory representatives, are now handled first by the Contact Center. The Contact Center is well equipped to handle most inquiries about benefits and claims.

If the Contact Center representative determines that an inquiry can best be handled by the TMHP Provider Relations department, the inquiry will be forwarded to Provider Relations. For example, providers who want to talk to their Provider Relations representative about a visit, in-service, or training, can call the Contact Center, and the Contact Center will forward the request to Provider Relations.

Resolution of more complex issues that are referred to Provider Relations for further analysis can take up to 30 days from the date of the referral. For these issues, Provider Relations will contact the provider by telephone or e-mail when the issue has been resolved.

For questions or information about Medicaid eligibility, clients are referred to their caseworker or the local HHSC office.

### A.12.3 Automated Inquiry System (AIS)

AIS provides the following information and services through the use of a touch-tone telephone:
- Claim status
- Patient eligibility
- Benefit limitations
- Medically Needy case status
- Family Planning
- Current weekly payment amount
- Claim appeals.

Eligibility and claim status information is available on AIS 23 hours a day, 7 days a week, with scheduled down time between 3 a.m. and 4 a.m., Central Time. All other AIS information is available from 7 a.m. until 7 p.m., Central Time, Monday through Friday. AIS offers 15 transactions per call.
A.12.4 TMHP Provider Relations

The TMHP Provider Relations Department comprises a staff of Austin- and field-based provider relations representatives whose goal is to serve the health-care community by furnishing a variety of services and activities designed to inform and educate health-care providers about Texas Medicaid activities and claim submission procedures.

Provider Relations activities include the following:

- **Provider education through planned events.** Provider Relations representatives conduct a planned program of educational workshops, in-services, webinars, computer-based training (CBT), and other training sessions designed to keep all actively-enrolled providers informed of the latest policies, claim processing procedures, and federal and state regulations affecting Texas Medicaid. Details of all available provider training can be found in the Provider Education section of the TMHP website at www.tmhp.com.

- **Problem identification and resolution.** A staff of research coordinators is available to assist providers with clarification of Medicaid policies and assist with in-depth problem claim submission issues after initial inquiries are made with the TMHP Contact Center. Coordinators work closely with field-based regional representatives to coordinate the educational needs of the community.

- **Relationship with professional health-care organizations.** To ensure that Texas associations that represent health-care professions have up-to-date information about the requirements for participation in Texas Medicaid, the Provider Relations Department maintains a work relationship with these organizations. Also, the Provider Relations Department participates in several events sponsored by Texas health-care associations, such as conventions and conferences.

Call the TMHP Contact Center at 1-800-925-9126 for assistance.

A.12.5 Reporting a Complaint to TMHP

Providers can report complaints by calling the TMHP Contact Center at 1-800-925-9126 or by submitting a written complaint to:

TMHP
Complaints Resolution Department
PO Box 2014270
Austin, TX 77872-4270

A complaint is defined as any dissatisfaction concerning any aspect of Texas Medicaid. The term “complaint” does not include a misunderstanding or a problem with misinformation that is resolved promptly by clearing up the misunderstanding or supplying the appropriate information to the provider’s satisfaction.

Providers should submit the following information when reporting the complaint:

- Point of contact name and phone number or email address
- Provider name
- Provider TPI, if available
- Description of the complaint situation
- Client name
- Client PCN
- Date of service

A.12.6 TMHP Electronic Data Interchange (EDI) Help Desk

The TMHP EDI Help Desk assists Medicaid providers with EDI transactions. The TMHP EDI Help Desk is available at 1-888-863-3638 from 7 a.m. to 7 p.m., Central Time, Monday through Friday.
TMHP EDI Help Desk activities and responsibilities include, but are not limited to, the following:

- Enrolling providers for electronic billing
- Qualifying vendors for TMHP EDI production through testing
- Diagnosing claim transmission problems through research
- Consulting with provider billing personnel, billing services, and software vendors regarding TMHP EDI

TMHP EDI Help Desk staff assists with questions about TMHP EDI, TexMedConnect, and electronic transmissions at 1-888-863-3638.

Providers who employ hardware or software vendors should contact those vendors for the resolution of technical problems.
APPENDIX B: HIV/AIDS

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B.1  CDC Revised Recommendations for HIV Testing of Adults, Adolescents, and Pregnant Women in Health-Care Settings

The revised Centers for Disease Control and Prevention (CDC) recommendations advocate routine voluntary human immunodeficiency virus (HIV) screening as a normal part of medical practice, similar to screening for other treatable conditions. Screening is a basic public health tool used to identify unrecognized health conditions so treatment can be offered before symptoms develop and, for communicable diseases, so interventions can be implemented to reduce the likelihood of continued transmission. HIV screening should be offered as an opt-out test in accordance with CDC testing guidelines, which may be viewed at www.cdc.gov/mmwr/preview/mmwrhtml/rr5514a1.htm.

B.1.1 Routine HIV Testing Procedure Codes

The following table lists the procedure codes for routine HIV testing and the corresponding modifiers that must be submitted for rapid testing. Routine HIV testing is covered as a preventative or screening benefit. Medical necessity is not required.

<table>
<thead>
<tr>
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</table>

B.2  Model Workplace Guidelines for Businesses, State Agencies, and State Contractors

B.2.1  Purpose

The purpose of this policy is to protect the employment rights and privileges of individuals infected with HIV and acquired immunodeficiency syndrome (AIDS) through compliance with federal, state, and local laws. This policy will provide Texas employers, especially state agencies, with a uniform approach to developing policies and education programs that address HIV/AIDS in the workplace. The Department of State Health Services (DSHS) encourages all employers to establish workplace policies...
concerning persons with HIV/AIDS. Employers can adapt this model to fit the particular needs of their organization, work force, and clients; however, the content and intent must remain consistent with this document and the Health and Safety Code (HSC).

**B.2.2 Authority**


The model workplace guidelines, developed by the DSHS HIV/Sexually Transmitted Disease (STD) Comprehensive Services Branch, as required by HSC §85.012, “Model Workplace Guidelines,” and adopted as HIV/STD Policy No. 090.021, are considered the minimum standards for the development of guidelines for state agencies. This policy also serves as the minimum standard for contractors of certain designated state agencies and organizations funded by those state agencies (HSC §85.113).

**Refer to:** subsection B.2.7, “State Agencies Listed Under Health and Safety Code (HSC) §85.113” in this section.

These guidelines are also the standard for health-care facilities licensed by DSHS and the Texas Health and Human Services Commission (HHSC) as stated in HSC §85.010, “Educational Course for Employees and Clients of Health Care Facilities.”

**B.2.3 Who Must Use Workplace Guidelines**

**B.2.3.1 State Agencies**

State law requires that each state agency adopt and carry out workplace guidelines. The agency’s workplace guidelines should incorporate, at a minimum, the DSHS model workplace guidelines in this policy.

**B.2.3.2 State Contractors**

A program that involves direct client contact and that contracts with or is funded by any of the state agencies listed in subsection B.2.7, “State Agencies Listed Under Health and Safety Code (HSC) §85.113” in this section will adopt and carry out workplace guidelines as stated in HSC §85.113.

**B.2.4 Why Have Guidelines**

Employers should develop and carry out policies and education programs concerning potentially limiting medical conditions before a crisis arises. Such policies and education programs help reduce employees’ fears and misconceptions about HIV/AIDS and help to:

- Provide current and accurate scientific evidence that people with HIV infection do not pose a risk of transmitting the virus to coworkers through ordinary workplace contact.
- Provide workers with current information about HIV risk reduction for employees and their families.
- Avoid conflict between the infected employee and the employer regarding discrimination or other employment issues.
- Prevent work disruption and rejection of the infected employee by coworkers.
- Inform employees that they have rights regarding work continuation, confidentiality of medical and insurance records, and general health and safety.
- Provide specific and ongoing education and equipment to employees in health-care settings who are at risk of exposure to HIV, and to assure that appropriate infection-control procedures are used.
- Reduce the financial impact, legal implications, and other possible effects of HIV/AIDS in the workplace.
B.2.5 Development of Workplace Policy Content

Individuals infected with HIV have the same rights and opportunities as other individuals. While some employers prefer a policy specific to HIV/AIDS and its unique issues, others prefer a general policy concerning illnesses and disabilities. A general policy should address HIV/AIDS in the same way as other major illnesses. Use of the following statements in agency policy is encouraged:

- Use of a person’s HIV status to decide employment status, service delivery, or to deny services to HIV-infected individuals is not acceptable. Employees who believe that they have been discriminated against because of HIV or AIDS should contact the personnel office to discuss the matter, or initiate action through the agency’s grievance procedure. Other legal options may also be available.

- This policy is consistent with current information from public health authorities, such as the CDC of the U.S. Public Health Service, and with state and federal laws and regulations.

While the approach and resolution of each employee’s situation may vary, similar issues may arise. A workplace policy should address the following issues about HIV/AIDS and other life-threatening illnesses or disabilities:

- **Discrimination.** The Americans with Disabilities Act of 1990 prohibits discrimination against people with disabilities, which includes HIV and AIDS, in employment, public accommodations, public transportation, and other situations. A specific policy statement that no one will be denied employment or employment opportunities because of a disability, satisfies the employer and employee’s need to address discrimination. Such a statement might be, “This agency complies with the Americans with Disabilities Act protections of all people with disabilities against discrimination in job application procedures, hiring, promotions, discharge, compensation, job training, and other terms or conditions of employment.” Managers may want to define ways in which they will deal with discriminatory actions.

- **Desire and Ability to Work.** A workplace policy should address the infected employee’s desire and need to work and the infected employee’s value to the workplace. Such a statement reassures employees that the employer supports them. The health status of someone with HIV may vary from healthy to critically ill. In the work setting, the ultimate concern is whether or not the employee can satisfy job expectations. A policy statement may say, for example, “Procedures may be adapted to provide reasonable accommodation so that people with disabilities may remain employed and productive for as long as possible. All employees, however, are expected to perform the essential functions of their job with or without reasonable accommodation.”

- **Performance Standards.** The Americans with Disabilities Act provides protections for disabled persons qualified to perform their jobs. And although an employer may be expected to provide reasonable accommodation to a disabled employee or applicant; employers may terminate employees and refuse to hire individuals who cannot perform the essential functions of the job, with or without the reasonable accommodation. One suggested statement is, “While the Americans with Disabilities Act does protect disabled employees from employment discrimination, all employees, those with and without disabilities, have the same performance and conduct standards regarding hiring, promotion, transfer, and dismissal.”

- **Reasonable Accommodation.** The Americans with Disabilities Act requires employers to provide reasonable accommodations for employees with disabilities. Employers do not have an obligation to provide any accommodation that imposes an undue hardship on the employer. Specific questions about the issue of reasonable accommodation and undue hardship should be directed to staff responsible for coordinating the requirements of the Americans with Disabilities Act. Such a policy statement might read, “The following options may be considered for people with HIV/AIDS: possible assignment or reassignment of job duties, working at home, leaves of absence, and flexible work schedules.”
• **Confidentiality and Privacy.** Organizations that receive funds from a state agency for residential or direct client services or programs shall develop and use confidentiality guidelines to protect their clients’ HIV/AIDS-related medical information (HSC §85.115, “Confidentiality Guidelines”). Organizations that fail to adopt and use confidentiality guidelines are ineligible to receive state funds. Employees are not required to reveal their HIV status to employers. All medical information that an HIV-infected employee provides to medical or management personnel is confidential and private. Employers may not reveal this information without the employee’s knowledge and written consent, except as provided by law (HSC §81.103, “Confidentiality; Criminal Penalty”). A suggested policy statement might be, “This agency will protect the confidentiality of employee medical records and information. Written consent of the employee must be obtained to share any confidential information with other staff. Those with access to confidential information must maintain strict confidentiality and privacy, separating this information from employees’ personnel records. Individuals who fail to protect these employee rights commit a serious offense, which may be cause for litigation resulting in both civil and criminal penalties, and may result in dismissal.”

• **Coworker Concerns.** Employers need to be aware of the concerns that coworkers may have about an HIV-infected coworker. A policy statement that acknowledges employee concerns and offers HIV/AIDS education helps to increase awareness and decrease fear. Equally important is a policy statement that clarifies the limits of an employer’s response to coworker concerns, e.g., “Employees do not have the right to refuse to work with someone who has any disability.”

• **Employee Education.** Any health-care facility licensed by DSHS or HHSC must require its employees to complete an educational course about HIV infection (HSC §85.010). A suggested policy statement may be: “All employees will receive education about methods of transmission and prevention of HIV infection and related conditions.” In response to HSC, §85.004, “Educational Programs,” DSHS developed model education program guidelines. These are available from DSHS, HIV/STD Comprehensive Services Branch, 1100 W. 49th St., Austin, TX. 78756-3199, 1-512-533-3000. Employers may also find the CDC’s educational kit, *Business Responds to AIDS*, useful in developing educational courses. HIV/AIDS education should address employee concerns about HIV communicability to themselves, their families, and coworkers. Experience shows that educated coworkers usually respond to persons with HIV/AIDS with support, rather than with fear and ostracism due to misconceptions. Education programs must stress that agency employees who provide direct client services may face occupational exposure to a client’s blood, semen, vaginal secretions, or other body fluids that are considered to be high-risk for transmission of blood born pathogens, including HIV/AIDS. All individuals receiving direct services are clients and include individuals who are physically or mentally impaired and individuals confined to correctional or residential facilities. All state agencies should have, as part of their employee education program, comprehensive policies and protocols based on universal precautions, body substance isolation, and barrier methods. These precautions prevent the spread of infection in clinical settings. The employer’s careful planning will reflect a commitment to the health and well-being of the work force and the community being served.

• **Assistance.** Some employers have designated benefits programs available to employees and family members with HIV infection. Such programs may:
  • Make referrals for testing, counseling, medical, and psychosocial services.
  • Provide HIV/AIDS workplace training for managerial staff.
  • Serve as a liaison between management and the employer’s clinical and occupational health programs.
  • Provide counseling for employees who irrationally fear coworkers or clients.
Employers who have no employee assistance program may consider working with other organizations that provide assistance. Some of these groups include local health departments, AIDS services organizations, American Red Cross chapters, community support groups, clinical treatment and counseling services, and the religious community.

A suggested policy statement might be: “An employee who wants assistance concerning a disability or a life-threatening illness should contact the Personnel Office. This agency offers the following resources to help employees and managers deal with these issues: education and information concerning HIV/AIDS; confidential referral to supportive services for employees and dependents affected by life-threatening illnesses; and benefits consultation to help employees effectively manage health, leave, and other benefits.”

**B.2.6 Where to Go for Help**

Employees may call 2-1-1 for HIV/STD testing locations in Texas. For questions related to issues such as transmission, signs and symptoms, or other concerns about HIV or other sexually transmitted infections, employees may call 1-800-CDC-INFO (English/Español) or 1-888-232-6348 (TTY).

**B.2.7 State Agencies Listed Under Health and Safety Code (HSC) §85.113**

HSC §85.113, “Workplace Guidelines for State Contractors” states “An entity that contracts with or is funded by… to operate a program involving direct client contact shall adopt and implement workplace guidelines similar to the guidelines adopted by the agency that funds or contracts with the entity.”

H.B. 2292, 78th Leg., abolished 10 of the 12 existing health and human services agencies and transferred their powers and duties to three new state agencies and to HHSC, which rendered the state agency list found in HSC §85.113 obsolete. The following list reflects the state agency consolidation brought about by H.B. 2292 and identifies the state agencies to which HSC §85.113 applies.

- DSHS
- HHSC
- Texas Department of Criminal Justice
- Texas Juvenile Probation Commission
- Texas Youth Commission
## C.1 Acronym Dictionary

<table>
<thead>
<tr>
<th>Term</th>
<th>Definition</th>
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<tbody>
<tr>
<td>A/EEG</td>
<td>Ambulatory Electroencephalogram</td>
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<tr>
<td>AAFP</td>
<td>American Academy of Family Physicians</td>
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<tr>
<td>AAP</td>
<td>American Academy of Pediatrics</td>
</tr>
<tr>
<td>AAPD</td>
<td>American Academy of Pediatric Dentistry</td>
</tr>
<tr>
<td>ABMG</td>
<td>American Board of Medical Geneticists</td>
</tr>
<tr>
<td>ABR</td>
<td>Auditory Brainstem Response</td>
</tr>
<tr>
<td>ACA</td>
<td>Affordable Care Act of 2010</td>
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<td>Augmentative Communication Device</td>
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<td>Automated Clearinghouse</td>
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<td>Advisory Committee on Immunization Practices</td>
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<tr>
<td>ADA</td>
<td>American Dental Association</td>
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<tr>
<td>ADL</td>
<td>Activity of Daily Living</td>
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<td>American Heart Association</td>
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<td>American National Standards Institute</td>
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<td>Atypical Provider Identifier</td>
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<td>Advanced Practice Registered Nurse</td>
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<td>Admission, Review, and Dismissal</td>
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<td>Compass21</td>
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<td>Critical Access Hospital</td>
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<td>Continuous Ambulatory Peritoneal Dialysis</td>
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<td>Community-Based Alternatives (Program)</td>
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<td>Community Based Organization</td>
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<td>Comprehensive Care Program</td>
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<td>Current Dental Terminology</td>
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1 General Information

The information in this handbook is intended for Texas Medicaid ambulance providers. The handbook provides information about Texas Medicaid’s benefits, policies, and procedures applicable to emergency and nonemergency ambulance transports.

Important: All providers are required to read and comply with “Section 1: Provider Enrollment and Responsibilities” (Vol. 1, General Information). In addition to required compliance with all requirements specific to Texas Medicaid, it is a violation of Texas Medicaid rules when a provider fails to provide health-care services or items to Medicaid clients in accordance with accepted medical community standards and standards that govern occupations, as explained in Title 1 Texas Administrative Code (TAC) §371.1659. Accordingly, in addition to being subject to sanctions for failure to comply with the requirements that are specific to Texas Medicaid, providers can also be subject to Texas Medicaid sanctions for failure, at all times, to deliver health-care items and services to Medicaid clients in full accordance with all applicable licensure and certification requirements including, without limitation, those related to documentation and record maintenance.

Refer to: “Section 1: Provider Enrollment and Responsibilities” (Vol. 1, General Information) for more information about enrollment procedures. The Medicaid Managed Care Handbook (Vol. 2, Provider Handbooks).

Subsection 5.1.8, “Prior Authorization for Nonemergency Ambulance Transport” in “Section 5: Fee-for-Service Prior Authorizations” (Vol. 1, General Information) for more information about nonemergency ambulance transport.

2 Ambulance Services

2.1 Enrollment

To enroll in Texas Medicaid, ambulance providers must operate according to the laws, regulations, and guidelines governing ambulance services under Medicare Part B; equip and operate under the appropriate rules, licensing, and regulations of the state in which they operate; acquire a license from the Texas Department of State Health Services (DSHS) approving equipment and training levels of the crew; and enroll in Medicare.

A hospital-operated ambulance provider must be enrolled as an ambulance provider and submit claims using the ambulance provider identifier, not the hospital provider identifier.

Refer to: Subsection 2.4.3, “Medicare and Medicaid Coverage” in this handbook.

Note: Air ambulance providers are not required to enroll with Medicare.

Reminder: When ambulance providers enroll in Texas Medicaid, they accept Medicaid payment as payment in full. They cannot bill clients for Texas Medicaid-covered benefits.

2.1.1 Subscription Plans

The Texas Insurance Code does not apply to ambulance providers who finance, in part or in whole, an ambulance service by subscription plan. DSHS’s license requirements do not permit providers of membership or subscription programs to enroll Medicaid clients. Emergency Medical Services (EMS) Subscription Programs are regulated by the DSHS-EMS Compliance Group. An EMS provider must have specific approval to operate a subscription program.
For more information, providers should contact the DSHS Office of EMS/Trauma Systems Coordination at 1-512-834-6700. A list of EMS office and contact information is available at http://dshs.texas.gov/emstraumasystems/contact.shtm.

### 2.2 Services, Benefits, Limitations, and Prior Authorization

Emergency and nonemergency ambulance transport services are a benefit of Texas Medicaid when the client meets the definition of emergency medical condition or meets the requirements for nonemergency transport.

Cardiopulmonary resuscitation (CPR) is included in ambulance transport when needed and is not a separately billable service. Claims for CPR during transport will be denied. If CPR is performed during a nonemergency transport, the advanced life support (ALS) procedure code must be billed.

Reimbursement for disposable supplies is separate from the established global fee for ambulance transports and is limited to one billable code per trip.

Providers must calculate the number of miles traveled by using the ambulance vehicle odometer reading or an Internet mapping tool. Mileage reported on the claim must be the actual number of miles traveled.

Claims for ground ambulance transports (procedure codes A0426, A0427, A0428, A0429, A0433, A0434, and A0999) must be submitted with mileage procedure code A0425.

Medical necessity and coverage of ambulance services are not based solely on the presence of a specific diagnosis. Medicaid payment for ambulance transportation may be made only for those clients whose condition at the time of transport is such that ambulance transportation is medically necessary. For example, it is insufficient that a client merely has a diagnosis such as pneumonia, stroke, or fracture to justify ambulance transportation. In each of those instances, the condition of the client must be such that transportation by any other means is medically contraindicated. In the case of ambulance transportation, the condition necessitating transportation is often an accident or injury that has occurred giving rise to a clinical suspicion that a specific condition exists (for instance, fractures may be strongly suspected based on clinical examination and history of a specific injury).

It is the requesting provider’s responsibility to supply the contractor with information describing the condition of the client that necessitated ambulance transportation. Medicaid recognizes the limitations of ambulance personnel in establishing a diagnosis, and recognizes therefore, that diagnosis coding of a client’s condition using *International Classification of Diseases, Tenth Revision, Clinical Modification* (ICD-10-CM) codes when reporting ambulance services may be less specific than those reported by other professional providers. Providers who submit diagnosis codes must choose the code that best describes the client’s condition at the time of transport. As a reminder to providers of ambulance services, “rule out” or “suspected” diagnoses must not be reported using specific ICD-10-CM codes. In such instances where a diagnosis is not confirmed, it is correct to use a symptom, finding, or injury code.

The ambulance provider may be sanctioned, including nonparticipation in the Medicaid Title XIX programs, for completing or signing a claim form that includes false or misleading representations of the client's condition or the medical necessity of the transport.

The inpatient hospital stay benefit includes medically necessary emergency and nonemergency ambulance transportation of the client during an inpatient hospital stay.

Ambulance transport during a client’s inpatient stay will not be reimbursed to the ambulance provider. One-time ambulance transports that occur immediately after the client’s discharge may be considered for reimbursement.

#### 2.2.1 Emergency Ambulance Transport Services

An emergency ambulance transport service is a benefit when the client has an emergency medical condition. An emergency medical condition is defined, according to 1 TAC §354.1111, as a medical condition (including emergency labor and delivery) manifesting itself by acute symptoms of sufficient
severity (including severe pain, psychiatric disturbances, or symptoms of substance use) such that a prudent layperson with an average knowledge of health and medicine, could reasonably expect the absence of immediate medical attention to result in one of the following:

- Placing the client’s health (or, with respect to a pregnant woman, the health of the woman or her unborn child) in serious jeopardy
- Serious impairment to bodily functions
- Serious dysfunction of any bodily organ or part

Facility-to-facility transport may be considered an emergency if emergency treatment is not available at the first facility and the client still requires emergency care. The transport must be to an appropriate facility, meaning the nearest medical facility equipped in terms of equipment, personnel, and the capacity to provide medical care for the illness or injury of the client involved.

Transports to out-of-locality providers (one-way transfers of 50 or more miles from the point of pickup to the point of destination) are covered if a local facility is not adequately equipped to treat the condition. Transports may be cut back to the closest appropriate facility.

When there are two responders to an emergency, the company that transports the client will be reimbursed for their services.

### 2.2.1.1 Prior Authorization for Emergency Out-of-State Transport

All emergency out-of-state (air, ground, and water) transports require authorization before the transport is considered for payment.

Prior authorization for emergency transport is required for out-of-state providers with the exception of those providers located within 200 miles of the Texas border.

**Refer to:** Subsection 2.6, “Out-of-State Medicaid Providers” in “Section 2: Texas Medicaid Fee-for-Service Reimbursement” (Vol. 1, General Information) for additional information on providers who are not considered out-of-state providers.

To initiate the prior authorization process, providers must call 1-800-540-0694.

Texas Medicaid & Healthcare Partnership (TMHP) is responsible for processing prior authorization requests for all Medicaid clients.

### 2.2.2 Nonemergency Ambulance Transport Services

According to 1 TAC §354.1111, nonemergency transport is defined as ambulance transport provided for a Medicaid client to or from a scheduled medical appointment, to or from a licensed facility for treatment, or to the client’s home after discharge from a hospital when the client has a medical condition such that the use of an ambulance is the only appropriate means of transportation (i.e., alternate means of transportation are medically contraindicated).

**Note:** In this circumstance, contraindicated means that the client cannot be transported by any other means from the origin to the destination without endangering the individual’s health.

According to Human Resource Code (HRC) §32.024 (t), a Medicaid-enrolled physician, nursing facility, health-care provider, or other responsible party is required to obtain authorization before an ambulance is used to transport a client in circumstances not involving an emergency.

Providers requesting prior authorization must document whether the client is currently an inpatient in a hospital when requesting prior authorization. Prior authorization will not be approved if the provider indicates the client is currently an inpatient in a hospital, except for one-time transports immediately after the client’s discharge from the hospital.
Prior authorization requests may be submitted to the TMHP Prior Authorization Department by mail, fax, or the electronic portal. Prescribing or ordering providers, dispensing providers, clients’ responsible adults, and clients may sign prior authorization forms and supporting documentation using electronic or wet signatures.

Medical necessity must be established through prior authorization for all nonemergency ambulance transports. Retrospective review may be performed to ensure that documentation supports the medical necessity of the transport.

Clients who do not meet medical necessity requirements for nonemergency ambulance transport may be able to receive transport through the Medical Transportation Program (MTP).

Transports must be limited to those situations where the transportation of the client is less costly than bringing the service to the client.

For non-emergency ambulance transportation services rendered to a client, ambulance providers may coordinate the nonemergency ambulance prior authorization request between the requesting provider, which may include a physician, nursing facility, healthcare provider, or other responsible party. Ambulance providers may assist in providing necessary information such as their National Provider Identifier (NPI) number, fax number, and business address to the requesting provider. However, the Non-emergency Ambulance Prior Authorization Request form must be signed and dated and submitted by the Medicaid-enrolled requesting provider, not the ambulance provider.

Refer to: Subsection 5.1.8, “Prior Authorization for Nonemergency Ambulance Transport” in “Section 5: Fee-for-Service Prior Authorizations” (Vol. 1, General Information) for more information about nonemergency ambulance transport prior authorization requests and appeals.

The Medical Transportation Program Handbook (Vol. 2, Provider Handbooks) for more information about the Medical Transportation Program.

2.2.3 Levels of Service

Levels of services as defined by Texas Medicaid:

- Basic Life Support (BLS) is emergency care that uses noninvasive medical acts and, if allowed by licensing jurisdiction, may include the establishment of a peripheral intravenous (IV) line.

- Advanced Life Support (ALS) is emergency care that uses invasive medical acts. For Medicaid purposes only, ALS services are divided into two categories, Level 1 and Level 2.
  - Level 1 ALS includes an ALS assessment or at least an ALS intervention.
  - Level 2 ALS includes either of the following:
    - At least three separate administration of one or more medications by intravenous push/bolus or by continuous infusion (excluding crystalloid fluids); or
    - At least one of the ALS 2 procedures: manual defibrillation/cardioversion; endotracheal intubation; central venous line; cardiac pacing; chest decompression; surgical airway; or intra-osseous line.

2.2.4 Oxygen

Reimbursement for oxygen (procedure code A0422) is limited to one billable code per transport.

2.2.5 Types of Transport

2.2.5.1 Multiple Client Transports

Multiple client transports occur when more than one client with Medicaid coverage is transported simultaneously in the same vehicle. A claim for each client must be billed with the transport procedure code and the mileage procedure code with the GM modifier that indicates multiple client transport.
Claims must include the names and Medicaid numbers of other Texas Medicaid clients who shared the transfer, or indicate "Not a Medicaid client" in Block 19 of the CMS 1500 paper form. Providers must enter charges on a separate claim for each client. TMHP adjusts the payment to 80 percent of the allowable base rate for each claim and divides mileage equally among the clients who share the ambulance.


2.2.5.2 Air or Specialized Vehicle Transports

Air ambulance transport services, by means of either fixed or rotary wing aircraft, and other specialized emergency medical services vehicles may be covered only if one of the following conditions exists:

- The client’s medical condition requires immediate and rapid ambulance transportation that could not have been provided by standard automotive ground ambulance.
- The point of client pick up is inaccessible by standard automotive ground vehicle.
- Great distances or other obstacles are involved in transporting the client to the nearest appropriate facility.

Claims for air ambulance transports procedure codes A0430 and A0431 must be submitted with the corresponding air mileage procedure code A0435 or A0436.

2.2.5.3 Specialty Care Transport (SCT)

SCT (procedure code A0434) is the interfacility transport of a critically injured or ill client by a ground ambulance vehicle, including the provision of medically necessary supplies and services, at a level of service beyond the scope of the emergency medical technician (EMT) or paramedic. SCT is necessary when a client’s condition requires ongoing care that must be furnished by one or more health professionals in an appropriate specialty area, for example, emergency or critical-care nursing, emergency medicine, respiratory care, cardiovascular care, or a paramedic with additional training.

2.2.5.4 Transports for Pregnancies

Transporting a pregnant woman may be covered as an emergency transfer if the client’s condition is documented as an emergency situation at the time of transfer.

Claims documenting an emergency home delivery or delivery en route are considered emergency transfers. Premature labor and early onset of delivery (less than 37 weeks gestation) may also be considered an emergency. Active labor without more documentation of an emergency situation is not payable as an emergency transport.

If the pregnant client is transported in an ambulance for a nonemergency situation, all criteria for nonemergency prior authorization must be met.

2.2.5.5 Transports to or from Prescribed Pediatric Extended Care Centers (PPECC)

Non-emergency ambulance transports between a client’s home and a PPECC are not covered.

2.2.5.6 Transports to or from State Institutions

Ambulance transports to or from a state-funded hospital for admission or following discharge are covered when nonemergency transfer criteria are met. Ambulance transfers of clients while they are inpatients of the institution are not covered. The institution is responsible for routine nonemergency transportation.
2.2.5.7 Not Medically Necessary Transports

Providers must use the GY modifier to submit claims for instances when the provider is aware no medical necessity existed. When billing for this type of transportation, ambulance providers must maintain a signed Client Acknowledgment Statement indicating that the client was aware, prior to service rendered, that the transport was not medically necessary. The Client Acknowledgment Statement is subject to retrospective review.

Refer to: Subsection 1.7.11.1, “Client Acknowledgment Statement” in “Section 1: Provider Enrollment and Responsibilities” (Vol. 1, General Information).

2.2.5.8 Transports for Nursing Facility Residents

Nursing facilities are responsible for providing or arranging transportation for their residents. Arranging transportation for Medicaid clients includes obtaining prior authorizations for nonemergency ambulance transports. The Nonemergency Ambulance Prior Authorization Request form must be filled out and submitted to TMHP by the facility or the physician’s staff that is most familiar with the client’s condition. The ambulance provider must not assist in completing or submitting any portion of this form.

Transports from a nursing facility to a hospital are covered if the client’s condition meets emergency criteria.

A return trip to a nursing facility following an emergency transport is not considered routine; therefore, transport back to the facility must be requested by the discharging hospital. Nonemergency transport for the purpose of required diagnostic or treatment procedures that are not available in the nursing facility (such as dialysis treatments at a freestanding facility) is also allowable only for clients whose medical condition is such that the use of an ambulance is the only appropriate means of transport (e.g., alternate means of transport are medically contraindicated).

The cost of routine nonemergency transportation is included in the nursing facility vendor rate. This nonemergency transport requires the nursing facility to request and obtain a Prior Authorization Number (PAN) from the TMHP Ambulance Unit before contacting the ambulance company for the transport.

Transports of nursing facility residents for rehabilitative treatment (e.g., physical therapy) to outpatient departments or physicians’ offices for recertification examinations for nursing facility care are not reimbursable ambulance services.

Claims for services to nursing facility residents must indicate the medical diagnosis or problem requiring treatment, the medical necessity for use of an ambulance for the transport, and the type of treatment rendered at the destination (e.g., admission or X-ray).

If a client is returned by ambulance to a nursing facility following inpatient hospitalization, the acute condition requiring hospitalization must be noted on the ambulance claim form. This transport is considered for payment only if the client’s medical condition is appropriate for transport by ambulance. This nonemergency transport requires the nursing facility to request and obtain a PAN from the TMHP Ambulance Unit before contacting the ambulance company for the transport.

Ambulance providers may bill a nursing facility or client for a nonemergency ambulance transport only under the following circumstances:

- Providers may bill the nursing facility when the nursing facility requests the nonemergency ambulance transport without a PAN.
- Providers may bill the client only when the client requests transport that is not an emergency and the client does not have a medical condition such that the use of an ambulance is the only appropriate means of transport (i.e., alternate means of transport are medically contraindicated). The provider
must advise the client of acceptance as a private pay patient at the time the service is provided, and the client is responsible for payment of all services. Providers are encouraged to have the client sign the Private Pay Agreement.

Providers may refer questions about a nursing facility's responsibility for payment of a transport to the TMHP Contact Center at 1-800-925-9126 or TMHP provider relations representatives.

### 2.2.5.9 Emergency Transports Involving a Hospital

Hospital-to-hospital transports that meet the definition of an emergency transport do not require prior authorization.

Providers must use modifier ET and one of the facility-to-facility transfer modifiers (HH, HI, or IH) on each procedure code listed on the claim.

<table>
<thead>
<tr>
<th>Modifier</th>
<th>Transport Type</th>
</tr>
</thead>
<tbody>
<tr>
<td>HH</td>
<td>From hospital to hospital</td>
</tr>
<tr>
<td>HI</td>
<td>From hospital to site of transfer</td>
</tr>
<tr>
<td>IH</td>
<td>From site of transfer to hospital</td>
</tr>
</tbody>
</table>

#### 2.2.5.10 No Transport

Texas Medicaid does not reimburse ambulance providers for services that do not result in a transport to a facility, regardless of whether any medical care was rendered. If a client contacts an ambulance provider, but the call does not result in a transport, the provider should have the client sign an acknowledgement statement and may bill the client for services rendered.

Texas Medicaid will not reimburse for the return trip of an empty ambulance. Texas Medicaid will not reimburse air or ground mileage when the client is not on board the ambulance.

### 2.3 Documentation Requirements

The requesting provider, which may include a physician, nursing facility, health-care provider, or other responsible party, is required to maintain the supporting documentation, physician's orders, the Non-emergency Ambulance Prior Authorization Request form and if applicable, the Nonemergency Ambulance Exception form.

The requesting provider (i.e., physician, nursing facility, healthcare provider, or other responsible party) must contact the transporting ambulance provider with the PAN and the dates of service that were approved. The transporting ambulance provider will submit claims for the nonemergency ambulance transportation services, using the approved PAN provided by the requesting provider.

An ambulance provider is required to maintain documentation that represents the client's medical condition and other clinical information to substantiate medical necessity, the level of service, and the mode of transportation requested. This supporting documentation is limited to documents developed or maintained by the ambulance provider.

Physicians, nursing facilities, health-care providers, or other responsible parties are required to maintain physician orders related to requests for prior authorization of nonemergency and out-of-state ambulance services. These providers must also maintain documentation of medical necessity for the ambulance transport.

In hospital-to-hospital transports or hospital-to-outpatient medical facility transports, the TMHP Ambulance Unit considers information by telephone from the hospital. Providers are not required to fax medical documentation to TMHP; however, in certain circumstances, TMHP may request that the hospital fax the supporting documentation. Hospitals are allowed to release a client's protected health information (PHI) to a transporting emergency medical services provider for treatment, payment, and health-care operations.
Providers must document whether the client is currently an inpatient in a hospital when requesting prior authorization. Prior authorization will not be approved if the provider indicates the client is currently an inpatient in a hospital, except for one-time transports immediately after the client’s discharge from the hospital.

The hospital must maintain documentation of medical necessity, including a copy of the authorization from TMHP in the client’s medical record for any item or service that requires prior authorization. The services provided must be clearly documented in the medical record with all pertinent information regarding the client’s condition to substantiate the need and medical necessity for the services.

### 2.3.1 Medicaid Surety Bond Requirements

Ambulance providers attempting to renew their Emergency Medical Services (EMS) license must submit a surety bond to TMHP for each license they are attempting to renew. A copy of the surety bond must also be attached to an application for renewal of an EMS license when submitted to DSHS.

**Refereto:** Subsection 1.1.7, “Surety Bond Enrollment Requirement” in “Section 1: Provider Enrollment and Responsibilities” (*Vol. 1, General Information*) for more information.

### 2.4 Claims Filing and Reimbursement

#### 2.4.1 Claims Information

Emergency and nonemergency claims may be billed electronically. For electronic billers, the hospital’s provider identifier must be entered in the Facility ID field. Providers should consult their software vendor for the location of this field on the electronic claim form.

The Healthcare Common Procedure Coding System (HCPCS)/Current Procedural Terminology (CPT) codes included in policy are subject to National Correct Coding Initiative (NCCI) relationships. Exceptions to NCCI code relationships that may be noted in the Texas Medicaid medical policy are no longer valid.

The CMS NCCI and MUE guidelines can be found in the [NCCI web page](https://www.cms.gov), which are available on the CMS website. Providers should refer to the CMS NCCI web page for correct coding guidelines and specific applicable code combinations.

In instances when Texas Medicaid medical policy quantity limitations are more restrictive than NCCI Medically Unlikely Edits (MUE) guidance, medical policy prevails.

#### 2.4.2 Reimbursement

Ground and air ambulance providers are reimbursed based on the lesser of a provider’s billed charges or the maximum fee established by the Texas Health and Human Services Commission (HHSC) in accordance with 1 TAC §355.8600. Providers can refer to the [Online Fee Lookup (OFL)](https://www.tmhp.com) or the applicable fee schedule on the TMHP website at [www.tmhp.com](http://www.tmhp.com).

**Refereto:** Subsection 2.2, “Fee-for-Service Reimbursement Methodology” in “Section 2: Texas Medicaid Fee-for-Service Reimbursement” (*Vol. 1, General Information*) for more information about reimbursement methodologies.

Subsection 1.12, “Texas Medicaid Limitations and Exclusions” in “Section 1: Provider Enrollment and Responsibilities” (*Vol. 1, General Information*) for information on Medicaid exclusions.

#### 2.4.2.1 Ambulance Disposable Supplies

Ambulance disposable supplies are included in the global fee for specialty care transport and must not be billed separately.
Reimbursement for BLS or ALS disposable supplies (procedure codes A0382 and A0398 respectively) is separate from the established fee for ALS and BLS ambulance transports and is limited to one billable procedure code per transport.

**2.4.2.2 Payment Window Reimbursement Guidelines for Services Preceding an Inpatient Admission**

The three-day and one-day payment window reimbursement guidelines do not apply for ambulance services.

*Refer to:* Subsection 3.7.4.14, “Payment Window Reimbursement Guidelines” in the *Inpatient and Outpatient Hospital Services Handbook* (Vol. 2, Provider Handbooks) for additional information about the payment window reimbursement guidelines.

**2.4.3 Medicare and Medicaid Coverage**

All ambulance claims are exempt from Medicare equalization, which pays the lesser of the coinsurance and deductible or the remainder of the amount that Medicaid would have paid for the same service minus what Medicare has already paid on Medicare crossover claims.

Ambulance providers can include copayment amounts as coinsurance for non-contracted Medicare Part C clients on dual eligible claims submitted to TMHP. Providers do not have to bill Medicare copay procedure codes to TMHP for ambulance services.

All claims for ambulance services provided to dual-eligible clients are reimbursed the full amount of the Medicare coinsurance and deductible for Part B claims and Part C claims from non-contracted Medicare Advantage Plans.

Medicaid prior authorization is not required for ambulance services for Qualified Medicare Beneficiary (QMB) clients because QMB clients are not eligible for Medicaid benefits. Providers can contact Medicare for the Medicare prior authorization guidelines.

Medicaid Qualified Medicare Beneficiary (MQMB) clients are eligible for all Medicaid benefits; therefore, the provider should simultaneously request prior authorization for the nonemergency transport from TMHP for the MQMB client in the event the service requested is denied by Medicare as a non-covered service.

*Refer to:* Subsection 4.9, “Medicare and Medicaid Dual Eligibility” in “Section 4: Client Eligibility” (Vol. 1, General Information).

Subsection 2.7, “Medicare Crossover Claim Reimbursement” in “Section 2: Texas Medicaid Fee-for-Service Reimbursement” (Vol. 1, General Information) for additional information about Medicare coinsurance and deductible payments and exceptions.

**2.4.3.1 Medicare Services Paid**

Assigned claims filed with and paid by Medicare should automatically transfer to TMHP for payment of the deductible and coinsurance liability. According to current guidelines, providers must submit Medicare-paid claims that do not cross over to TMHP for the coinsurance and deductible. Providers must send the Medicare Remittance Advice Notice (MRAN) with the client information circled in black ink.

**2.4.3.2 Medicare Services Denied**

A Medicare ambulance claim that has been denied must go through the appropriate Medicare claim appeals process with a decision by the administrative law judge before TMHP will process the ambulance claim. MQMB ambulance claims that have exhausted the Medicare third level of appeal by the administrative law judge (ALJ) must be submitted to TMHP with the disposition letter from the ALJ along with all other required documents for an appeal.
An assigned claim that was denied by Medicare because the client has no Part B benefits or because the transport destination is not allowed can be submitted to TMHP for consideration. Providers must send claims to TMHP on a CMS-1500 paper claim form with the ambulance provider identifier, unless they are a hospital-based provider. Hospital-based ambulance providers must send Medicare denied claims to TMHP on a CMS-1500 paper claim form with the ambulance provider identifier and a copy of the MRAN.

**Note:** All claims for STAR+PLUS clients with Medicare and Medicaid must follow the same requirements used for obtaining prior authorization for Medicaid-only services from TMHP. The STAR+PLUS HMO is not responsible for reimbursement of these services.

### 2.4.4 Ambulance Claims Coding

Providers must submit claims for emergency transport with the ET modifier on each procedure code submitted. Any procedure code submitted on the claim for emergency transport without the ET modifier will be subject to prior authorization requirements.

#### 2.4.4.1 Place of Service Codes

The place of service (POS) for all ambulance transports is considered the destination.

POS codes 41 and 42 (other) are national POS codes that are accepted by Texas Medicaid only for electronic claims. POS code 9 is accepted by Texas Medicaid for ambulance claims submitted on paper.

#### 2.4.4.2 Origin and Destination Codes

All claims submitted on paper or electronically must include the two-character origin and destination codes for every claim line. The origin is the first character, and the destination is the second character.

The following are the origin and destination codes accepted by Texas Medicaid:

<table>
<thead>
<tr>
<th>Origin and Destination Code</th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td>D</td>
<td>Diagnostic or therapeutic site/freestanding facility (e.g., radiation therapy center) other than P or H</td>
</tr>
<tr>
<td>E</td>
<td>Residential/domiciliary/custodial facility (e.g., nonskilled facility)</td>
</tr>
<tr>
<td>G</td>
<td>Hospital-based dialysis facility (hospital or hospital-related)</td>
</tr>
<tr>
<td>H</td>
<td>Hospital (e.g., inpatient or outpatient)</td>
</tr>
<tr>
<td>I</td>
<td>Site of transfer (e.g., airport or helicopter pad) between modes of ambulance transport</td>
</tr>
<tr>
<td>J</td>
<td>Non-hospital-based dialysis facility</td>
</tr>
<tr>
<td>N</td>
<td>Skilled Nursing Facility (SNF) (swingbed is considered an SNF)</td>
</tr>
<tr>
<td>P</td>
<td>Physician’s office (includes HMO and nonhospital facility)</td>
</tr>
<tr>
<td>R</td>
<td>Residence (client’s home or any residence)</td>
</tr>
<tr>
<td>S</td>
<td>Scene of accident or acute event</td>
</tr>
<tr>
<td>X</td>
<td>Intermediate stop at physician’s office en route to the hospital (destination code only)</td>
</tr>
</tbody>
</table>

Nonemergency claims filed electronically must include the PAN in the appropriate field. For nonemergency hospital-to-hospital transfers, indicate the services required from the second facility and unavailable at the first facility in Block 19 of the CMS-1500 paper claim form. If the destination is a hospital, enter the name and address and the provider identifier of the facility in Block 32.
For nonemergency transports, ambulance providers must enter the ICD-10-CM diagnosis code to the highest level of specificity available for each diagnosis observed in Block 21 of the claim form.

**Reminder:** Providers must submit multiple transports for the same client on the same date of service through one claim submission. Additional claims information can be found within individual topics in this section.

Providers should consult their software vendor for the location of the field on the electronic claim form. Providers must submit ambulance services to TMHP on a CMS-1500 paper claim form. Providers may purchase CMS-1500 paper claim forms from a vendor of their choice. TMHP does not supply the forms.

When completing a CMS-1500 paper claim form, all required information must be included on the claim, as TMHP does not key information from claim attachments. Superbills, or itemized statements, are not accepted as claim supplements.

**Refer to:** “Section 3: TMHP Electronic Data Interchange (EDI)” (*Vol. 1, General Information*) for information on electronic claims submissions.

“Section 6: Claims Filing” (*Vol. 1, General Information*) for general information about claims filing.

Subsection 6.5, “CMS-1500 Paper Claim Filing Instructions” in “Section 6: Claims Filing” (*Vol. 1, General Information*). Blocks that are not referenced are not required for processing by TMHP and may be left blank.

### 2.4.4.3 Transports Billed Without Mileage

Ambulance transport claims with a billed mileage amount of $0.00 will be reimbursed. To qualify for reimbursement, the transport claim must include a mileage quantity that is greater than zero.

Providers may not include a mileage charge as part of the transport charge or as part of any other charges on the claim.

Payments for ambulance transports are only made if the client is actually transported and the mileage quantity billed is greater than zero. Mileage charges greater than zero will be considered for reimbursement when a transport procedure code is included on the claim.

### 2.4.5 Air or Specialized Vehicle Transports

Procedure codes A0430 and A0435, or A0431 and A0436 are used to bill air transport. Procedure code A0999 is used to bill for specialized vehicle transports. Transport claims may be submitted electronically with a short description of the client’s physical condition in the comment field. If the client’s condition cannot be documented, providers must file a paper claim with supporting documentation.

**Refer to:** Subsection 2.2.5.2, “Air or Specialized Vehicle Transports” in this handbook for more information about how to meet the specific criteria for reimbursement consideration for air or specialized transport claims.

### 2.4.6 Emergency Transport Billing

Emergency transport is a benefit when billed with the ET modifier and the most appropriate emergency medical condition codes. The ET modifier is required for every detail on an emergency transport claim, but is not required to be listed in the first position on the claim line.

The following procedure codes are for emergency transport:

<table>
<thead>
<tr>
<th>Procedure Codes</th>
</tr>
</thead>
<tbody>
<tr>
<td>A0382</td>
</tr>
<tr>
<td>A0434</td>
</tr>
</tbody>
</table>

*A0425 is denied if it is billed without procedure code A0427, A0429, A0433, or A0434.*
One of the following emergency medical condition code is required on all emergency ambulance claims and must be listed in Box 21 of the CMS-1500 claim form:

<table>
<thead>
<tr>
<th>Emergency Medical Condition Codes</th>
</tr>
</thead>
<tbody>
<tr>
<td>B9689</td>
</tr>
<tr>
<td>G8929</td>
</tr>
<tr>
<td>R002</td>
</tr>
<tr>
<td>R109</td>
</tr>
<tr>
<td>R569</td>
</tr>
<tr>
<td>T1491XA</td>
</tr>
<tr>
<td>T68XXXA</td>
</tr>
<tr>
<td>T82519A</td>
</tr>
<tr>
<td>Z9981</td>
</tr>
</tbody>
</table>

While ICD-10-CM codes are not precluded from use on ambulance claims, they are currently not required (per the Health Insurance Portability and Accountability Act [HIPAA] of 1996) on most ambulance claims and the use of these codes generally does not trigger a payment or a denial of a claim.

Claims for emergency transports that are denied for not meeting the emergency criteria will be considered on appeal with additional documentation to support the emergency nature of the transport. Claims that have denied for not meeting emergency transport criteria cannot be appealed for reimbursement as a nonemergency claim.

Refer to: Subsection 2.2.1, “Emergency Ambulance Transport Services” in this handbook.

### 2.4.7 Nonemergency Transport Billing

The following procedure codes are used when billing for nonemergency ambulance services:

<table>
<thead>
<tr>
<th>Procedure Codes</th>
</tr>
</thead>
<tbody>
<tr>
<td>A0382</td>
</tr>
<tr>
<td>A0433</td>
</tr>
</tbody>
</table>

*A0425 is denied if it is billed without procedure code A0426, A0428, A0433, or A0434.

### 2.4.8 Extra Attendant

The use of additional attendants (procedure code A0424) must be related to extraordinary circumstances when the basic crew is unable to transport the client safely.

An extra attendant on a nonemergency transport must be prior authorized. On an emergency transport, the billing provider’s medical documentation must clearly indicate the services the attendant performs along with a rationale for the services to indicate medical necessity of the attendant.

The information supporting medical necessity must be kept in the billing provider’s medical record and is subject to retrospective review.

Situations when an extra attendant may be required beyond the basic crew include, but are not limited to:

- Necessity of additional special medical equipment or treatment en route to destination (describe what special treatment and equipment is required and why it requires an attendant).
- Client behavior that may be a danger to self or ambulance crew or that requires, or may require, restraints.
- Extreme obesity of client (provide weight and client’s functional limitations).
• The extra attendant must be certified by DSHS to provide emergency medical services.

• The use of an extra attendant for air transport is not a benefit of Texas Medicaid. Claims submitted with procedure code A0424 will be denied if billed with air transports (procedure code A0430 or A0431).

2.4.8.1 Emergency Transports
Emergency transports that use an extra attendant do not require prior authorization. Modifier ET must be billed with the extra attendant procedure code A0424.

The billing provider’s medical documentation must clearly indicate the services the attendant performed along with rationale for the services to indicate the medical necessity of having the attendant. The billing provider must keep the information that supports medical necessity in the client’s medical record, which will be subject to retrospective review.

When more than one client is transported at the same time in the same vehicle, the use of an extra attendant may be required when each client who is being transported requires medical attention or close monitoring.

2.4.8.2 Nonemergency Transports
Prior authorization is required when an extra attendant is needed for any nonemergency transport. When a client’s condition changes, such as a need for oxygen or an extra attendant for transport, the prior authorization request must be updated.

To receive prior authorization, the requesting provider must prove medical necessity and identify attendant services that could not be provided by the basic crew. The information supporting medical necessity must be kept in the requesting provider’s medical record and is subject to retrospective review.

Texas Medicaid does not reimburse for an extra attendant based only on an ambulance provider’s internal policy.

2.4.9 Night Call
Texas Medicaid does not reimburse an extra charge for a night call.

2.4.10 Waiting Time
Procedure code A0420 may be billed when it is the general billing practice of local ambulance companies to charge for unusual waiting time (longer than 30 minutes). Providers must use the following procedures:

• Separate charges must be billed for all clients, Medicaid and non-Medicaid, for unusual waiting time.

• The circumstances requiring waiting time and the exact time involved must be documented in Block 24 of the CMS-1500 paper claim form.

• The amount charged for waiting time must not exceed the charge for a one-way transfer.

Important: Waiting time is reimbursed up to one hour.

2.4.11 Appeals
Only a denial of prior authorization may be appealed. Clients may appeal prior authorization request denials by contacting TMHP Client Notification at 1-800-414-3406. The Non-emergency Ambulance Prior Authorization Request form is not considered to be documentation after the service has been rendered.

Claims denied due to an inappropriate emergency medical condition code may be resubmitted with the appropriate emergency medical condition code.
On appeal, supporting documentation is critical for determining the client's condition at the time of transport. Ambulance providers who file paper claims must include all information that supports the reason for the transport and attach a copy of the run sheet to the claim. The EMT who transported the client must sign the documentation.

**Refer to:** Subsection 2.3, “Documentation Requirements” in this handbook.

Medicaid clients have the right to request a Fair Hearing within 90 days of the date of the denial action.

### 2.4.12 Relation of Service to Time of Death

Medicaid benefits cease at the time of the client’s death. However, if the client dies in the ambulance while en route to the destination, Texas Medicaid covers the transport. If a physician pronounces the client dead after the ambulance is called, Texas Medicaid covers the ambulance service (base rate plus mileage) to the point of pick up. Providers must indicate the date and time the client died in Block 19 of the CMS-1500 paper claim form. If a physician or coroner pronounces the client dead before the ambulance is called, the service is not covered.

Equipment and nondisposable supplies are included in the base rate. These items are not separately reimbursable and are considered part of another procedure. Therefore, equipment and supplies cannot be billed to the client.

### 2.5 Claims Resources

Providers may refer to the following sections or forms when filing claims:

<table>
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<th>Location</th>
</tr>
</thead>
<tbody>
<tr>
<td>Acronym Dictionary</td>
<td>“Appendix C: Acronym Dictionary” (Vol. 1, General Information)</td>
</tr>
<tr>
<td>Automated Inquiry System (AIS)</td>
<td>Subsection A.10, “TMHP Telephone and Fax Communication” in “Appendix A: State, Federal, and TMHP Contact Information”</td>
</tr>
<tr>
<td>CMS-1500 Paper Claim Filing Instructions</td>
<td>Subsection 6.5, “CMS-1500 Paper Claim Filing Instructions” in “Section 6: Claims Filing” (Vol. 1, General Information)</td>
</tr>
<tr>
<td>State, federal, and TMHP contact information</td>
<td>“Appendix A: State, Federal, and TMHP Contact Information” (Vol. 1, General Information)</td>
</tr>
<tr>
<td>TMHP Electronic Data Interchange (EDI)</td>
<td>“Section 3: TMHP Electronic Data Interchange (EDI)” (Vol. 1, General Information)</td>
</tr>
</tbody>
</table>

### 2.6 Contact TMHP

The TMHP Contact Center at 1-800-925-9126 is available Monday through Friday from 7 a.m. to 7 p.m., Central Time.

### 3 Forms

The following linked forms can also be found on the Forms page of the Provider section of the TMHP website at [www.tmhp.com](http://www.tmhp.com):

<table>
<thead>
<tr>
<th>Forms</th>
</tr>
</thead>
<tbody>
<tr>
<td>Non-emergency Ambulance Prior Authorization Request</td>
</tr>
</tbody>
</table>
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The following linked claim form examples can also be found on the Claim Form Examples page of the Provider section of the TMHP website at www.tmhp.com:

<table>
<thead>
<tr>
<th>Claim Form Examples</th>
</tr>
</thead>
<tbody>
<tr>
<td>Ambulance Non-emergency Transport</td>
</tr>
<tr>
<td>Ambulance Emergency Transport from Residence to Hospital</td>
</tr>
<tr>
<td>Ambulance Emergency Transport from Scene of Accident to Hospital</td>
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1 General Information

The information in this handbook is intended for the Case Management for the Blind Children’s Vocational Discovery and Development Program (BCVDDP), Case Management for Children and Pregnant Women, and services provided by a licensed clinical social worker (LCSW), licensed marriage and family therapist (LMFT), licensed professional counselor (LPC), psychologist, physician, advanced practice registered nurse (APRN), physician assistant (PA), or providers of intellectual and developmental disability (IDD) case management, mental health targeted case management, and mental health rehabilitative services.

All providers are required to report suspected child abuse or neglect as outlined in subsection 1.7.1.2, “Reporting Child Abuse or Neglect” in “Section 1: Provider Enrollment and Responsibilities” (Vol. 1, General Information) and subsection 1.7.1.5, “Training” in “Section 1: Provider Enrollment and Responsibilities” (Vol. 1, General Information).

Important: All providers are required to read and comply with “Section 1: Provider Enrollment and Responsibilities” (Vol. 1, General Information). In addition to required compliance with all requirements specific to Texas Medicaid, it is a violation of Texas Medicaid rules when a provider fails to provide health-care services or items to Medicaid eligible persons in accordance with accepted medical community standards and standards that govern occupations, as explained in Title 1 Texas Administrative Code (TAC) §371.1659. Accordingly, in addition to being subject to sanctions for failure to comply with the requirements that are specific to Texas Medicaid, providers can also be subject to Texas Medicaid sanctions for failure to deliver, at all times, health-care items and services to Medicaid eligible persons in full accordance with all applicable licensure and certification requirements including, without limitation, those related to documentation and record maintenance.

Refer to: “Section 1: Provider Enrollment and Responsibilities” (Vol. 1, General Information).

1.1 Payment Window Reimbursement Guidelines for Services Preceding an Inpatient Admission

According to the three-day and one-day payment window reimbursement guidelines, most professional and outpatient diagnostic and nondiagnostic services that are rendered within the designated timeframe of an inpatient hospital stay and are related to the inpatient hospital admission will not be reimbursed separately from the inpatient hospital stay if the services are rendered by the hospital or an entity that is wholly owned or operated by the hospital.

These reimbursement guidelines do not apply in the following circumstances:

- The professional services are rendered in the inpatient hospital setting.
- The hospital and the physician office or other entity are both owned by a third party, such as a health system.
- The hospital is not the sole or 100-percent owner of the entity.

Refer to: Subsection 3.7.4.14, “Payment Window Reimbursement Guidelines” in the Inpatient and Outpatient Hospital Services Handbook (Vol. 2, Provider Handbooks) for additional information about the payment window reimbursement guidelines.
2 Blind Children’s Vocational Discovery and Development Program (BCVDDP)

2.1 Overview
BCVDDP services are provided to help children who are blind and visually impaired to develop their individual potential. This program offers a wide range of services that are tailored to each child and their family’s needs and circumstances. By working directly with the entire family, this program can help children develop the concepts and skills needed to realize their full potential.

BCVDDP services include the following:
- Assisting the child in developing the confidence and competence needed to be an active part of their community
- Providing support and training to children in understanding their rights and responsibilities throughout the educational process
- Assisting family and children in the vocational discovery and development process
- Providing training in areas like food preparation, money management, recreational activities, and grooming
- Supplying information to families about additional resources

2.2 Enrollment
Texas Health and Human Services Commission (HHSC) Blind Children’s Vocational Discovery and Development Program (BCVDDP) is the Medicaid provider of case management for persons who are 22 years of age and younger and blind or visually impaired. Providers must meet educational and work experience requirements that are commensurate with their job responsibilities and must be trained in BCVDDP case management activities.

Refer to: Subsection 1.1, “Provider Enrollment” in “Section 1: Provider Enrollment and Responsibilities” (Vol. 1, General Information) for more information about procedures for enrolling as a Medicaid provider.

2.3 Services, Benefits, Limitations, and Prior Authorization
Services eligible for reimbursement are limited to one contact per month, per person, regardless of the number of contacts that are made during the month. HHSC BCVDDP providers should bill procedure code G9012.

A contact is defined as “an activity performed by a case manager with the person or organization on behalf of the person to locate, coordinate, and monitor necessary services.”

Refer to: Subsection A.8, “Texas Health and Human Services Commission Blind Children’s Vocational Discovery and Development Program (BCVDDP)” in “Appendix A: State, Federal, and TMHP Contact Information” (Vol. 1, General Information).

2.3.1 Prior Authorization
Prior authorization is not required for BCVDDP case management services.

2.4 Documentation Requirements
All services require documentation to support the medical necessity of the service rendered, including BCVDDP services.

BCVDDP services are subject to retrospective review and recoupment if documentation does not support the service billed.
2.5 Claims Filing and Reimbursement

BCVDDP case management services must be submitted to the Texas Medicaid & Healthcare Partnership (TMHP) in an approved electronic format or on the CMS-1500 paper claim form. Providers may purchase CMS-1500 paper claim forms from the vendor of their choice. TMHP does not supply the forms.

When completing a CMS-1500 paper claim form, all required information must be included on the claim, as TMHP does not key any information from claim attachments. Superbills or itemized statements are not accepted as claim supplements. Providers must not submit a claim when or after the person turns 21 years of age.

Claims may be submitted up to 365 days from the date of service in accordance with 1 TAC §354.1003. Any child who has a suspected or diagnosed visual impairment may be referred to BCVDDP. HHSC BCVDDP assesses the impact the visual impairment has on the child’s development and provides blindness-specific services to increase the child’s skill level in the areas of independent living, communication, mobility, social, recreational, and vocational discovery and development. For more information, visit the HHS website at https://hhs.texas.gov/.

Providers can refer to the Online Fee Lookup (OFL) or the applicable fee schedule on the TMHP website at www.tmhp.com.

Texas Medicaid implemented mandated rate reductions for certain services. The OFL and static fee schedules include a column titled “Adjusted Fee” to display the individual fees with all mandated percentage reductions applied.

Additional information about rate changes is available on the TMHP website at www.tmhp.com/resources/rate-and-code-updates/rate-changes.

Refer to: “Section 3: TMHP Electronic Data Interchange (EDI)” (Vol. 1, General Information) for information about electronic claims submissions.

Subsection 6.1, “Claims Information” in “Section 6: Claims Filing” (Vol. 1, General Information) for general information about claims filing.


Subsection 2.9, “Federal Medical Assistance Percentage (FMAP)” in “Section 2: Texas Medicaid Fee-for-Service Reimbursement” (Vol. 1, General Information) for federal matching percentage.

3 Case Management for Children and Pregnant Women

3.1 Overview

Case management services are provided to help Medicaid eligible persons gain access to necessary medical, social, educational, and other services. Case managers assess a person’s need for these services and then develop a service plan to address those needs.

3.1.1 Eligibility

To be eligible for services, a person must:

- Be eligible for Texas Medicaid.
- Be a pregnant woman who has a high-risk condition or a child (birth through 20 years of age) who has a health condition or health risk.
• Need assistance in gaining access to necessary medical, social, educational and other services related to their health condition, health risk, or high-risk condition.

• Want to receive case management services.

Pregnant women who have a high-risk condition are defined as those who have a medical or psychosocial condition that places them and their fetuses at a greater than average risk for complications, either during pregnancy, delivery, or following birth. Children with a health condition are defined as children who have a health condition or health risk or children who have or are at risk for a medical condition, illness, injury, or disability that results in the limitation of function, activities, or social roles in comparison with healthy same-age peers in the general areas of physical, cognitive, emotional, or social growth and development.

3.1.2 Referral Process
To refer a Medicaid eligible person for Case Management for Children and Pregnant Women services, providers may do one of the following:

• Visit https://hhs.texas.gov/doing-business-hhs/provider-portals/health-services-providers/case-management-providers-children-pregnant-women to obtain a referral form.

• Call THSteps toll free at 1-877-847-8377 from 8 a.m. to 8 p.m., Central Time, Monday through Friday.

• Contact a Case Management for Children and Pregnant Women provider directly at https://hhs.texas.gov/doing-business-hhs/provider-portals/health-services-providers/case-management-providers-children-pregnant-women. A case management provider will contact the family to offer a choice of providers and obtain information necessary to request prior authorization for case management services.

A referral for Case Management for Children and Pregnant Women services can be received from any source.

3.2 Enrollment
Enrollment for Case Management for Children and Pregnant Women providers is a two-step process.

Step 1
Potential providers must submit a Health and Human Services Commission (HHSC) Case Management for Children and Pregnant Women provider application to the HHSC Health Screening and Case Management Unit.

Both registered nurses who have an associate’s, bachelor’s, or advanced degree and social workers who have a bachelor’s or advanced degree are eligible to become case managers if they are currently licensed by their respective Texas licensure boards and the license is not temporary in nature. Registered nurses with associate degrees must also have at least two years of cumulative, paid, full-time work experience or two years of supervised full-time, educational, internship/practicum experience in the past ten years. The experience must be with pregnant women or with children who are 20 years of age and younger. The experience must include assessing psychosocial and health needs and making community referrals for these populations. Registered nurses with bachelor or advance degrees and social workers do not have to meet any experience requirements.
For more information about provider qualifications and enrollment, contact HHSC at 1-512-458-7111, ext. 2168, visit the case management website at https://hhs.texas.gov/doing-business-hhs/provider-portals/health-services-providers/case-management-providers-children-pregnant-women, or write to the following address:

Health and Human Services Commission  
Case Management for Children and Pregnant Women  
PO Box 149347, MC 1938  
Austin, TX 78714-9347

**Note:** Before providing services, each case manager must attend HHSC case manager training. Training is conducted by DSHS regional staff.

**Step 2**

Upon approval by HHSC, potential providers must enroll as a Medicaid provider for Case Management for Children and Pregnant Women and submit a copy of their HHSC approval letter. Facility providers must enroll as a Case Management for Children and Pregnant Women group, and each eligible case manager must enroll as a performing provider for the group. Federally Qualified Health Center (FQHC) facilities that provide Case Management for Children and Pregnant Women services will use their FQHC number and should not apply for an additional provider number for Case Management for Children and Pregnant Women.

**Refer to:** Subsection 1.1, “Provider Enrollment” in “Section 1: Provider Enrollment and Responsibilities” (Vol. 1, General Information) for more information about the procedures for enrolling as a Medicaid provider.

### 3.3 Services, Benefits, Limitations, and Prior Authorization

Case Management for Children and Pregnant Women services are limited to one contact per day per person. Additional provider contacts on the same day are denied as part of another service rendered on the same day.

Procedure code G9012 is to be used for all Case Management for Children and Pregnant Women services. Modifiers are used to identify which service component is provided.

<table>
<thead>
<tr>
<th>Service</th>
<th>Contact Code</th>
</tr>
</thead>
<tbody>
<tr>
<td>Comprehensive visit</td>
<td>G9012 with modifier U5 and modifier U2</td>
</tr>
<tr>
<td>Follow-up face-to-face</td>
<td>G9012 with modifier U5 and modifier TS</td>
</tr>
<tr>
<td>Follow-up telephone</td>
<td>G9012 with modifier TS</td>
</tr>
</tbody>
</table>

Providers must adhere to Case Management for Children and Pregnant Women program rules, policies, and procedures.

**Note:** Case Management for Children and Pregnant Women providers are not required to file claims with other health insurance before filing with Medicaid.

**Reminder:** Billable services are defined in program rule 25 TAC §27.11.

Case Management for Children and Pregnant Women services are not billable when a person is an inpatient at a hospital or other treatment facility.

Reimbursement will be denied for services rendered by providers who have not been approved by HHSC.
3.3.1 Prior Authorization

All services must be prior authorized. One comprehensive visit is approved for all Medicaid eligible persons. Follow-up visits are authorized based on contributing factors. Additional visits can be requested and may be authorized based on a continuing need for services. A prior authorization number is required on all claims for Case Management for Children and Pregnant Women services.

Note: Prior authorization is a condition of reimbursement, not a guarantee of payment.

Approved case management providers may submit requests for prior authorization from HHSC on the Department of State Health Services (DSHS) website at https://www.cpwforms.dshs.state.tx.us/cpw/.

3.4 Technical Assistance

Providers may contact HHSC program staff as needed for assistance with program concerns. Providers should contact TMHP provider relations staff as needed for assistance with claims problems or concerns.

3.4.1 Assistance with Program Concerns

Providers who have questions, concerns, or problems with program rule, policy, or procedure may contact HHSC program staff. Contact names and numbers can be obtained from the case management website at https://hhs.texas.gov/doing-business-hhs/provider-portals/health-services-providers/case-management-providers-children-pregnant-women, or by calling 1-512-458-7111, Ext. 2168.

Regional DSHS staff make routine contact with providers to ensure providers are delivering services as required.

3.5 Documentation Requirements

All services require documentation to support the medical necessity of the service rendered, including Case Management for Children and Pregnant Women services.

Case Management for Children and Pregnant Women services are subject to retrospective review and recoupment if documentation does not support the service billed.

3.6 Claims Filing and Reimbursement

3.6.1 Claims Information

Case Management for Children and Pregnant Women services must be submitted to TMHP in an approved electronic format or on the CMS-1500 paper claim form. Providers may purchase CMS-1500 paper claim forms from the vendor of their choice. TMHP does not supply the forms.

When completing a CMS-1500 paper claim form, all required information must be included on the claim, as TMHP does not key any information from claim attachments. Superbills, or itemized statements, are not accepted as claim supplements.

Case Management for Children and Pregnant Women providers are reimbursed in accordance with 1 TAC §355.8401. Providers can refer to the OFL or the applicable fee schedule on the TMHP website at www.tmhp.com.

Texas Medicaid implemented mandated rate reductions for certain services. The OFL and static fee schedules include a column titled “Adjusted Fee” to display the individual fees with all mandated percentage reductions applied.

Additional information about rate changes is available on the TMHP website at www.tmhp.com/resources/rate-and-code-updates/rate-changes.
3.6.2 Managed Care Clients

Case Management for Children and Pregnant Women services are carved out of Medicaid managed care and must be billed to TMHP for payment consideration. Carved-out services are those that are rendered to Medicaid managed care clients, but are administered by TMHP and not the client’s managed care organization (MCO).

4 Outpatient Mental Health Services

Outpatient mental health services are used for the treatment of mental illness and emotional disturbances in which the clinician establishes a professional contract with the person and, utilizing therapeutic interventions, attempts to alleviate the symptoms of mental illness or emotional disturbance, and reverse, change, or ameliorate maladaptive patterns of behavior.

Outpatient mental health services include psychiatric diagnostic evaluation, psychotherapy (including individual, group, or family psychotherapy), psychological, neurobehavioral, or neuropsychological testing, pharmacological management services, and electroconvulsive therapy (ECT).

Outpatient mental health services are benefits when provided in the office, home, skilled nursing or intermediate care facility (SNF/ICF), outpatient hospital, extended care facility (ECF), or in other locations.

Outpatient mental health services are benefits of Texas Medicaid when provided to persons who are experiencing a mental health issue that is causing distress, dysfunction, and/or maladaptive functioning as a result of a confirmed or suspected psychiatric condition as defined in the current edition of the American Psychiatric Association’s Diagnostic and Statistical Manual of Mental Disorders (DSM).

Note: Claims will require the corresponding diagnosis code(s) from the current edition of the International Classification of Diseases (ICD).

4.1 Provider Enrollment

Mental health service providers include physicians, PAs, APRNs, LCSWs, LMFTs, LPCs, psychologists, licensed psychological associates (LPAs), provisionally licensed psychologists (PLPs), post-doctoral fellows, and pre-doctoral psychology interns.

Refer to: Subsection 1.1, “Provider Enrollment” in “Section 1: Provider Enrollment and Responsibilities” (Vol. 1, General Information) for more information about procedures for enrolling as a Medicaid provider.
4.1.1 Physicians
To enroll in Texas Medicaid to provide medical services, physicians (doctor of medicine [MD] or doctor of osteopathy [DO]) and doctors (doctor of dental medicine [DMD], doctor of dental surgery [DDS], doctor of optometry [OD], and doctor of podiatric medicine) must be authorized by the licensing authority of their profession to practice in the state where the services are performed at the time they are provided.

Providers cannot be enrolled in Texas Medicaid if their licenses are due to expire within 30 days. A current Texas license must be submitted.

All physicians except gynecologists, pediatricians, pediatric subspecialists, pediatric psychiatrists, and providers performing only THSteps medical or dental checkups must be enrolled in Medicare before enrolling in Medicaid. TMHP may waive the Medicare enrollment prerequisite for pediatricians or physicians whose type of practice and service may never be billed to Medicare.

4.1.2 Physician Assistants (PAs)
To enroll in Texas Medicaid, a PA must be licensed as a PA and be recognized as a PA by the Texas Physician Assistant Board. All PAs are enrolled within the categories of practice as determined by the Texas Medicaid Board. PAs can enroll as an individual, group, or as a performing provider into a clinic/group practice. If enrolling into a Medicare enrolled clinic/group practice, Medicare enrollment is required.

4.1.3 Advanced Practice Registered Nurses (APRNs)
To enroll in Texas Medicaid, whether as an individual or as part of a group, a nurse practitioner (NP) or clinical nurse specialist (CNS) recognized as an APRN must be licensed by the Texas Board of Nursing (TBON). NP/CNSs must also be enrolled in Medicare or obtain a pediatric practice exemption from TMHP Provider Enrollment. If a pediatric-based NP/CNSs is enrolling as part of a Medicare-enrolled group, then the NP/CNSs must also be enrolled in Medicare.

Providers that hold a temporary license are not eligible to enroll in Medicaid. NP/CNSs cannot be enrolled if their license is due to expire within 30 days. A current license must be submitted.

4.1.4 Licensed Clinical Social Workers (LCSWs)
To enroll in Texas Medicaid, whether as an individual or as part of a group, an LCSW must be licensed by the Texas State Board of Social Worker Examiners. LCSWs must also be enrolled in Medicare or obtain a pediatric practice exemption from TMHP Provider Enrollment. If a pediatric-based LCSW is enrolling as part of a Medicare-enrolled group, then the LCSW must also be enrolled in Medicare.

Providers that hold a temporary license are not eligible to enroll in Medicaid. LCSWs cannot be enrolled if their license is due to expire within 30 days. A current license must be submitted.

4.1.5 Licensed Marriage and Family Therapists (LMFTs)
To enroll in Texas Medicaid, whether as an individual or as part of a group, an LMFT must be licensed by the Texas State Board of Examiners of Licensed Marriage and Family Therapists. LMFTs are covered as Medicaid-only providers; therefore, enrollment in Medicare is not a requirement. LMFTs can enroll as part of a clinic/group practice whether or not they are enrolled in Medicare. Providers that hold a temporary license are not eligible to enroll in Medicaid. LMFTs cannot be enrolled if their license is due to expire within 30 days. A current license must be submitted.

4.1.6 Licensed Professional Counselors (LPCs)
To enroll in Texas Medicaid, whether as an individual or as part of a group, an LPC must be licensed by the Texas Board of Examiners of Professional Counselors. LPCs are covered as Medicaid-only providers; therefore, enrollment in Medicare is not a requirement. LPCs can enroll as part of a clinic/group practice
whether or not they are enrolled in Medicare. Providers that hold a temporary license are not eligible to enroll in Medicaid. LPCs cannot be enrolled if their license is due to expire within 30 days. A current license must be submitted.

4.1.7 Psychologists
To enroll in Texas Medicaid, whether as an individual or as part of a group, a psychologist must be licensed by the Texas State Board of Examiners of Psychologists (TSBEP). Psychologists must also be enrolled in Medicare or obtain a pediatric practice exemption from TMHP Provider Enrollment. If a pediatric-based psychologist is enrolling as part of a Medicare-enrolled group, then the psychologist must also be enrolled in Medicare. Psychologists cannot be enrolled if they have a license that is due to expire within 30 days. A current license must be submitted. Texas Medicaid accepts temporary licenses for psychologists.

4.1.8 Licensed Psychological Associates (LPAs)
LPAs must be licensed by TSBEP. LPAs are expected to abide by their scope and standards of practice. Services performed by an LPA are a Medicaid-covered benefit when the following conditions are met:

- The services must be performed under the required supervision of a licensed, Medicaid-enrolled psychologist.
- The supervising psychologist must be in the same office, building, or facility when the service is provided and must be immediately available to furnish assistance and direction.
- The LPA performing the service must be an employee of either the licensed psychologist or the legal entity that employs the licensed psychologist.

Psychological services provided by an LPA must be billed under the supervising psychologist’s Medicaid identifier or the Medicaid identifier of the legal entity employing the supervising psychologist.

4.1.9 Provisionally Licensed Psychologists (PLPs)
PLPs must be licensed by TSBEP. A PLP may perform all of the services that are benefits of Texas Medicaid when the services are performed by a psychologist.

PLPs are expected to abide by their scope and standards of practice. Services performed by a PLP are a Medicaid-covered benefit when the following conditions are met:

- The services must be performed under the required supervision of a licensed psychologist in accordance with the TSBEP guidelines.
- The supervising psychologist must be in the same office, building, or facility when the service is provided and must be immediately available to furnish assistance and direction.
- The PLP who is performing the service must be an employee of either the licensed psychologist or the legal entity that employs the licensed psychologist.

The TSBEP requires a PLP to work under the required supervision of a licensed psychologist and does not allow a PLP to engage in independent practice. Therefore, a PLP will not be independently enrolled in the Medicaid program and must provide services under the delegating psychologist’s provider identifier.

Psychological services provided by a PLP must be billed under the supervising psychologist’s Medicaid identifier or the Medicaid identifier of the legal entity employing the supervising psychologist.

4.1.10 Post-Doctoral Fellows
Post-doctoral psychology fellows who satisfy the provisional licensure examination requirements but have not yet been awarded the PLP designation are eligible to perform delegated psychological services within their scope of practice and under the required supervision of a licensed psychologist.
Psychology interns are not independently enrolled in Texas Medicaid; therefore, they do not have a provider identifier.

Psychological services provided by an intern must be billed under the supervising psychologist’s Medicaid identifier or the Medicaid identifier of the legal entity employing the supervising psychologist.

### 4.1.11 Pre-doctoral Psychology Interns

Pre-doctoral psychology interns who are participating in a pre-doctoral psychology internship at a site that is a member of the Association of Psychology Postdoctoral and Internship Centers (APPIC) are eligible to perform delegated psychological services within their scope of practice and under the required supervision of a licensed psychologist.

Psychology interns are not independently enrolled in Texas Medicaid; therefore, they do not have a provider identifier.

### 4.2 Services, Benefits, Limitations

The following procedure codes may be reimbursed for outpatient mental health services:

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*Note: Add-on procedure codes indicated with asterisk must be billed with the appropriate primary procedure code.*

The following psychotherapy procedure codes are limited to 30 visits per calendar year. Additional services require prior authorization:

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*Add-on procedure code must be billed with the appropriate E/M code*

The following add-on procedure codes may be used for prolonged psychotherapy services:

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Procedure codes 90791, 90792, 90832, 90833, 90834, 90836, 90837, 90838, 90846, 90847, and 90853 are limited to the following diagnosis codes:

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<td>T7621XS</td>
<td>T7622XA</td>
<td>T7622XD</td>
<td>T7622XS</td>
<td>T7631XA</td>
</tr>
<tr>
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<td>T7631XS</td>
<td>T7632XA</td>
<td>T7632XD</td>
<td>T7632XS</td>
<td>T7651XA</td>
<td>T7651XD</td>
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</tr>
<tr>
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<td>T7652XD</td>
<td>T7652XS</td>
<td>T7661XA</td>
<td>T7661XD</td>
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<td>T7662XA</td>
<td>T7662XD</td>
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<td>Z634</td>
<td>Z644</td>
<td>Z658</td>
<td>Z72810</td>
<td>Z72811</td>
<td>Z736</td>
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<td>Z73810</td>
<td>Z73811</td>
<td>Z73812</td>
<td>Z73819</td>
<td>Z781</td>
<td>Z818</td>
<td>Z8651</td>
<td>Z8659</td>
</tr>
</tbody>
</table>

*Add-on procedure codes must be billed with the appropriate primary code*
4.2.1 Telemedicine and Telehealth

Certain outpatient mental health services may be provided by distant site providers through telemedicine or telehealth when billed with modifier 95.

Mental health services delivered through telemedicine or telehealth do not require a patient site presenter unless the person is experiencing a mental health emergency.

Refer to: The Telecommunication Services Handbook (Vol. 2, Provider Handbooks) for more information about telemedicine and telehealth.

4.2.2 Psychotherapy

Individual psychotherapy is therapy that focuses on a single person.

Group psychotherapy is a type of psychotherapy that involves one or more therapists working with several persons at the same time.

Family psychotherapy is therapy that focuses on the dynamics of the family unit where the goal is to strengthen the family’s problem solving and communication skills.

Providers must bill a modifier to identify a separate and distinct service when performing individual psychotherapy (procedure codes 90832, 90834, and 90837) and family psychotherapy (procedure codes 90846 or 90847) on the same day for the same person. When billing for these services, providers must submit the family psychotherapy procedure code with the modifier on the claim to indicate that the procedure or service was distinct or independent from other services performed on the same day for the same person. Documentation that supports the provision of distinct or independent services must be maintained in the person’s medical record and made available to Texas Medicaid upon request.

Prolonged psychotherapy services delivered in addition to procedure code 90837 should be billed using the appropriate prolonged services add-on code (procedure code 99354 or 99355)

Note: The add-on codes may only be billed by physician, APRN, or PA providers.

Refer to: Subsection 9.2.56.5, “Prolonged Physician Services” in the Medical and Nursing Specialists, Physicians, and Physician Assistants Handbook (Vol. 2, Provider Handbooks) for more information about prolonged physician services.

Psychotherapy (individual, family, or group) is limited to 4 hours per person, per day.

Psychotherapy is limited to 30 individual, group, or family psychotherapy visits per person, per calendar year. Additional psychotherapy services must be prior authorized. Prior authorization requests in increments of up to 10 additional visits may be considered. The request must be submitted on an Outpatient Mental Health Services Request Form and include the following information:

- Identifying information for the person receiving services
- Provider name and identifier
- Current DSM diagnosis(es)
- Current psychotropic medications
- Current symptoms requiring additional psychotherapy
- Treatment plan, including measurable short term goals, specific therapeutic interventions utilized, and measurable expected outcomes of therapy
- Number and type of services requested and anticipated dates that the services will be provided
• Indication of court-ordered or DFPS-directed services

Providers with an established relationship with a person receiving services must request prior authorization when they determine the person is approaching 30 psychotherapy visits for the calendar year. If the person changes providers during the year and the new provider is unable to obtain complete information on the person’s previous treatment history, providers are encouraged to obtain prior authorization before rendering services. Requests submitted on the same day as the initial session with a new provider will be considered based on medical necessity criteria.

Providers must bill the preponderance of each half hour of psychotherapy and indicate the number of units on the claim form.

LMFTs must bill with modifier U8 to differentiate from LPCs.

Supporting documentation for individual, family, or group psychotherapy must include:

• Start and end time of session
• Modality or modalities utilized
• Frequency of psychotherapy sessions
• Clinical notes for each encounter must include: diagnosis; symptoms; functional status; focused mental status examination, if indicated; treatment plan, prognosis, and progress; name, signature and credentials of person performing the service

4.2.2.1 Family Psychotherapy

Family psychotherapy may be provided to Medicaid eligible persons 20 years of age and younger using procedure code 90846 or persons of any age using procedure code 90847.

Family psychotherapy is only reimbursable for one Medicaid eligible person per session regardless of the number of family members present per session.

Family psychotherapy for Medicaid eligible persons 20 years of age and younger may be provided to the child’s parent(s), foster parent(s), or legal guardian without the child present, as clinically appropriate, using procedure code 90846. Parent- or guardian-only sessions may be indicated when addressing sensitive topics such as parenting challenges or related stressors that would be inappropriate to discuss with the child present at the session.

Only the following specific relatives are allowed to participate in family psychotherapy services:

• Biological parent, foster parent, or legal guardian
• Child
• Grandfather or grandmother
• Sibling (biological, foster, or kinship)
• Uncle, aunt, nephew, or niece
• First cousin or first cousin once removed
• Stepfather, stepmother, stepbrother, or stepsister

4.2.2.2 Treatment for Alzheimer’s Disease and Dementia

Psychotherapy for persons with Alzheimer’s disease or dementia may be a benefit of Texas Medicaid for persons with very mild or mild cognitive decline.

Documentation to support the treatment for Alzheimer’s disease or dementia must be maintained in the person’s medical record and may be subject to retrospective review. Psychotherapy services must not be continued if no longer beneficial to the person due to diminished cognitive functioning.
4.2.3 Delegated Services

Services provided by a psychologist, LPA, PLP, psychology intern, or post-doctoral fellow must be billed with a modifier on each detail. Psychological services provided by an LPA, PLP psychology intern, or post-doctoral fellow must be billed under the supervising psychologist’s Medicaid provider identifier or the Medicaid identifier of the legal entity employing the supervising psychologist.

Services performed by a LPA or PLP will be reimbursed at 70 percent of the psychologist rate. Services performed by the psychology intern or post-doctoral fellow will be reimbursed at 50 percent of the psychologist rate.

The following modifiers are to be used with procedure codes for licensed psychologist and delegated services:

<table>
<thead>
<tr>
<th>Modifier</th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td>AH</td>
<td>Identifies service provided by a clinical psychologist</td>
</tr>
<tr>
<td>UB</td>
<td>Identifies service provided by a pre-doctoral psychology intern or post-doctoral psychology fellow</td>
</tr>
<tr>
<td>UC</td>
<td>Identifies service provided by an LPA</td>
</tr>
<tr>
<td>U9</td>
<td>Identifies service provided by a PLP</td>
</tr>
</tbody>
</table>

Claims submitted without a modifier or with two of these modifiers on the same detail will be denied.

Only the LCSW, LMFT, LPC, APRN, or PA actually performing the mental health service may bill Texas Medicaid. The LCSW, LMFT, LPC, APRN, or PA must not bill for services performed by people under his or her supervision.

4.2.4 Pharmacological Management

Pharmacological management is the in-depth management of psychopharmacological agents to treat a person’s mental health symptoms.

Pharmacological management is a physician service and cannot be provided by a non-physician or “incident to” a physician service, with the exception of APRNs and PAs whose scope of license in this state permits them to prescribe.

Pharmacological management is limited to one service per day, per person, by any provider in any setting.

The treating provider should use the most appropriate E/M code for the pharmacological management visit depending on the place of service and complexity of the person’s condition, along with modifier UD to designate the visit as primarily focused on pharmacological management.

Supporting documentation for pharmacological management must include:

- A complete diagnosis utilizing diagnostic criteria from the current edition of the DSM
- Current list of medications
- Current psychiatric symptoms and problems, to include presenting mental status
- Problems, reactions, and side effects, if any, to medications
- Any medication modifications made during visit and the reasons for medication adjustments, changes, or discontinuation
- Desired therapeutic drug levels, if applicable, for medications requiring blood level monitoring, e.g. Lithium
- Current laboratory values, if applicable, for medications requiring monitoring for potential side effects, e.g. hyperglycemia caused by anti-psychotic medications
• Treatment goals

4.2.5 Electroconvulsive Therapy
Electroconvulsive therapy (ECT) is the induction of convulsions by the passage of an electric current through the brain used in the treatment of certain psychiatric disorders.

Individual psychotherapy, psychological testing, neurobehavioral testing, or neuropsychological testing billed in addition to ECT on the same day, by any provider will be denied as part of another procedure on the same day.

ECT billed in addition to psychiatric diagnostic evaluation, group psychotherapy, or family psychotherapy on the same day, by the same provider will be denied as part of another procedure.

4.2.6 Psychiatric Diagnostic Evaluation
Psychiatric diagnostic evaluation is an integrated biopsychosocial assessment, including history, mental status, and recommendations. Psychiatric diagnostic evaluation with medical services also includes a medical assessment, other physical examination elements as indicated, and may also include prescription of medications, and laboratory or other diagnostic studies.

A psychiatric diagnostic evaluation (without medical services) (procedure code 90791) may be reimbursed to physicians, psychologists, APRNs, PAs, LCSWs, LPCs, LMFTs, PLPs, psychology interns, and post-doctoral fellows.

A psychiatric diagnostic evaluation (with medical services) (procedure code 90792) may be reimbursed to physicians, APRNs, and PAs.

Psychiatric diagnostic evaluations (procedure codes 90791 or 90792) are limited to once per person, per rolling year, same provider in the office, home, outpatient hospital, or other settings, regardless of the number of professionals involved in the interview. Additional psychiatric diagnostic evaluations may be considered for prior authorization on a case-by-case basis when submitted on an Outpatient Mental Health Services Request Form with supporting documentation, including but not limited to:

• A court order or a Department of Family and Protective Services (DFPS) directive
• If a major change of status occurs

Supporting documentation for psychiatric diagnostic evaluations must include:
• Reason for referral and/or presenting problem
• Prior diagnoses and any prior treatment
• Other pertinent medical, social, and family history
• Clinical observations and results of mental status examination
• A complete diagnosis utilizing diagnostic criteria from the current edition of the DSM
• Recommendations, including expected long term and short term goals

4.2.7 Psychological, Neurobehavioral, and Neuropsychological Testing
Psychological, neurobehavioral, and neuropsychological testing involves the use of formal tests and other assessment tools to measure and assess a person’s emotional, and cognitive functioning in order to arrive at a diagnosis and guide treatment.

Neurobehavioral testing (procedure codes 96116 and 96121*) is limited to four hours per person, per day and eight hours per person, per calendar year.
Psychological testing (procedure codes 96130, 96131*, 96136, and 96137*) and neuropsychological testing (procedure codes 96132, 96133*, 96136, and 96137*) are limited to eight hours per person, per calendar year. Additional hours require prior authorization when medically necessary. The request must be submitted on an Outpatient Mental Health Services Request Form and include the following information:

- Identifying information for the person receiving services
- Provider name and identifier
- Current DSM diagnoses
- Indication of court-ordered or DFPS-directed services
- Type of testing requested (psychological, neurobehavioral, or neuropsychological) including specific procedure codes
- Rationale for requested testing, to include the current symptoms of the person receiving services
- Previous history and testing results

Psychological, neurobehavioral, and neuropsychological testing will not be reimbursed to an APRN or a PA. The most appropriate office encounter/visit procedure code must be billed. Mental health screening may be performed during an assessment by an APRN or a PA, but will not be reimbursed separately.

Psychological testing (procedure codes 96130, 96131*, 96136, and 96137*) or neuropsychological testing (procedure codes 96132, 96133*, 96136, and 96137*) may be reimbursed on the same date of service as an initial psychiatric diagnostic evaluation (procedure code 90791 or 90792).

Neurobehavioral testing (procedure code 96116) may not be reimbursed on the same date of service as an initial psychiatric diagnostic evaluation (procedure code 90791 or 90792) to the same provider.

Neurobehavioral testing (procedures codes 96116 and 96121*) will not be paid for the same date of service to the same provider as psychological testing (procedure codes 96130, 96131*, 96136 and 96137*) or neuropsychological testing (procedure codes 96132, 96133*, 96136, and 96137*). All documentation must be maintained by the provider in the person’s medical record.

The reimbursement for procedure codes 96116, 96121*, 96130, 96131*, 96132, 96133*, 96136, and 96137* includes the face-to-face testing and the scoring and interpretation of the results. The number of units of testing on the claim for procedure codes 96116, 96121*, 96130, 96131*, 96132, 96133*, 96136, and 96137* must be in accordance with the allowable activities outlined in each code description.

**Note:** Add-on procedure codes indicated with asterisk must be billed with the appropriate primary procedure code.

Assessment, treatment planning, and documentation time, including time to document test results in the person’s medical record, is not reimbursed separately. Reimbursement is included in the covered procedure codes.

### 4.2.7.1 Testing in Facilities

Psychological testing, neurobehavioral testing, or neuropsychological testing may be reimbursed when provided in a skilled nursing facility (SNF), intermediate care facility (ICF), or extended care facility (ECF) as clinically indicated. Testing may be indicated, for example, when a person has experienced significant change in mental status requiring specialized testing, or to evaluate a person’s competency to return to a community-based setting. Persons with well-established mental or cognitive issues do not require additional testing.
Psychological, neurobehavioral, or neuropsychological testing will not be reimbursed in a SNF, ICF, or ECF when conducted prior to the performance of initial intake assessments such as the Minimum Data Set or Preadmission Screening and Resident Review (PASRR) (a completed Level I Screening and a Level II Evaluation, as applicable).

Supporting documentation for psychological, neurobehavioral, or neuropsychological testing must include:

- Reason for referral and/or presenting problem
- The name of the tests (e.g., WAIS-R, Rorschach, MMPI) performed
- The scoring of the test
- Location the testing is performed
- The name and credentials of each provider involved in administering, interpreting, and preparing the report
- Interpretation of the test to include narrative descriptions of the test findings
- Length of time spent by each provider, as applicable, in face-to-face administration, interpretation, integrating the test interpretation, and documenting the comprehensive report based on the integrated data
- Recommended treatment, including how test results affect the prescribed treatment
- Recommendations for further testing to include an explanation to substantiate the necessity for retesting, if applicable
- Rationale or extenuating circumstances that impact the ability to complete the testing, such as, but not limited to, the person’s condition requires testing over two days and the person does not return, or the person’s condition precludes completion of the testing.

The original testing material must be maintained by the provider and must be readily available for retrospective review by HHSC.

When psychological, neurobehavioral, or neuropsychological testing is performed in a SNF, ICF, or ECF, a copy of the test and the resulting report must also be maintained in the person’s medical record at the facility.

### 4.3 Prior Authorization

Prior authorization requests may be submitted to the TMHP Prior Authorization Department via mail, fax, or the electronic portal. Performing providers may sign prior authorization forms and supporting documentation using electronic or wet signatures.

Refer to: Subsection 5.5.1.2, “Document Requirements and Retention” in “Section 5: Fee-for-Service Prior Authorizations” (Vol. 1, General Information) for additional information about electronic signatures.

All providers are required to adhere to prior authorization requirements.

#### 4.3.1 Services Requiring Prior Authorization

Prior Authorization is required for the following services:

- Psychiatric diagnostic evaluation (procedure codes 90791 and 90792) after the one evaluation per person, per provider, per rolling year limitation has been met
- Individual, family, or group psychotherapy (procedure codes 90832, 90833, 90834, 90836, 90837, 90838, 90846, 90847, and 90853) after the 30 visit per calendar year limitation has been met
• Neurobehavioral testing (procedure codes 96116 and 96121*) after the 4 hour per day limitations have been met

• Psychological testing (procedure codes 96130, 96131*, 96136, and 96137*) or neuropsychological testing (procedure codes 96132, 96133*, 96136, and 96137*) after the 8 hour per calendar year limitations have been met

• Unlisted psychiatric service or procedure (procedure code 90899)

Requests for prior authorization for procedure code 90899 must be submitted by the provider to the Special Medical Prior Authorization (SMPA) department using the Special Medical Prior Authorization (SMPA) Request Form with documentation supporting medical necessity including:

• Diagnosis(es)

• Prior treatment for this diagnosis and the medical necessity of the requested procedure

• A clear, concise description of the evidence-based service or procedure to be performed, and the intended fee for the service or procedure

• The reason for recommending this particular service or procedure

• A procedure code that is comparable to the service or procedure being requested

• Documentation that this service or procedure is not investigational or experimental

4.3.2 Prior Authorization Not Required

Prior authorization is not required for the following services:

• One psychiatric diagnostic evaluation (procedure codes 90791 and 90792) per person, per rolling year, per provider (same provider)

• 30 individual, family, or group psychotherapy (procedure codes 90832, 90833, 90834, 90836, 90837, 90838, 90846, 90847, and 90853) visits per person per calendar year

• 4 hours of psychotherapy services per person per day

• 4 hours of neuropsychological testing (procedure codes 96116 and 96121*) per person per day

• 8 hours of psychological, neurobehavioral, or neuropsychological testing (procedure codes 96130, 96131*, 96132, 96133*, 96136, or 96137*) per person, per calendar year

• Electroconvulsive therapy (procedure code 90870)

4.4 Documentation Requirements

All services require documentation to support the medical necessity of the service rendered, including mental health services. The documentation must support the medical necessity of the treatment for its entire duration.

Mental health services outlined in this handbook are subject to retrospective review to ensure that the documentation in the person’s medical record supports the medical necessity of the services provided.

4.5 Twelve Hour System Limitation

The following provider types are limited to a maximum combined total of 12 hours per provider, per day, regardless of the number of persons seen for outpatient mental health services:

• Psychologist

• APRN

• PA
The following table lists the procedure codes for mental health services included in the system limitation, along with the time increments the system will apply based on the billed procedure code. The time increments applied will be used to calculate the 12-hour per day system limitation.

<table>
<thead>
<tr>
<th>Procedure Code</th>
<th>Time Applied</th>
</tr>
</thead>
<tbody>
<tr>
<td>90791</td>
<td>60 minutes</td>
</tr>
<tr>
<td>90792</td>
<td>60 minutes</td>
</tr>
<tr>
<td>90832</td>
<td>30 minutes</td>
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<tr>
<td>90833*</td>
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<td>45 minutes</td>
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<td>90836*</td>
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<td>90837</td>
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<td>90838*</td>
<td>60 minutes</td>
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<tr>
<td>90846</td>
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<td>30 minutes</td>
</tr>
<tr>
<td>96137*</td>
<td>30 minutes</td>
</tr>
</tbody>
</table>

* Add-on procedure codes to be billed with the most appropriate E/M procedure code.

Court-ordered and DFPS directed services are not subject to the 12-hour per provider, per day system limitation when billed with modifier H9.

Physicians are not subject to the 12-hour system limitation since they can delegate and may submit claims in excess of 12 hours per day.

Psychologists can delegate to multiple LPAs, PLPs, interns, or post-doctoral fellows and therefore delegated services are not subject to the 12-hour system limitation since they may submit claims for delegated services in excess of 12 hours per day.

### 4.6 Court-Ordered Services

The court-ordered services listed below for persons who are age 20 years of age and younger or 65 years of age and older are not subject to utilization management reviews, including prior authorization, concurrent reviews, or retrospective reviews that have the effect of denying, reducing, or controverting the court-ordered service. In these situations, the court order is considered the determination of medical necessity.

When billed with modifier H9, court-ordered services are not subject to the 12-hour system limitation per provider, per day.
Federal law prohibits the use of federal Medicaid funding for medical care provided to persons who are considered incarcerated. A person is considered incarcerated when a criminal justice facility has custody of the person. Examples include:

- A person who is currently residing in a criminal justice facility and receiving treatment through a program at the criminal justice facility.
- A person who is committed under Title 1 Texas Code of Criminal Procedure §46(B), which addresses persons ages 18 and older who have been ordered to receive competency restoration.
- A person who is committed under Title 3 Texas Family Code §§55.01-55.45. These sections refer to persons committed to inpatient psychiatric care because they are deemed unfit to proceed.

The following court-ordered services are required to be provided to Medicaid eligible persons who are not considered incarcerated:

- Emergency detention ordered by a judge or magistrate under Title 7 Texas Health and Safety Code §§573.011-573.026.
- Mental health services ordered under Title 7 Texas Health and Safety Code §§574.01-574.110. Mental health services may include:
  - A mental health examination.
  - Inpatient or outpatient treatment.
  - Detention under protective custody and temporary mental health services.
- Treatment of persons who are found not guilty based on lack of responsibility under Title 3 Texas Family Code §55.
- Treatment that is a condition of probation.
- Treatment of persons with chemical dependencies ordered under Title 6 Texas Health and Safety Code §462.042.

For authorization of court-ordered services, the provider must submit documentation that includes:

- The court-order.
- Information about the statute under which the court is ordering the services.
- Verification of the person’s incarceration status.

For court-ordered inpatient admissions, providers must submit documentation that includes:

- A copy of the doctor’s certificate.
- All court-ordered commitment papers signed by the judge.

For persons with fee-for-service benefits, this supporting documentation must be submitted with the Psychiatric Inpatient Extended Stay Request Form.

Requested services beyond those that are court-ordered are subject to medical necessity review.

4.7 Exclusions

The following services are not benefits of Texas Medicaid:

- Psychoanalysis
- Multiple Family Group Psychotherapy
- Marriage or couples counseling
- Narcosynthesis
• Biofeedback training as part of psychophysiological therapy
• Psychiatric Day Treatment Programs
• Services provided by a psychiatric assistant, psychological assistant (excluding Master’s level LPA), or a licensed chemical dependency counselor

4.8 Claims Filing and Reimbursement

Providers must bill Medicare before Medicaid when a person is eligible for services under both programs. Medicaid’s responsibility for the coinsurance or deductible is determined in accordance with Medicaid benefits and limitations. Providers must check the person’s Medicare card for Part B coverage before billing Medicaid. When Medicare is primary, it is inappropriate to bill Medicaid without first billing Medicare.

Note: Texas Medicaid may reimburse the full amount of the Medicare coinsurance and deductible for services rendered by LCSW providers.

Refer to: Subsection 2.7.2, “Part B” in “Section 2: Texas Medicaid Fee-for-Service Reimbursement” (Vol. 1, General Information).

Subsection 4.9.2, “Medicare Part B Crossovers” in “Section 4: Client Eligibility” (Vol. 1, General Information) for information about how coinsurance and deductibles may be reimbursed by Texas Medicaid.

LCSW, LMFT, and LPC services must be submitted to TMHP in an approved electronic format or on the CMS-1500 paper claim form. Providers may purchase CMS-1500 paper claim forms from the vendor of their choice. TMHP does not supply the forms. When completing a CMS-1500 paper claim form, all required information must be included on the claim, as TMHP does not key any information from claim attachments. Superbills, or itemized statements, are not accepted as claim supplements.

Refer to: “Section 3: TMHP Electronic Data Interchange (EDI)” (Vol. 1, General Information) for information on electronic claims submissions.

Subsection 6.1, “Claims Information” in “Section 6: Claims Filing” (Vol. 1, General Information) for general information about claims filing.

Subsection 6.5, “CMS-1500 Paper Claim Filing Instructions” in “Section 6: Claims Filing” (Vol. 1, General Information). Blocks that are not referenced are not required for processing by TMHP and may be left blank.

According to 1 TAC §355.8091, the Texas Medicaid rate for LCSWs, LMFTs, and LPCs is 70 percent of the rate paid to a psychiatrist or psychologist for a similar service per 1 TAC §355.8085. Providers can refer to the OFL or the applicable fee schedule on the TMHP website at www.tmhp.com. Under 1 TAC §355.8261, an FQHC is reimbursed according to its specific prospective payment system (PPS) rate per visit for LCSW services.

Texas Medicaid implemented mandated rate reductions for certain services. The OFL and static fee schedules include a column titled “Adjusted Fee” to display the individual fees with all mandated percentage reductions applied.

Additional information about rate changes is available on the Reimbursement Rate Changes page of the TMHP website at www.tmhp.com.

Note: Texas Medicaid may reimburse the full amount of the Medicare coinsurance and deductible for services rendered by LCSW providers.
Refer to: Subsection 2.2, “Fee-for-Service Reimbursement Methodology” in “Section 2: Texas Medicaid Fee-for-Service Reimbursement” (Vol. 1, General Information) for more information about reimbursement.

Subsection 4.5, “Twelve Hour System Limitation” in this handbook for details about the 12-hours-per-day behavioral health services limitation.

4.9 NCCI and MUE Guidelines
The HCPCS and CPT codes included in the Texas Medicaid Provider Procedures Manual are subject to NCCI relationships, which supersede any exceptions to NCCI code relationships that may be noted in the manual. The CMS NCCI and MUE guidelines can be found in the NCCI Policy and Medicaid Claims Processing manuals, which are available on the CMS website. Providers should refer to the CMS NCCI web page for correct coding guidelines and specific applicable code combinations.

Whenever Texas Medicaid limitations are more restrictive than NCCI MUE guidance, Texas Medicaid limitations prevail.

5 Intellectual Disability Service Coordination, Mental Health Targeted Case Management, and Mental Health Rehabilitative Services

5.1 Enrollment

5.1.1 Local Intellectual and Developmental Disability Authority (LIDDA) Providers
A LIDDA provider who is authorized by HHSC to provide service coordination must be enrolled as a Long Term Care provider, and must submit claims through the Long Term Care system.

LIDDA are the only entities that provide case management (service coordination) services to persons who have an intellectual disability.

Refer to: The TMHP website at www.tmhp.com for additional information about Long Term Care enrollment and billing requirements.

5.1.2 Local Mental Health Authority (LMHA) Providers
LMHA providers are authorized by the DSHS to provide targeted case management services and mental health rehabilitative services. To enroll in Texas Medicaid, LMHA providers must contact DSHS at 1-512-206-5288 to be approved.

5.1.3 Non-Local Mental Health Authority (Non-LMHA) Providers
Non-LMHAs are private providers of both mental health (MH) case management and MH rehabilitative services, but they are not LMHAs. They must comply with all applicable federal and local laws and all of the regulations that are related to the services they provide. After receiving approval for enrollment in Texas Medicaid, the Non-LMHA provider must be credentialed by a Texas Medicaid managed care organization (MCO) to provide services to Texas Medicaid eligible persons.

Non-LMHA providers also must register to use the DSHS Clinical Management for Behavioral Health Services (CMBHS) clinical record-keeping system before providing services to Texas Medicaid eligible persons.
5.1.4 Provider Credentials for Facilities Delivering MHTCM and Mental Health Rehabilitative Services

Community Services Specialist (CSSP), Qualified Mental Health Professional - Community Services (QMHP-CS), family partners, and peer providers are eligible to deliver some or all of the mental health rehabilitative services and mental health targeted case management services. The credentialing requirements and services each provider may deliver are listed in the following sections.

Staff administering the assessment instruments must have documentation of current certification in the CANS or ANSA. Certification must be updated annually through an approved entity.

5.1.4.1 Community Services Specialist (CSSP)

CSSP providers are eligible to deliver Mental Health Targeted Case Management (MHTCM) and Mental Health (MH) Rehabilitative services and must meet the following minimum credentialing requirements:

- High school diploma or high school equivalency
- Three continuous years of documented full-time experience in the provision of MH rehabilitative services prior to August 30, 2004
- Demonstrated competency in the provision and documentation of MHTCM and MH rehabilitative services

A CSSP performing MHTCM and MH rehabilitative services must:

- Be an employee of the facility where the case management is delivered.
- Be clinically supervised by at least a QMHP-CS.

5.1.4.2 Qualified Mental Health Professional - Community Services (QMHP-CS)

QMHP-CS providers are eligible to deliver MHTCM and MH rehabilitative services and must meet the following minimum credentialing requirements:

- Completed a standardized training curriculum
- Demonstrated competency in the work to be performed
- Obtained one of the following:
  - A bachelor’s degree from an accredited college or university with a minimum number of hours that are equivalent to a major in psychology, social work, medicine, nursing, rehabilitation, counseling, sociology, human growth and development, physician assistant, gerontology, special education, educational psychology, early childhood education, or early childhood intervention
  - A license as a registered nurse (RN)

Staff administering the assessment instruments must have documentation of current certification in the Child and Adolescent Needs and Strengths Assessment (CANS) or the Adult Needs and Strengths Assessment (ANSA). Certification must be updated annually through an approved entity.

An individual who possesses any of the following licenses is considered a Licensed Practitioner of the Healing Arts (LPHA) and is automatically certified as a QMHP-CS:

- Physician
- Physician Assistant
- Advanced Practice Registered Nurse
- Psychologist
- Licensed Clinical Social Worker (LCSW)
- Licensed Marriage and Family Therapist (LMFT)
• Licensed Professional Counselor (LPC)

A QMHP-CS must be clinically supervised by another QMHP-CS. If a QMHP-CS is clinically supervised by another QMHP-CS, the supervising QMHP-CS must be clinically supervised by an LPHA.

5.1.4.3 Peer Provider

Peer providers must have a high school diploma or high school equivalency, one cumulative year of receiving mental health services, and be clinically supervised by an LPHA. The supervising LPHA must conduct at least monthly documented meetings with the peer provider and conduct an additional monthly documented observation of the peer providing services.

A peer provider must satisfy all staff credentialing, competency, training, and clinical supervision requirements.

Services provided by a peer provider must be included in the treatment plan.

5.1.4.4 Family Partner

A certified family partner must have a high school diploma or high school equivalency and one cumulative year of participating in mental health services as the parent or legally authorized representative (LAR) of a child receiving mental health services.

A family partner must be supervised by at least a QMHP-CS and must satisfy all staff credentialing, competency, training, and clinical supervision requirements.

Services provided by a family partner must be included in the treatment plan.

Family partners must be credentialed as a certified family partner within one year of their hire date.

The family partner service is provided to parents or LARs for the benefit of the Medicaid eligible child.

5.1.4.5 Certifications for Mental Health Rehabilitative Services

The following provider certifications are required for mental health rehabilitative services:

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<tr>
<th>Service</th>
<th>Provider Types</th>
<th>Peer Provider</th>
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5.2 Services, Benefits, Limitations, and Prior Authorization

5.2.1 Intellectual and Developmental Disabilities Service Coordination

Texas Medicaid provides the following:

- Service coordination for persons who have an intellectual disability or a related condition (adult or child). Persons who have a related condition are eligible if they are being enrolled into the home and community based waiver (HCS); the Texas Home Living Waiver; or an Intermediate Care Facility for Individuals with Intellectual Disabilities (ICF-IID).

- Service coordination for persons who have an intellectual disability or a related condition who are enrolled in HCS or Texas Home Living waiver programs.

Service coordination funded by Medicaid as TCM is reimbursed by encounter. There are two types of encounters:

- **Comprehensive encounter (Type A):** A face-to-face contact with a person to provide service coordination. The comprehensive encounter is limited to one billable encounter per person per calendar month. HHSC will not authorize payment for a comprehensive encounter that exceeds the cap of one encounter per person per calendar month.

- **Supportive encounter (Type B):** A face-to-face, telephone, or telemedicine contact with a person or with a collateral on the person’s behalf to provide service coordination.

A LIDDA is allowed up to three Type B encounters per calendar month for each Type A encounter that has occurred within the calendar month.

The Type B encounters are not limited to three per person. Rather, the allowed Type B encounters may be delivered to any person who needs a Type B encounter. These Type B encounters are allowable as long as the person who received the Type B encounter also received a Type A encounter that same month.

For example, Sam and Mary receive a Type A encounter in June. It is allowable for the LIDDA to bill for one Type B encounter for Sam in June and five Type B encounters for Mary in June.

Payment for a person’s Type B encounter is contingent on that the person having a Type A encounter within the same calendar month.

Within the calendar month, the Type A encounter does not have to occur on a date before any of the Type B encounters occur.

Prior authorization is not required for IDD coordination services.
5.2.2 Mental Health Targeted Case Management (MHTCM)

Targeted case management services are case management services to persons within targeted groups. The target population that may receive Mental Health Targeted Case Management (MHTCM) as part of the Texas Medicaid Program are persons, regardless of age, with a single diagnosis of chronic mental illness or a combination of chronic mental illnesses as defined in the latest edition of the American Psychiatric Association’s DSM, and who have been determined via a uniform assessment process to be in need of MHTCM services. Persons of any age with a single diagnosis of intellectual and developmental disabilities (IDD) and related conditions, or a single diagnosis of substance use disorder (SUD) are not eligible for MHTCM services.

MHTCM services are furnished to assist persons in gaining access to needed medical, social/behavioral, educational, and other services and supports. MHTCM activities and services include:

- A comprehensive assessment and periodic reassessment, as medically necessary, of persons needs to determine the need for any medical, educational, social/behavioral, or other services.

- The development (and periodic revision, as medically necessary) of a specific care plan that:
  - Is based on the information collected through the assessment;
  - Specifies the goals and actions to address the medical, social/behavioral, educational, and other services and supports needed by the person;
  - Includes activities such as ensuring the active participation of the eligible person and working with the person (or the person’s authorized health care decision maker) and others to develop these goals; and
  - Identifies a course of action to respond to the assessed needs of the eligible person.

- Making referrals and performing other related activities, such as scheduling an appointment on behalf of the person, to help an eligible person obtain needed services and supports, including activities that help link a person with:
  - Medical, social/behavioral, and educational providers; and
  - Other programs and services that are capable of providing needed services to address identified needs and achieve goals in the care plan.

- Monitoring and performing the necessary follow-up that is necessary to ensure the care plan is implemented and adequately addresses the person’s needs.

MHTCM activities may be with the person, family members, LAR, providers, or other entities or individuals and conducted as frequently as necessary, and at least once annually, to determine whether the following conditions are met:

- Services are being furnished in accordance with the person’s care plan;
- Services in the care plan are adequate in amount, scope, and duration to meet the needs of the person; and
- The care plan and service arrangements are modified when the person’s needs or status change.

MHTCM is a benefit for persons transitioning to a community setting for up to 180 consecutive days prior to leaving a nursing facility; however, MHTCM services are coordinated with and do not duplicate activities provided as part of nursing facility services and discharge planning activities.

MHTCM consists of intensive case management and routine case management. Intensive case management services are predominantly community-based case management activities provided to the person or to the LAR on behalf of the person (who may or may not be present) to assist a person and caregiver or LAR in obtaining and coordinating access to necessary care and services appropriate to the
person’s needs. Routine case management services are primarily office-based case management activities that assist a person, caregiver, or LAR in obtaining and coordinating access to necessary care and services appropriate to the person’s needs.

Intensive case management and routine case management are benefits for persons who are 20 years of age and younger. Intensive case management and routine case management are not payable on the same day.

Routine case management is a benefit for persons who are 21 years of age and older.

Providers must use procedure code T1017 and the appropriate modifier for MHTCM:

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<thead>
<tr>
<th>Modifier</th>
<th>Description</th>
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<tr>
<td>TF</td>
<td>Routine Case Management</td>
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<td>TG</td>
<td>Intensive Case Management</td>
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<td>HA</td>
<td>Child/Adolescent Program</td>
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<td>HZ</td>
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Procedure code T1017 is limited to the following diagnosis codes:

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An MHTCM reimbursable session is the provision of a case management activity by an authorized case manager during a face-to-face meeting with a person who is authorized to receive that specific type of case management. A billable unit of MHTCM is 15 continuous minutes of contact.

MHTCM is not payable when delivered on the same day as psychosocial rehabilitative services.

The following activities are included in the MHTCM rate and will not be reimbursed separately:

- Documenting the provision of MHTCM services.
- On-going administration of the Uniform Assessment to determine amount, duration, and type of MHTCM.
- Travel time required to provide MHTCM services at a location not owned, operated, or under arrangement with the provider.

Texas Medicaid must not be billed for MHTCM services provided before the establishment of a diagnosis of mental illness and the authorization of services.

### 5.2.2.1 Collateral Contacts

MHTCM may include contacts with non-eligible individuals who are directly related to identifying the eligible person’s needs and care for the purposes of helping the eligible person access services, identifying needs and supports to assist the eligible person in obtaining services, providing case managers with useful feedback, and alerting case managers to changes in the eligible person’s needs.

MHTCM services involving collateral contacts are only payable when the person or LAR is also present during the case management session.

### 5.2.2.2 Intensive Case Management

Intensive case management incorporates a wraparound approach to care planning and treatment plan implementation. The wraparound process is a strengths-based course of action involving a child or youth and their family, including any additional people identified by the child or youth, LAR, primary caregiver, and family, that results in a unique set of community services and natural supports that are individualized for the child or youth to achieve a positive set of identified outcomes.

Intensive case management is primarily community-based, meaning that services are provided in whatever setting is clinically appropriate and person-centered.

A case manager assigned to a child or youth who is authorized to receive intensive case management services must have completed training in the National Wraparound Implementation Center’s Wraparound Practice model and must incorporate wraparound process planning or other approved models in developing a plan that addresses the child’s or youth’s unmet needs across life domains.
The case manager must develop an intensive case management treatment plan based on the child’s or youth’s needs that may include information across life domains from relevant sources.

The case manager must meet face-to-face with the child or youth and the LAR or primary caregiver:

- Within seven days after the case manager is assigned to the child or youth;
- Within seven days after discharge from an inpatient psychiatric setting, whichever is later; or
- Document the reasons the meeting did not occur and meet at the soonest available opportunity.

The case manager must identify the child’s or youth’s strengths, service needs, and assistance that will be required to address the identified needs in the plan.

The case manager must take steps that are necessary to assist the child or youth in gaining access to the needed services and service providers, including:

- Making referrals to potential service providers.
- Initiating contact with potential service providers.
- Arranging, and if necessary to facilitate linkage, accompanying the child or youth to initial meetings and non-routine appointments.
- Arranging transportation to ensure the child or youth attendance.
- Advocating with service providers.
- Providing relevant information to service providers.
- Monitoring the child’s or youth’s progress toward the goals set forth in the plan.

5.2.2.2.1 Authorization Requirements

LMHAs delivering services to persons with fee-for-service benefits must obtain authorization from their internal utilization management department. When providing care to persons enrolled in managed care, LMHAs and other providers contracted with MCOs must submit authorization requests to the MCO with whom the person is enrolled. The MCO may choose to waive this authorization submission requirement. Additionally, MCOs must follow the requirements set forth in the Uniform Managed Care Manual regarding utilization management for targeted case management and mental health rehabilitative services.

Eligibility determinations occur at the facility providing targeted case management services using the Clinical Management of Behavioral Health Services (CMBHS) software system.

Criteria used to make these service determinations are from the recommended Level of Care (LOC) of the person as derived from the Uniform Assessment (UA), the needs of the person, and the Texas Resilience and Recovery Utilization Management Guidelines.

In determining service, the Qualified Mental Health Professional-Community Services (QMHP-CS) or Licensed Practitioner of the Healing Arts (LPHA) performs a screening for eligibility utilizing the UA. The LPHA gives a diagnosis and determines if the services are medically necessary.

The LPHA determination of diagnosis shall include an interview with the person conducted either in-person or via telemedicine or telehealth.

Refer to: The Telecommunication Services Handbook (Vol. 2, Provider Handbooks).

A facility that provides MHTCM must ensure that at minimum a QMHP-CS administers the uniform assessment to the person at specified intervals (every 90 calendar days for persons who are 20 years of age and younger and every 180 calendar days for persons who are 21 years of age and older), and obtains a recommended LOC for the person.
The facility must evaluate the clinical needs of the person to determine if the amount of MHTCM services associated with the recommended LOC described in the utilization management guidelines is sufficient to meet those needs and ensure that an LPHA reviews the recommended LOC and verifies whether the services are medically necessary.

If the facility determines that the type of MHTCM services associated with the recommended LOC is sufficient to meet the person's needs, the facility must submit a request for service authorization according to the recommended LOC.

If the facility determines that a LOC other than the recommended LOC is more appropriate for the person, the provider must submit a deviation request that includes:

- A request for an authorization of an LOC that is higher or lower than initially recommended; and
- The clinical justification for the request.

The clinical justification must include the specific reasons why the person requires interventions outside the recommended LOC. Refusal of recommended LOC by the person receiving services may be noted as part of the justification.

All plans of care are subject to retrospective review by the state.

### 5.2.2.3 Eligibility and Service Determinations for Persons Who are 20 Years of Age and Younger

MHTCM is available to persons who are 20 years of age and younger with a diagnosis of mental illness (excluding a single diagnosis of IDD and related disorders, or a single diagnosis of SUD) or serious emotional disturbance and who:

- Have a serious functional impairment; or
- Are at risk of disruption of a preferred living or child care environment due to psychiatric symptoms; or
- Are enrolled in a school system's special education program because of serious emotional disturbance.

The initial assessment is the clinical process of obtaining and evaluating historical, social/behavioral, functional, psychiatric, developmental, or other information from the person seeking services to determine specific treatment and support needs.

Functioning is assessed using one of the following standardized assessment tools:

- The Child and Adolescent Needs and Strengths Assessment (CANS) for persons who are 17 years of age and younger
- The Adult Needs and Strengths Assessment (ANSA) and any necessary supplemental assessments for persons who are 18 years of age and older

Services and supports to be provided to the person are determined jointly by the person receiving services, family, and the provider.

MHTCM services authorized for care by the provider through a clinical override are eligible for the duration of the authorization.

Continued eligibility for MHTCM services is based on a reassessment every 90 calendar days by the provider and reauthorization of services by the facility. Assignment of diagnosis in the CMBHS is required at any time the DSM diagnosis changes and at least annually from the last diagnosis entered into CMBHS.
5.2.2.4 Eligibility and Service Determinations for Persons who are 21 Years of Age and Older

MHTCM is available to persons who are 21 years of age and older and who have severe and persistent mental illnesses such as schizophrenia, major depression, bipolar disorder, post-traumatic stress disorder, or other severely disabling mental disorders (excluding a single diagnosis of IDD and related disorders, or a single diagnosis of SUD) which require crisis resolution or ongoing and long-term support and treatment.

Adults with schizophrenia and bipolar disorder are automatically eligible for services. Adults with any other mental health diagnoses require evidence of significant difficulty functioning across one or more domains, such as work or school, to be eligible for services.

Functioning is assessed using a standardized assessment tool called the Adult Needs and Strengths Assessment (ANSA). Adults are reassessed every 180 calendar days for continued need for services. Assignment of diagnosis in the CMBHS is required at any time the DSM diagnosis changes and at least annually from the last diagnosis entered into CMBHS.

Adults with a diagnosis of schizophrenia or bipolar disorder are automatically eligible for continued services. Adults with major depressive disorder whose level of functioning qualified them initially are also automatically eligible for continued services, regardless of whether their level of functioning has improved or not.

Adults with any other mental health diagnoses are eligible should their level of functioning continue to be significantly impaired, as evidenced by the results of a standardized assessment tool.

5.2.2.5 Documentation Requirements

A comprehensive diagnosis must be included in the person’s medical record, including documentation of applicable diagnostic criteria according to the latest edition of the DSM, as well as the specific justification of need for services.

MHTCM services, including attempts to provide MHTCM services, must be documented in the person’s medical record.

For routine case management, the case manager must document the person’s strengths, service needs, and assistance required to address the service needs as well as the steps that are necessary to accomplish the goals required to meet the person’s service needs.

For intensive case management, the assigned case manager must include the intensive case management treatment plan in the child’s or youth’s medical record and document steps taken to meet the child’s or youth’s goals and needs in the child’s or youth’s progress notes.

As a result of the face-to-face meetings, assessments, and reassessments conducted, the case manager must document the person’s identified strengths, service needs, and assistance given to address the identified need, and specific goals and actions to be accomplished.

The case manager must document the following for all services provided:

- The event or behavior that occurs while providing the MHTCM service or the reason for the specific case management encounter
- The person, persons, or entity, including other case managers, with whom the encounter or contact occurred
- Collateral contacts such as contacts with non-eligible individuals that are directly related to identifying the needs and supports for helping the person access services and managing the person’s care, including coordination with other case managers
- The recovery plan goal(s) that was the focus of the service, including the progress or lack of progress in achieving recovery plan goal(s)
• The time line for obtaining the needed services
• The specific intervention that is being provided
• The date the MHTCM service was provided
• The start and end time of the MHTCM service
• The location where the MHTCM service was provided and whether it was a face-to-face or telephone contact
• The name of the provider agency and the signature of the employee providing the MHTCM service, including their credentials
• The time line for reevaluating the needed service

If the person refuses MHTCM services, the case manager must document the reason for the refusal in the most appropriate area of the person’s medical record and request that the person sign a waiver of MHTCM services that is filed in the person’s medical record.

The provider must retain documentation in compliance with applicable records retention requirements in federal and state laws, rules, and regulations.

5.2.2.6 Exclusions

The following services are not covered by MHTCM:

• Case management activities that are an integral component of another covered Medicaid service
• The provision of a medical, educational, social/behavioral, or other service to which a person has been referred, including for foster care programs, services such as, but not limited to, the following:
  • Research gathering and completion of documentation required by the foster care program
  • Assessing adoption placements
  • Recruiting or interviewing potential foster care parents
  • Serving legal papers
  • Home investigations
  • Providing transportation, including transporting the person to his/her LAR/primary caregiver
  • Administering foster care subsidies
  • Making placement arrangements
• Performing an activity that does not directly assist a person in gaining or coordinating access to needed services
• Providing medical or nursing services
• Performing preadmission or intake activities
• Monitoring the person’s general health status
• Performing outreach activities
• Performing quality oversight of a service provider
• Conducting utilization review or utilization management activities
• Conducting quality assurance activities
• Authorizing services or authorizing the provision of services
• Services to inmates of public institutions
5.2.3 Mental Health Rehabilitative Services

Mental health rehabilitative services are defined as providing assistance in maintaining or improving functioning and may be considered rehabilitative when necessary to help a person achieve a rehabilitation goal as defined in the treatment plan.

Mental health rehabilitative services may be provided to a person with a serious mental illness as defined in the latest edition of the American Psychiatric Association’s DSM.

Mental health rehabilitative services are age-appropriate, individualized, and designed to ameliorate functional impairments that negatively affect any of the following:

- Community integration
- Community tenure
- Behaviors resulting from serious mental illness (SMI) or severe emotional disturbance (SED) that interfere with a person’s ability to remain in the community as a fully integrated and functioning member of that community

Mental health rehabilitative services may include:

- Medication training and support services
- Psychosocial rehabilitative services
- Skills training and development
- Crisis intervention services
- Day programs for acute needs

Mental health rehabilitative services may only be provided by a member of the person’s therapeutic team. The therapeutic team includes a sufficient number of staff to adequately address the rehabilitative needs of persons assigned to the team.

Team members must be appropriately credentialed and have completed required trainings to provide the full array of component services, have regularly scheduled team meetings either in person or by teleconference, and every member of the team must be knowledgeable of the needs and the services available to the specific persons assigned to the team.

Mental health rehabilitative services may be a benefit for persons residing in a nursing facility (NF) when medically necessary as determined via a uniform assessment protocol and determined through preadmission screening and resident review (PASRR) to require specialized services.

The following procedure codes are a benefit for mental health rehabilitation:

<table>
<thead>
<tr>
<th>Service Category</th>
<th>Procedure Codes</th>
<th>Modifiers</th>
</tr>
</thead>
<tbody>
<tr>
<td>Day Program for Acute Needs</td>
<td>H2012</td>
<td></td>
</tr>
<tr>
<td>Medication Training and Support</td>
<td>H0034</td>
<td>HA/HQ: group services for adults</td>
</tr>
<tr>
<td></td>
<td></td>
<td>HQ: group services for child/youth</td>
</tr>
<tr>
<td>Crisis Intervention</td>
<td>H2011</td>
<td>HA: child/youth</td>
</tr>
<tr>
<td>Skills Training and Development</td>
<td>H2014</td>
<td>HA: individual services for child/youth</td>
</tr>
<tr>
<td></td>
<td></td>
<td>HA/HQ: group services for child/youth</td>
</tr>
<tr>
<td>Psychosocial Rehabilitation Services</td>
<td>H2017</td>
<td>TD: individual services provided by RN</td>
</tr>
<tr>
<td></td>
<td></td>
<td>HQ: group services</td>
</tr>
<tr>
<td></td>
<td></td>
<td>HQ/TD: group services provided by RN</td>
</tr>
<tr>
<td></td>
<td></td>
<td>ET: individual crisis services</td>
</tr>
</tbody>
</table>
Psychosocial rehabilitation is not reimbursable on the same day as mental health targeted case management or skills training and development.

Reimbursement for procedure codes H0034, H2012, H2014, and H2017 are limited to the following diagnosis codes:

<table>
<thead>
<tr>
<th>Diagnosis Codes</th>
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<tbody>
<tr>
<td>F060 F061 F062 F0630 F0631 F0632 F0633 F0634</td>
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<tr>
<td>F064 F068 F070 F0789 F09 F200 F201 F202</td>
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<tr>
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<tr>
<td>T7602XS T7612XA T7612XD T7612XS T7622XA T7622XD T7622XS T7652XA</td>
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<tr>
<td>T7652XD T7652XS T7662XA T7662XD T7662XS</td>
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</table>
No diagnosis is required for crisis intervention services (procedure code H2011); however, all CMS 1500 claim forms require a diagnosis.

A Medicaid provider may only bill for medically necessary mental health rehabilitative services that are provided face-to-face to:

- A Medicaid-eligible person;
- The LAR of a Medicaid-eligible person who is 21 years of age and older (on behalf of the person); or
- The LAR or primary caregiver of a Medicaid-eligible person who is 20 years of age and younger (on behalf of the person).

Rehabilitative services delivered via group modality are limited to an 8-person maximum for adults and a 6-person maximum for children or adolescents (not including LARs or caregivers).

### 5.2.3.1 Eligibility and Service Determinations for Persons Who are 20 Years of Age and Younger

Certain mental health rehabilitative services (crisis intervention services, medication training and support, and skills training and development) are available to persons who are 20 years of age and younger with a diagnosis of mental illness or serious emotional disturbance and who:

- Have a serious functional impairment; or
- Are at risk of disruption of a preferred living or child care environment due to psychiatric symptoms; or
- Are enrolled in a school system’s special education program because of serious emotional disturbance.

Functioning is assessed using a standardized assessment tool, the Child and Adolescent Needs and Strengths Assessment (CANS) for persons who are 17 years of age and younger and the ANSA for persons who are 18 years of age and older.

Continued eligibility for mental health rehabilitative services for persons who are 17 years of age and younger is based on a reassessment at least every 90 calendar days, or more frequently if clinically indicated by the provider. Persons who are 18 years of age and older are reassessed every 180 calendar days, or more frequently if clinically indicated by the provider.

Assignment of diagnosis in the Clinical Management for Behavioral Health Services (CMBHS) is required at any time the mental illness diagnosis changes and at least annually from the last diagnosis entered into CMBHS.

The LPHA determination of diagnosis shall include an interview with the child or youth conducted either in person or via telemedicine or telehealth.

In order to complete a comprehensive diagnosis for a child or youth, documentation of the required diagnostic criteria according to the latest version of the DSM, as well as the specific level of functioning, shall be included in the child’s or youth’s record. This information shall be included as part of the required assessment information.

### 5.2.3.2 Eligibility and Service Determinations for Persons Who are 21 Years of Age and Older

Persons who are 21 years of age and older with serious mental illness, determined to be medically necessary via a uniform assessment protocol, are eligible for Mental Health Rehabilitative Services if the adult is:

- A resident of the state of Texas.
- Determined by a uniform assessment and clinician observation to require mental health rehabilitative services.
• An LPHA has made a determination that such services are medically necessary.

Mental health rehabilitative services are available to persons who are 21 years of age and older who have serious mental illnesses and significant functional impairments which require crisis resolution or ongoing treatment. Functioning is assessed using a standardized assessment tool, the Adult Needs and Strengths Assessment (ANSA).

Adults with schizophrenia and bipolar disorder are automatically eligible for services. Adults with any other mental health diagnoses require evidence of significant difficulty functioning across one or more domains, such as work or school, to be eligible for services.

Persons who are 18 years of age and older are reassessed for continued need for services at least every 180 calendar days, or more frequently if clinically indicated by the provider.

Assignment of diagnosis in the Clinical Management for Behavioral Health Services (CMBHS) is required at any time the mental illness diagnosis changes and at least annually from the last diagnosis entered into CMBHS.

The LPHA determination of diagnosis shall include an interview with the adult conducted either in person or via telemedicine or telehealth.

In order to complete a comprehensive diagnosis for an adult, documentation of the required diagnostic criteria according to the latest version of the American Psychiatric Association’s DSM, as well as the specific level of functioning, shall be included in the adult’s record. This information shall be included as part of the required assessment information.

Adults with a diagnosis of schizophrenia or bipolar disorder are automatically eligible for continued services. An adult with major depressive disorder whose level of functioning qualified them initially also is automatically eligible for continued services, regardless of whether their level of functioning has improved or not. Adults with any other mental health diagnoses are eligible should their level of functioning continue to be significantly impaired, as evidenced by the results of a standardized assessment tool called the Adult Needs and Strengths Assessment (ANSA).

5.2.3.3 Treatment Planning

Mental health rehabilitative services are part of a person’s treatment plan and are intended to:

• Reduce a person’s functional impairments resulting from serious mental illness (SMI) for adults.
• Reduce serious emotional disturbance in children.
• Restore a person to his/her optimal functioning level in the community.

The treatment planning process for mental health rehabilitative services requires the active participation of the Medicaid eligible person or LAR when necessary due to the person’s age or legal status. Treatment plans are based on a comprehensive assessment and must address the person’s strengths, areas of need, the person’s preferences, and descriptions of the person’s treatment goals.

5.2.3.4 Medication Training and Support

Medication training and support services consist of education and guidance about medications and their possible side effects. It is curriculum-based training and guidance that serves as an initial orientation for the person in understanding the nature of his/her mental illnesses or emotional disturbances and the role of medications in ensuring symptom reduction and increased tenure in the community.

Medication training and support includes:

• Assisting the person to manage symptomology and maximize functioning.
• Understanding the concepts of recovery and resilience within the context of the serious mental illness.
• Developing an understanding of the relationship between mental illness and the medications prescribed to treat the illness.
• The interaction of medication with other medications, diet, and mood altering substances.
• Understanding the overdose precautions of the person’s medication.
• The identification and management of potential side effects.
• Learning self-administration of the person’s medication.
• Necessity of taking medications prescribed and following the physician’s or other qualified health care professional’s orders.

Medication training and support is available to eligible children, youth, and adults. The LAR or primary caregiver may receive medication training and support services on behalf of an eligible adult, child or youth.

5.2.3.5 **Psychosocial Rehabilitative Services**

Psychosocial rehabilitative services are social, behavioral, and cognitive interventions provided by members of a person’s therapeutic team that build on strengths and focus on restoring the person’s ability to develop and maintain social relationships, occupational or educational achievement, and other independent living skills that are affected by or the result of a serious mental illness in persons who are 17 years of age and older.

Psychosocial rehabilitative services include independent living services, coordination services, and employment, housing, and medication-related services. Psychosocial rehabilitative services may also address the impact of co-occurring disorders upon the person’s ability to reduce symptomology and increase daily functioning.

If psychosocial rehabilitation is in the treatment plan, the treatment plan cannot simultaneously include skills training and development or targeted case management services.

Psychosocial rehabilitative services may not be provided to a person who is currently admitted to a crisis stabilization unit.

5.2.3.5.1 **Independent Living Services**

Independent living services assist a person in acquiring the most immediate, fundamental functional skills needed to enable the person to reside in the community and avoid more restrictive levels of treatment or reducing behaviors or symptoms that prevent successful functioning in the person’s environment of choice.

Independent living services include skills training and/or supportive interventions that focus on the improvement of communication skills, appropriate interpersonal behaviors, and other skills necessary for independent living or, when age appropriate, functioning effectively with family, peers, and teachers.

Training for independent living includes skills related to:

• Personal hygiene.
• Transportation utilization.
• Money management.
• The development of natural supports.
• Access to needed services in the community (e.g., medical care, substance use services, legal services, living accommodations).
• Social skills (e.g., communicating one’s needs to strangers and making appropriate choices for the use of leisure time).
5.2.3.5.2 Coordination Services

Coordination services are training activities that assist a person in improving his or her ability to gain and coordinate access to necessary care and services appropriate to the needs of the person.

Training for coordination skills includes instruction and guidance in such areas as:

- Identifying areas of need across all life domains.
- Prioritizing needs and setting goals.
- Identifying potential service providers and support systems.
- Initiating contact with providers and support systems.
- Participating in the development and subsequent revisions of their plan of care.
- Coordinating their services and supports.
- Advocating for necessary changes and improvements to ensure that they obtain maximum benefit from their services and supports.

5.2.3.5.3 Employment-Related Services

Employment-related services provide supports and skills training that are not job-specific and focus on developing skills to reduce or manage the symptoms of serious mental illness that interfere with a person’s ability to make vocational choices or obtain or retain employment.

Included in employment-related services are activities such as:

- Skills training related to task focus, task completion, planning and managing activities to achieve outcomes, personal hygiene, grooming and communication, and skills training related to securing appropriate clothing, developing natural supports, and arranging transportation.
- Establishing supportive contacts related to the school or work-site situation to reduce or manage behaviors or symptoms related to the person’s mental illness or emotional disturbance that interfere with job performance or progress towards the development of skills that would enable the person to obtain or retain employment.

5.2.3.5.4 Housing-Related Services

Housing-related services develop a person’s strengths and abilities to manage the symptoms of the person’s serious mental illness that interfere with the person’s capacity to obtain or maintain tenure in independent integrated housing.

Included in housing-related services are activities such as:

- Skills training related to home maintenance and cleanliness.
- Problem solving with landlord and other residents.
- Maintaining appropriate interpersonal boundaries.
- Establishing supportive contacts related to the housing situation to reduce or manage behaviors or symptoms related to the person’s mental illness or emotional disturbance that interfere with maintaining independent integrated housing.

5.2.3.5.5 Medication-Related Services

Medication-related services provide individualized training regarding the person’s medication adherence and is different from medication-training and support.

Services consist of training and supportive interventions that focus on person-specific needs and goals regarding the administration of medication, monitoring efficacy and side effects of medication, and other nursing services that enable the person to attain or maintain an optimal level of functioning.
Medication-related services do not include services or activities that are incidental to services performed by a physician (or other qualified health care professional) during an evaluation and management services visit.

### 5.2.3.6 Skills Training and Development

Skills training and development is training provided to an eligible person, the LAR, or primary caregiver on behalf of an eligible adult, child, or youth.

The training addresses:

- Serious mental illness or SED and symptom-related problems that interfere with the person’s functioning and living, working, and learning environment.
- Provides opportunities for the person to acquire and improve skills needed to function as appropriately and independently as possible in the community.
- The person’s community integration and increases his or her community tenure.

Skills training and supportive interventions focus on the improvement of communication skills, appropriate interpersonal behaviors and other skills necessary for independent living or, when age appropriate, functioning effectively with family, peers, and teachers.

Skills training and development may include:

- Skills related to personal hygiene.
- Pro-social skills.
- Assertiveness skills.
- Anger management skills.
- Stress reduction techniques.
- Communication skills.
- Transportation utilization.
- Money management.
- The development of natural supports.
- Access to needed services in the community, e.g., medical care, substance use services, legal services, living accommodations.
- Social skills (e.g., communicating one’s needs to strangers and making appropriate choices for the use of leisure time).

Skills training and development services consist of increasing the LAR’s or primary caregiver’s understanding of and ability to respond to the person’s needs identified in the uniform assessment or documented in the treatment plan.

Persons receiving skills training and development are not eligible to simultaneously receive psychosocial rehabilitative services and both services should not be simultaneously listed in the person’s treatment plan.

### 5.2.3.7 Crisis Intervention

Crisis intervention services are intensive community-based one-to-one services provided to persons who require services to control acute symptoms that place the person at immediate risk of hospitalization, incarceration, or placement in a more restrictive treatment setting.

This service includes assessment, behavioral skills training, problem-solving, and reality orientation to help a person identify and manage their symptoms of mental illness, and cope with stressors.
Crisis intervention services may be provided in extended observation or crisis residential units. Crisis intervention services may not be provided to a person who is currently admitted to a crisis stabilization unit.

Crisis intervention services consist of the following interventions:

- An assessment of dangerousness of the person to self or others
- The provision of emergency care services that include crisis screening and response, telephone access, emergency case services, urgent care services, routine care services, and access to emergency medical/crisis services
- Behavior skills training to assist the person in reducing distress and managing symptoms
- Problem-solving
- Reality orientation to help the person identify and manage his or her symptoms of serious mental illness or SED
- Providing instruction, structure, and emotional support to the person in adapting to and coping with immediate stressors

Crisis intervention services are available to eligible children youth and adults.

5.2.3.8 Day Programs for Acute Needs

Day programs for acute needs provide short term, intensive treatment to an eligible persons who is 18 years of age or older and who requires multidisciplinary treatment to stabilize acute psychiatric symptoms or prevent admission to a more restrictive setting. Day program services are a site-based treatment provided in a group modality.

Day programs for acute needs are provided in a highly structured and safe environment with constant supervision and ensure an opportunity for frequent interaction between the adult and staff members.

Day programs for acute needs must at all times have sufficient staff to ensure safety and program adequacy according to an established staffing ratio and staff response times. This service focuses on intensive, medically-oriented, multidisciplinary interventions such as behavior skills training, crisis management, and nursing services that are designed to stabilize acute psychiatric symptoms.

These services may be provided in a residential facility; however, none of the residential facilities can contain greater than 16 beds.

Day programs for acute needs include:

- Psychiatric nursing services.
- Pharmacological instruction that addresses medication issues related to the crisis precipitating the need for provision of day programs for acute needs.
- Symptom management training.
- Functional skills training.

Day programs for acute needs must, at all times, have a sufficient number of staff members to ensure safety and program adequacy, and, at a minimum, include:

- One RN for every 16 persons at the day program’s location,
- One physician to be available by phone, with a response time not to exceed 15 minutes,
- Two staff members who are QMHP-CSs, CSSPs, or peer providers at the day program’s location,
- One additional QMHP-CS who is not assigned full-time to another day program to be physically available, with a response time not to exceed 30 minutes,
Additional QMHP-CSs, CSSPs, or peer providers at the day program’s location sufficient to maintain a ratio of one staff member to every four persons receiving care.

5.2.3.9 **Authorization Requirements**

Providers must obtain prior authorization for mental health rehabilitative services, with the exception of crisis intervention services.

LMHAs delivering services to persons eligible for fee-for-service benefits must obtain authorization from their internal utilization management department. When providing care to persons enrolled in managed care, LMHAs and other providers contracted with MCOs must submit authorization requests to the MCO with whom the person is enrolled. The MCO may choose to waive this authorization submission requirement. Additionally, MCOs must follow the requirements set forth in the Uniform Managed Care Manual regarding utilization management for targeted case management and mental health rehabilitative services.

Eligibility determinations occur at the facility providing mental health rehabilitative services using the Clinical Management of Behavioral Health Services (CMBHS) software system.

Criteria used to make these service determinations are from the recommended LOC of the person as derived from the Uniform Assessment (UA), the needs of the person, and the Texas Resilience and Recovery Utilization Management Guidelines.

A facility that provides mental health rehabilitative services must ensure that at minimum a QMHP-CS administers the uniform assessment to the person at specified intervals (every 90 calendar days for persons who are 20 years of age and younger and every 180 calendar days for persons who are 21 years of age and older), and obtains a recommended LOC for the person.

The provider must evaluate the clinical needs of the person to determine if the amount of services associated with the recommended LOC described in the utilization management guidelines is sufficient to meet those needs and ensure that an LPHA reviews the recommended LOC and verifies whether the services are medically necessary.

Changes to the treatment plan with regard to type, amount, or duration of services must be approved by an LPHA practicing within the scope of his/her licensure.

If the facility determines that an LOC other than the recommended LOC is more appropriate for the person, the provider must submit a deviation request that includes:

- A request for an authorization of an LOC that is higher or lower than initially recommended.
- The clinical justification for the request.

The clinical justification must include the specific reason(s) why the person requires interventions outside the recommended LOC. Refusal of recommended LOC by the person receiving services may be noted as part of the justification.

All treatment plans are subject to retrospective review by the state.

5.2.3.9.1 **Reauthorization Requirements**

A QMHP-CS must conduct the uniform assessment at specified intervals (every 90 calendar days for children/youth and every 180 calendar days for adults) to determine the type, amount, and duration of mental health rehabilitative services.

Prior to the expiration of the authorization period or depleting the amount of services authorized, the provider must make a determination of whether the person continues to need mental health rehabilitative services. An LPHA must also determine whether the continuing need for mental health rehabilitative services meets the definition of medical necessity.
If the determination is that the person continues to need mental health rehabilitative services and that such services are medically necessary, the provider must:

- Request another authorization for the same type and amount of mental health rehabilitative service previously authorized; or
- Submit a request, with documented clinical reasons for such request, to change the type or amount of mental health rehabilitative services previously authorized if:
  - The provider determines that the type or amount of mental health rehabilitative services previously authorized is inappropriate to address the person’s needs.
  - The criteria described in the utilization management guidelines for changing the type or amount of mental health rehabilitative services has been met.

### 5.2.3.10 Documentation Requirements

All services require documentation to support the medical necessity of the service rendered. An LPHA must document in the person’s medical record that mental health rehabilitative services are medically necessary when the services are authorized and reauthorized.

Persons determined to need mental health rehabilitative services must have a treatment plan developed by the Medicaid enrolled provider of mental health rehabilitative services that describes in writing the type, amount, and duration of mental health rehabilitative services determined to be medically necessary to meet the needs of the person.

A rehabilitative services provider must document the following for all mental health rehabilitative services:

- The name of the person to whom the service was provided
- The type of service provided
- The specific goal or objective addressed, and the modality and method used to provide the service
- The date the service was provided
- The start and end time of the service
- The location where the service was provided
- The signature of the staff member providing the service and a notation of their credentials
- Any pertinent event or behavior relating to the person’s treatment which occurs during the provision of the service
- The outcome or progress in achieving treatment plan goals

In addition to the general requirements described above, when providing crisis services, a provider must document the following information:

- Risk of suicide or homicide
- Substance use
- Trauma, abuse, or neglect
- The outcome of the crisis (e.g., person in hospital, person with friend and scheduled to see doctor at 9:00 a.m. the following day)
- All actions (including rehabilitative interventions and referrals to other agencies) used by the provider to address the problems presented
- The response of the person, and if appropriate, the response of the LAR and family members
• Any pertinent event or behavior relating to the person’s treatment that occurs during the provision of the service
• Follow up activities that may include referral to another provider

Documentation for day programs for acute needs must be made daily. Documentation must be made after each face-to-face contact occurs to provide the mental health rehabilitative service for all other services.

An LPHA must, within two business days after crisis intervention services are provided, determine whether the crisis intervention services met the definition of medical necessity. If medical necessity is met then the LPHA must document the medical necessity.

Services are subject to retrospective review and recoupment if documentation does not support the service billed.

A provider must retain documentation in compliance with applicable federal and state laws, rules, and regulations.

5.2.3.11 Exclusions

Persons receiving psychosocial rehabilitation services are not eligible to simultaneously receive skills training and development or targeted case management services.

Mental health rehabilitative services do not include any of the following services that must be billed to Texas Medicaid:

• Rehabilitative services provided:
  • Before the establishment of a diagnosis of mental illness and authorization of services
  • To persons who reside in an institution for mental diseases
  • To general acute care hospital inpatients
• Services to residents of institutions that furnish food, shelter, and treatment to four or more unrelated persons
• Services to nursing facility residents who have not been identified through the PASSR process as needing specialized mental health services
• Services to inmates of public institutions
• Job task-specific vocational services
• Educational services
• Room and board residential costs
• Services that are an integral and inseparable part of another Medicaid-reimbursable service, including targeted case management services, residential rehabilitative behavioral health services, institutional and waiver services
• Services that are covered elsewhere in the state Medicaid plan
• Services to persons with a single diagnosis of intellectual or developmental disability or substance use disorder who do not have a co-occurring diagnosis of mental illness in adults or serious emotional disturbance in children
• Inpatient hospital services
• Respite services
• Family support services
5.2.3.12 Non-reimbursable Activities

A Medicaid provider will not be reimbursed for a mental health rehabilitative service:

- That is not included in the person’s treatment plan (except for crisis intervention services and psychosocial rehabilitative services provided in a crisis situation).
- That is not authorized, except for crisis intervention services.
- Provided in excess of the amount authorized.
- Provided outside of the duration authorized.
- Provided to a person receiving MH case management services.
- That is not documented.
- Provided to a person who does not meet the eligibility criteria.
- Provided to a person who does not have a current uniform assessment (except for crisis intervention services).
- Provided to a person who is not present, awake, and participating during such service.
- Provided via electronic media.

A Medicaid provider will not be reimbursed for a crisis service provided to a person who does not have a serious mental illness.

The cost of the following activities is included in the Medicaid mental health rehabilitative services reimbursement rate(s) and may not be directly billed by the Medicaid provider:

- Developing and revising the treatment plan and interventions that are appropriate to a person’s needs.
- Staffing and team meetings to discuss the provision of mental health rehabilitative services to a specific person.
- Monitoring and evaluating outcomes of interventions, including contacts with a person other than the person receiving services.
- Documenting the provision of mental health rehabilitative services.
- A staff member’s travel time to and from a location to provide mental health rehabilitative services.
- All services provided within a day program for acute needs that are delivered by a staff member, including services delivered in response to a crisis or an episode of acute psychiatric symptoms.
- Administering the uniform assessment to persons who are receiving mental health rehabilitative services.

5.3 Documentation Requirements

All services require documentation to support the medical necessity of the service rendered, including MH and IDD services.

MH and IDD services are subject to retrospective review and recoupment if documentation does not support the service billed.

5.4 Claims Filing and Reimbursement

IDD service coordination, MHTCM, and mental health rehabilitative services must be submitted to TMHP in an approved electronic claims format or on the CMS-1500 paper claim form. Providers may purchase CMS-1500 paper claim forms from the vendor of their choice. TMHP does not supply them.
When completing a CMS-1500 paper claim form, all required information must be included on the claim, as TMHP does not key any information from claim attachments. Superbills, or itemized statements, are not accepted as claim supplements.

Services are cost reimbursed in accordance with 1 TAC §§355.743, 355.746, and 355.781. Providers can refer to the OFL or the applicable fee schedule on the TMHP website at www.tmhp.com.

### 5.4.1 Managed Care Clients

Claims for persons in managed care must be submitted to the client’s MCO. Mental health targeted case management and mental health rehabilitative services that are funded by a criminal justice agency (submitted with modifier HZ) are carved out and must be submitted to TMHP.

### 5.4.2 Reimbursement Reductions

Texas Medicaid implemented mandated rate reductions for certain services. The OFL and static fee schedules include a column titled “Adjusted Fee” to display the individual fees with all mandated percentage reductions applied.

Additional information about rate changes is available on the TMHP website at www.tmhp.com/resources/rate-and-code-updates/rate-changes.

**Refero:** "Section 3: TMHP Electronic Data Interchange (EDI)” *(Vol. 1, General Information)* for information on electronic claims submissions.


Subsection 6.5, “CMS-1500 Paper Claim Filing Instructions” in “Section 6: Claims Filing” *(Vol. 1, General Information)*. Blocks that are not referenced are not required for processing by TMHP and may be left blank.

Subsection 2.2, “Fee-for-Service Reimbursement Methodology” in “Section 2: Texas Medicaid Fee-for-Service Reimbursement” *(Vol. 1, General Information)* for more information about reimbursement and the federal matching percentage.

### 6 Peer Specialist Services

#### 6.1 Services, Benefits, Limitations, and Prior Authorization

Peer specialist services (procedure code H0038) for mental health conditions or substance use disorders are a benefit of Texas Medicaid for persons who are 21 years of age and older, and who have peer specialist services included as a component of their person-centered recovery plan.

Peer specialist services are recovery-oriented, person-centered, relationship-focused, voluntary, and trauma-informed.

Peer specialist services may include the following:

- Recovery and wellness support services, which include providing information and support for recovery planning.
- Mentoring, which includes serving as a role model and helping find needed community resources and services.
- Advocacy, which includes providing support during stressful or urgent situations and helping to ensure that the person’s rights are respected. Advocacy may also include encouraging the person to advocate for him or herself to obtain services.
Peer specialist services are based on a mutual relationship between the peer specialist and the Medicaid eligible person. A peer specialist uses his or her lived experience to support the person with the following:

- Achieving the goals and objectives of the person’s individualized recovery plan
- Skill development
- Problem solving strategies
- Coping mechanisms for stressors and barriers encountered when recovering from a mental health condition or a substance use disorder

Peer specialist services can be delivered individually or in a group setting.

### 6.2 Peer Specialist Requirements

Peer specialist services may be delivered as part of a coordinated, comprehensive, and individualized approach to treating a person’s mental health condition, substance use disorder, or both, if the peer specialist is employed by one of the following Medicaid-enrolled provider types:

- Clinic/group practices that treat behavioral health conditions
- Physicians (M.D.s), osteopaths (D.O.s), and nurse practitioners (NPs), clinical nurse specialists (CNSs), and physicians assistants (PAs) that treat behavioral health conditions
- Psychologists, licensed clinical social workers, licensed marriage and family therapists, and licensed professional counselors
- Comprehensive provider agencies of targeted case management and mental health rehabilitative services
- Local mental and behavioral health authorities
- Chemical dependency treatment facilities
- Federally qualified health clinics (FQHCs)
- Rural health clinics (RHCs)
- Opioid Treatment Providers (OTPs)

Only clinic/group practices or behavioral health care individual providers (M.D., D.O., NP, CNS, and PA) with a behavioral health focus may be reimbursed for peer specialist services.

Peer specialists coordinate with all behavioral health service providers involved in the person’s care and utilize a person-centered, recovery-oriented approach to treatment planning and service delivery.

Non-Medicaid enrolled providers that employ peer specialists can contract with one of the listed Medicaid-enrolled provider types to furnish peer specialist services as part of a continuum of comprehensive treatment services. Subcontracted peer specialist services must also be part of the coordinated, comprehensive, and individualized person-centered recovery plan.

A peer specialist must meet all of the following criteria:

- Be at least 18 years of age.
- Have lived experience with a mental health condition, substance use disorder, or both.
- Have a high school diploma or General Equivalency Diploma (GED).
- Be willing to appropriately share his or her own recovery story with the person receiving services.
- Demonstrate current self-directed recovery.
- Pass criminal history and registry checks as described in 1 TAC §354.3201.
A peer specialist must not:

- Practice psychotherapy.
- Make clinical or diagnostic assessments.
- Dispense expert opinions.
- Engage in any service that requires a license.
- Falsify any documentation related to application, training, testing, certification, or services provided.

### 6.2.1 Certification

A peer specialist must complete all required training and certification before providing services. To be certified as a peer specialist as specified in 1 TAC §354.3155, a candidate must complete the following training:

- Required orientation
- Self-assessment activities
- Core training delivered by a certified training entity
- Supplemental training in one of two specialty areas:
  - Mental health peer specialist
  - Recovery support peer specialist

The candidate can apply for initial certification after successful completion of core and one supplemental training and a knowledge assessment.

A peer specialist who is initially certified may begin to deliver Medicaid billable services, if participating in a supervised internship at their place of employment. The internship consists of 250 hours of supervised work experience to be completed within a 6-month period. An extension may be granted by the certification entity should a peer be unable to complete the required hours within the 6-month time frame.

Independent study, such as reading or watching instructional videos, does not count toward the required supervised work experience hours. Time spent receiving supervision, other than the observation of the peer specialist providing services, does not count toward the required hours.

After completing the required internship hours, certified peer specialists can apply for renewed certification through the approved certification entity. Peer specialists must renew their certification every two years, which requires continuing education hours.

Certified peer specialists should only deliver services in their specialty area.

### 6.2.2 Supervision

As defined in 1 TAC §354.3003, providers may be reimbursed for peer specialist services rendered under the supervision of one of the following:

- Qualified credentialed counselor (QCC)
- Licensed practitioner of the healing arts (LPHA)
- Qualified mental health professional (QMHP), with a QCC or LPHA supervising the QMHP
- Qualified peer supervisor (QPS), with a QCC or LPHA supervising the QPS

Supervision must focus on a peer specialist’s provision of services, including:

- Review of cases and activities
• Skill building
• Problem resolution
• Professional growth

Supervision may also include aspects specific to the organization, such as following organizational policy or other administrative matters.

Peer specialist supervision may be provided as follows:
• Individually
• In a group setting
• Face-to-face
• By teleconference
• Through observation of the peer specialist providing services

Supervision must occur at least once weekly for a peer specialist with an initial certification, at least once monthly for a peer specialist with a two-year certification, or more frequently at the request of the peer specialist.

A QCC or LPHA who supervises a QMHP or QPS must provide individual or group supervision at least once monthly and conduct an observation of the QMHP or QPS supervising the peer specialist at a self-determined frequency based on the QMHP’s or QPS’s skill level.

A supervisor must successfully complete supervisory training for peer specialist services and the recovery model from a certified training entity before supervising a peer specialist. Supervisor training must include instruction about:
• The distinction between peer support and therapy.
• The role of peer support in building and sustaining recovery goals.
• Advocacy for peer specialists and peer specialist services.
• Job performance review, including strengths-based, timely, and respectful feedback.
• Supervisory skills, such as how to work with a variety of personality types and communication styles.

After completing training, each candidate must successfully complete a knowledge assessment before receiving approval to supervise a peer specialist from a certified training entity. Peer specialist supervisor certification must be renewed every two years, which requires continuing education hours.

### 6.3 Prior Authorization Requirements

Prior authorization is not required for the first 104 units of peer specialist services in a rolling 6-month period. Prior authorization is required once a person exceeds 104 units of individual or group peer specialist services in a rolling 6-month period.

Prior authorization requests for procedure code H0038 must be submitted to TMHP using the Special Medical Prior Authorization (SMPA) Request Form. Requests for continued services must demonstrate all of the following:
• The person continues to meet eligibility criteria as outlined in the statement of benefits above, including current DSM diagnoses
• The current person-centered recovery plan and goals
• The progress made relative to the goals outlined in the person-centered recovery plan
• The need for continued services

Requests must indicate how many additional units of service are being requested (up to 30 units are allowed per request) and the type (individual or group), as well as the expected time frame when services will be delivered.

*Note:* The requesting provider may be asked for additional information to clarify or complete a request.

Retrospective review may be performed to ensure that the documentation supports the medical necessity of the requested service.

*Refer to:* Refer to “Section 5: Fee-for-Service Prior Authorizations” (Vol. 1, General Information) for more information on submitting prior authorization methods.

### 6.4 Documentation Requirements

The Medicaid-enrolled provider must ensure proper documentation of all peer specialist services that are rendered. Documentation of peer specialist services must:

- Indicate the date, time, and place of service.
- Summarize the purpose and content of the services.
- Include the specific strategies and activities utilized as related to the goals of the person’s plan of care.

Peer specialist supervisors must document all supervisory sessions and maintain the records in the peer specialist’s employee personnel file.

#### 6.4.1 Reimbursement

Reimbursement for procedure code H0038 is limited to substance use disorders and mental health conditions, including, but not limited to:

- Schizophrenia spectrum and other psychotic disorders.
- Bipolar and related disorders.
- Depressive disorders.
- Anxiety disorders.
- Obsessive-compulsive and related disorders.
- Trauma and stressor related disorders.
- Feeding and eating disorders.

Procedure code H0038 is limited to 104 units in a rolling six-month period. This limitation may be exceeded with documentation of medical necessity for the additional services.

Peer specialist services will also be limited as follows:

- Must not be delivered simultaneously with other behavioral health services that are delivered to the person or group of persons receiving services
- Must be delivered in person and not through advanced telecommunications technology
- Limited to 12 total persons per group session

#### 6.4.2 Claim Filing

Procedure code H0038 must be submitted with one of the following specialty modifiers:

- Modifier HE-mental health
• Modifier HF-substance use

If services are provided in a group setting, procedure code H0038 must also be submitted with modifier HQ.

Mental health rehabilitative services must be billed separately from peer specialist services.

FQHCs and RHCs should submit claims using procedure code H0038 for informational purposes only.

### 6.5 Exclusions

The following services are not a benefit of Texas Medicaid:

- Record keeping or documentation activities
- Services provided without the person present

### 7 Psychiatric Services for Hospitals

Inpatient admissions to acute care hospitals for adults and children for psychiatric conditions are a benefit of Texas Medicaid. Admissions must be medically necessary and are subject to Texas Medicaid’s retrospective utilization review (UR) requirements. The UR requirements are applicable regardless of the hospital’s designation of a unit as a psychiatric unit versus a medical or surgical unit.

Persons who are 20 years of age and younger may be admitted to a freestanding psychiatric facility or a state psychiatric facility. Persons who are 21 years of age and older may be admitted only to an acute care facility. Providers should use the most appropriate revenue code when billing for inpatient psychiatric services in an acute care facility. A certification of need must be completed and placed in the person’s medical record within 14 days of the admission or once the person becomes Medicaid-eligible while in the facility.

Inpatient psychiatric treatment is a benefit of Texas Medicaid if all the following apply:

- The person has a psychiatric condition that requires inpatient treatment.
- The inpatient treatment is directed by a psychiatrist.
- The inpatient treatment is provided in a nationally accredited facility or hospital.
- The provider is enrolled in Texas Medicaid.

Persons of all ages may be admitted to an acute care facility. Inpatient admissions for the single diagnosis of chemical dependency or abuse (such as alcohol, opioids, barbiturates, and amphetamines) without an accompanying medical complication are not benefits of Texas Medicaid. Additionally, admissions for chronic diagnoses such as intellectual disability, organic brain syndrome, or chemical dependency or abuse are not covered benefits for acute care hospitals without an accompanying medical complication or medical condition. The UB-04 CMS-1450 paper claim form must indicate all relevant diagnoses that necessitate the inpatient stay.

Supporting documentation (certification of need) must be documented in the person’s medical record. This documentation must be maintained by each facility for a minimum of five years and be readily available for review when requested by HHSC or its designee.

Additional coverage through the Comprehensive Care Program (CCP) may be allowed for Medicaid-eligible persons who are 20 years of age and younger. Providers should use revenue code 124 when billing for inpatient psychiatric services in freestanding and state psychiatric facilities.

**Refer to:** Subsection 2.16, “Inpatient Psychiatric Hospital or Facility (Freestanding) (CCP)” in the Children’s Services Handbook (Vol. 2, Provider Handbooks).
7.1 Prior Authorization Requirements

Prior authorization is not required for persons with fee-for-service benefits who are admitted to psychiatric units in acute care hospitals. Out-of-network admissions require notification within the next business day and submission of clinical information to determine appropriateness for transfer to a contracted facility.

Prior authorization is not required for initial admission to freestanding psychiatric facilities or state psychiatric hospitals for persons who are 20 years of age and younger for a maximum of five days based on Medicaid eligibility and documentation of medical necessity. Extended stay requests beyond the initial 5 days require prior authorization.

Refer to: Subsection 2.17.3, “Prior Authorization and Documentation Requirements” in the Children’s Services Handbook (Vol. 2, Provider Handbooks) for more information about inpatient psychiatric services.

7.2 Documentation Requirements

Documentation of medical necessity for inpatient psychiatric care must specifically address the following issues:

- Why the ambulatory care resources in the community cannot meet the treatment needs of the person receiving the services.
- Why inpatient psychiatric treatment under the care of a psychiatrist is required to treat the acute episode of the person receiving the services
- How the services can reasonably be expected to improve the person’s condition or prevent further regression of the person’s condition in a proximate time period.

7.3 Inpatient Hospital Discharge

Procedure codes 99238 and 99239 must be submitted when billing for a hospital discharge.

8 Screening, Brief Intervention, and Referral to Treatment (SBIRT)

SBIRT is a comprehensive, public health approach to the delivery of early intervention and treatment services for persons who are 10 years of age and older and who have alcohol or substance use disorders or are at risk of developing such disorders. SBIRT is used for intervention directed to a person and not for group intervention.

SBIRT services can be provided by physicians, registered nurses, advanced practice nurses, physician assistants, psychologists, licensed clinical social workers, licensed professional counselors, certified nurse midwives, outpatient hospitals, federally qualified health centers (FQHCs), and rural health clinics (RHCs). Non-licensed providers may deliver SBIRT under the supervision of a licensed provider if such supervision is within the scope of practice for that licensed provider. The same SBIRT training requirements apply to non-licensed providers.

A person may have a maximum of two screening only sessions per rolling year, and up to four combined screening and brief intervention sessions per rolling year. Providers must refer the person to treatment if the screening results reveal severe risk of alcohol or substance use.

Refer to: Section 9, “Substance Use Disorder (SUD) Services” in this handbook for additional information on SUD treatment.
8.1  **SBIRT Training**

Providers that perform SBIRT must be trained in the correct practice of this method and will be required to complete at least four hours of training. Proof of completion of SBIRT training must be maintained in an accessible manner at the provider’s place of service.

Information regarding available trainings and standardized screening tools can be found through the Substance Abuse and Mental Health Services Administration.

8.2  **Screening**

Screening persons for problems related to alcohol or substance use identifies the person’s level of risk and determines the appropriate level of intervention indicated for the person. Providers must explain the screening results to the person, and if the results are positive, be prepared to subsequently deliver, or delegate to another provider, brief intervention services. Screening must be conducted using a standardized screening tool. Standardized tools that may be used include, but are not limited to, the following:

- Alcohol, Smoking, and Substance Involvement Screening Test (ASSIST)
- Drug Abuse Screening Test (DAST)
- Alcohol Use Disorders Identification Test (AUDIT)
- Cut-down, Annoyed, Guilty, Eye-opener (CAGE) questionnaire
- Car, Relax, Alone, Forget, Family or Friends, Trouble (CRAFFT) questionnaire
- Binge drinking questionnaire

Results obtained through blood alcohol content (BAC) or through toxicology screening may also be used to screen for alcohol or substance use risk.

8.3  **Brief Intervention**

Brief intervention is performed following a positive screen or a finding of at least a mild to moderate risk for alcohol or substance use. During the session, brief intervention involves motivational interviewing techniques (such as the Brief Negotiated Interview) that is focused on raising the person’s awareness of his or her alcohol or substance use and its consequences. The session is also focused on motivating the person toward behavioral change.

Subsequent screening and brief intervention sessions within the allowable annual limitations may be indicated to assess for behavior change and further explore a person’s readiness to make behavioral changes related to their alcohol or substance use.

*Note:* Providers may choose to schedule multiple screening and brief intervention sessions in a rolling year in order to provide ongoing support to a person at risk for substance use who is receptive to behavior change.

8.4  **Referral to Treatment**

If the provider determines that the person is in need of more extensive treatment or has a severe risk for alcohol or substance use, the person must be referred to an appropriate substance use treatment provider.
Referral to more extensive treatment is a proactive process that facilitates access to care for persons who require a more extensive level of service than SBIRT provides. Referral is an essential component of the SBIRT intervention because it ensures that all persons who are screened have access to the appropriate level of care.

**Note:** If the person is currently under the care of a behavioral health provider, the person must be referred back to that provider.

### 8.5 Reimbursement and Limitations

SBIRT is limited to persons who are 10 years of age and older. SBIRT is limited to up to two screening sessions per rolling year. A screening that results in a negative result does not require a brief intervention. In these instances procedure code H0049 should be used. A provider may re-screen a person within the same rolling year to determine whether their substance use behavior has changed.

Procedure code 99408 or G2011 should be used when a brief intervention follows an SBIRT screening. Procedure code 99408 is limited to once per day. SBIRT is limited to four sessions per rolling year when it constitutes a screening followed by a brief intervention.

If a person requires more than four combined screening and brief intervention sessions per rolling year, the person must be referred for substance use disorder treatment.

SBIRT is not reimbursable to providers (whether licensed or non-licensed) who have not completed the required number of training hours in SBIRT methodology.

Procedure codes 99408, G2011, and H0049 will be denied if billed for the same date of service as any of the following procedure codes:

<table>
<thead>
<tr>
<th>Procedure Codes</th>
</tr>
</thead>
<tbody>
<tr>
<td>90791 90792 90832 90833* 90834 90836* 90837 90838* 90847 90853</td>
</tr>
<tr>
<td>90865 90870 96130 96131* 96132 96133* 96136 96137*</td>
</tr>
</tbody>
</table>

Procedure codes 99408 and H0049 cannot both be billed on the same date.

Physicians and other qualified health care professionals that bill an Evaluation and Management (E/M) code for a visit where SBIRT occurred must use modifier 25 to identify a significant, separately identifiable E/M service rendered by the same provider on the same date of service.

**Note:** FQHCs and RHCs should submit claims using SBIRT procedure codes for informational purposes only.

### 8.6 Documentation Requirements

A person’s record documentation must support medical necessity for the SBIRT services provided and must be maintained by the SBIRT provider and made readily available for review when requested by the Health and Human Services Commission (HHSC) or its designee. SBIRT documentation for screening must include the following:

- The provider who performed the SBIRT screening
- Screening results from a standardized screening tool or laboratory results such as BAC, toxicology screen, or other measures showing risk for alcohol or substance use and the specific screening tool used.
Documentation for SBIRT brief intervention sessions must include a person-centered plan for the delivery of medically necessary services that supports the use of procedure code 99408. The plan must include all of the following:

- The provider who performed the SBIRT brief intervention, if different from the provider who screened the person.
- Start and stop time of the session, or the total face-to-face time spent providing SBIRT services to the person.
- Goals established.
- Specific strategies to achieve the goals.
- The person’s support system such as family members, a legal guardian, or friends.

Note: If subsequent sessions are indicated, the provider who performed the SBIRT session must document that a follow up SBIRT appointment was made and with whom, or document another mechanism established to reassess progress.

- The name, address, and phone number of the provider that the person has been referred to for substance use disorder treatment.

Services are subject to retrospective review to ensure that the documentation in the person’s medical record supports the medical necessity of the services provided.

### 8.7 Claims Filing and Reimbursement

SBIRT services must be submitted to TMHP in an approved electronic format or on the CMS-1500 paper claim form. Providers may purchase CMS-1500 paper claim forms from the vendor of their choice. TMHP does not supply the forms. When completing a CMS-1500 paper claim form, all required information must be included on the claim, as TMHP does not key any information from claim attachments. Superbills, or itemized statements, are not accepted as claim supplements.

**Refer to:** “Section 3: TMHP Electronic Data Interchange (EDI)” (Vol. 1, General Information) for information on electronic claims submissions.

Subsection 6.1, “Claims Information” in “Section 6: Claims Filing” (Vol. 1, General Information) for general information about claims filing.


Texas Medicaid rates for Hospitals are calculated according to 1 TAC §355.8061.

According to 1 TAC §355.8091, the Medicaid rate for LCSWs, LMFTs, and LPCs is 70 percent of the rate paid to a psychiatrist or psychologist for a similar service per 1 TAC §355.8085.

The Medicaid rates for psychologists are calculated in accordance with 1 TAC §355.8085.

Texas Medicaid rates for physicians and certain other practitioners are calculated in accordance with TAC §355.8085.

Texas Medicaid rates for Nurse Practitioners and Clinical Nurse Specialists are calculated in accordance with TAC §355.8281.
According to 1 TAC §355.8093, the Medicaid rate for PAs is 92 percent of the rate paid to a physician (MD or DO) for the same professional service and 100 percent of the rate paid to physicians for laboratory services, X-ray services, and injections. Services performed by a PA and billed under a physician’s or RHC’s provider identifier are reimbursed according to the Texas Medicaid Reimbursement Methodology (TMRM) for physician services.

**Note:** For more information about Texas Medicaid rates for the provider types above, refer to the OFL or the applicable fee schedule on the TMHP website at www.tmhp.com.

Texas Medicaid implemented mandated rate reductions for certain services. The OFL and static fee schedules include a column titled “Adjusted Fee” to display the individual fees with all mandated percentage reductions applied.

Additional information about rate changes is available on the TMHP website at www.tmhp.com/resources/rate-and-code-updates/rate-changes.

**Note:** Certain rate reductions including, but not limited to, reductions by place of service, client type program, or provider specialty may not be reflected in the Adjusted Fee column.

### 9 Substance Use Disorder (SUD) Services

#### 9.1 Overview

SUDs are chronic, relapsing medical illnesses that require an array of best practice medical and psychosocial interventions of sufficient intensity and duration to achieve and maintain remission and support progress toward recovery. SUD may include problematic use of alcohol, prescription drugs, illegal drugs (e.g., cannabis, opioids, stimulants, inhalants, hallucinogens, “club” drugs, other synthetic euphoriants), and other substances that may be identified in the future.

Treatment for SUD is a benefit of Texas Medicaid for persons who meet the criteria for a substance-related disorder, as outlined in the current edition of the American Psychiatric Association’s DSM.

SUD treatment services are individualized, age-appropriate medical and psychosocial interventions designed to treat a person’s problematic use of alcohol or other drugs, including prescription medication.

SUD services may include:

- Withdrawal management services.
- Individual and group SUD counseling in an outpatient setting.
- Residential treatment services.
- Medication assisted treatment.
- Evaluation and treatment (or referral for treatment) for co-occurring physical and behavioral health conditions.

Level of care (e.g., outpatient, residential, inpatient hospital) and specific services provided must adhere to current evidence-based industry standards and guidelines for SUD treatment, such as those outlined in the current edition of the American Society of Addiction Medicine’s Treatment Criteria for Addictive Substance-Related and Co-Occurring Conditions, as well as the licensure requirements outlined in 25 TAC §448 pertaining to standards of care.

SUD treatment services (outpatient or residential) may only be delivered in a licensed chemical dependency treatment facility (CDTF). Medication assisted treatment (MAT) may also be delivered in the office setting by appropriately trained physicians, physician assistants (PAs), and advanced practice registered nurses (APRNs) who are recognized by the Texas Board of Nursing as either nurse practi-
tioners (NPs), clinical nurse specialists (CNSs), nurse anesthetists (CRNAs), or nurse midwives (CNMs), provided that the APRN is a qualifying practitioner and possesses the Drug Addiction Treatment Act (DATA) waiver.

SUD withdrawal management in an inpatient hospital setting may be provided for persons who meet hospital level of care requirements as a result of the severity of their withdrawal syndrome or the severity of their co-occurring conditions. These services may be reimbursed as general hospital inpatient services.

The treatment setting and the intensity or level of services will vary depending on the severity of the person’s SUD and what is clinically appropriate. The intensity or level of services refers to the number of hours of services per week, as well as the types of services the person receives. Early Intervention services are part of the spectrum of SUD treatment and are a benefit in Texas Medicaid. Early intervention services target persons who are at risk of developing a substance related problem but may not have a diagnosed SUD.

Refer to: Subsection 8, “Screening, Brief Intervention, and Referral to Treatment (SBIRT)” in this handbook for further information on early intervention services.

Upon admission into a treatment setting, a face-to-face multi-dimensional assessment (procedure code H0001) must be conducted by a qualified credentialed counselor (QCC) or intern as defined in DSHS TAC §441.101 to determine a course of treatment that is medically necessary and clinically appropriate. The assessment must be signed off by a QCC.

9.2 Evaluation, Treatment, or Referral for Co-Occurring Conditions
CDTFs shall facilitate access to physical health, mental health, and ancillary services if those services are not available through the program and are necessary to meet treatment goals or needs of the person receiving services.

Persons in residential CDTFs commonly require medications unrelated to their SUD treatment for which costs are not covered in the reimbursement for SUD or MAT services. These medications, if included in the Medicaid formulary, may be obtained and reimbursed through the person’s Medicaid pharmacy benefit.

Persons in residential CDTFs also commonly require other services that are benefits of Texas Medicaid, but not included in the CDTF rate. Claims for these services can be submitted by the appropriate providers.

CDTFs should screen each person for risk for contracting tuberculosis, Hepatitis B and C, HIV antibody, and sexually transmitted infections, and if appropriate, provide access to testing and follow up. Testing may be performed on site and billed by the ordering provider if appropriate testing facilities are available that are compliant with the rules and regulations for the Clinical Laboratory Improvement Amendments (CLIA). Providers that do not comply with CLIA are not reimbursed for laboratory services.

9.3 Withdrawal Management Services
Withdrawal management, formerly known as detoxification, is the medical and behavioral treatment of persons experiencing or potentially experiencing withdrawal symptoms as a result of ceasing or reducing substance use.

Withdrawal management involving opioids, alcohol, sedatives, hypnotics, or anxiolytics will vary depending on the severity of the withdrawal symptoms experienced but will typically involve medications to treat symptoms in addition to supportive care, observation, and monitoring. Withdrawal management involving stimulants, inhalants, and cannabis typically involves supportive care, observation and monitoring, and medications to treat withdrawal symptoms as required.
Withdrawal management may be performed in an outpatient setting for persons experiencing mild to moderate withdrawal symptoms that can be successfully, as well as safely, managed outside of a residential setting or an inpatient hospital. Withdrawal management in a residential setting may be required for persons whose multidimensional assessment indicates one or more of the following circumstances that would make outpatient withdrawal management unsafe or unsuccessful:

- A level of severity of withdrawal, medical, or mental health complication
- Sufficient challenges with readiness to change, ability to stop using, or social support

Withdrawal management in an inpatient hospital setting may be required for persons whose severity of medical withdrawal (e.g., impending delirium tremens, severe withdrawal seizures), comorbid medical conditions (e.g., severe liver impairment, acute pneumonia, endocarditis, dementia), or comorbid psychiatric conditions (e.g., severe suicidality, acute and unstable psychosis or mania) requires a hospital level of care.

### 9.4 Individual and Group SUD Counseling in an Outpatient Setting

Counseling for substance use disorders is designed to assist persons in developing a better understanding of their SUD, help to establish treatment goals and plans for achieving those goals, and provide interventions to assist persons in accordance with the plan. The overall intent of the service is to assist persons in understanding their SUD and developing the skills and supports needed to address their SUD over time. Counseling may be done individually or in a group setting with multiple members. Group counseling sessions are limited to a total of 16 persons per session.

Outpatient counseling services are appropriate for the following:

- Persons with less severe disorders
- Persons who are in the early stages of change
- As a step down from more intensive services
- Persons who are stable but for whom ongoing monitoring is appropriate

**Note:** For persons unable or unwilling to access SUD treatment services at a CDTF, psychotherapy delivered by a licensed practitioner of the healing arts (LPHA) may be an alternative treatment option to address a person’s SUD.

Outpatient services may be appropriate at the start of treatment, throughout treatment, or after an episode of residential or inpatient treatment, depending on the person’s acuity, severity, comorbidity, needs, or preferences. Outpatient services can address active symptoms as well as provide ongoing treatment for persons in partial or full remission who need continuing help to maintain progress.

Abstinence should not necessarily be a requirement for participation in outpatient services.

### 9.5 Residential Treatment Services

Residential treatment programs provide a structured therapeutic environment where persons reside with staff support and deliver comprehensive substance use disorder treatment with attention to co-occurring conditions as appropriate. The frequency and duration of services should be based on meeting the person’s needs and achieving the person’s treatment goals.

Residential services are appropriate for persons who require a structured therapeutic environment to stabilize SUD and develop coping and recovery skills. Residential treatment programs may specialize in the unique needs of a specific population such as adolescents, or pregnant or parenting women with children.
Episodes of residential treatment may be required for persons with more severe SUD, more significant medical or psychiatric comorbidities, more significant challenges with sustaining motivation, maintaining control in an outpatient setting, or a living environment that jeopardizes their current ability to be successful in outpatient treatment.

Residential SUD treatment services may only be provided by a licensed CDTF.

**9.6 Outpatient Treatment Services**

Outpatient treatment services must be billed with procedure codes H0004 or H0005.

Procedure codes H0004 and H0005 are limited to the following diagnosis codes:

<table>
<thead>
<tr>
<th>Diagnosis Codes</th>
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</thead>
<tbody>
<tr>
<td>F1010</td>
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<tr>
<td>F10139</td>
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<tr>
<td>F10188</td>
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<td>F10231</td>
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<tr>
<td>F14282</td>
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<td>F1493</td>
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<td>F14988</td>
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</tbody>
</table>
9.7 Medication Assisted Treatment

MAT is the use of FDA-approved medications in combination with psychosocial treatment to treat substance use disorders, particularly alcohol and opioid use disorders.

MAT is a recognized best practice for alcohol use disorder (AUD) and opioid use disorder (OUD). All persons with AUD and OUD should be educated about the availability of MAT and the evidence supporting MAT, and have the opportunity to receive MAT regardless of where they are receiving SUD services. This could be accomplished on site or through a written agreement with a collaborating opioid treatment program (OTP) or office-based opioid treatment (OBOT) program.

Initiation or induction of MAT can appropriately occur in lieu of withdrawal management for opioid use disorders, may begin early in withdrawal management for either AUD or OUD, and can be initiated as appropriate at any point in time during the course of treatment. Duration of MAT is determined on an person basis, depending on the person’s unique needs and treatment goals.

Determination of which MAT medication to use is also an individualized treatment decision based on provider assessment and the person’s needs and treatment goals. Providers are encouraged to offer as many treatment options as possible (within the parameters of their licensing and scope of practice) to maximize the person’s choice and access to care.

MAT may be utilized as appropriate, as part of the service array delivered by outpatient providers or residential treatment services programs at CDTFs.
Opioid treatment programs (also referred to as narcotic treatment programs) are the only settings permitted by law to provide methadone for OUD and must comply with additional federal and state requirements, rules on licensure and scope of practice, including physician delegation, supervision, and prescriptive authority. Opioid treatment programs can also provide or administer other forms of MAT.

CDTFs, physicians, NPs, and PAs may prescribe and provide for the administration of long acting injectable naltrexone (Vivitrol) to treat cravings associated with either opioid use disorder or alcohol use disorder.

Physicians, PAs, and APRNs who are recognized by the Texas Board of Nursing as either NPs, CNSs, APRNs, or CNMs who have received a federal waiver to dispense buprenorphine may choose to incorporate this form of MAT into their medical practice while also providing or referring for other types of treatment services (also referred to as OBOT).

Certain MAT medications to treat alcohol and opioid use disorders (such as buprenorphine, disulfiram, acamprosate, and naltrexone), are available as a pharmacy benefit and may be prescribed to a person by their physician or other qualified health care professional. Providers may refer to the Vendor Drug Program Formulary for additional information on covered medications.

Prescribing of certain MAT medications may be done via telemedicine presuming all other applicable state and federal laws are followed.

A prescription for an opioid antagonist (e.g., naloxone) should be given to all persons receiving treatment for opioid use disorder, and instruction should be provided on how to administer if needed.

Claims for urinalysis drug screens ordered by a physician, NP, or CNS to monitor compliance with MAT may be submitted by the individually-enrolled physician or APRN.

The following MAT procedure codes may be separately reimbursed from withdrawal management and treatment services in the outpatient or residential setting:

<table>
<thead>
<tr>
<th>Procedure Codes</th>
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<tbody>
<tr>
<td>H0020</td>
<td>H0033</td>
</tr>
<tr>
<td>J0570</td>
<td>J2315</td>
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<tr>
<td>Q9991</td>
<td>Q9992</td>
</tr>
</tbody>
</table>

### 9.7.1 Opioid Treatment Providers

Substance Abuse and Mental Health Services Administration certified (SAMHSA-certified) opioid treatment providers (OTPs) that are also licensed as narcotic treatment programs in Texas are required to enroll in Medicare before enrolling with Texas Medicaid as OTPs. Providers billing claims for persons who have dual eligibility for Medicaid and Medicare must first submit their claims to Medicare.

The following procedure codes may be reimbursed to Opioid Treatment Providers:

<table>
<thead>
<tr>
<th>Procedure Codes</th>
<th></th>
</tr>
</thead>
<tbody>
<tr>
<td>H0001</td>
<td>H0004</td>
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<tr>
<td>H0005</td>
<td>H0020</td>
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<td>H0033</td>
<td>Q9991</td>
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<tr>
<td>J0570</td>
<td>J2315</td>
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</tbody>
</table>

**Important:** CDTFs cannot bill for OTP services through Medicare, as CDTFs are not Substance Abuse and Mental Health Services Administration certified (SAMHSA-certified) OTPs.

A comprehensive assessment (procedure code H0001) is limited to once per day, any provider. An assessment is also limited to once per episode of care and should be performed at the start of each new episode of care.


9.8 Exclusions
SUD treatment services for tobacco use disorder as the primary diagnosis are not a covered benefit, although a comprehensive SUD treatment approach should address tobacco use if reducing or eliminating this substance is part of the person’s treatment goal.

With the exception of prescribing MAT medications via telemedicine, SUD treatment services may not be delivered via telemedicine or telehealth.

9.9 Prior Authorization
The following services do not require prior authorization:

- Assessment
- Outpatient treatment services
- MAT

**Exception:** Outpatient treatment services require prior authorization if the calendar year hours/units are exceeded. Those limits are 135 units of group services and 26 hours of individual services per calendar year.

The following services require prior authorization:

- Outpatient withdrawal management services
- Outpatient treatment for persons who exceed the benefit limitation
- Residential withdrawal management services
- Residential treatment services

Providers must submit the appropriate prior authorization request form for the initial or continuation of outpatient or residential withdrawal management treatment and residential treatment services. A QCC (as defined by the DSHS licensure standard) must complete and sign the prior authorization request forms.

Providers must submit one of the following forms to obtain prior authorization:

- Outpatient Withdrawal Management Authorization Request Form
- Outpatient Substance Use Disorder Counseling Extension Request Form
- Residential Withdrawal Management Authorization Request Form
- Residential Substance Use Disorder Treatment Request Form

Prior authorization will be considered for the least restrictive environment appropriate to the person’s medical need as determined in the person’s plan of care (POC), based on national standards.

Prior authorization requests for services beyond the limitations outlined in this section, may be considered with documentation supporting the medical necessity for continuation of the treatment.

9.9.1 Prior Authorization to Persons with Fee-For-Service Benefits
Prior authorization requests for persons with fee-for-service benefits may be submitted to the TMHP Prior Authorization Unit online at www.tmhp.com, by fax at 1-512-514-4211, or by mail to:

Texas Medicaid & Healthcare Partnership
TMHP Prior Authorization Department
12357-B Riata Trace Parkway, Suite 100
Austin, TX 78727
To complete the prior authorization process electronically, the provider must complete the prior authorization requirements through any approved electronic methods. Providers must retain a copy of the signed and dated prior authorization form in the person’s medical record.

Providers may contact the TMHP Prior Authorization Unit by telephone at 1-800-213-8877, Option 2, to obtain information about substance use disorder benefits, the prior authorization process, or the status of a prior authorization request. Prior authorization for substance use disorder services cannot be obtained through this line.

Prior authorization for outpatient withdrawal management, residential treatment, or residential withdrawal management services will be considered when requested within three business days after the date of admission.

Prior authorization may be considered for persons who are enrolled in a Medicaid MCO when they are admitted to SUD services and whose eligibility changes to fee-for-service during treatment. Requests must be submitted within three business days of the date on which the fee-for-service eligibility started.

**9.9.2 Prior Authorization for Outpatient Withdrawal Management Treatment Services**

Outpatient withdrawal management services may be prior authorized for up to 21 days. The level of service and number of days that are prior authorized will be based on the substances that are used, level of intoxication and withdrawal potential, and the person’s medical needs. Providers may submit requests for services using the Outpatient Withdrawal Management Authorization Request Form.

**9.9.2.1 Admission Criteria for Outpatient Withdrawal Management Treatment Services**

The admission criteria for outpatient withdrawal management treatment services follow the existing Texas Department of Insurance licensure requirements and standards that are specified in 28 TAC §3.8001–§3.8030.

To be considered eligible for treatment for outpatient withdrawal management services, the person must meet the following conditions:

- **Chemical Substance Withdrawal**—The person must meet all of the following criteria with regard to chemical substance withdrawal:
  - The person is expected to have a stable withdrawal from alcohol or drugs.
  - The diagnosis must meet the criteria for the definition of substance use disorder or the most current revision of the DSM accompanied by evidence that some of the symptoms have persisted for at least one month or have occurred repeatedly over a longer period of time.

- **Medical Functioning**—The person must meet all of the following criteria with regard to medical functioning:
  - No history of recent seizures or past history of seizures during withdrawal.
  - No clinical evidence of altered mental state as manifested by disorientation to self, alcoholic hallucinations, toxic psychosis, or altered level of consciousness (clinically significant obtundation, stupor, or coma).
  - The symptoms are due to withdrawal and not due to a general medical condition. Absence of any presumed new asymmetric or focal findings (i.e., limb weakness, clonus, spasticity, unequal pupils, facial asymmetry, eye ocular movement paresis, papilledema, or localized cerebellar dysfunction, as reflected in asymmetrical limb coordination).
  - Stable vital signs as interpreted by a physician. The person must also be without a previous history of complications from acute chemical substance withdrawal and judged to be free of a health risk as determined by a physician.
• No evidence of a coexisting serious injury or systemic illness either newly discovered or progressive in nature.
• Absence of serious disulfiram-alcohol (Antabuse) reaction with hypothermia, chest pains, arrhythmia, or hypotension.
• Clinical condition that allows for a comprehensive and satisfactory assessment.

• Family, Social, or Academic Dysfunction—The person must meet at least one of the following criteria with regard to family, social, or academic dysfunction:
  • The person’s social system and significant others are supportive of recovery to the extent that the person can adhere to a treatment plan and treatment service schedules without substantial risk of reactivating the person’s substance use disorder.
  • The person’s family or significant others are willing to participate in the outpatient withdrawal management treatment program.
  • The person may or may not have a primary or social support system to assist with immediate recovery, but the person has the social skills to obtain such a support system or to become involved in a self-help fellowship.
  • The person does not live in an environment where licit or illicit mood altering substances are being used. A person living in an environment where licit or illicit mood altering substances are being used may not be a candidate for this level of care.

• Emotional and Behavioral Status—The person must meet all of the following criteria with regard to emotional and behavioral status:
  • The person is coherent, rational, and oriented for treatment.
  • The mental state of the person does not preclude the person’s ability to comprehend and understand the materials presented, and the person is able to participate in the outpatient withdrawal management treatment process.
  • Documentation exists in the medical record that the person expresses an interest to work toward outpatient withdrawal management treatment goals.
  • The person has no neuropsychiatric condition that places the person at imminent risk of harming self or others (e.g. pathological intoxication or alcohol idiosyncratic intoxication).
  • The person has no neurological, psychological, or uncontrolled behavior that places the person at imminent risk of harming self or others (depression, anguish, mood fluctuations, overreactions to stress, lower stress tolerance, impaired ability to concentrate, limited attention span, high level of distractibility, negative emotions, or anxiety).
  • The person has no documented DSM condition or disorder that, in combination with alcohol or drug use, compounds a pre-existing or concurrent emotional or behavioral disorder and presents a major risk to the person.
  • The person has no mental confusion or fluctuating orientation.

• Chemical Substance Use—The person must meet the criteria in at least one of the following conditions with regard to recent chemical substance use:
  • The person’s chemical substance use is excessive, and the person has attempted to reduce or control it but has been unable to do so (as long as chemical substances are available).
  • The person is motivated to stop using alcohol or drugs and is in need of a supportive, structured treatment program to facilitate withdrawal from chemical substances.
9.9.2.2 Continued Stay Criteria for Outpatient Withdrawal Management Treatment Services

A person is considered eligible for continued stay in the outpatient withdrawal management treatment service when the person meets at least one of the conditions for either chemical substance withdrawal or psychiatric or medical complications. Requests for extension of services must be received on or before the last date authorized or denied. The prior authorization unit will notify the provider by fax or electronic portal. If the date of the prior authorization unit determination letter is on or after the last date authorized or denied or if the last date falls on a holiday or weekend, the request for extension of services is due by 5 p.m. of the next business day. Documentation in the person’s medical record must support either Chemical Substance Withdrawal or Psychiatric or Medical Complications.

Chemical Substance Withdrawal
The person must meet at least one of the following conditions with regard to chemical substance withdrawal complications:

- The person, while physically abstinent from chemical substance use, is exhibiting incomplete stable withdrawal from alcohol or drugs, as evidenced by psychological and physical cravings.
- The person, while physically abstinent from chemical substance use, is exhibiting incomplete stable withdrawal from alcohol or drugs, as evidenced by significant drug levels.

Psychiatric or Medical Complications
The person must meet both of the following psychiatric or medical complication conditions:

- The intervening medical or psychiatric event was serious enough to interrupt the outpatient withdrawal management treatment.
- Evidence that the person is progressing in treatment again.

9.9.3 Prior Authorization for Residential Withdrawal Management Treatment Services

 Withdrawal management services may be prior authorized for up to 21 days. The level of service and number of prior authorized days will be based on the substances that are used, level of intoxication and withdrawal potential, and the person’s medical needs. Providers may submit requests for services using the Residential Withdrawal Management Authorization Request form.

Requests for withdrawal management services for persons who need more than 21 days of residential withdrawal management require review of documentation of medical necessity from a provider who is familiar with the person.

9.9.3.1 Admission Criteria for Residential Withdrawal Management Treatment Services

The admission criteria for residential withdrawal management treatment services follow the existing Texas Department of Insurance licensure requirements and standards that are specified in 28 TAC §3.8001–§3.8030.

A person is eligible for admission to a residential withdrawal management service when they have failed two previous individual treatment episodes of outpatient withdrawal management or when they have a diagnosis that meets the criteria for the definition of substance use disorder or the most current revision of the DSM.

The person must also meet at least one of the following criteria for admission to residential withdrawal management treatment:

- Chemical Substance Withdrawal—The person must have impaired neurological functions as evidenced by:
- Extreme depression (e.g., suicidal).
- Altered mental state with or without delirium as manifested by disorientation to self; alcoholic hallucinosis, toxic psychosis, altered level of consciousness, as manifested by clinically significant obtundation, stupor, or coma.
- History of recent seizures or past history of seizures on withdrawal.
- The presence of any presumed new asymmetric or focal findings (i.e., limb weakness, clonus, spasticity, unequal pupils, facial asymmetry, eye ocular movement paresis, papilledema, or localized cerebellar dysfunction, as reflected in asymmetrical limb incoordination).
- Unstable vital signs combined with a history of past acute withdrawal syndromes that are interpreted by a physician to be indication of acute alcohol or drug withdrawal.
- Evidence of coexisting serious injury or systemic illness, newly discovered or progressive.
- Clinical condition (e.g., agitation, intoxication, or confusion) that prevents satisfactory assessment of the above conditions and indicates placement in residential withdrawal management service may be justified.
- Neuropsychiatric changes of such severity and nature that they put the person at imminent risk of harming self or others (e.g., pathological intoxication or alcohol idiosyncratic intoxication).
- Serious disulfiram-alcohol (Antabuse) reaction with hypothermia, chest pains, arrhythmia, or hypotension.
- Major Medical Complications—The person must present a documented condition or disorder that, in combination with alcohol or drug use, presents a determined health risk (e.g., gastrointestinal bleeding, gastritis, severe anemia, uncontrolled diabetes mellitus, hepatitis, malnutrition, cardiac disease, hypertension).
- Major Psychiatric Illness—The person must meet at least one of the following conditions with regard to major psychiatric illness:
  - Documented DSM condition or disorder that, in combination with alcohol or drug use, compounds a pre-existing or concurrent emotional or behavioral disorder and presents a major risk to the person.
  - Severe neurological and psychological symptoms: (e.g., anguish, mood fluctuations, overreactions to stress, lowered stress tolerance, impaired ability to concentrate, limited attention span, high level of distractibility, extreme negative emotions, or extreme anxiety).
  - Danger to others or homicidal.
  - Uncontrolled behavior that endangers self or others, or documented neuropsychiatric changes of a severity and nature that place the person at imminent risk of harming self or others.
  - Mental confusion or fluctuating orientation.

### 9.9.3.2 Continued Stay Criteria for Residential Withdrawal Management Treatment Services

Eligibility for continued stay for residential withdrawal management services is based on the person meeting at least one of the criteria for chemical substance withdrawal, major medical complications, or major psychiatric complications.
Chemical Substance Withdrawal
The person must exhibit one of the following conditions with regard to chemical substance withdrawal complications:

- Incomplete medically stable withdrawal from alcohol or drugs, as evidenced by documentation of at least one of the following conditions:
  - Unstable vital signs
  - Continued disorientation
  - Abnormal laboratory findings related to chemical dependency
  - Continued cognitive deficit related to withdrawal so that the person is unable to recognize alcohol or drug use as a problem
  - Laboratory finding that, based on the judgment of a physician, indicates that a drug has not sufficiently cleared the person’s system

Major Medical Complications
For major medical complications, the person must have documentation in the medical record that indicates that a medical condition or disorder (e.g., uncontrolled diabetes mellitus) continues to present a health risk and is being actively treated.

Major Psychiatric Complications
The person must meet at least one of the following with regard to major psychiatric complications:

- Documentation in the medical record that a psychiatric condition or disorder that, in combination with alcohol or drug use, continues to present a major health risk, is actively being treated.
- Documentation in the medical record that severe neurological or psychological symptoms have not been satisfactorily reduced but are actively being treated.

9.9.4 Prior Authorization for Residential Treatment Services
Residential treatment may be prior authorized for up to 35 days per episode of care, with a maximum of two episodes of care per rolling six-month period and four episodes of care per rolling year.

Providers can use the Residential Substance Use Disorder Treatment Request form to submit authorization requests for persons who require additional episodes within the 6- or 12-month time frame.

9.9.4.1 Admission Criteria for Residential Treatment Services
The admission criteria for residential treatment services follow the existing Texas Department of Insurance licensure requirements and standards that are specified in 28 TAC §3.8001-§3.8030.

The diagnosis must meet the criteria for the definition of substance use disorder or the most current version of the DSM.

All persons must meet the following conditions in order to receive treatment in a residential treatment service program:

- Medical Functioning—The following must be present with regard to medical functioning:
  - Documented medical assessment following admission (except in instances where the person is being referred from an inpatient service) indicates that the person is medically stable and not in acute withdrawal.
  - The person is not bed-confined and has no medical complications that would hamper participation in the residential service.
- Family, Social, or Academic Dysfunction and Logistic Impairments—At least one of the following must be present with regard to family, social, or academic dysfunction and logistic impairments:
• The person manifests severe social isolation or withdrawal from social contacts.

• The person lives in an environment (social and interpersonal network) in which treatment is unlikely to succeed (e.g., a chaotic family dominated by interpersonal conflict, which undermines person’s efforts to change).

• The person’s family or significant others are opposed to the person’s treatment efforts and are not willing to participate in the treatment process.

• Family members or significant others living with the person manifest current substance use disorders and are likely to undermine treatment.

• Logistic impairments (e.g., distance from treatment facility or mobility limitations) preclude participation in an outpatient treatment setting.

• Emotional and Behavioral Status—The person must meet all three of the following criteria with regard to emotional and behavioral status:
  
  • The person is coherent, rational, and oriented for treatment.
  
  • Mental state of the person does not preclude the person’s ability to comprehend and understand the materials presented and participate in rehabilitation or the treatment process.
  
  • The medical record contains documentation that with continued treatment the person will be able to improve or internalize the person’s motivation toward recovery within the recommended length of stay time frames (e.g., becoming less defensive, verbalizing, and working on alcohol or drug related issues). Interventions, treatment goals, or contracts are in place to help the person deal with or confront the blocks to treatment (e.g., family intervention or employee counseling confrontation).

• Chemical Substance Use—The person must meet at least one of the following criteria with regard to chemical substance use:
  
  • The person’s chemical substance use is excessive, and the person has attempted to reduce or control it but has been unable to do so (as long as chemical substances are available).
  
  • Virtually all of the person’s daily activities revolve around obtaining, using, or recuperating from the effects of chemical substances, and the person requires a secured environment to control the person’s access to chemical substances.

9.9.4.2 Residential Treatment Services for Adolescents

Adolescents who are 13 through 17 years of age must meet all above conditions and the following conditions in order to receive treatment in an adolescent residential treatment service program:

• At the maturation level, the adolescent must meet both of the following criteria:
  
  • The adolescent is assessed as manifesting physical maturation at least in middle adolescent range (i.e., post-pubescent).
  
  • The history of the adolescent reflects cognitive development of at least 11 years of age.

• The adolescent must display at least one of the following with regard to developmental status:
  
  • Documented history of inability to function within the expected age norms despite normal cognitive and physical maturation (e.g., refusal to interact with family members, overt prostitution, felony, or other criminal charges).
  
  • A recent history of moderate to severe conduct disorder, as defined in the DSM, or impulsive disregard for social norms and rights of others.
Documented difficulty in meeting developmental expectations in a major area of functioning (e.g., social, academic, or psychosexual) to an extent that interferes with the capacity to remain behaviorally stable.

9.9.4.3 Continued Stay Criteria for Residential Treatment Services

At least one of the following conditions must be present for continued stay in a residential treatment program:

- Chemical Dependency Rehabilitation or Treatment Complications:
  - The person recognizes or identifies with the severity of the alcohol or drug problem but demonstrates minimal insight into the person’s defeating the use of alcohol or drugs. However, documentation in the medical record indicates that the person is progressing in treatment.
  - The person identifies with the severity of the alcohol or drug problem and manifests insight into the person’s personal relationship with mood-altering chemicals, yet does not demonstrate behaviors that indicate the development of problem-solving skills that are necessary to cope with the problem.
  - The person would predictably relapse if moved to a lesser level of care.

- Psychiatric or Medical Complications:
  - Documentation in the medical record indicates an intervening medical or psychiatric event that was serious enough to interrupt rehabilitation or treatment, but the person is again progressing in treatment.
  - Documentation in the medical record indicates that the person is being held pending an immediate transfer to a psychiatric, acute medical service, or inpatient withdrawal management alcohol or drug service.

9.9.5 Prior Authorization for Outpatient Treatment Services

Prior authorization for outpatient treatment services beyond the annual limitation of 135 units of group services and 26 hours of individual services per calendar year, may be considered with documentation supporting medical necessity for continued treatment services. Providers may submit requests requiring additional services using the Outpatient Substance Use Disorder Counseling Extension request form.

Requests must be submitted before providing the extended services. The documentation must include the following information:

- The person is meeting treatment goals.
- The person demonstrates insight and understanding into relationship with mood altering chemicals, but continues to present with issues addressing the life functions of work, social, or primary relationships without the use of mood-altering chemicals.
- The person is physically abstinent from chemical substance use, but remains mentally preoccupied with such use to the extent that the person is unable to adequately address primary relationships, or social or work tasks, but there are indications that, with continued treatment, the person will effectively address these issues.
- Although other psychiatric or medical complications exist that affect the person’s treatment, there is documentation to support the person continues to show treatment progress and there is evidence to support the benefits of continued treatment.
9.10 Documentation Requirements
To facilitate determination of medical necessity and avoid unnecessary denials, the provider must provide correct and complete information, including documentation for medical necessity for the services requested. The provider must maintain documentation of medical necessity in the person’s medical record.

The requesting provider may be asked for additional information to clarify or complete a request.

Retrospective review may be performed to ensure documentation supports the medical necessity of the requested services.

9.11 Reimbursement and Limitations
9.11.1 Withdrawal Management Services
Inpatient hospital-based withdrawal management is reimbursed by the reimbursement methodology specific to the inpatient hospital. Separate reimbursement may be provided for physician services performed during an inpatient stay.

Residential withdrawal management and treatment services are considered outpatient services for the purposes of reimbursement and should be billed accordingly.

Residential withdrawal management services (procedure codes H0012, H0031, S9445, and T1007) are limited to once per day, any provider.

Residential withdrawal management services (procedure codes H0031, H0047, S9445, or T1007) will be denied if billed without lead procedure code H0012 on the same day, same provider.

Room and board for residential withdrawal management and treatment (procedure code H0047) is limited to once per date of service, any provider. Procedure code H0047 is reimbursed for persons who are 21 years of age and older as an access-based fee and as an informational detail for persons who are 20 years of age and younger.

Outpatient withdrawal management (procedure codes H0016, H0050, and S9445) is limited to once per day, any provider and may be reimbursed on the same date of service as outpatient SUD treatment by the same or different provider when medically necessary and identified in the person’s treatment plan.

Outpatient withdrawal management (procedure codes H0050 and S9445) will be denied if billed without lead procedure code H0016 on the same day, same provider.

Separate reimbursement may be provided for physician services during a residential stay.

9.11.1.1 Treatment Services
Outpatient treatment services are limited to 135 units of group counseling and 26 hours of individual counseling per calendar year when provided by a CDTF. Providers may submit requests requiring additional services using the Outpatient Substance Use Disorder Counseling Extension request form.

Residential treatment services (procedure code H2035) are limited to one per day and are allowed up to a maximum of 35 days.

Outpatient treatment (procedure codes H0004 and H0005) will be denied if billed on the same date of service as residential withdrawal management (procedure codes H0012, H0031, H0047, S9445, and T1007) or residential treatment (procedure code H2035).

Procedure code H0047 will be denied if billed without lead procedure code H2035 or H0012 on the same day, same provider.

Refer to: Subsection 6.4.1, “National Correct Coding Initiative (NCCI) Guidelines” in “Section 6: Claims Filing” (Vol. 1, General Information) for information about NCCI MUE guidelines.
9.11.2 MAT Services

Claims billed for MAT must include the person’s substance use disorder diagnosis. MAT billing may include billing for induction as well as maintenance.

Methadone administration (procedure code H0020) for opioid disorder must be submitted with the following modifiers:

- When methadone is administered with supervision in a facility the provider must submit claims using the UA modifier to indicate the facility administered doses
- When methadone is dispensed without supervision as a take home dose the provider must submit claims using the U1 modifier to indicate take home doses

Methadone provided in an outpatient setting (procedure code H0020) is limited to once per date of service, by any provider and is reimbursed at a fixed daily rate. Reimbursement for procedure code H0020 with modifier U1 is limited to a quantity of 30 per 30 days, any provider.

Providers that allow take-home doses must submit procedure code H0020 with modifier U1 for each date of service for which a take-home dose is dispensed. Methadone that is dispensed for unsupervised take-home use should be dispensed in alignment with the federal opioid treatment standards in Title 42 Code of Federal Regulations (CFR) §8.12.

Methadone administration (procedure code H0020) submitted without a modifier will be denied.

Non-methadone (e.g., buprenorphine) administration (procedure code H0033) for opioid disorder must be submitted with the following modifiers:

- When non-methadone is administered with supervision in a facility the provider must submit claims using the modifier UA to indicate opioid disorder treatment facility doses or claims will be denied
- When non-methadone is dispensed without supervision as a take home dose the provider must submit claims using the modifier U1 to indicate opioid disorder take home doses or claims will be denied

Non-methadone provided in an outpatient setting (procedure code H0033) is limited to once per date of service, by any provider. Reimbursement for procedure code H0033 with modifier U1 is limited to a quantity of 30 per 30 days, any provider.

Providers that allow take-home doses must submit procedure code H0033 with modifier U1 for each date of service for which a take-home dose is dispensed. Non-methadone that is dispensed for unsupervised take-home use should be dispensed in alignment with the federal opioid treatment standards in Title 42 Code of Federal Regulations (CFR) §8.12.

When non-methadone is administered in a facility for a non-opioid treatment, providers must use procedure code H0033 to indicate non-opioid treatment in a facility.

Non-methadone administration (procedure code H0033) submitted without a modifier will be denied. Physician and physician extenders may be reimbursed separately using the appropriate evaluation and management procedure codes.

Injectable administration is considered part of MAT and is not reimbursed separately. Procedure code 96372 will be denied when billed for the same date of service by any provider as procedure code H0020 or H0033.

Texas Medicaid implemented mandated rate reductions for certain services. The OFL and static fee schedules include a column titled “Adjusted Fee” to display the individual fees with all mandated percentage reductions applied.
Additional information about rate changes is available on the TMHP website at www.tmhp.com/resources/rate-and-code-updates/rate-changes.

Note: Certain rate reductions including, but not limited to, reductions by place of service, client type program, or provider specialty may not be reflected in the Adjusted Fee column.

9.12 Claims Filing

Claims for SUD services must be submitted to TMHP in an approved electronic format or on the CMS-1500 paper claim form. Providers may purchase CMS-1500 paper claim forms from the vendor of their choice. TMHP does not supply the forms. When completing a CMS-1500 paper claim form, all required information must be included on the claim, as TMHP does not key any information from claim attachments. Superbills, or itemized statements, are not accepted as claim supplements.

Refer to: “Section 3: TMHP Electronic Data Interchange (EDI)” (Vol. 1, General Information) for information about electronic claims submissions.

Subsection 6.1, “Claims Information” in “Section 6: Claims Filing” (Vol. 1, General Information) for general information about claims filing.


10 Claims Resources

Refer to the following sections or forms when filing claims:

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11 Contact TMHP

Providers can call the TMHP Contact Center at 1-800-925-9126 from Monday through Friday, 7 a.m. to 7 p.m., Central Time.
12 Forms

The following linked forms can also be found on the Forms page of the Provider section of the TMHP website at www.tmhp.com:

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13 Claim Form Examples

The following linked claim form examples can also be found on the Claim Form Examples page of the Provider section of the TMHP website at www.tmhp.com:

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1 General Information

The information in this handbook is intended for dentists, school districts, physicians, physician assistants (PAs), rural health clinics (RHCs), federally qualified health centers (FQHCs), advanced practice registered nurses (APRNs), home health agencies (HHAs), durable medical equipment (DME) suppliers, hospitals, and clinics. The handbook provides information about Texas Medicaid’s benefits, policies, and procedures applicable to these providers.

**Important:** All providers are required to read and comply with “Section 1: Provider Enrollment and Responsibilities” (Vol. 1, General Information). In addition to required compliance with all requirements specific to Texas Medicaid, it is a violation of Texas Medicaid rules when a provider fails to provide health-care services or items to Medicaid clients in accordance with accepted medical community standards and standards that govern occupations, as explained in 1 Texas Administrative Code (TAC) §371.1659. Accordingly, in addition to being subject to sanctions for failure to comply with the requirements that are specific to Texas Medicaid, providers can also be subject to Texas Medicaid sanctions for failure, at all times, to deliver health-care items and services to Medicaid clients in full accordance with all applicable licensure and certification requirements including, without limitation, those related to documentation and record maintenance.

All providers are required to report suspected child abuse or neglect as outlined in subsection 1.7, “Provider Responsibilities” in “Section 1: Provider Enrollment and Responsibilities” (Vol. 1, General Information).

1.1 Medical Transportation Program

The Medical Transportation Program (MTP) is funded with federal and state dollars to arrange nonemergency transportation to medical or dental appointments for eligible clients and their attendants.

The Health and Human Services Commission (HHSC) administrative rules govern parental accompaniment of children who receive Medicaid screenings, treatments, and MTP services.

Titles 1 Texas Administrative Code (TAC), Part 15, §380.207 allows parents or guardians to authorize one adult and one alternate adult to accompany their children on MTP rides when the parent or guardian is unable to do so. The parent or guardian is required to designate the other adult on a form prescribed by HHSC in accordance with section §380.207(4).

An adult who is authorized by a parent or guardian may not be a provider or an employee or affiliate of a provider that submits claims for services.

**Refer to:** The Medical Transportation Program Handbook (Vol. 2, Provider Handbooks) for more information.

1.2 Rates Reduction

Texas Medicaid implemented mandated rate reductions for certain services. The Online Fee Lookup (OFL) and static fee schedules include a column titled “Adjusted Fee” to display the individual fees with all mandated percentage reductions applied. Additional information about rate changes is available on the Texas Medicaid & Healthcare Partnership (TMHP) website at [www.tmhp.com/resources/rate-and-code-updates/rate-changes](http://www.tmhp.com/resources/rate-and-code-updates/rate-changes).
1.3 NP, CNS, PA, and CNM Claims Submitted by a Physician

Physicians will be reimbursed 92 percent of the established reimbursement rate for services provided by an NP, CNS, PA, or CNM if the physician does not make a decision regarding the client’s care or treatment on the same date of service as the billable medical visit. Physicians who submit a claim using the physician’s own provider identifier for the services that were provided by the NP, CNS, PA, or CNM must submit one of the following modifiers on each claim detail:

- SA—Services were provided by an NP or CNS
- U7—Services were provided by a PA
- SB—Services were provided by a CNM

Exception: The 92 percent reimbursement rate does not apply to laboratory services, radiology services, or injections provided by an NP, CNS, PA, or CNM.

1.4 Payment Window Reimbursement Guidelines for Services Preceding an Inpatient Admission

According to the three-day and one-day payment window reimbursement guidelines, most professional and outpatient diagnostic and nondiagnostic services that are rendered within the designated time frame of an inpatient hospital stay and are related to the inpatient hospital admission will not be reimbursed separately from the inpatient hospital stay if the services are rendered by the hospital or an entity that is wholly owned or operated by the hospital.

These reimbursement guidelines do not apply in the following circumstances:

- Services are rendered at a federally qualified health center (FQHC) or rural health clinic (RHC).
- Services are for a THSteps medical checkup.
- Professional services are rendered in the inpatient hospital setting.
- The hospital and the physician office or other entity are both owned by a third party, such as a health system.
- The hospital is not the sole or 100-percent owner of the entity.

These reimbursement guidelines do not apply for FQHC, RHC, THSteps, and professional services that are rendered in the inpatient hospital setting.

Refer to: Subsection 3.7.4.14, "Payment Window Reimbursement Guidelines" in the Inpatient and Outpatient Hospital Services Handbook (Vol. 2, Provider Handbooks) for additional information about the payment window reimbursement guidelines.

2 Medicaid Children’s Services Comprehensive Care Program (CCP)

2.1 CCP Overview

CCP is an expansion of the Early Periodic Screening, Diagnosis, and Treatment (EPSDT) service as mandated by the Omnibus Budget Reconciliation Act (OBRA) of 1989, which requires all states to provide all medically necessary treatment for correction of physical or mental problems to Texas Health Steps (THSteps)-eligible clients when federal financial participation (FFP) is available, even if the services are not covered under the state’s Medicaid plan.

Under the Early Periodic Screening, Diagnostic, and Treatment (EPSDT) regulation, known in Texas as Texas Health Steps (THSteps), Section 1905(r) of the Social Security Act mandates that all Medicaid-eligible beneficiaries who are birth through 20 years of age receive medically necessary services to treat, correct and ameliorate illnesses and conditions identified if the service is covered in the state’s Medicaid plan.
plan or is an optional Medicaid service. It is the responsibility of the state to determine medical necessity on a case specific basis. No arbitrary limitations on services are allowed (e.g., one pair of eyeglasses or 10 therapy sessions per year) if determined to be medically necessary.

Services not covered under this section include:

- Experimental or investigational treatment.
- Services or items not generally accepted as effective and/or not within the normal course and duration of treatment.
- Services for the caregiver or provider convenience.

All EPSDT requirements must be adhered to for beneficiaries who receive services under managed care arrangements.

The following CCP provider sections describe the specific requirements of each area of responsibility:

- Subsection 2.5, “Clinician-Directed Care Coordination Services (CCP)” in this handbook.
- Subsection 2.10, “Medical Nutrition Counseling Services (CCP)” in this handbook.
- Subsection 2.8, “Early Childhood Intervention (ECI) Services” in this handbook.
- Subsection 2.11, “Personal Care Services (PCS) (CCP)” in this handbook.
- Subsection 2.16, “Inpatient Psychiatric Hospital or Facility (Freestanding) (CCP)” in this handbook.
- Subsection 2.17, “Inpatient Rehabilitation Facility (Freestanding) (CCP)” in this handbook.

Refer to:

- The Physical Therapy, Occupational Therapy, and Speech Therapy Services Handbook (Vol. 2, Provider Handbooks) for information about comprehensive outpatient rehabilitation facilities (CORFs) and outpatient rehabilitation facilities (ORFs).
- The Home Health Nursing and Private Duty Nursing Services Handbook (Vol. 2, Provider Handbooks) for more information about private duty nursing (PDN) services.
- The Physical Therapy, Occupational Therapy, and Speech Therapy Services Handbook (Vol. 2, Provider Handbooks) for more information about CCP therapy services.
- The Certified Respiratory Care Practitioner (CRCP) Services Handbook (Vol. 2, Provider Handbooks) for more information about CRCP CCP services.

2.1.1 Client Eligibility

The client must be birth through 20 years of age and eligible for THSteps on the date of service. If the client’s Your Texas Benefits Medicaid card states “Emergency,” “PE,” or “QMB,” the client is not eligible for CCP benefits.

Clients are ineligible for CCP services beginning the day of their 21st birthday.

2.1.2 Enrollment

Refer to: Subsection 1.7.18, “Children’s Services Comprehensive Care Program (CCP)” in “Section 1: Provider Enrollment and Responsibilities” (Vol. 1, General Information) for enrollment information.

2.1.3 Services, Benefits, and Limitations

Payment is considered for any health-care service that is medically necessary and for which FFP is available. CCP benefits are allowable services not currently covered under Texas Medicaid (e.g., speech-language pathology [SLP] services for nonacute conditions, PDN, prosthetics, orthotics, apnea monitors and some DME, some specific medical nutritional products, medical nutrition services, inpatient
rehabilitation, travel strollers, and special needs car seats). CCP benefits also include expanded coverage of current Texas Medicaid services where services are subject to limitations (e.g., diagnosis restrictions for total parenteral nutrition [TPN] or diagnosis restrictions for attendant care services).

Requests for services that require a prior authorization must be submitted to TMHP. Prior authorization is a condition for reimbursement, not a guarantee of payment. For information about specific benefits, providers can refer to provider-specific sections of this manual.

Payment cannot be made for any service, supply, or equipment for which FFP is not available. The following are some examples:

- Vehicle modification, mechanical, or structural (such as wheelchair lifts).
- Structural changes to homes, domiciles, or other living arrangements.
- Environmental equipment, supplies, or services, such as room dehumidifiers, air conditioners, filters, space heaters, fans, water purification systems, vacuum cleaners, and treatments for dust mites, rodents, and insects.
- Ancillary power sources and other types of standby equipment (except for technology-dependent clients such as those who are ventilator-dependent for more than six hours per day).
- Educational programs, supplies, or equipment (such as a personal computer or software).
- Equine or hippotherapy.
- Exercise equipment, home spas or gyms, toys, therapeutic balls, or tricycles.
- Tennis shoes.
- Respite care (relief to caregivers).
- Aids for daily living (toothbrushes, spoons, reachers, and foot stools).
- Take-home drugs from hospitals (Eligible hospitals may enroll in and bill Vendor Drug Program (VDP). Pharmacies that want to enroll should call 1-512-491-1429.
- Therapy involving any breed of animal.

### 2.1.4 Prior Authorization and Documentation Requirements

Prior authorization is a condition for reimbursement; it is not a guarantee of payment. A prior authorization number (PAN) is a TMHP-assigned number establishing that a service or supply has been determined to be medically necessary and for which FFP is available. It is each provider’s responsibility to verify the client’s eligibility at the time each service is provided. Any service provided while the client is not eligible cannot be reimbursed by TMHP. The responsibility for payment of services is determined by private arrangements made between the provider and client.

Prior authorization of CCP services may be requested in writing by completing the appropriate request form, attaching any necessary supportive documentation, and submitting them by mail, fax, or the electronic portal to the TMHP-CCP department. Prescribing or ordering providers, dispensing providers, clients’ responsible adults, and clients may sign prior authorization forms and supporting documentation using electronic or wet signatures.

Providers who fax new prior authorization requests, resubmitted requests, or additional information to complete a request must include:

- A working fax number on the prior authorization form, so that they can receive faxed responses and correspondence from TMHP.
- The last four digits of the client’s Medicaid identification number on the fax coversheet.
Prior authorization may also be requested through the TMHP website. (Providers can refer to Subsection 5.5.1, “Prior Authorization Requests Through the TMHP Website” in “Section 5: Fee-for-Service Prior Authorizations” (Vol. 1, General Information) for additional information to include mandatory documentation and retention requirements). All requested information on the form must be completed, or the request is returned to the provider. Incomplete forms are not accepted. If prior authorization is granted, the potential service provider (such as the DME supplier, pharmacy, registered nurse (RN), or physical therapist) receives a letter that includes the PAN, the procedures prior authorized, and the length of the authorization. Providers are notified in writing when additional information is needed to process the request for services.

Providers must submit a CCP Prior Authorization Request Form and documentation to support medical necessity to the CCP department before providing services. Providers must submit the CCP Prior Authorization Request Form when requesting a medically necessary service if the service is not addressed in the Texas Medicaid Provider Procedures Manual and the client is 20 years of age or younger.

Important: Documentation to support medical necessity of the service, equipment, or supply (such as a prescription, letter, or medical records) must be current, signed, and dated by a physician (M.D. or D.O.) before services are performed. Providers must keep the information on file.

Refer to: CCP provider-specific sections for prior authorization requirements of specific services, including the appropriate prior authorization request forms.

2.1.4.1 Incomplete Prior Authorization Requests

Providers must respond to an incomplete prior authorization request within 14 business days of the request receipt date. Incomplete prior authorization requests are requests that are received by TMHP with missing, incomplete, or illegible information.

Prior to denying an incomplete request, TMHP’s Prior Authorization department will attempt to get the correct information from the requesting provider. The Prior Authorization department will make a minimum of three attempts to contact the requesting provider before sending a letter to the client about the status of the request and the need for additional information.

If the necessary information to make a prior authorization determination is not received within 14 business days of the request receipt date, the request will be denied as “incomplete.” To ensure timely processing, providers should respond to requests for missing or incomplete information as quickly as possible.

For fee-for-service (FFS) Medicaid requests that require a physician review before a final determination can be made, TMHP’s Physician Reviewer will complete the review within three business days of receipt of the completed prior authorization request. An additional three business days will be allowed for requests that require a peer-to-peer review with the client’s prescribing physician.

2.1.4.2 Diagnosis Coding

All providers must obtain the client’s medical diagnosis from the physician. This information must be reflected on each claim submitted to TMHP.

2.1.4.3 Drug and Medical Device Approval

Manufacturers may request to have drug or medical device products added as a CCP benefit by sending the information in writing to the following address:

HHSC
1100 West 49th Street
Austin, TX 78756-3179

HHSC reviews the information. Requests for consideration must not be sent to TMHP.
2.1.4.4 Physician Signature

The dated signature of the physician (M.D. or D.O.) on a prescription or CCP Authorization Request Form must be current to the service date(s) of the request, i.e., the signature must always be on or before the service start date and no older than three months before the current date(s) of service requested. Physician signatures dated after the service start date on initial requests cannot be accepted as documentation supporting medical necessity for dates of service prior to the signature date. A request for prior authorization must include documentation from the provider to support the medical necessity of the service, equipment, or supply. If services begin as a result of a verbal order before the physician’s dated signature, proof of the verbal order must be submitted with the request.

Stamped signatures and dates are not accepted on CCP Authorization Request Forms or prescriptions for CCP prior authorized services, supplies, or equipment. Verbal orders must be cosigned and dated by a physician (M.D. or D.O.) within two weeks, per provider policy. Signatures of chiropractors or doctors of philosophy (PhDs) are not accepted on CCP Authorization Request Forms or prescriptions for CCP prior authorized services.

Certified nurse midwife (CNM), clinical nurse specialist (CNS), nurse practitioner (NP), and PA providers may sign on behalf of the physician for private duty nursing, physical, occupational and speech therapy services when the physician delegates this authority.

Physician prescriptions must be specific to the type of service requested.

2.2 Managed Care Organization (MCO) Clients Who Transition to Medicaid Fee-For-Service (FFS)

When clients transition from an MCO to FFS, providers can request that previously approved authorizations for Comprehensive Care Program (CCP) services, occupational therapy (OT), physical therapy (PT), private duty nursing (PDN), and speech therapy (ST) be transferred from the MCO to FFS.

2.2.1 Submission Guidelines

TMHP will consider the reimbursement of claims for services that were rendered on or after the MCO’s disenrollment date only when the provider submits a request to TMHP to transfer the previously approved authorization for CCP services.

The request to TMHP must be received on or before the end date of the previously approved MCO authorization. Any requests submitted after the MCO’s authorization end date will have to meet the regular submission guidelines for the specific service type.

2.2.2 Documentation Requirements

All of the requests to transfer the authorizations from the MCO to FFS must include:

- A copy of the previously approved authorization letter.
- All of the documentation that was sent in the original authorization request, including any physician orders that were used to determine the start of care. TMHP will accept the physician orders as the required documentation for the requested services.
- The completed CCP Prior Authorization Request form, Special Medical Prior Authorization (SMPA) form, Home Health Plan of Care, or Texas Medicaid Physical, Occupational, or Speech Therapy (PT, OT, ST) Prior Authorization Form, whichever is applicable for the requested service. The form must include the dates of service and quantities that are being requested from TMHP, which must match the dates of service and quantities that were approved in the original authorization.

Note: It is not necessary to obtain signatures or dates on the forms listed above when submitted to TMHP for the purpose of transferring an authorization from an MCO to FFS Medicaid.
Authorizations for services transferred from an MCO to FFS Medicaid are subject to retrospective review.

TMHP will verify the client’s eligibility, the dates of service, and the quantities requested.

TMHP will process reimbursement claims as follows:

- Claims for services that were rendered before the date on which the transfer request was received will be denied as a late submission, and the provider will be notified of their administrative appeal rights through the Health and Human Services Commission (HHSC).

- Claims for services that were rendered on or after the date of receipt use the required information from the transferred authorization and will be processed as if the request was received in a timely manner.

- Claims for services that were paid by an MCO and then recouped must contain the recoupment EOB from the MCO for consideration of payment. The claims must meet the 95-day deadline from the recoupment disposition date.

  Note: Letter requests for refunds will not be accepted. A recoupment EOB with a disposition date is required.

If a request to transfer an MCO authorization is submitted after the end date of the MCO authorization or the provider does not have an authorization letter from the MCO, TMHP will process the request to transfer the authorization based on established TMHP authorization submission guidelines for CCP services, PDN, OT, PT, and ST.

All new requests for rendered services must meet the documentation requirements.

### 2.2.3 New Services and Extension of Services

For new services that occur after the client’s MCO disenrollment change date, the provider is responsible for submitting all TMHP required paperwork and meeting all established submission guidelines for prior authorization.

Requests for the extension of services that occur after the MCO disenrollment change date must include all of the paperwork that is required by TMHP and meet all established submission guidelines for prior authorization.

### 2.2.4 Loss of Eligibility

If an MCO disenrolled a client and the client also loses Medicaid eligibility, providers must anticipate, if and when Medicaid eligibility is restored, that the client will initially be considered a Medicaid FFS client and will have a retroactive eligibility period.

All requests for services that require prior authorization and that occur during the client’s retroactive eligibility period, must be submitted to TMHP following the process that is outlined in subsection 5.1.1, “Prior Authorization Requests for Clients with Retroactive Eligibility” in “Section 5: Fee-for-Service Prior Authorizations” (Vol. 1, General Information).

If a client is retroactively disenrolled by an MCO, all of the services that are rendered by the provider during this retroactive disenrollment period (specifically from the date on which the client was eligible for FFS to the date of the client’s MCO eligibility change) will be denied by TMHP, and the provider will be notified of their administrative appeal rights.

TMHP may consider services for the MCO transition beginning on the date of the client’s MCO eligibility change date and going forward. TMHP uses the MCO transition process for the submission of paperwork and the processing of provider requests.
2.3 **Breastfeeding Support Services**

*Refer to:* Section 3, “Breastfeeding Support Services” in the *Gynecological, Obstetrics, and Family Planning Title XIX Services Handbook (Vol. 2, Provider Handbooks)* for information about breastfeeding support services.

2.4 **Certified Respiratory Care Practitioner Services (CCP)**

*Refer to:* The *Certified Respiratory Care Practitioner (CRCP) Services Handbook (Vol. 2, Provider Handbooks)* for information about CRCP (CCP) services.

2.5 **Clinician-Directed Care Coordination Services (CCP)**

2.5.1 **Services, Benefits, and Limitations**

Clinician-directed (physician, NP, CNS, and PA) care coordination services are a benefit of CCP for eligible clients who are birth through 20 years of age and have special health needs. These services are payable only to the clinician (primary care, specialist, or sub-specialist) who provides the medical home for the client.

To provide a medical home for the client, the primary care clinician directs care coordination together with the client and family. Care coordination consists of managing services and resources for clients with special health needs and their families to maximize the clients’ potential and provide them with optimal health care.

Clinician-directed care coordination services (face-to-face and non-face-to-face) must include the following components:

- A written care plan (either a formal document or documentation contained in the client’s progress notes) developed and revised by the medical home clinician, in partnership with the client, family, and other agreed-upon contributors. This plan is shared with other providers, agencies, and organizations involved with the care of the client, including educational and other community organizations with permission of the client or family. The care plan must be maintained by the medical home clinician and reviewed every six months or more frequently as necessary for the client’s needs.

- Care among multiple providers that are coordinated through the clinician.

- A central record or database maintained by the medical home clinician containing all pertinent medical information, including hospitalizations and specialty care.

- Assistance for the client or family in communicating clinical issues when a client is referred for a consultation or additional care, such as evaluation, interpretation, implementation, and management of the consultant recommendations for the client or family in partnership and collaboration with other providers, the client, or family.

Clinician-directed care coordination services must also include the supervision of the development and revision of the client’s emergency medical plan in partnership with the client, the family, and other providers for use by emergency medical services (EMS) personnel, utility service companies, schools, other community agencies, and caregivers.

Face-to-face care coordination services are encompassed within the various levels of evaluation and management (E/M) encounters and prolonged services.

Non-face-to-face clinician-directed care coordination services include:

- Prolonged services (procedure codes 99358 and 99359).

- Medical team conference (procedure code 99367).
• Care plan oversight and supervision, including telephone consultations with a specialist or subspecialist (procedure codes 99339, 99340, 99374, 99375, 99377, 99378, 99379, and 99380).
• Specialist or subspecialist telephone consultations (procedure code 99499 with modifier U9).

Non-face-to-face clinician-directed care coordination services are not considered case management by Texas Medicaid.

Specifically, non-face-to-face medical home clinician oversight and supervision of the development or revision of a client’s care plan may include the following activities, which do not have to be contiguous:
• Review of charts, reports, treatment plans, and lab or study results, except for the initial interpretation or review of lab or study results ordered during, or associated with, a face-to-face encounter.
• Telephone calls with other Medicaid-enrolled health-care professionals (not employed in the same practice) involved in the care of the client.
• Telephone or face-to-face discussions with a pharmacist about pharmacological therapies (not just ordering a prescription).
• Medical decision-making.
• Activities to coordinate services, if the coordination activities require the skill of a clinician.
• Documenting the services provided, which includes writing a note in the client’s chart describing the services provided, decision-making performed, and the amount of time spent performing the countable services, including the start and stop times and time spent by the physician working on the care plan after the nurse has conveyed pertinent information from agencies and facilities to the physician.

The following activities are not covered as non-face-to-face clinician supervision of the development or revision of the client’s care plan (care plan oversight services):
• Time that the staff spends getting or filing charts, calling home health agencies or clients, and similar administrative actions.
• Clinician telephone calls to client or family, except when necessary to discuss changes in client’s care plan.
• Clinician time spent telephoning prescriptions to a pharmacist (does not require clinician work and does not require a clinician to perform).
• Clinician time getting or filing the chart, dialing the telephone, or time on hold (does not require clinician work and does not meaningfully contribute to the treatment of the illness or injury).
• Travel time.
• Time spent preparing claims and for claims processing.
• Initial interpretation or review of lab or study results that were ordered during, or associated with, a face-to-face encounter.
• Services included as part of other E/M services.
• Consultations with health professionals not involved in the client’s case.

2.5.1.1 Non-Face-to-Face Services
2.5.1.1.1 Non-Face-to-Face Medical Conferences
Procedure code 99367 must be used when billing for medical team conferences.
2.5.1.1.2 Non-Face-to-Face Clinician Supervision of a Home Health Client
Procedure code 99374 or 99375 must be used when billing for services requiring interaction with a home health agency.

2.5.1.1.3 Non-Face-to-Face Clinician Supervision of a Hospice Client
Procedure code 99377 or 99378 must be used when billing for services requiring interaction with a hospice.

2.5.1.1.4 Non-Face-to-Face Clinician Supervision of a Nursing Facility Client
Procedure code 99379 or 99380 must be used when billing for services requiring interaction with a nursing facility.

2.5.1.1.5 Other Non-Face-to-Face Supervision
Procedure code 99339 or 99340 must be used when billing for services requiring interaction with an independently-enrolled nurse or other provider (e.g., not a home health agency, nursing facility, or hospice provider).

2.5.1.1.6 Non-Face-to-Face Prolonged Services
Procedure code 99358 or 99359 must be used when billing for prolonged services without face-to-face contact. This service is to be reported in addition to other clinician services, including E/M services at any level, or health-care professionals outside of a home health agency, hospice, or nursing facility.

Non-face-to-face prolonged services are limited to a maximum of 90 minutes once per client by the same provider unless one of the following significant changes in the client’s clinical condition occurs:

- The client will soon be, or has recently been, discharged from a prolonged and complicated hospitalization that required coordination of complex care with multiple providers in order for the client to be adequately cared for in the home.
- The client has experienced recent trauma resulting in new medical complications that require complex interdisciplinary care.
- The client has a new diagnosis of a medically complex condition requiring additional interdisciplinary care with additional specialists.

Procedure code 99359 must be billed on the same date of service as procedure code 99358. Additional prolonged non-face-to-face services may be authorized if the provider submits supporting documentation for authorization.

Procedure code 99358 must be used to report the first hour of prolonged services and must be billed with the appropriate physician E/M procedure code listed in the following table. Prolonged services of less than 30 minutes are considered part of the physician's E/M service being provided.

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<th>Procedure Codes</th>
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<td>99221</td>
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Procedure code 99359 is used to report an additional 15 to 30 minutes of prolonged non-face-to-face services beyond the first hour. Prolonged services of less than 15 minutes beyond the first hour are considered part of the first hour.
Non-face-to-face prolonged services procedure codes 99358 and 99359 may be used when billing for completion of paperwork required by a judge to determine guardianship of a client. Required paperwork may include a certified medical examination form for clients who are birth through 20 years of age that are eligible for the Texas Health Steps program on the date of service.

### 2.5.1.1.7 Non-Face-to-Face Specialist or Subspecialist Telephone Consultation

Telephone consultations are limited to two every six months to the same provider and will not be reimbursed to the clinician providing the medical home.

The clinician providing the medical home must have an authorization on file for one of the following procedure codes before the specialist or subspecialist can be reimbursed:

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<th>Procedure Codes</th>
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<td>99339</td>
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Because the specialist or sub-specialists cannot be reimbursed without the medical home clinician’s current prior authorization information, the clinician providing the medical home should provide their information to the specialist or subspecialist.

The specialist or subspecialist will not be separately reimbursed for the telephone consultation if he or she is the medical home clinician because care plan oversight by the medical home provider includes telephone consultations. The referring provider identifier and prior authorization number must be submitted on the claim.

### 2.5.1.1.8 General Requirements for Non-Face-to-Face Clinician-Directed Care Coordination Services

These services may be reimbursed for the medical home clinician time involved in this coordination. The clinician billing the services must personally perform the services. Care coordination services delegated to, or performed by others, do not count towards care coordination reimbursement. Care coordination provided during post-surgical care is a benefit if the care is unrelated to the surgery.

### 2.5.1.1.9 Non-Face-to-Face Care Plan Oversight

The medical home clinician who bills for the care plan oversight must be the clinician who signed the plan of care (POC) in the home or domiciliary (procedure codes 99339 and 99340), home health agency (procedure codes 99374 and 99375), hospice (procedure codes 99377 and 99378), or nursing facility (procedure codes 99379 and 99380).

Procedure code 99339 is denied when billed on the same date of service by the same provider as procedure code 99340.

Procedure code 99374 is denied when billed on the same date of service by the same provider as procedure code 99375.

Procedure code 99377 is denied when billed on the same date of service by the same provider as procedure code 99378.

Procedure code 99379 is denied when billed on the same date of service by the same provider as procedure code 99380.

Care plan oversight services may be reimbursed for the clinician time involved in this coordination. The clinician billing the services must personally perform the services. Care coordination services delegated to or performed by others do not count towards care coordination reimbursement.

Only one clinician-directed care plan oversight service (procedure codes 99339, 99340, 99374, 99375, 99377, 99378, 99379 or 99380) will be reimbursed per client, per calendar month to any provider.

The medical home clinician may not have a significant financial or contractual relationship with the home health agency as defined in 42 Code of Federal Regulations (CFR) §424.
The medical home clinician may not be the medical director or employee of the hospice and may not furnish services under arrangements with the hospice, including volunteering.

### 2.5.1.10 Medical Team Conference

One medical team conference (procedure code 99367) may be reimbursed once every six months when the medical home coordinating clinician attests that they are providing the medical home for the client. The coordinating clinician may be the client’s primary care provider or a specialist.

Additional medical team conferences may be considered with documentation of a change in the client’s medical home.

The medical team conference time must be documented in the client’s record.

### 2.5.1.2 Face-to-Face Services

#### 2.5.1.2.1 General Requirements for Face-to-Face Clinician-Directed Care Coordination Services

Providers must use the most appropriate face-to-face E/M procedure codes to bill for care coordination services.

- When counseling or care coordination requires more than 50 percent of the client or family encounter (face-to-face time in the office or other outpatient setting, or floor/unit time in the hospital), then time may be considered the key or controlling factor to qualifying for a particular level of E/M service.
- Counseling is a discussion with the client or family concerning diagnostic studies or results, prognosis, risks and benefits, management options, importance of adhering to the treatment regimen, and client and family education.

Modifiers must be used as appropriate for billing.

Any face-to-face inpatient or outpatient E/M procedure code that is a benefit of Texas Medicaid may be billed on the same day as the following non-face-to-face clinician-directed care coordination procedure codes when the procedure requires significant, separately-identifiable E/M services by the same physician on the same day.

<table>
<thead>
<tr>
<th>Procedure Codes</th>
</tr>
</thead>
<tbody>
<tr>
<td>99339</td>
</tr>
<tr>
<td>99380</td>
</tr>
</tbody>
</table>

### 2.5.2 Prior Authorization and Documentation Requirements

Non-face-to-face clinician-directed care coordination services provided by the medical home require prior authorization. Providers must submit a request for prior authorization within seven business days of the date of service. Prior authorization is limited to a maximum of six months. Prior authorization is required to recertify the client for additional six-month periods and requires submission of a new request with documentation supporting medical necessity for ongoing services.

Prior authorization for initial non-face-to-face clinician-directed care coordination requires documentation of at least one covered face-to-face inpatient or outpatient E/M visit by the medical home clinician directing the care coordination during the six months preceding the provision of the first non-face-to-face care coordination service.

Prior authorization for subsequent non-face-to-face clinician-directed care coordination services requires at least one covered face-to-face inpatient or outpatient E/M visit by the medical home clinician directing the care coordination during the previous 12 months or more frequently as indicated by the client’s condition.
Prior authorization of CCP services may be requested in writing by completing a CCP Prior Authorization Request Form, attaching the necessary supportive documentation as detailed below, and mailing or faxing it to the TMHP-CCP department:

Texas Medicaid & Healthcare Partnership
Comprehensive Care Program
PO Box 200735
Austin, TX 78720-0735
Fax: 1-512-514-4212

For prior authorization to be considered, clients must require complex and multidisciplinary care modalities involving regular clinician development or revision of care plans, review of subsequent reports of client status, and review of related laboratory and other studies:

- **Medically complex.** The health care needed by a Medicaid client achieves the designation of medically complex when the approved POC necessitates a clinical professional practicing within the scope of his or her license and in the context of a medical home to coordinate ongoing treatment to ensure its safe and effective delivery. The diagnosis must be covered under Texas Medicaid and be characterized by one of the following:
  - Significant and interrelated disease processes that involve more than one organ system (including behavioral health diagnoses) and require the services of two or more licensed clinical professionals, specialists, or subspecialists.
  - Significant physical or functional limitations that require the services of two or more therapeutic or ancillary disciplines, including, but not limited to, nursing, nutrition, OT, PT, ST, orthotics, and prosthetics.
  - Significant physical, developmental, or behavioral impairment that requires the integration of two or more medical or community-based providers, including, but not limited to, educational, social, and developmental professionals, that impact the care of the client.

- **Multidisciplinary Care.** Care is multidisciplinary when the medically necessary covered services of an approved POC include the need to coordinate the assessment, treatment, or services of a Medicaid-enrolled clinical provider with two or more additional medical, educational, social, developmental, or other professionals impacting the health care of the client.

Prior authorization is effective for care coordination services provided over a period of six months. Medical home clinicians must submit a revised care plan for subsequent periods of prior authorization. Documentation of the following components must be submitted with the prior authorization form to obtain an initial authorization or renewal:

- A current medical summary, encompassing all disciplines and all aspects of the client’s care, and containing key information about the client’s health, including conditions, complexity, medications, allergies, past surgical procedures, and so on.
- A current list of the main concerns, issues, and problems as well as key strengths and assets and the related current clinical information including a list of all diagnosis codes.
- Planned action steps and interventions to address the concerns and to sustain and build strengths, with the expected outcomes.
- Disciplines involved with the client’s care and how the multiple disciplines will work or are working together to meet the client’s need. Providers must explain how the multidisciplinary approach will or do benefit the client’s needs.
- Short-term and long-term goals with timeframes.
The supporting documentation can be any of the following:

- A formal written care plan
- Progress note detailing the care coordination planning
- A letter of medical necessity detailing the care plan oversight and care coordination

Clinician-directed care coordination services must be documented in the client’s medical record. Documentation must support the services being billed and must include a record of the medical home clinician’s time spent performing specific care coordination activities, including start and stop times. The documentation must also include a formal care plan and an emergency services plan. The supporting documentation maintained in the client’s medical records must be dated and include the following components and requirements:

- Problem list
- Interventions
- Short-term and long-term goals
- Responsible parties

Client medical records are subject to retrospective review.

Documentation for care coordination provided during post-surgical care must clearly indicate the care coordination is unrelated to the surgery.

2.5.2.1 Documentation Requirements for the Medical Home Clinician for a Telephone Consult with a Specialist

The clinician providing the medical home must maintain the following documentation in the client’s medical record:

- Start and stop times showing that the consultation was at least 15 minutes
- The reason for the call
- The specialist’s or subspecialist’s medical opinion
- The recommended treatment or laboratory services
- The name of the specialist or subspecialist

2.5.2.2 Documentation Requirements for the Specialist or Subspecialist for a Telephone Consult with the Medical Home Clinician

Specialists or subspecialists must complete and retain the Specialist or Subspecialist Telephone Consultation Form for Non-Face-to-Face Clinician Directed Care Coordination Services-CCP. These records are subject to retrospective review. The supporting documentation must include, but is not limited to the following:

- The client’s name, date of birth, and Medicaid identification number
- Start and stop times indicating the consultation lasted at least 15 minutes
- The reason for the call
- The specialist’s or subspecialist’s medical opinion
- The recommended treatment or laboratory services
- The name and telephone number of the clinician providing the medical home
- Provider information for the specialist’s or subspecialist’s and the clinician providing the medical home
2.5.3  Claims Information
Claims for clinician-care coordination services must be submitted to TMHP in an approved electronic claims format or on a CMS-1500 paper claim form. Providers may purchase CMS-1500 paper claim forms from the vendor of their choice. TMHP does not supply the forms.

*Refer to:* “Section 3: TMHP Electronic Data Interchange (EDI)” (Vol. 1, General Information) for information on electronic claims submissions.

"Section 6: Claims Filing” (Vol. 1, General Information) for general information about claims filing.

Subsection 6.5, “CMS-1500 Paper Claim Filing Instructions” in “Section 6: Claims Filing” (Vol. 1, General Information) for instructions on completing paper claims.

2.5.4  Reimbursement
Clinician-directed care coordination services are reimbursed in accordance with 1 TAC §355.8441.

2.6  Comprehensive Outpatient Rehabilitation Facilities (CORFs) and Outpatient Rehabilitation Facilities (ORFs)

*Refer to:* The Physical Therapy, Occupational Therapy, and Speech Therapy Services Handbook (Vol. 2, Provider Handbooks) for more information about CCP therapy services.

2.7  Durable Medical Equipment (DME) Supplier (CCP)

2.7.1  Enrollment
To be eligible to participate in CCP, providers of DME (including customized or non-basic medical equipment) and expendable medical supplies must be enrolled in Medicare.

Home health agencies that provide DME and supplies should refer to subsection 2.1, “Enrollment” in the Durable Medical Equipment, Medical Supplies, and Nutritional Products Handbook (Vol. 2, Provider Handbooks).

2.7.1.1  Pharmacies (CCP)
Pharmacy providers are eligible to participate in CCP. To be enrolled in CCP, the pharmacy must also be enrolled in VDP.

This enrollment allows pharmacy providers to bill for those medications and supplies payable by Medicaid for clients who are birth through 20 years of age but not covered by VDP (e.g., some over-the-counter drugs, some nutritional products, diapers, and disposable or expendable medical supplies). Pharmacy providers must continue to bill HHSC for drugs covered under VDP.

To locate a pharmacy CCP provider, use the Online Provider Lookup (OPL) at http://opl.tmhp.com/ProviderManager/AdvSearch.aspx.

*Refer to:* Subsection 2.1.2, "Enrollment” in this handbook for more information about CCP enrollment procedures.

Section 1.1, “About the Vendor Drug Program” in the Outpatient Drug Services Handbook (Vol. 2, Provider Handbooks) for information about this program.

Section 2, “Texas Medicaid (Title XIX) Home Health Services” in the Durable Medical Equipment, Medical Supplies, and Nutritional Products Handbook (Vol. 2, Provider Handbooks) for details about coverage through Texas Medicaid (Title XIX) Home Health Services.
2.7.2 Services, Benefits, and Limitations

DME is defined as medical equipment that is manufactured to withstand repeated use, ordered by a physician for use in the home, and required to correct or ameliorate the client’s disability, condition, or illness.

Because there is no single authority (such as a federal agency) that confers the official status of “DME” on any device or product, HHSC retains the right to make such determinations with regard to DME covered by Texas Medicaid. DME covered by Texas Medicaid must either have a well-established history of efficacy or, in the case of novel or unique equipment, valid peer-reviewed evidence that the equipment serves a medical purpose, can withstand repeated use, and is appropriate and safe for use in the home.

Requested DME may be a benefit of Texas Medicaid when it meets the Medicaid definition of DME.

The majority of DME and expendable medical supplies are covered through Texas Medicaid (Title XIX) Home Health Services.

If a service cannot be provided through Texas Medicaid (Title XIX) Home Health Services, the service may be covered through CCP if it is determined to be medically necessary for the client and if FFP is available.

If a DME provider is unable to deliver a piece of equipment, the provider must allow the client the option of obtaining the DME or expendable medical supplies from another provider.

Periodic rental payments are made only for the lesser of the following:

- The period of time the equipment is medically necessary
- The total monthly rental payments equal the reasonable purchase cost for the DME

DME will be purchased when a purchase is determined to be medically necessary and more cost effective than leasing the device with supplies. Only new, unused equipment will be purchased. When a provider is replacing a piece of rental DME with purchased DME, the provider must supply a new piece of DME to the client.

Purchase is justified when the estimated duration of need multiplied by the rental payments would exceed the reasonable purchase cost of the equipment or it is otherwise more practical to purchase the equipment.

DME repair will be considered based on the age of the item and cost to repair it. A request for repair of DME must include an itemized estimated cost list from the vendor or DME provider who will make the repairs.

Rental equipment may be provided to replace purchased medical equipment for the period of time it will take to make necessary repairs to purchased medical equipment.

All adjustments and modifications that are made within the first six months after delivery are considered part of the purchase price. However, DME that has been delivered to the client’s home and then found to be inappropriate for the client’s condition will not be eligible for an upgrade within the first six months following purchase unless there had been a significant change in the client’s condition, as documented by the physician familiar with the client.

Rental reimbursement to the same provider cannot exceed the purchase price, except as addressed in specific policies.

All DME purchased for a client becomes the Medicaid client’s property upon receipt of the item. Delivered equipment will become the Medicaid client’s property in the following instances even though it will not be prior authorized or reimbursed:

- Equipment delivered to the client before the physician signature date on the CCP Prior Authorization Request Form or prescription.
• Equipment delivered more than three business days before obtaining prior authorization from TMHP that meets the criteria for purchase.

As long as the client is eligible for CCP services on the date the custom equipment is ordered from the manufacturer, the provider must use the order date as the date of service since custom equipment is client specific and cannot be used for another client.

To establish medical necessity of the equipment for the client, the provider must have on file in the client’s records current documentation that is signed by a physician (e.g., a signed and dated prescription) showing the following:

• A diagnosis relative to each item requested.
• The specific type of supply needed.
• The length of time needed.

2.7.2.1 Purchase Versus Equipment Rental

When providing equipment not prior authorized under Texas Medicaid (Title XIX) Home Health Services for CCP clients with long-term or chronic conditions, it is more cost-effective, in many cases, to purchase the equipment rather than rent it. The client’s condition and length of time the equipment will be used must be carefully assessed before prior authorization for rental or purchase is requested. CCP nurses determine whether the equipment will be rented, purchased, repaired, or modified based on the client’s needs, the duration of use, and the age of the equipment.

CCP does not pay for the purchase of certain types of equipment; consequently, long-term rental may be considered. Most other equipment is rented for only four months initially. During this time, the provider must assess whether the equipment should be purchased before the rental lapses. Rentals and purchases must be prior authorized.

After prior authorization is obtained for purchase, new equipment must be provided and the rental discontinued. CCP does not purchase used equipment.

Providers of customized or nonbasic medical equipment also must be enrolled as Medicare DME providers.

2.7.3 Prior Authorization and Documentation Requirements

Providers can request prior authorization for most DME through the TMHP website. Providers that make written requests for prior authorization must complete the CCP Prior Authorization Request Form on the TMHP website at www.tmhp.com, and they must attach the documentation necessary to support the request. The documentation must include a current prescription that has been signed and dated by a physician (M.D. or D.O.), and it must be mailed or faxed to TMHP with the prior authorization request. For specific policy information not contained in this manual related to the purchase of DME, providers can call TMHP-CCP Customer Service at 1-800-846-7470.

A completed CCP Prior Authorization Request Form prescribing the DME or medical supplies must be signed and dated by the prescribing physician familiar with the client before requesting prior authorization. The completed CCP Prior Authorization Request Form must be maintained by the requesting provider and the prescribing physician. The original signature copy must be kept in the physician’s medical record for the client.

To complete the prior authorization process by paper, the DME provider must fax or mail the completed CCP Prior Authorization Request Form to the CCP prior authorization unit and retain a copy of the signed and dated CCP form in the client’s medical record at the provider’s place of business.

To complete the prior authorization process electronically, the DME provider must complete the prior authorization requirements through any approved electronic methods and retain a copy of the signed and dated CCP Prior Authorization Request form in the client’s medical record at the provider’s place of business.
To avoid unnecessary denials, the physician must provide correct and complete information, including accurate documentation of the medical necessity for the equipment and services requested. The physician must maintain documentation of medical necessity in the client’s medical record. The requesting provider may be asked for additional information to clarify or complete a request.

Retrospective review may be performed to ensure documentation supports the medical necessity of the requested equipment or supplies.

A determination as to whether the equipment will be rented, purchased, repaired, or modified will be made by HHSC or its designee based on the client’s needs, duration of use, and age of the equipment.

Equipment that has been purchased may be considered for replacement when loss or irreparable damage has occurred outside the warranty terms, conditions, and limitations. A copy of the police or fire report, when appropriate, and the measures to be taken to prevent reoccurrence must be submitted with the prior authorization request.

A request for prior authorization must include documentation from the provider to support the medical necessity of the service, equipment, or expendable medical supply. Physician prescriptions must be specific to the item requested. For example, if the provider is requesting a customized wheelchair, the prescription must request a customized wheelchair, not just a wheelchair. Providers must submit a CCP Prior Authorization Request Form and documentation to support medical necessity to the CCP department before providing services. Providers must obtain prior authorization within three business days of the requested date of service.

Refer to: Section 2.2.30, ”Procedure Codes That Do Not Require Prior Authorization” in the Durable Medical Equipment, Medical Supplies, and Nutritional Products Handbook (Vol. 2, Provider Handbooks) for details about specific procedure codes that do not require prior authorization through Texas Medicaid (Title XIX) Home Health Services.

2.7.3.1 Equipment Accessories
CCP may consider prior authorization of equipment accessories, such as ventilator and oxygen trays and positioning inserts, when supporting documentation takes into account all the client’s needs, capabilities, and physical or mental status.

2.7.3.2 Equipment Modifications
A modification is the replacement of a component due to changes in the client’s condition, not the replacement of a component that is no longer functioning.

DME that has been delivered to the client’s home and then found to be inappropriate for the client’s condition will not be eligible for an upgrade within the first six months following purchase. All modifications that are made within the first six months after delivery are considered part of the purchase price.

However, CCP may consider prior authorization of modifications to custom equipment if a change occurs in the client’s needs, capabilities, or physical or mental status that cannot be anticipated. Documentation must include:

- All projected changes in the client’s needs.
- The age of the current equipment, and the cost of purchasing new equipment versus modifying current equipment.

2.7.3.3 Equipment Adjustments
Adjustments do not require supplies.

Labor for adjustments within the first six months after delivery are not prior authorized because these are considered part of the purchase price.
Up to one hour of labor for adjustments may be considered for reimbursement with prior authorization through CCP as needed after the first six months. Providers must use procedure code K0739 for adjustments.

### 2.7.3.4 Repair to Client-Owned Equipment

Repairs to client-owned equipment may be prior authorized as needed with documentation of medical necessity. Technician fees are considered part of the cost of the repair.

HHSC or its designee reserves the right to request additional documentation about the need for repairs when there is evidence of abuse or neglect to equipment by the client, client’s family, or caregiver. When there is documented proof of abuse or neglect, requests for repairs will not be prior authorized.

Providers are responsible for maintaining documentation in the client’s medical record that specifies the repairs and supporting medical necessity.

Documentation must include all of the following:

- The date of purchase
- The serial number of the current equipment (as applicable)
- The cause of the damage or need for repairs
- What steps the client or caregiver will take to prevent further damage if repairs are due to an accident
- When requested, the cost of purchasing new equipment as opposed to repairing current equipment

Temporary replacement of client-owned respiratory equipment during the repair may be considered for prior authorization for one month using procedure code K0462.

Labor for repair of client-owned respiratory equipment may be considered for prior authorization using procedure code K0739 up to a maximum of two hours per day (maximum quantity of eight units).

Routine maintenance of rental equipment is the provider’s responsibility.

### 2.7.3.5 DME Certification and Receipt Form

The DME Certification and Receipt Form is required and must be completed before reimbursement can be made for any DME delivered to a client. The certification form must include the name of the item, the date the client received the DME, and the signatures of the provider and the client or primary caregiver.

The DME provider must maintain the signed and dated form in the client’s medical record.

DME claims and appeals that meet or exceed a billed amount of $2,500 for the same date of service will suspend for verification of client receipt of the DME item(s). The DME Certification and Receipt Form must be faxed to 1-512-506-6615. If the claim is submitted without the form or if receipt of the DME item(s) cannot be verified, the DME item(s) on the claim will be denied. TMHP may contact the client that received the product for verification of services rendered.

Refer to: [DME Certification and Receipt Form](www.tmhp.com) on the TMHP website.

### 2.7.3.6 Documentation of Supply Delivery

Providers must retain individual delivery slips or corresponding invoices for each date of service to document the date of delivery for all supplies provided to a client. Providers must disclose this documentation to HHSC or its designee upon request. These records and claims must be retained for a minimum of five years from the date of service (DOS) or until all audit questions, appeals, hearings, investigations, or court cases are resolved. The DOS is the date on which supplies are delivered to the client or shipped by a carrier to the client as evidenced by the dated tracking document attached to the corresponding invoice for that date.
Documentation of delivery must include one of the following:

- Delivery slip or invoice signed and dated by the client or caregiver.
- A dated carrier tracking document that includes the shipping date and delivery date must be printed from the carrier’s website as confirmation that the supplies were shipped and delivered. The dated carrier tracking document must be attached to the delivery slip or corresponding invoice.

The dated delivery slip or invoice must include the client’s full name and address to where supplies were delivered, and an itemized list of goods that includes the descriptions and numerical quantities of the supplies delivered to the client, and the corresponding tracking number from the carrier. This document could also include prices, shipping weights, shipping charges, and any other description.

All claims submitted for DME supplies must include the same quantities or units that are documented on the delivery slip or corresponding invoice and on the CCP Prior Authorization Request form. They must reflect the number of units by which each product is measured. For example, diapers are measured as individual units. If one package of 300 diapers is delivered, the delivery slip or invoice and the claim must reflect that 300 diapers were delivered and not that one package was delivered. Diaper wipes are measured as boxes or packages. If one box of 200 wipes is delivered, the delivery slip or invoice and the claim must reflect that one box was delivered and not that 200 individual wipes were delivered. There must be one dated delivery slip or invoice for each claim submitted for each patient. All claims submitted for DME supplies must reflect either one business day before or one business day after the date of service as documented on the delivery slip or corresponding invoice and the same timeframe covered by the CCP Prior Authorization Request form. The DME Certification and Receipt Form is still required for all equipment delivered.

### 2.7.3.7 Specific CCP Policies

Most DME and expendable medical supplies are available under Texas Medicaid (Title XIX) Home Health Services. If the service is not available under Texas Medicaid (Title XIX) Home Health Services, CCP may cover the requested service, if the client is CCP-eligible and the service is medically necessary, requested by a physician, and for which FFP is available.

**Refer to:** [DME Certification and Receipt Form](https://www.tmhp.com) on the TMHP website at [www.tmhp.com](http://www.tmhp.com).

The *Durable Medical Equipment, Medical Supplies, and Nutritional Products Handbook (Vol. 2, Provider Handbooks)* for DME services.

### 2.8 Early Childhood Intervention (ECI) Services

The Texas Health and Human Services (HHS) ECI program is available statewide to all children who have been determined to be eligible for ECI services by ECI contractors. To be eligible for ECI services, children must be 35 months of age and younger (i.e., before their third birthday) and have disabilities or developmental delays as defined by ECI criteria. Texas Medicaid covers the ECI claims for children who are Medicaid clients.

All health-care professionals are required by federal and state regulations to refer children who are 35 months of age and younger (i.e., before their third birthday) to the Texas HHS ECI program as soon as possible, but no longer than 7 days after identifying a disability or suspected delay in development. Referrals can be based on professional judgment or a family’s concern. A medical diagnosis or a confirmed developmental delay is not required for referrals.

To refer families for services, providers can call their local ECI program, or they can call the HHS Inquiry Line at 1-877-787-8999. For additional ECI information, providers can visit the [Early Childhood Intervention Services page](https://hhs.texas.gov) of the HHS website at [https://hhs.texas.gov](http://https://hhs.texas.gov). Persons who are deaf or hard of hearing may use the relay option of their choice or dial 7-1-1 to connect with Relay Texas.
2.8.1 Enrollment
The Texas HHS ECI program contracts with local non-profit entities to take referrals, determine clients’ eligibility, and provide services to ECI-eligible children and their families. The non-profit entities must contract with the Texas HHS ECI program and must comply with all of the applicable federal and state laws and regulations that govern the Texas HHS ECI program.

ECI contractors are eligible to enroll as Texas Medicaid ECI providers to render services to eligible Medicaid clients. After providers meet the criteria of the Texas HHS ECI program, they must complete a Medicaid application.

To participate in Texas Medicaid, an ECI contractor must submit a copy of the current contract award from the Texas HHS ECI program.

*Refer to:* Subsection 1.1, “Provider Enrollment” in “Section 1: Provider Enrollment and Responsibilities” (*Vol. 1, General Information*) for more information about the procedures for enrolling as a Medicaid provider.

2.8.2 Services, Benefits, and Limitations
Prior authorization is not required for evaluations, re-evaluations, seating assessments, therapy services, SST, and TCM. The IFSP Services Pages identify the amount, duration, and scope for the provision of SRS treatment services and serves as the prior authorization for ECI services. The IFSP is retained in the client’s record and is subject to retrospective review.

ECI services include targeted case management (TCM) and specialized rehabilitative services (SRS), which includes occupational therapy (OT), physical therapy (PT), speech therapy (ST), and specialized skills training (SST).

ECI SRS services may be provided in the following places of service (POS): office/facility (POS 1), home (POS 2), outpatient (POS 5 applicable only to ECI services rendered in a Prescribed Pediatric Extended Care Center [PPECC]), and other locations (POS 9). In addition to these places of service, TCM may be provided in inpatient hospital (POS 3) and outpatient hospital (POS 5).

ECI services of OT, PT, ST, SST and TCM are provided to Medicaid-eligible clients who are birth through 35 months of age and have a documented developmental delay or a medically diagnosed condition as established by HHSC (40 TAC, Part 2, Chapter 108), or an auditory or visual impairment as defined by the Texas Education Agency (19 TAC §89.1040).

To the maximum extent appropriate, ECI services are delivered in the client’s natural environment, as defined in 40 TAC, Part 2, Chapter 108, and are family-centered.

The interdisciplinary team must document ECI eligibility decisions in accordance with 40 TAC, Part 2, Chapter 108. The eligibility statement must be in the child’s record and updated when eligibility changes or is re-determined.

All documentation of ECI services, including the plan of care specified in the Individualized Family Service Plan (IFSP) must be retained in the client’s record and available upon request. The IFSP is a written plan of care for providing early childhood intervention services and other medical, health, and social services to an eligible child and the child’s family when necessary to enhance the child’s development.

ECI service providers are employees and subcontractors of non-profit entities that have contracts with the State of Texas for the provision of Individuals with Disabilities Education Act (IDEA) Part C Early Childhood Intervention services.
Medically necessary services may be provided by other Medicaid-enrolled providers in addition to the services provided by the ECI contractor. For example, the family may choose to receive speech therapy from the ECI contractor and physical therapy from a home health provider. Or, outpatient clinic personnel may have expertise that will enhance the services of the ECI provider resulting in ECI providers and other Medicaid-enrolled providers providing services within the same discipline.

Only the services provided to ECI enrolled children by ECI contracted entities must comply with the Medicaid medical guidelines for ECI services.

Services provided by other Medicaid-enrolled providers, including other providers of physical, occupational, and speech therapy, must comply with Medicaid medical guidelines that apply to those provider types (e.g., outpatient rehabilitation facility, home health agency).

### 2.8.2.1 Physical, Occupational, and Speech Therapies and Specialized Skills Training (PT, OT, ST, and SST)

ECI services use techniques by which the ECI service provider engages the family or caregiver in activities to meet the developmental needs of the child.

ECI services are performed in accordance with 40 TAC, Part 2, Chapter 108.

To the maximum extent possible, ECI services are provided in the client’s natural environment, as defined in 34 CFR Part 303, unless the IFSP team determines the identified outcomes cannot be achieved in a natural environment. Natural environments are defined as settings that are natural or typical for the same-aged infant or toddler without a disability, and may include the home and community settings such as daycare, playgrounds, stores, and restaurants.

Justification for providing services in other settings (e.g., office, clinic, Prescribed Pediatric Extended Care Center (PPECC)) must be documented in the client’s record.

PT, OT, ST, and SST are benefits for clients with an acute or a chronic condition when documented on the IFSP. Documentation on the IFSP is evidence that services are developed and recommended by the child’s interdisciplinary team, including the parents and a licensed practitioner of the healing arts (as defined in 40 TAC, Part 2, Chapter 108).

PT, OT, ST, and SST must be performed and delivered as identified in the IFSP.

Missed visits may be rescheduled within the authorization period as long as the total number of visits or units provided does not exceed the amount authorized in the client’s IFSP. The ECI contractor must document the reason for visits outside of the weekly or monthly frequency in the client’s record.

A single identified need and treatment goal (outcome on the IFSP) may be addressed by more than one discipline.

More than one discipline can evaluate a child at the same time to facilitate compliance with the federal requirement for multidisciplinary evaluation (34 CFR, Part 303).

A client may receive a combination of PT, OT, ST, or SST with any other IFSP service when the IFSP indicates necessity for co-visits or co-treatment (i.e., two or more services to be provided at the same time).

PT, OT, ST, and SST may be delivered to a client individually or in a group setting according to 40 TAC, Part 2, Chapter 108 and when documented in the IFSP.

Documentation of each PT, OT, ST, and SST contact must be entered into the child’s record in accordance with 40 TAC, Part 2, Chapter 108.

### 2.8.2.2 Physical, Occupational, and Speech Therapy (PT, OT, and ST)

Physical and occupational therapy treatment services require orders from a referring provider once a year.
Speech therapy treatment services do not require an order from a referring provider.

Therapy goals for acute or chronic conditions include, but are not limited to the following:

- Improving function
- Maintaining function
- Slowing the deterioration of function

### 2.8.2.2.1 Physical Therapy (PT)

PT includes services that address the promotion of sensory and motor function through enhancement of musculoskeletal status, neurobehavioral organization, perceptual and motor development, cardiopulmonary status, and effective environmental adaptation.

All services must be performed in accordance with 42 CFR 440.110.

A PT evaluation, re-evaluation, or seating assessment may be performed without an order from a referring provider as allowed by 22 TAC Part 16, Chapter 322, §322.1(a)(2)(A).

PT services must be provided by one of the following:

- A licensed physical therapist who meets the requirements of 42 CFR 440.110(a)
- A licensed physical therapy assistant (PTA) when the assistant is acting under the direction of a licensed physical therapist in accordance with 42 CFR 440.110 and all other applicable state and federal law

### 2.8.2.2.2 Occupational Therapy (OT)

OT includes services that address the functional needs of a child related to adaptive development, adaptive behavior and play, and sensory, motor, and postural development. These services are designed to improve the client’s functional ability to perform tasks in the home and community settings.

All services must be performed in accordance with 42 CFR 440.110.

An OT evaluation, re-evaluation, or seating assessment may be performed without an order from a referring provider as allowed by §454.213 of the Texas Occupations Code.

OT services must be provided by one of the following:

- A licensed occupational therapist who meets the requirements of 42 CFR 440.110(b)
- A licensed or licensed and certified occupational therapist assistant (OTA) when the assistant is acting under the direction of a licensed occupational therapist in accordance with 42 CFR 440.110 and all other applicable state and federal law

### 2.8.2.2.3 Speech Therapy (ST)

Speech and language therapy includes services designed to promote rehabilitation and remediation of delays or disabilities in language-related symbolic behaviors, communication, language, speech, emergent literacy, or feeding and swallowing behavior.

All services must be delivered in accordance with 42 CFR 440.110 and §401.001(6) of the Texas Occupations Code.

A ST evaluation, re-evaluation, and treatment service may be performed without a physician order as allowed by Chapter 401 of the Texas Occupations Code.

ST services must be provided by one of the following:

- A licensed speech-language pathologist (SLP) who meets the requirements of 42 CFR 440.110(c) and all other applicable state and federal law
A licensed assistant in SLP when the assistant is acting under the direction of a licensed SLP in accordance with 42 CFR 440.110

A licensed intern when the intern is acting under the direction of a licensed SLP in accordance with 42 CFR 440.110 and all other applicable state and federal law

2.8.2.3 Physical Therapy, Occupational Therapy, and Speech Therapy Procedure Codes

Clients who are eligible for ongoing PT, OT, and ST through the ECI program may request additional therapy under the Early & Periodic Screening, Diagnosis, & Treatment (EPSDT) benefit of Medicaid (also known as Texas Health Steps) when medically necessary.

Refero: Section 5, “Children’s Therapy Services Clients birth through 20 years of age” in the Physical Therapy, Occupational Therapy, and Speech Therapy Services Handbook (Vol. 2, Provider Handbooks) for information about physical, occupational, and speech therapy procedure codes outside of the ECI benefit that are not defined in this section.

2.8.2.3.1 Evaluation and Re-evaluation Procedure Codes

The following encounter-based evaluation and re-evaluation procedure codes for PT, OT, and ST are benefits of Texas Medicaid:

<table>
<thead>
<tr>
<th>Procedure Codes</th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td>97165, 97166, or 97167</td>
<td>OT Evaluation</td>
</tr>
<tr>
<td>97168</td>
<td>OT Re-evaluation</td>
</tr>
<tr>
<td>97161, 97162, or 97163</td>
<td>PT Evaluation</td>
</tr>
<tr>
<td>97164</td>
<td>PT Re-Evaluation</td>
</tr>
<tr>
<td>92521, 92522, 92523, or 92524</td>
<td>ST Evaluation</td>
</tr>
<tr>
<td>S9152</td>
<td>ST Re-Evaluation</td>
</tr>
<tr>
<td>92610</td>
<td>ST Evaluation swallowing function</td>
</tr>
</tbody>
</table>

2.8.2.3.2 Time-Based Procedure Codes

The following time-based PT and OT treatment procedure codes may be a benefit of Texas Medicaid and must be billed in 15-minute increments (units).

<table>
<thead>
<tr>
<th>Procedure Codes</th>
</tr>
</thead>
<tbody>
<tr>
<td>97032 97033 97034 97035 97036 97110 97112 97113 97116 97124</td>
</tr>
<tr>
<td>97140 97530 97535 97542 97750 97760 97761 97763</td>
</tr>
</tbody>
</table>

2.8.2.3.3 Untimed PT and OT Procedure Codes

The following untimed PT and OT treatment procedure codes representing supervised modalities are limited to one encounter each, per date of service per discipline, must be delivered on the same day as one or more time-based codes listed above, and are subject to the CMS NCCI relationships.

<table>
<thead>
<tr>
<th>Procedure Codes</th>
</tr>
</thead>
<tbody>
<tr>
<td>97012 97014 97016 97018 97022 97024 97026 97028</td>
</tr>
</tbody>
</table>

The following PT and OT group therapy code may be reimbursed as an untimed procedure code, payable per encounter, and reimbursed once per date of service per discipline.

<table>
<thead>
<tr>
<th>Procedure Codes</th>
</tr>
</thead>
<tbody>
<tr>
<td>97150</td>
</tr>
</tbody>
</table>
2.8.2.3.4  Encounter-Based Speech Therapy Procedure Codes

The following speech therapy individual treatment codes must be billed per encounter and are limited to once per day per provider. Only one ST treatment procedure code 92507 or 92526 may be reimbursed per date of service.

<table>
<thead>
<tr>
<th>Procedure Codes</th>
</tr>
</thead>
<tbody>
<tr>
<td>92507</td>
</tr>
<tr>
<td>92526</td>
</tr>
</tbody>
</table>

The following ST group treatment code may be reimbursed as an untimed procedure code, payable per encounter, and reimbursed once per date of service.

<table>
<thead>
<tr>
<th>Procedure Codes</th>
</tr>
</thead>
<tbody>
<tr>
<td>92508</td>
</tr>
</tbody>
</table>

2.8.2.3.5  Modifier Requirements for PT, OT, or ST Services

The following modifiers must be submitted for PT, OT, and ST treatment services:

<table>
<thead>
<tr>
<th>Modifier</th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td>GO</td>
<td>Services delivered under an outpatient occupational therapy plan of care</td>
</tr>
<tr>
<td>GP</td>
<td>Services delivered under an outpatient physical therapy plan of care</td>
</tr>
<tr>
<td>GN</td>
<td>Services delivered under an outpatient speech therapy plan of care</td>
</tr>
<tr>
<td>UB</td>
<td>Services delivered by a therapy assistant under supervision of a licensed therapist</td>
</tr>
<tr>
<td>U5</td>
<td>Services delivered by a licensed therapist or a physician</td>
</tr>
</tbody>
</table>

Modifier UB or U5 is required on all claims for therapy treatment procedure codes to designate whether treatment was provided by a licensed therapist or a licensed assistant.

Modifier U3 is not used by an ECI contractor for co-visits or co-treatment services.

<table>
<thead>
<tr>
<th>Modifier</th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td>U3</td>
<td>Not used by an ECI contractor</td>
</tr>
</tbody>
</table>

2.8.2.3.6  Seating Assessments

Seating assessments are reimbursed in 15-minute increments (units) and must be billed with the following procedure code:

<table>
<thead>
<tr>
<th>Procedure Codes</th>
</tr>
</thead>
<tbody>
<tr>
<td>97542</td>
</tr>
</tbody>
</table>

The PT completing the assessment must submit procedure code 97542 with modifiers GP and UC in order to bill for the seating assessment.

<table>
<thead>
<tr>
<th>Modifier</th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td>GP</td>
<td>Services delivered under an outpatient physical therapy plan of care</td>
</tr>
<tr>
<td>UC</td>
<td>Assessment performed by an OT or PT</td>
</tr>
</tbody>
</table>
The OT completing the assessment must submit procedure code 97542 with modifiers GO and UC in order to bill for the seating assessment:

<table>
<thead>
<tr>
<th>Modifier</th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td>GO</td>
<td>Services delivered under an outpatient occupational therapy plan of care</td>
</tr>
<tr>
<td>UC</td>
<td>Assessment performed by an OT or PT</td>
</tr>
</tbody>
</table>

2.8.2.3.7 Specialized Skills Training (SST) Services

SST services are rehabilitative services to promote age-appropriate development by providing skills training to correct deficits and teach compensatory skills for deficits that directly result from medical, developmental, or other health-related conditions.

Services must include all the following:

- Be designed to create learning environments and activities that promote the client’s acquisition of skills in one or more of the following developmental areas: physical or motor, communication, adaptive, cognitive, and social or emotional.
- Skills training and anticipatory guidance for family members, or other significant caregivers, to ensure effective treatment and to enhance the client’s development.

SST services do not require an order from a referring provider. The ECI contractor ensures that SST services are provided by a certified early intervention specialist. SST services must be provided by an early intervention specialist who meets the criteria established in 40 TAC Part 2, Chapter 108.

SST services must be submitted with the following procedure codes and modifiers, and they must be billed in 15-minute increments:

<table>
<thead>
<tr>
<th>Procedure Code</th>
<th>Description</th>
<th>Modifier</th>
</tr>
</thead>
<tbody>
<tr>
<td>T1027</td>
<td>Individual setting</td>
<td>U1</td>
</tr>
<tr>
<td>T1027</td>
<td>Group setting</td>
<td></td>
</tr>
</tbody>
</table>

2.8.2.3.8 Reimbursement Guidelines for PT, OT, ST, and SST

Claims may be submitted to Medicaid when the interaction is directly with the client and the client’s parent(s) as defined in 20 U.S.C. §1401, or the client and the routine caregiver(s) as defined in 40 TAC, Part 2, Chapter 108.

ECI services must be billed under the ECI contractor’s Texas Provider Identifier, National Provider Identifier, and benefit code of EC1 as the insured’s policy group when submitting claims.

Refer to: “Section 6: Claims Filing” (Vol. 1, General Information) for more information about benefit codes.

Physical therapy, occupational therapy, and speech-language pathology evaluations are performed for the purposes of initial determination of need for rehabilitative services and annually to verify the child’s ongoing need for rehabilitative services. To ensure there are no gaps in rehabilitative services, the annual evaluation should occur prior to the child’s annual IFSP meeting.

Physical, occupational, and speech therapy evaluation and re-evaluation services are benefits through the ECI Medicaid benefit and do not require an order from a referring provider.

Physical therapy, occupational therapy, and speech-language pathology re-evaluations may be performed periodically during the child’s annual enrollment in ECI services, and without a physician’s order, to determine if changes to the IFSP are necessary.
Evaluations, re-evaluations, and seating assessments are not required to be listed on the IFSP Service Pages. A physical or occupational therapist may provide a seating assessment that is required to order a wheeled mobility system. A seating assessment does not require an order from a referring provider.

**Refer to:** Subsection 2.2.17, “Mobility Aids” in the *Durable Medical Equipment, Medical Supplies, and Nutritional Products Handbook* (Vol. 2, Provider Handbooks) for information about mobility aids.

Reimbursement is available to two or more of the ECI contractor’s service providers when the client receives a combination of any Medicaid-covered service identified on the IFSP and the IFSP indicates necessity for co-visits or co-treatments (i.e., two or more services to be provided at the same time). For example, the child may receive both PT and ST at the same time. Another example, the child may receive counseling and SST at the same time.

Reimbursement is available to two or more of the ECI contractor’s service providers when they are conducting an evaluation at the same time.

When an evaluation and treatment service within the same discipline occur on the same day, only the evaluation will pay.

When a re-evaluation and treatment service within the same discipline occur the same day, only the treatment will pay.

PT, OT, and ST equipment and supplies used during therapy visits are not reimbursed separately.

Reimbursement under Medicaid benefit guidelines applies to only the services provided to ECI enrolled children by ECI contracted entities.

Reimbursement for services provided to ECI enrolled children by other Medicaid-enrolled providers (e.g., home health, CORF) is available under the Medicaid medical policies that apply to those provider types.

### 2.8.2.4 Targeted Case Management (TCM)

TCM services are provided to assist an eligible client and his or her family in gaining access to the rights and procedural safeguards under Part C of IDEA, and to needed medical, social, educational, developmental, and other appropriate services.

TCM services are performed in accordance with the ECI Medicaid benefit guidelines and 40 TAC, Part 2, Chapter 108.

TCM services do not require an order from a referring provider, but must be delivered by a qualified ECI contractor. The ECI contractor ensures that TCM services are provided by the assigned Service Coordinator who meets the criteria established in 40 TAC Part 2, Chapter 108.

TCM is provided in the natural environment (including office, home, daycare, and other community locations), outpatient, PPECC, and inpatient hospital setting.

The documentation for each TCM contact must be in accordance with 40 TAC, Part 2, Chapter 108. The place of service is the location of the service coordinator at the time of service delivery.

TCM services must be submitted with the following procedure codes and modifiers, and they must be billed in 15-minute increments:

<table>
<thead>
<tr>
<th>Procedure Code</th>
<th>Description</th>
<th>Modifier</th>
</tr>
</thead>
<tbody>
<tr>
<td>T1017</td>
<td>Face-to-face interaction</td>
<td>U1</td>
</tr>
<tr>
<td>T1017</td>
<td>Telephone interaction</td>
<td></td>
</tr>
</tbody>
</table>

TCM services may be delivered face-to-face or by telephone.
2.8.2.4.1 Guidelines for TCM Services

Claims may be submitted to Texas Medicaid when the interaction is directly with the client or the client’s parent(s) as defined in 20 United States Code (U.S.C.) §1401), or other routine caregiver(s) as defined in 40 TAC, Part 2, Chapter 108.

Contacts may be made with other individuals when directly related to identifying the eligible client’s needs, helping the eligible client access services, identifying needs and support to assist the eligible client in obtaining services, providing the service coordinator with useful feedback, and alerting the service coordinator to changes in the eligible client’s needs. These contacts must be documented in the client’s record, but are not submitted as claims to Medicaid if they took place outside of the presence of the client or the client’s parent or routine caregivers.

2.8.2.5 Guidelines for ECI Services Performed in a Prescribed Pediatric Extended Care Center (PPECC)

When ECI services are rendered in a PPECC, the place of service will be outpatient hospital (used for a PPECC). The PPECC’s NPI must appear on the claim, in addition to the ECI contractor’s NPI. The ECI contractor and PPECC must have a written agreement for the provision of ECI services at the PPECC. The written agreement must address responsibilities of both parties, and how the parties will coordinate related to the client’s IFSP or plan of care, which includes documentation of coordination with the PPECC. The written agreement must be maintained in the client’s record.

2.8.3 Claims Filing and Reimbursement

2.8.3.1 Claims Information

Claims for SST and TCM services that have been rendered by an ECI contractor must be submitted to TMHP in an approved electronic format or on the CMS-1500 paper claim form. Contractors may purchase CMS-1500 paper claim forms from the vendor of their choice; TMHP does not supply the forms. When completing a CMS-1500 paper claim form, all required information must be included on the claim, as TMHP does not key any information from claim attachments. Superbills or itemized statements are not accepted as claim supplements.

Refer to: “Section 3: TMHP Electronic Data Interchange (EDI)” (Vol. 1, General Information) for information on electronic claims submissions.

Subsection 6.1, “Claims Information” in “Section 6: Claims Filing” (Vol. 1, General Information) for general information about claims filing.

Subsection 6.5, “CMS-1500 Paper Claim Filing Instructions” in “Section 6: Claims Filing” (Vol. 1, General Information) to find the instructions for completing paper claims.

Subsection 2.2, “Fee-for-Service Reimbursement Methodology” in “Section 2: Texas Medicaid Fee-for-Service Reimbursement” (Vol. 1, General Information) for more information about reimbursement.

2.8.3.1.1 Billing Units Based on 15 Minutes

All claims for reimbursement are based on the actual amount of billable time associated with the service. For those services for which the unit of service is 15 minutes (1 unit = 15 minutes), partial units should be rounded to the nearest quarter hour.

The following table shows the time intervals for 1 through 8 units:

<table>
<thead>
<tr>
<th>Units</th>
<th>Number of Minutes</th>
</tr>
</thead>
<tbody>
<tr>
<td>0 units</td>
<td>0 minutes through 7 minutes</td>
</tr>
<tr>
<td>1 unit</td>
<td>8 minutes through 22 minutes</td>
</tr>
<tr>
<td>2 units</td>
<td>23 minutes through 37 minutes</td>
</tr>
</tbody>
</table>
2.8.3.1.2 Managed Care Clients

If the child is enrolled in a Medicaid managed care organization (MCO), claims for PT, OT, and ST are submitted to the MCO.

TCM services are carved-out of Medicaid managed care and must be billed to TMHP for payment consideration.

SST services are carved-out of Medicaid managed care and claims must be billed to TMHP for payment consideration.

2.8.3.2 Reimbursement

ECI therapy, SST, and TCM services are reimbursed according to a maximum allowable fee established by HHSC. See the applicable fee schedule on the TMHP website at www.tmhp.com.

- ECI therapy services are reimbursed in accordance with 1 TAC §355.8441.
- SST services are reimbursed in accordance with 1 TAC §355.8422.
- TCM services are reimbursed in accordance with 1 TAC §355.8421.

2.9 Health and Behavior Assessment and Intervention

2.9.1 Services, Benefits, and Limitations

Health and Behavior Assessment and Intervention (HBAI) services are a benefit of Texas Medicaid for clients who are 20 years of age and younger when the services are provided by a licensed practitioner of the healing arts (LPHA) who is co-located in the same office or building complex as the physician, PA, NP, or CNS who is treating the client.

In many cases, the treating physician, PA, NP, or CNS will be the client’s primary care provider; however, a specialist seeing a client regularly may function in a similar role to a primary care provider and may also make HBAI referrals to a co-located LPHA.

These services are designed to identify the psychological, behavioral, emotional, cognitive and social factors important to prevention, treatment or management of physical health symptoms.

HBAI services are a benefit when the client meets all of the following criteria:

- The client has an underlying physical illness or injury.
- There are indications that biopsychosocial factors may be significantly affecting the treatment or medical management of an illness or an injury.
- The client is alert, oriented, and, depending on the client’s age, has the capacity to understand and to respond meaningfully during the in-person evaluation.
- The client has a documented need for psychological evaluation or intervention to successfully manage his or her physical illness, and activities of daily living.
- The assessment is not duplicative of other provider assessments.

<table>
<thead>
<tr>
<th>Units</th>
<th>Number of Minutes</th>
</tr>
</thead>
<tbody>
<tr>
<td>3 units</td>
<td>38 minutes through 52 minutes</td>
</tr>
<tr>
<td>4 units</td>
<td>53 minutes through 67 minutes</td>
</tr>
<tr>
<td>5 units</td>
<td>68 minutes through 82 minutes</td>
</tr>
<tr>
<td>6 units</td>
<td>83 minutes through 97 minutes</td>
</tr>
<tr>
<td>7 units</td>
<td>98 minutes through 112 minutes</td>
</tr>
<tr>
<td>8 units</td>
<td>113 minutes through 127 minutes</td>
</tr>
</tbody>
</table>
HBAI services that include the client’s family are a benefit when the family member directly participates in the overall care of the client.

Family is defined as a responsible adult. This adult individual has agreed to accept the responsibility for providing food, shelter, clothing, education, nurturing, and supervision for the client. Responsible adults include, but are not limited to, biological parents, adoptive parents, foster parents, guardians, court-appointed managing conservators, and other family members by birth or marriage.

HBAI services may be reimbursed when billed with the following procedure codes:

<table>
<thead>
<tr>
<th>Procedure Codes</th>
<th>Limitations</th>
</tr>
</thead>
<tbody>
<tr>
<td>96156</td>
<td>Limited to a maximum of 2 per client, per rolling 180 days, any provider</td>
</tr>
<tr>
<td>96158, 96164, 96167</td>
<td>Limited to a maximum of 8 per client, per rolling 180 days, any provider</td>
</tr>
<tr>
<td>96159, 96165, 96168</td>
<td>Limited to a maximum of 14 per client, per rolling 180 days, any provider</td>
</tr>
</tbody>
</table>

These services may be rendered by physician, nurse practitioner (NP), clinical nurse specialist (CNS), physician assistant (PA), licensed professional counselor (LPC), licensed clinical social worker (LCSW), licensed marriage family therapist (LMFT), Comprehensive Care Program (CCP) LCSW, or psychologist provider in the office or outpatient setting.

LMFTs must bill with state defined modifier U8 to identify services billed.

For services that are rendered by physician, NP, CNS, or PA providers, claims must be submitted with the appropriate evaluation and management (E/M) procedure codes (99202, 99203, 99204, 99205, 99211, 99212, 99213, 99214, or 99215). The physician, NP, CNS, or PA may bill the HBAI procedure codes for an LPHA that is in the medical practice.

HBAI services are limited to a total of three units per day, by any provider, as follows:

- The initial 30 minutes of health behavior intervention (procedure codes 96158, 96164, and 96167) is limited to one unit per day.
- Each additional 15 minutes of health behavior intervention (procedure codes 96159, 96165, and 96168) is limited to two units per day.

Procedure codes 96167 and 96168 which include the client’s family, are a benefit when the family member directly participates in the overall care of the client.

An in-person evaluation is defined as a patient evaluation conducted by a provider who is at the same physical location as the client. These services are considered acute per rolling 180 days from the initiation of services and are limited as shown in the following table:

<table>
<thead>
<tr>
<th>Procedure Codes</th>
<th>Limitations</th>
</tr>
</thead>
<tbody>
<tr>
<td>96156</td>
<td>Limited to a maximum of 2 per client, per rolling 180 days, any provider</td>
</tr>
<tr>
<td>96158, 96164, 96167</td>
<td>Limited to a maximum of 8 per client, per rolling 180 days, any provider</td>
</tr>
<tr>
<td>96159, 96165, 96168</td>
<td>Limited to a maximum of 14 per client, per rolling 180 days, any provider</td>
</tr>
</tbody>
</table>

Rural Health Clinics and Federally Qualified Health Centers may be reimbursed for client in-person evaluation visits based on encounter rates.

Documentation must be maintained in the client’s medical record that details the change in the mental or medical status warranting reassessment of the client’s capacity to understand and cooperate with the medical interventions that are necessary to the client’s health and well-being.

Clients must be referred for psychiatric evaluation or psychotherapy as soon as the need is identified. Providers cannot use all 16 units if the need for psychiatric or psychological intervention is identified earlier.
After the initial HBAI assessment, if the client is receiving behavioral health services from another health-care provider, the HBAI provider should coordinate with the external behavioral health provider and establish the most appropriate course of treatment for the client.

**Refer to:** Section 4, “Outpatient Mental Health Services” in the *Behavioral Health and Case Management Services Handbook* (Vol. 2, Provider Handbooks) for more information about behavioral health services beyond the acute care limitations outlined in this section.

The initial clinical interview, reassessment, psychophysiological monitoring, observation, and intervention do not include the following:

- Conversations about educating the family or caregivers outside of the in-person evaluation sessions
- Psychotherapy

After the initial 180 days of HBAI services, the client may receive another episode of HBAI with the same medical diagnosis if there is a newly identified behavioral health issue. The client may have two episodes of HBAI per rolling year.

HBAI services are adjunct to other services and are to be used as a non-intensive means to identify specific needs. As appropriate, the client should be referred for those additional services that would meet the client’s biopsychosocial needs.

### 2.9.2 Prior Authorization and Documentation Requirements

Prior authorization is not required for HBAI services.

Documentation is required for HBAI services to support the medical necessity of the initial assessment, reassessment, and intervention.

For the initial assessment, documentation must support the medical necessity of the assessment and must include the following information:

- The date of initial diagnosis of physical illness
- A clear rationale for assessment
- Outcome of assessment, which includes mental status and the client’s or caregiver’s ability to understand and respond meaningfully
- Goals and expected duration of specifically recommended psychological intervention(s)

For reassessment, documentation must support the reassessment is necessary and include the following information:

- The date of change in mental or physical status
- Rationale for re-assessment with a clear indication of precipitating events.

For the intervention, documentation must support the necessity of the intervention and include the following information:

- Evidence that the client or caregiver has the capacity to understand and respond meaningfully,
- Clear outline of planned psychological intervention
- Goals of the psychological intervention identifying expected improvement in compliance with the medical treatment plan
- The client’s response to the intervention
- Rationale for frequency and duration of acute care services

All documentation must include the amount of time spent in the HBAI assessment or intervention and must be documented in the client’s medical record.
All services are subject to retrospective review to ensure that the documentation in the client’s medical record supports the medical necessity of the services provided.

### 2.9.3 Claims Information

Claims for HBAI services must be submitted to TMHP in an approved electronic claims format or on a CMS-1500 paper claim form. Providers may purchase CMS-1500 paper claim forms from the vendor of their choice. TMHP does not supply the forms.

*Refer to:* “Section 3: TMHP Electronic Data Interchange (EDI)” *(Vol. 1, General Information)* for information on electronic claims submissions.

“Section 6: Claims Filing” *(Vol. 1, General Information)* for general information about claims filing.


### 2.9.4 Reimbursement

Providers may refer to the OFL or the applicable fee schedule on the TMHP website at [www.tmhp.com](http://www.tmhp.com) for reimbursement rates.

### 2.10 Medical Nutrition Counseling Services (CCP)

#### 2.10.1 Enrollment

Independently practicing licensed dietitians may enroll in Texas Medicaid to provide services to CCP-eligible clients. Dieticians who provide nutrition assessments and counseling must be currently licensed by the Texas Department of Licensing and Regulation (TDLR) in accordance with the Licensed Dietitians Act, Chapter 701, Texas Occupations Code.

*Refer to:* Subsection 2.1.2, “Enrollment” in this handbook for more information about CCP enrollment procedures.

#### 2.10.2 Services, Benefits, and Limitations

Medical nutrition therapy (assessment, re-assessment, and intervention) and medical nutrition counseling may be beneficial for treating, preventing, or minimizing the effects of illness, injuries, or other impairments. A case manager, school counselor, or school nurse may refer a client for medical nutrition counseling services.

Medical nutrition counseling services are a benefit when all of the following criteria are met:

- The client is 20 years of age or younger
- The client is eligible for CCP
- The services are prescribed by a physician
- The services are performed by a Medicaid-enrolled licensed dietitian
- Clinical documentation supports medical necessity and medical appropriateness
- FFP is available

Medical nutrition therapy and nutrition counseling may be considered beneficial for disease states for which dietary adjustment has a therapeutic role. Such disease states include, but are not limited to, the following conditions:

- Abnormal weight gain
- Cardiovascular disease
- Diabetes or alterations in blood glucose
• Eating disorders
• Gastrointestinal disorders
• Gastrostomy or other artificial opening of gastrointestinal tract
• Hypertension
• Inherited metabolic disorders
• Kidney disease
• Lack of normal weight gain
• Multiple food allergies
• Nutritional deficiencies

Nutrition intervention for the following conditions is considered experimental and investigational and is not a benefit:
• Attention-deficit hyperactivity disorder
• Chemical sensitivities
• Chronic fatigue syndrome
• Idiopathic environmental intolerance

Medical nutrition counseling services for the diagnosis of obesity without a comorbid condition is not a benefit.

Medical nutrition therapy (procedure code 97802) is a more comprehensive service than medical nutrition counseling and is provided to individual clients for assessment and intervention. Procedure code 97802 is limited to one session per day and four units per rolling year.

Medical nutrition therapy (procedure code 97803) is provided to individual clients for a reassessment and intervention, after the initial assessment and intervention. Procedure code 97803 may be used for direct therapy sessions with clients. These sessions are limited to 1 session per day and 12 units per rolling year.

Nutrition assessments and re-assessments are in-depth evaluations of both objective and subjective data related to an individual’s food and nutrient intake, lifestyle, and medical history. Nutrition assessments and re-assessments are performed as part of medical nutrition therapy. Nutrition assessments and re-assessments may be required as a result of a medical diagnosis and may be performed in conjunction with other therapies for treatment or as a goal to help clients make and maintain dietary changes. Documentation must include the following:
• Objective and subjective data obtained
• Height, weight, body mass index (BMI), and correlating percentiles on the growth curves
• Estimated caloric needs
• Nutritional diagnosis
• Intervention and plan
• Evaluation

Medical nutrition counseling (procedure code S9470) is provided to individual clients after an initial assessment and is less comprehensive than medical nutrition therapy. Nutritional counseling may be used to discuss the plan of care or intervention and to determine whether modifications are needed. Procedure code S9470 is limited to one visit per day and four visits per rolling year.
Medical nutrition group therapy (procedure code 97804) is not a benefit in the home setting, and does not include an individual nutrition assessment. Medical nutrition group therapy is limited to eight units per rolling year.

Medical nutrition group therapy may be provided to a group of clients with the same condition. While medical nutrition group therapy must be led by a Medicaid-enrolled dietitian licensed by the TDLR, other health-care providers may participate in the group sessions. The focus of the therapy is on nutrition and health for chronic conditions such as the following:

- Acquired acanthosis nigricans
- Diabetes
- Dysmetabolic syndrome X
- Eating disorder
- Hyperlipidemia
- Other specified hypoglycemia
- Pure hypercholesterolemia
- Pure hyperglyceridemia

Medical nutrition group therapy sessions must last at least 30 minutes, have a minimum of two clients and a maximum of ten clients, and must include the following:

- An age-appropriate presentation on nutrition issues related to the chronic condition. (The presentation may include information about prevention of disease exacerbation or complications and living with chronic illness. The presentation may also offer suggestions for making healthy food choices or changing ideas about food.)
- A question-and-answer period.

Client participation in medical nutrition group therapy is optional. Providers must obtain an informed consent from a client’s parent or guardian before rendering services. The medical documentation maintained in a client’s medical record must include the following:

- Physician prescription
- Referral, if applicable
- Location where the services were provided
- Services that were provided during medical nutrition group therapy
- Goals or objectives for the group therapy
- Client participation
- Beginning and ending time of the group therapy session

In the following table, the procedure codes in Column A will be denied as part of another service if they are submitted by any provider for the same date of service as the corresponding procedure codes in Column B:

<table>
<thead>
<tr>
<th>Column A: Procedure Codes Denied When Submitted With…</th>
<th>Column B: Procedure Codes</th>
</tr>
</thead>
<tbody>
<tr>
<td>S9470</td>
<td>97802, 97803, or 97804</td>
</tr>
</tbody>
</table>
Claims for medical nutrition therapy and counseling services should be submitted as follows:

<table>
<thead>
<tr>
<th>Procedure Code</th>
<th>Time Unit</th>
<th>Limitation</th>
</tr>
</thead>
<tbody>
<tr>
<td>97802 Initial assessment</td>
<td>15 minutes</td>
<td>4 units per rolling year</td>
</tr>
<tr>
<td>97803 Reassessment</td>
<td>15 minutes</td>
<td>12 units per rolling year</td>
</tr>
<tr>
<td>97804 Group</td>
<td>30 minutes</td>
<td>8 units per rolling year</td>
</tr>
<tr>
<td>S9470 Dietitian visit</td>
<td>Per visit</td>
<td>1 visit per day/ 4 visits per rolling year</td>
</tr>
</tbody>
</table>

### 2.10.3 Prior Authorization and Documentation Requirements

Prior authorization is required for services that exceed the limitations for medical nutrition therapy (assessment, re-assessment, and intervention), medical nutrition group therapy, and nutrition counseling visits.

Prior authorization is also required for consideration of other health conditions that are not addressed.

The following documentation must be submitted to the CCP Prior Authorization Unit for prior authorization:

- Completed CCP Prior Authorization Request Form
- Treatment plan
- Diagnosis of a condition for which there is medical necessity for the service
- Obstacles for not meeting goals
- Interventions planned to meet goals

The prescribing physician and provider must maintain documentation of medical necessity, including the completed CCP Prior Authorization Request Form, in a client’s medical record. The physician must maintain the original signed copy of the CCP Prior Authorization Request Form. The completed CCP Prior Authorization Request Form is valid for a period of up to six months from the date of the physician’s signature.

### 2.10.4 Claims Information

Providers must submit services provided by licensed dietitians in an approved electronic claims format or on a CMS-1500 paper claim form from the vendor of their choice. TMHP does not supply the forms.

Claims for services that have been prior authorized must reflect the PAN in Block 23 of the CMS-1500 paper claim form or its equivalent.

Refer to: “Section 3: TMHP Electronic Data Interchange (EDI)” (Vol. 1, General Information) for information on electronic claims submissions.

“Section 6: Claims Filing” (Vol. 1, General Information) for general information about claims filing.

Subsection 6.5, “CMS-1500 Paper Claim Filing Instructions” in “Section 6: Claims Filing” (Vol. 1, General Information) for instructions on completing paper claims.

Medical Nutrition Counseling (CCP Only) on the TMHP website at [www.tmhp.com](http://www.tmhp.com) for a claim form example.
2.10.5 Reimbursement
Dietitian services are reimbursed in accordance with 1 TAC §355.8441.

2.11 Personal Care Services (PCS) (CCP)

2.11.1 Enrollment
CCP providers that want to participate in the delivery of PCS to Medicaid clients must be enrolled with TMHP and have the appropriate HHSC licensure or certification.

All PCS providers must have a TPI and a National Provider Identifier (NPI).

Providers that are currently contracted with HHSC to administer consumer-directed services (CDS) or provide PCS through the service responsibility option (SRO), including providers currently enrolled in Texas Medicaid, are required to enroll or re-enroll separately as a CDS or SRO provider. Texas Medicaid enrolls only new providers that are currently contracted with HHSC to provide PCS through CDS and SRO.

Providers (other than those discussed above) that want to provide PCS to Medicaid clients must enroll through TMHP. Texas Medicaid enrollment rules for PCS participation require providers to have one of the following categories of HHSC licensure prior to enrollment:

- Personal Assistance Services (PAS)
- Licensed Home Health Services (LHHS)
- Licensed and Certified Home Health Services (LCHHS)

LCHH and LHH agencies that are currently enrolled through TMHP do not need to enroll as CCP-PCS providers to provide PCS. Providers must have a TPI in one of the following enrollment categories: LHHS agency, LCHHS agency, or PCS provider.

Providers that are enrolled as any entity other than an LHHS agency or LCHHS agency are required to meet the provider enrollment rules in order to participate in the delivery of PCS through Texas Medicaid.

Refer to: Subsection 2.1.2, "Enrollment" in this handbook for more information about CCP enrollment procedures.

2.11.2 Services, Benefits, and Limitations
PCS is a benefit of CCP for Texas Medicaid clients who are birth through 20 years of age. PCS may not be authorized in hospitals, nursing facilities, or intermediate care facilities for individuals with intellectual or developmental disabilities (ICF-IID). PCS will be denied when billed on the same date of service as an inpatient stay service. The provider may appeal the denied claim with documentation supporting that PCS was performed while the client was not in a hospital setting. PCS are support services provided to clients who meet the definition of medical necessity and require assistance with the performance of ADLs, instrumental activities of daily living (IADLs), and health maintenance activities (HMAs) due to a physical, cognitive, or behavioral limitation related to a client’s disability or chronic health condition. PCS are provided by someone other than the responsible adult of the client who is a minor child or the legal spouse of the client.

A responsible adult is an individual, 18 years of age or older, who has agreed to accept the responsibility for providing food, shelter, clothing, education, nurturing, and supervision for the client. Responsible adults include, but are not limited to, biological parents, adoptive parents, step parents, foster parents, legal guardians, court-appointed managing conservators, and the primary adult who is acting in the role of parent.
PCS are those services that assist eligible clients in performing ADLs, IADLs, and HMAs. The scope of ADLs, IADLs, and HMAs includes a range of activities that healthy, nondisabled adults can perform for themselves. Typically, developing children gradually and sequentially acquire the ability to perform these ADLs, IADLs, and HMAs for themselves. If a typically developing child of the same chronological age could not safely and independently perform an ADL, IADL, or HMA without adult supervision, then the client’s responsible adult ensures that the client’s needs for the ADLs, IADLs, and HMAs are met.

PCS include direct intervention (assisting the client in performing a task) or indirect intervention (cuing or redirecting the client to perform a task). ADLs, IADLs, and HMAs include, but are not limited to, the following:

<table>
<thead>
<tr>
<th>ADLs</th>
<th>IADLs</th>
</tr>
</thead>
<tbody>
<tr>
<td>Bathing</td>
<td>Escort or Assistance with Transportation Services</td>
</tr>
<tr>
<td>Dressing</td>
<td>Grocery or Household Shopping</td>
</tr>
<tr>
<td>Eating</td>
<td>Laundry</td>
</tr>
<tr>
<td>Locomotion or Mobility</td>
<td>Light housework</td>
</tr>
<tr>
<td>Personal Hygiene</td>
<td>Meal preparation</td>
</tr>
<tr>
<td>Positioning</td>
<td>Medication Assistance</td>
</tr>
<tr>
<td>Toileting</td>
<td>Money management</td>
</tr>
<tr>
<td>Transferring</td>
<td>Telephone Use or Other Communication</td>
</tr>
</tbody>
</table>

* Escort or Assistance with Transportation Services includes the coordination of transportation to medical appointments and accompaniment to appointments to assist with needed ADLs. PCS does not include the payment for transportation or transportation vehicles since these services are available through MTP.

Note: Health maintenance activities (HMAs) and nurse-delegated tasks that fall within the scope of the task listed above are allowable in PCS.

Note: Exercise and range of motion are not available through PCS, but are services that could be provided through PT, PDN, or home health SN.

PCS does not include the following:

- ADLs, IADLs, or HMAs that a typically developing child of the same chronological age could not safely and independently perform without adult supervision
- Services that provide direct intervention when the client has the physical, behavioral, and cognitive abilities to perform an ADL, IADL, or health-related function without adult supervision
- Services used for or intended to provide respite care, child care, or restraint of a client
- Stand-by supervision related to safety
- Potty training
- Grocery shopping for members of the client’s family or household
- Cleaning for members of the client’s family or household (exception: light housework is approved if the client shares a room with a person)
- Cleaning the entire house (exception: a need for clean environment is approved if related to the client’s diagnosis or condition [e.g., asthma, allergies, or autoimmune deficiencies])

Note: Cleaning an area or equipment that is used to complete a task may be included in the light housework IADL, as appropriate.
• Laundry services for members of the client’s family or household (exception: laundry is approved when related to the client’s diagnosis or condition that results in soiled bedding or clothing for the client beyond the norm [e.g., incontinence, feeding tube, trachea, an ostomy, diapers, or skin condition])

• Waiting time for the laundry machine to complete a cycle in the home setting (exception: the time an attendant is at a laundromat completing the laundry task for the client is covered for PCS)

• Meal preparation for members of the client’s family or household

• Time of a PCS attendant while acting as the responsible adult for the receipt of medical care or providing medical transportation

An escort is approved if it is related to the client’s diagnosis or condition, such as using the toilet at the appointment or assistance carrying equipment (e.g., feeding pump, oxygen tank).

An escort is approved if it is related to the client’s diagnosis or condition and the responsible adult is occupied during the transport. For example, a child’s condition might include behaviors that create an unsafe situation for the child during transport, such as removing a seatbelt, attempting to open the car door while the car is in motion, or elopement.

PCS does cover the entire time that an attendant is away from the home performing this task.

PCS is considered for reimbursement when providers use procedure code T1019 in conjunction with the appropriate modifier listed in the following table. PCS provided by a home health agency or PCS-only provider, including PCS being provided under the SRO defined in 40 TAC Part 1, Chapter 41, must be billed in 15-minute increments. PCS provided by a financial management services agency (FMSA) under the CDS option defined in 40 TAC Part 1, Chapter 41, must submit the attendant fee in 15-minute increments. FMSAs must bill the administration fee once per calendar month per client for any month in which the client receives PCS under the CDS option and regardless of the number of PCS units of service the client receives under the CDS option during the month. PCS claims are considered for reimbursement only when TMHP has issued a valid PAN to a PCS provider.

<table>
<thead>
<tr>
<th>PCS Procedure Codes</th>
</tr>
</thead>
<tbody>
<tr>
<td><em><em>All PCS Providers</em> (except FMSA)</em>*</td>
</tr>
<tr>
<td>Procedure Code: T1019</td>
</tr>
<tr>
<td>Modifier: U6 (PCS each 15 minutes)</td>
</tr>
<tr>
<td>UA (Behavioral health condition, each 15 minutes)</td>
</tr>
</tbody>
</table>

| **FMSA Under CDS Option*** |
| Procedure Code: T1019 |
| Modifier: U7 (Attendant fee each 15 minutes) |
| U8 (Administration fee once a month) |
| UB (Behavior health condition, each 15 minutes) |

* 40 TAC Part 1, Chapter 41

Home health agencies and Personal Care Services (PCS) providers that provide PCS and Community First Choice (CFC) Services in the home setting may be reimbursed for nurse evaluation and supervision using procedure code G0162.

The following limitations apply for procedure code G0162:

• For a registered nurse (RN) assessment, procedure code G0162 (without modifier) is limited to three hours per day (12 to 15 minute increments) and two occurrences per rolling year for any provider.
• For training and supervision of the attendant, procedure code G0162 must be billed with modifier U1 and is limited to three hours (12 to 15 minute increments) per 30 days for any provider.

    **Note:** *Training and supervision and an RN assessment may be billed on the same day.*

Prior authorization is not required for procedure code G0162.

### 2.11.2.1 Place of Service

PCS may be provided in the following settings if medically necessary:

- The client’s home
- The client’s school
- The client’s daycare facility
- Other community setting in which the client is located

    **Note:** *For claims filing purposes, the PCS provider must bill POS 2 (home) when submitting claims to TMHP.*

Texas Medicaid does not reimburse providers for PCS that duplicate services that are the legal responsibility of school districts. The school district, through the School Health and Related Services (SHARS) program, is required to meet the client’s personal care needs while the client is at school. If those needs cannot be met by SHARS or the school district, the school district must submit documentation to the Texas Department of State Health Services (DSHS) case manager indicating the school district is unable to provide all medically necessary services. When clients are receiving both PCS and PDN services from an individual person over the same span of time, the combined total number of hours for PCS and PDN are reimbursed according to the maximum allowable rate.

### 2.11.2.2 Client Eligibility

The PCS benefit is available to Texas Medicaid clients who:

- Are birth through 20 years of age.
- Are enrolled with Texas Medicaid.
- Are eligible for CCP.
- Have physical, cognitive, or behavioral limitations related to a disability or chronic health condition that inhibits the client’s ability to accomplish ADLs, IADLs, or HMAs.

Whether the client has a physical, cognitive, or behavioral limitation related to a disability or chronic health condition that inhibits the client’s ability to accomplish ADLs, IADLs, or HMAs, the following needs and conditions of the responsible adult will be considered:

- The responsible adult’s need to sleep, work, attend school, and meet their own medical obligations
- The responsible adult’s legal obligation to care for, support, and meet the medical, educational, and psychosocial needs of other dependents
- Whether requiring the responsible adult to perform the PCS will put the client’s health or safety in jeopardy
- The time periods during which the PCS tasks are required by the client, as they occur over the course of a 24-hour day and a seven-day week.
- Whether or not the need to help the family perform PCS on behalf of the client is related to a medical, cognitive, or behavioral condition that results in a level of functional ability that is below that expected of a typically developing child of the same chronological age
- Whether services are needed based on:
  - The Practitioner Statement of Need (PSON)
• The client’s personal care assessment form (PCAF)

Clients who are enrolled in a HHSC waiver program may also receive PCS if they are eligible for it, as long as the services that are provided through the waiver program and PCS are not duplicated. Clients who are enrolled in the following HHSC waiver programs may access the PCS benefits if they meet the PCS eligibility requirements:

- Community Living Assistance and Support Services (CLASS)
- Deaf/Blind Multiple Disabilities (DBMD)
- Community-Based Alternatives (CBA)
- Consolidated Waiver Program (CWP)
- Medically Dependent Children Program (MDCP)
- Texas Home Living Waiver (TxHmL)
- Home and Community Services (HCS)

Note: Clients who receive HCS Residential Support Services, Supervised Living Services, or Foster/Companion Care Services are not eligible to receive attendant care services through PCS.

Clients must choose the program through which they receive attendant care, if they meet the eligibility requirements of both programs. Clients will be given the following options for the delivery of attendant care services:

- A client can receive all attendant care services through PCS.
- A client can decline PCS and receive all attendant care service through a waiver program, if the waiver program offers attendant care.

Clients who participate in the CDS option for PCS and for a waiver program are required to choose one FMSA to provide services through both programs. FMSAs will only be permitted to file the financial management services (FMS) fee, also known as the monthly administrative fee, through one program. The FMSA must file the FMS claim through the program that provides the highest reimbursement rate.

2.11.2.2.1 Accessing the PCS Benefit

Clients must be referred to DSHS before receiving the PCS benefit. A referral can be made by any person who recognizes a client may have a need for PCS, including, but not limited to, the following:

- The client, a parent, a guardian, or a responsible adult
- A primary practitioner, primary care provider, or medical home
- A licensed health professional who has a therapeutic relationship with the client and ongoing clinical knowledge of the client
- A family member
- Home health, personal assistance, or FMSA providers

Referrals to DSHS can be made to the appropriate DSHS Health Service Region, based on the client’s place of residence in the state. Clients, parents, or guardians may also call the TMHP PCS Client Line at 1-888-276-0702 for more information on PCS. PCS providers must provide contact information for the client or responsible adult to DSHS or the TMHP PCS Client Contact Line when making a referral.

Upon receiving a referral, DSHS assigns the client a case manager, who then conducts an assessment in the client’s home with the input and assistance of the client or responsible adult. Based on the assessment, the case manager identifies whether the client has a need for PCS. If the case manager identifies a need for PCS, the client or responsible adult is asked to select a Medicaid-enrolled PCS provider in their area.
Once a provider is selected, the DSHS case manager prior authorizes a quantity of PCS based on the assessment and requests TMHP to issue a PAN to the selected PCS provider. The PCS provider uses the PAN to submit claims to TMHP for the services provided.

2.11.2.2.2 The Primary Practitioner’s Role in the PCS Benefit

A client who is assessed for the PCS benefit must have a primary practitioner (a licensed physician, APRN, or PA) or a primary care provider who has personally examined the client within the last 12 months and reviewed all of the appropriate medical records. The primary practitioner or primary care provider must have established a diagnosis for the client and must provide continuing care and medical supervision of the client. Prior to authorizing PCS, HHSC requires the completion of an HHSC-approved Practitioner Statement of Need (PSON) by a primary practitioner. The PSON must be on file with HHSC prior to the initiation of PCS and will only accept the PSON from an individual who is a physician, APRN, or PA.

The PSON certifies that the client is 20 years of age or younger and has a physical, cognitive, or behavioral limitation related to a disability or chronic health condition. The primary practitioner or primary care provider must mail or fax the completed PSON to the appropriate DSHS Health Services Region. DSHS keeps the signed and dated PSON and the client’s PCAF in the client’s case management record for the duration of the client’s participation in the benefit.

When a behavioral health condition exists, the primary practitioner may be a behavioral health provider. If the client’s medical record does not include the primary practitioner’s documentation and a PSON that certifies that the client has a physical, cognitive or behavioral health condition that impacts the client’s ability to perform an ADL or IADL, then PCS payments may be recouped.

Note: If a client is entering or is already in the conservatorship of the state, PCS may be provisionally initiated for up to 60 days once eligibility has been established through the assessment.

HHSC requires the reassessment of the client’s need for PCS every 12 months or when requested due to a change in the client’s health or living condition. A new PSON will be required at each annual reassessment and when there is a change in the client’s medical condition that may increase the need for services.

2.11.2.3 PCS Provided in Group Settings

PCS may be provided in a provider to client ratio other than one-to-one. Settings in which providers can provide PCS in a provider to client ratio other than one-to-one include homes with more than one client needing PCS, foster homes, and independent living arrangements.

A PCS provider may provide PCS to more than one client over the span of the day as long as:

- Each client’s care is based on an individualized service plan.
- Each client’s needs and service plan do not overlap with another client’s needs and service plan.

Example: If the prior authorized PCS hours for Client A is four hours, Client B is six hours, and the actual time spent with both clients is eight hours, the provider must bill for the actual one-on-one time spent with each client, not to exceed the client’s prior authorized hours or total hours worked. It would be acceptable to bill four hours for Client A and four hours for Client B, or three hours for Client A and five hours for Client B. It would not be acceptable to bill five hours for Client A and three hours for Client B. It would be acceptable to bill ten hours if the individual person actually spent ten hours onsite providing prior authorized PCS split as four hours for Client A and six hours for Client B. A total of ten hours cannot be billed if the individual person worked only eight hours.
• PCS may be delivered in a client-to-provider ratio other than one-on-one as long as each client’s care is based on an individualized POC and each client’s needs are being met. Only the time spent on authorized PCS tasks for each client is eligible for reimbursement. Total PCS billed for all clients cannot exceed an individual attendant’s total number of hours at the place of service.

When there is more than one client within the same household receiving PCS, the DSHS case manager will synchronize authorizations within the households for all eligible clients. The DSHS case manager will assess all eligible clients in the home and submit authorizations for all eligible clients in the household for the same authorization period. DSHS case managers will communicate with the provider the actions that are being taken using the existing Communication Tool.

Note: There should be no lapse in services to the client.

2.11.3 Prior Authorization and Documentation Requirements

Prior authorization is required before services are provided. All PCS must be prior authorized by a DSHS case manager based upon client need, as determined by the client assessment. DSHS prior authorizes PCS for eligible clients. The DSHS case manager notifies TMHP of the authorized quantity of PCS. TMHP sends a notification letter with the PAN to the client or responsible adult and the selected PCS provider if PCS is approved or modified. Only the client or responsible adult receives a notification letter with an explanation of denied services. PCS is prior authorized for periods of up to twelve months. PCS providers must provide services from the start of care date agreed to by the client or responsible adult, the case manager, and the PCS provider.

PCS may be authorized in the same day as PPECC, if medically necessary. However, they must be rendered in a home setting, before or after PPECC services.

A PCS provider may obtain prior authorization to provide enhanced PCS to clients with a behavioral health condition when the following criteria are met:

• The DSHS case manager completes the Personal Care Assessment Form (PCAF) and identifies the behavioral health condition.
• The PCAF indicates that the identified behavioral health condition impacts the client’s ability to perform an ADL or IADL.
• The PCAF indicates which ADL(s) or IADL(s) cannot be performed by the client without assistance.
• The DSHS case manager submits the appropriate modifier on the authorization request.

When a client experiences a change in condition, the client or responsible adult must notify the DSHS Health Service Office in the client’s region. A new assessment is required when a client’s physician orders services in a PPECC. A DSHS case manager must perform a new assessment and prior authorize any revisions in the quantity of PCS based on the new assessment. TMHP issues a revised authorization and notifications are sent to the client or responsible adult and the selected PCS provider. If the change is made during a current prior authorization period, the new prior authorization will maintain the same end date as the original prior authorization period. The revised authorization period will begin on the SOC date stated in the new assessment.

For continuing and ongoing PCS needs beyond the initial prior authorization period of up to twelve months, a DSHS case manager must conduct a new assessment and submit a new authorization request to TMHP. TMHP sends a notification letter updating the prior authorization to the client, responsible adults, and the selected PCS provider.

HHSC or its designee may suspend an authorization for PCS when either:

• The client or the client’s family creates an unsafe environment for the attendant’s health and safety.
• The provider requests suspension for the reasons outlined in 40 TAC Part 1, Chapter 41.
Providers can call a toll-free PCS Provider Inquiry Line at 1-888-648-1517 for assistance with inquiries about the status of a PCS prior authorization. Providers should direct inquiries about other Medicaid services to the TMHP Contact Center at 1-800-925-9126. PCS providers should encourage the client or responsible adult to contact the appropriate DSHS Health Service Region with inquiries or concerns about the PCS assessment.

**Note:** Any organization that employs attendants who provide PCS, and any organization serving as an FMSA, must comply with all documentation requirements as specified by the PCS program.

### 2.11.3.1 PCS Provider Responsibilities

PCS providers must comply with all applicable federal, state, and local laws and regulations.

All PCS providers must maintain written policies and procedures for obtaining consent for medical treatment in the absence of the responsible adult. The procedure and policy must meet the standards of the Texas Family Code, Chapter 32.

Providers must accept clients only when there is a reasonable expectation and evidence that the client’s needs can be adequately met in the POS. The POS must be able to support the client’s health and safety needs and adequately support the use, maintenance, and cleaning of all required medical devices, equipment, and supplies. Necessary primary and backup utility, communication, and fire safety systems must be available in the POS.

The PCS provider is responsible for the supervision of the PCS attendant as required by the PCS provider’s licensure requirements.

### 2.11.3.2 Documentation of Services Provided and Retrospective Review

Documentation elements are routinely assessed for compliance in retrospective review of client records, including the following:

- All entries are legible to people other than the author, dated (month, day, year, time), and signed by the author.
- Each page of the record documents the client’s name and Medicaid identification number.
- All attendants’ arrival and departure times are documented with signature and time.
- Documentation of services correlates with, and reflects medical necessity for, the services provided on any given day.
- Client’s arrival or departure from the home setting is documented with the time of arrival, departure, mode of transportation, and who accompanied the client.

### 2.11.4 Coordination with PPECC Provider

When a DSHS case manager is notified by the client, client’s responsible adult, or client’s physician that PPECC services have been initiated, revised, or recertified, the DSHS case manager must conduct a PCS reassessment, and submit all documentation required for a revision, modification, or denial of the original PCS authorization request, including a Physician’s Statement of Need if there is a change in client condition. The new authorization request must be submitted within ten (10) business days of notification.

DSHS case managers must provide documentation to support medical necessity if PCS service hours do not decrease when PPECC services are initiated.

DSHS case managers must also document coordination with the PPECC provider, maintaining documentation that the client or the client’s responsible adult has participated in the development of the plan of care.
When a client receives both PCS and PPECC in a single day, and decides to receive fewer hours of service in a PPECC (i.e., shifts more services to the home setting), or terminates PPECC services, the PCS case manager must conduct a reassessment, and submit all documentation required for a revision, modification, or denial of the original PCS authorization request, including a Physician’s Statement of Need if there is a change in condition within ten (10) business days of notification by the client, client’s responsible adult, or client’s physician.

Similarly, if the client decides to receive fewer hours of PCS services in the home, and increase PPECC hours, the DSHS case manager must conduct a reassessment, and submit all documentation required for an modification of the authorization request, within ten (10) business days of notification by the client, client’s responsible adult, or client’s physician.

Note: Coordination requirements may be different in a Medicaid managed care environment.

PCS services rendered in a client’s home may be billed before or after PPECC services on the same day, but not at the same time as PPECC services. PCS services required while a client is in a PPECC are considered part of the PPECC billable rate.

2.11.5 Claims Information

TMHP processes PCS claims. PCS providers must submit claims for services in an approved electronic claims format or on the appropriate claim form based on their provider type. PCS providers, other than home health agencies, that are enrolled as PAS-only providers, FMSAs, or SRO providers must file PCS claims using a CMS-1500 paper claim form. Home health agencies, including those enrolled as an FMSA, or an SRO provider, must file PCS claims using the UB-04 CMS-1450 paper claim form. TMHP does not supply the forms.

Home health agencies and consumer-directed agencies that bill for PCS using procedure code T1019 must include the prior authorization number on claims submitted for reimbursement. Additionally, providers utilizing paper, TexMedConnect, or billing through EDI must include the prior authorization number with all claims submissions.

2.11.5.1 Managed Care Clients

PCS services are carved-out of the Medicaid Managed Care Program for State of Texas Access Reform (STAR) clients and must be billed to TMHP for payment consideration. Carved-out services are those that are rendered to Medicaid Managed Care clients but are administered by TMHP and not the client’s MCO. Claims for STAR Health, STAR Kids, and STAR+PLUS are not carved out and must be submitted to the client’s MCO for payment consideration.

2.11.5.2 PCS for STAR Health Clients

PCS for eligible STAR Health clients are authorized and processed by Superior HealthPlan.

Medicaid providers that want to provide PCS services to clients in the STAR Health program should contact Superior HealthPlan for information regarding the contracting and credentialing process at:

Superior HealthPlan - Network Development
Telephone: 1-866-615-9399 Ext. 22534
Email: shp-networkdevelopment@centene.com

2.11.6 Reimbursement

Providers of PCS are reimbursed in accordance with 1 TAC §355.8441.
2.12 Community First Choice (CFC) Services

2.12.1 Enrollment

CFC providers, including providers offering the Service Responsibility Option (SRO), must be licensed and enrolled in Texas Medicaid and comply with all applicable federal, state, and local laws and regulations. When CFC is provided through the Consumer Directed Services (CDS) option by a Financial Management Services Agency (FMSA), the FMSA must be certified and enrolled in Texas Medicaid as a FMSA and must comply with all applicable federal, state, and local laws and regulations.

Note: CDS, FMSA, and SRO are defined in the Title 40 Texas Administrative Code (TAC), Part 1, Chapter 41. Licensure requirements for FMSA and SRO providers are defined in the Title 40 TAC, Part 1, Chapter 49.

Note: Any organization that employs attendants who provide CFC, and any organization serving as an FMSA, must comply with all documentation requirements as specified in CFC program policy.

All CFC providers must maintain written policies and procedures for obtaining consent for medical treatment for clients in the absence of a responsible adult that meet the standards of the Texas Family Code, Chapter 32.

Providers must only accept clients when there is a reasonable expectation and evidence that the client’s CFC needs can be adequately met in the place of service.

The CFC provider is responsible for supervising the CFC attendant in accordance with the provider’s licensure requirements.

Note: For CFC services delivered through a HHSC 1915(c) waiver or through a managed care organization (MCO), providers must refer to HHSC or the MCO for information about benefits, limitations, prior authorization, reimbursement, and specific claim processing procedures.

2.12.2 Services, Benefits, and Limitations

CFC services may be rendered by a Home Health Agency, PCS-only provider, Financial Management Services Agency (FMSA) under the CDS option, or by a Service Responsibility Option (SRO) Provider.

CFC is a benefit for Texas Medicaid fee-for-service clients who are birth through 20 years of age and who are:

- Eligible for medical assistance under the state plan
- Need help with activities of daily living; and
- Need an institutional level of care (to include hospital, nursing facility, intermediate care facility for clients with intellectual disabilities or institution of mental disease)

Services included are CFC personal assistance services, CFC habilitation, and CFC support management. CFC personal assistance services is the assistance with activities of daily living (ADLs) and instrumental activities of daily living (IADLs) through hands-on assistance, supervision, and/or cueing. CFC habilitation services are the acquisition, maintenance, and enhancement of skills necessary for the client to accomplish ADLs, IADLs, and health maintenance activities (HMAs). CFC support management is voluntary training on how to select, manage, and dismiss attendants.

CFC includes assistance with ADLs, IADLs and HMAs through hands-on assistance, supervision, or cueing. CFC also includes training on the acquisition, maintenance, and enhancement of skills necessary for the client to accomplish ADLs, IADLs, and HMAs.
ADLs are activities that include:

- **Bathing**—Assisting the client with any or all parts of bathing; selecting appropriate water temperature and flow speed, turning water on and off; laying out and putting away supplies; transferring in and out of bathtub or shower; washing and drying hair and body; clean up after task is completed.

- **Dressing**—Assisting the client with any or all parts of getting dressed; putting on, fastening, and taking off all items of clothing; donning and removing shoes or prostheses; choosing and laying out weather-appropriate clothing.

- **Eating**—Assisting the client with some or all parts of eating and drinking; feeding the client; assistance with utensils or special or adaptive eating devices; clean up after task is completed.

- **Personal hygiene**—Assisting the client with some or all parts of personal hygiene; routine hair care; oral care; ear care; shaving; applying makeup; managing feminine hygiene; washing and drying face, hands, perineum; basic nail care; applying deodorant; routine skin care; clean up after task is completed.

- **Toileting**—Assisting the client with some or all parts of toileting; using commode, bedpan, urinal, toilet chair; transferring on and off; cleansing; changing diapers, pad, incontinence supplies; adjusting clothing; clean up after task is completed.

- **Locomotion or mobility**—Assisting the client with moving between locations; assisting the client with walking or using wheelchair, walker, or other mobility equipment.

- **Positioning**—Assisting the client with positioning their body while in a chair, bed, or other piece of furniture or equipment; changing and adjusting positions; moving to or from a sitting position; turning side-to-side; assisting the client to sit upright.

- **Transferring**—Assisting the client with moving from one surface to another with or without a sliding board; moving from bed, chair, wheelchair, or vehicle to a new surface; moving to or from a standing or sitting position; moving the client with lift devices.

IADLs are activities that include:

- **Telephone use or other communication**—Assisting the client in making or receiving telephone calls; managing and setting up communication devices; making and receiving the call for the client.

- **Grocery or household shopping**—Shopping for or assisting clients in shopping for grocery and household items; preparing a shopping list; putting food and household items away; picking up medication and supplies.

- **Light housework**—Performing or assisting the client in performing light housework such as: cleaning and putting away dishes; wiping countertops; dusting; sweeping, vacuuming or mopping; changing linens and making bed; cleaning bathroom; taking out trash.

- **Laundry**—Assisting the client with doing laundry; gathering, sorting, washing, drying, folding, and putting away personal laundry, bedding, and towels; removing bedding to be washed and remaking the bed; using a laundry facility.

- **Meal preparation**—Assisting clients in preparing meals and snacks; cooking; assembling ingredients; cutting, chopping, grinding, or pureeing food; setting out food and utensils; serving food; preparing and pouring a predetermined amount of liquid nutrition; cleaning the feeding tube; cleaning area after meal; washing dishes.

- **Money management**—Assisting the client with managing their day-to-day finances; paying bills; balancing checkbook; making deposits or withdrawals; assisting in preparing and adhering to a budget.
- Medication assistance or administration—Assisting the client with oral medications that are normally self-administered, including administration through a permanently placed feeding tube with irrigation.

- Escort or assistance with transportation services—Assisting the client in making transportation arrangements for medical and other appointments; accompanying the client to a health care appointment to assist with needed ADLs.

HMAs include tasks that may be exempt from delegation based on the Registered Nurse (RN) assessment that enables the client to remain in an independent living environment and go beyond ADLs because of the higher skill level in which they are required to perform. HMAs will be limited to those within the scope of CFC that include:

- Administering oral medications that are normally self-administered, including administration through a permanently placed feeding tube with irrigation.

- Topically applied medications.

- Insulin or other injectable medications prescribed in the treatment of diabetes mellitus administered subcutaneously, nasally, or via an insulin pump.

- Unit dose medication administration by way of metered dose inhaler (MDIs) including medications administered as nebulizer treatments for prophylaxis or maintenance.

- Routine administration of a prescribed dose of oxygen.

- Noninvasive ventilation (NIV) such as continuous positive airway pressure (CPAP) and bi-level positive airway pressure (BiPAP) therapy.

- The administering of a bowel and bladder program, including suppositories, enemas, manual evacuation, intermittent catheterization, digital stimulation associated with a bowel program, tasks related to external stoma care including but not limited to pouch changes, measuring intake and output, and skin care surrounding the stoma area.

- Routine preventive skin care and care of Stage 1 pressure ulcers.

- Feeding and irrigation through a permanently placed feeding tube inserted in a surgically created orifice or stoma.

- Those tasks that an RN may reasonably conclude as safe to exempt from delegation based on an assessment consistent with 22 Texas Administrative Code (TAC) §225.6 of this title (relating to the RN Assessment of the client).

- Reporting as to the client’s condition, including changes to the client’s condition or needs and completing appropriate record.

- Skin care: Maintenance of the hygienic state of the client’s skin under optimal conditions of cleanliness and comfort.

- Use of durable medical equipment (DME).

- Such other tasks as the Board of Nursing may designate.

Whether the client has a physical, cognitive, or behavioral limitation related to a disability or chronic health condition that inhibits the client’s ability to accomplish ADLs, IADLs, or HMAs, the following needs and conditions of the responsible adult will be considered in the determination of hours for CFC personal assistance services:

- The responsible adult’s need to sleep, work, attend school, and meet their own medical obligations

- The responsible adult’s legal obligation to care for, support, and meet the medical, educational, and psychosocial needs of other dependents
• Whether requiring the responsible adult to perform the CFC personal assistance services will put the client's health or safety in jeopardy
• The time periods during which the CFC personal assistance services tasks are required by the client, as they occur over the course of a 24-hour day and a seven-day week.
• Whether or not the need to help the family perform CFC personal assistance services on behalf of the client is related to a medical, cognitive, or behavioral condition that results in a level of functional ability that is below that expected of a typically developing child of the same chronological age

CFC also includes training on the acquisition, maintenance, and enhancement of the following additional habilitation needs:

• Community integration—Client may need assistance finding, participating in and accessing community activities or community services such as free meal programs, churches, parks or self-advocacy training or events.
• Use of adaptive equipment—Client may need assistance operating, learning to use, or accessing adaptive equipment.
• Personal decision-making—Client may need assistance making decisions for him or herself, including assistance in assessing what is important to that client, pros and cons, as well as consequences.
• Reduce challenging behaviors to allow clients to accomplish ADLs, IADLs, and HMAs—Client may need assistance in increasing positive social encounters and engagement in preferred activities. Client may have challenging behaviors that can be reduced through behavior support plans, prompting, rewards, or redirection among others.
• Socialization/relationship development—Client may need assistance with development and maintenance of relationships or appropriate social behaviors.
• Accessing leisure and recreational activities—Client may need assistance identifying, finding, or accessing activities they would like to participate in during leisure time.

CFC does not include the following:

• Direct intervention to perform a task the client has the physical, behavioral, and cognitive abilities to perform;
• Skilled nursing services, or the supervision of delegated nursing tasks as described in the Texas Nurse Practice Act, the Act’s implementing regulations, the Texas Medicaid Provider Procedures Manual sections for Private Duty Nursing (PDN) Services - THSteps - Comprehensive Care Program (CCP) and Home Health Skilled Nursing and Home Health Aide Services;
• Costs associated with purchasing products for ADLs or IADLs;
• Services used for or intended to provide respite care, child care, or restraint of a client;
• Duplication of services provided by another program;
• Tasks that a typically developing child of the same chronological age could not safely and independently perform without adult supervision;
• Services provided in an institutional setting including hospitals, nursing facilities, psychiatric hospitals, or intermediate care facilities for clients with intellectual or developmental disabilities.
CFC is considered for reimbursement when billed with procedure code T1019 in conjunction with the appropriate modifier listed in the following table. CFC provided by a home health agency or PCS-only provider, including CFC being provided under the SRO defined in 40 TAC Part 1, Chapter 41, must be billed in 15-minute increments. CFC provided by a FMSA under the CDS option defined in 40 TAC Part 1, Chapter 41, must submit the attendant fee in 15-minute increments.

### CFC Procedure Codes

<table>
<thead>
<tr>
<th>Procedure Code</th>
<th>Modifier</th>
<th>Description</th>
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<tbody>
<tr>
<td>T1019</td>
<td>UD</td>
<td>UD (CFC—client needs attendant care only, each 15 minutes)</td>
</tr>
<tr>
<td></td>
<td>U9</td>
<td>U9 (CFC—client needs habilitation only, or attendant and habilitation, each 15 minutes)</td>
</tr>
<tr>
<td>T1019</td>
<td>U3</td>
<td>U3 (CFC attendant care for PCS - CDS Option, each 15 minutes)</td>
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<tr>
<td></td>
<td>U4</td>
<td>U4 (CFC habilitation for PCS - CDS Option, each 15 minutes)</td>
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<tr>
<td></td>
<td>U5</td>
<td>U5 (CFC CDS, per month)</td>
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*Note:* This modifier will be used for individuals receiving attendant care only.

*Note:* This modifier will be used for individuals receiving attendant care and habilitation.

*Note:* This modifier is used for the administrative fee for CFC provider under the CDS option.

FMSAs must bill the administration fee once per calendar month per client for any month in which the client receives CFC under the CDS option and regardless of the number of CFC units of service the client receives under the CDS option during the month. CFC claims are considered for reimbursement only when TMHP has issued a valid PAN to a CFC provider.

Home health agencies and Personal Care Services (PCS) providers that provide PCS and Community First Choice (CFC) Services in the home setting may be reimbursed for nurse evaluation and supervision using procedure code G0162.

The following limitations apply for procedure code G0162:

- For a registered nurse (RN) assessment, procedure code G0162 (without modifier) is limited to three hours per day (12 to 15 minute increments) and two occurrences per rolling year for any provider.
- For training and supervision of the attendant, procedure code G0162 must be billed with modifier U1 and is limited to three hours (12 to 15 minute increments) per 30 days for any provider.

*Note:* Training and supervision and an RN assessment may be billed on the same day.

Prior authorization is not required for procedure code G0162.

### 2.12.2.1 Place of Service

CFC may be provided in the following settings:

- Client’s home;
- Client’s school;
• Client’s daycare facility; or
• Other community setting in which the client is located.

Note: For claims filing purposes, the CFC provider must bill POS 2 (home) when submitting claims to TMHP.

Texas Medicaid does not reimburse providers for CFC services that duplicate services that are the legal responsibility of school districts. The school district, through the School Health and Related Services (SHARS) program, is required to meet the client’s personal care needs while the client is at school. If those needs cannot be met by SHARS or the school district, the school district must submit documentation to the DSHS case manager indicating the school district is unable to provide all medically necessary services.

2.12.3 CFC Attendant and Habilitation Services in Group Settings

CFC may be provided in a provider to client ratio other than one-to-one. Settings in which providers can provide CFC in a provider to client ratio other than one-to-one include homes with more than one client needing CFC, foster homes, and independent living arrangements. A CFC provider may provide CFC to more than one client over the span of the day as long as:

• Each client’s care is based on an individualized service plan.
• Each client’s needs and service plan do not overlap with another client’s needs and service plan.
• Only the time spent on authorized CFC tasks for each client is eligible for reimbursement. Total CFC billed for all clients cannot exceed an individual attendant’s total number of hours at the place of service.

When there is more than one client within the same household receiving CFC, the Department of State Health Services (DSHS) case manager will synchronize authorizations within the households for all eligible clients. The DSHS case manager will assess all eligible clients in the home and submit authorizations for all eligible clients in the household for the same authorization period. DSHS case managers will communicate with the provider the actions that are being taken using the existing Communication Tool.

2.12.4 Prior Authorization

Prior authorization is required before services are provided. All CFC must be prior authorized by a DSHS case manager based upon client need, as determined by the client assessment. DSHS prior authorizes CFC for eligible clients. The DSHS case manager notifies TMHP of the authorized quantity of CFC. TMHP sends a notification letter with the prior authorization number (PAN) to the client or responsible adult and the selected CFC provider if CFC is approved or modified. Only the client or responsible adult receives a notification letter with an explanation of denied services. CFC is prior authorized for periods of up to twelve months. CFC providers must provide services from the start of care date agreed to by the client or responsible adult, the case manager, and the CFC provider.

When DSHS has approved CFC services, DSHS will send the client’s selected CFC provider: A CFC Communication tool, specifying the approved hours and CFC tasks and a copy of the Personal Care Assessment Form (PCAF) CFC Addendum, which documents that client’s goals and preferences for the delivery of CFC services. The CFC provider may receive a Practitioner’s Statement of Need (PSON) for the client, but this form is not required documentation for CFC and is intended merely for informational purposes.

When a client experiences a change in condition, the client or responsible adult must notify the DSHS Health Service Office in the client’s region. A DSHS case manager must perform a new assessment and prior authorize any revisions in the quantity of CFC based on the new assessment. TMHP issues a revised authorization and notifications are sent to the client or responsible adult and the selected CFC provider. If the change is made during a current prior authorization period, the new prior authorization will maintain the same end date as the original prior authorization period. The revised authorization period will begin on the start of care date stated in the new assessment.
For ongoing CFC needs beyond the initial prior authorization period of up to twelve months, a DSHS case manager must conduct a new assessment and submit a new authorization request to TMHP. A new in-home assessment must be conducted every twelve months with the client. TMHP will send a notification letter updating the prior authorization to the client, responsible adults, and the selected CFC provider. HHSC or its designee may suspend an authorization for CFC when either:

- The client or the client’s family creates an unsafe environment for the attendant’s health and safety; or
- The provider requests suspension for the reasons outlined in 40 TAC Part 1, Chapter 41.

Providers can call a toll-free Provider Inquiry Line at 1-888-648-1517 for assistance with inquiries about the status of a CFC prior authorization. Providers should direct inquiries about other Medicaid services to the TMHP Contact Center at 1-800-925-9126. CFC providers should encourage the client or responsible adult to contact the appropriate DSHS Health Service Region with inquiries or concerns about the CFC assessment.

2.12.4.1 CFC Provider Responsibilities

CFC providers must comply with all applicable federal, state, and local laws and regulations. All CFC providers must maintain written policies and procedures for obtaining consent for medical treatment in the absence of the responsible adult. The procedure and policy must meet the standards of the Texas Family Code, Chapter 32. Providers must accept clients only when there is a reasonable expectation and evidence that the client’s needs may be adequately met in the place of service (POS). The POS must be able to support the client’s health and safety needs and adequately support the use, maintenance, and cleaning of all required medical devices, equipment, and supplies. Necessary primary and backup utility, communication, and fire safety systems must be available in the POS. The CFC provider is responsible for the supervision of the CFC attendant as required by the CFC provider’s licensure requirements.

2.12.4.2 Documentation Requirements

Documentation elements are routinely assessed for compliance in retrospective review of client records, including the following:

- All entries are legible to people other than the author, dated (month, day, year, time), and signed by the author.
- Each page of the record documents the client’s name and Medicaid identification number.
- All attendants’ arrival and departure times are documented with signature and time.
- Documentation of services correlates with, and reflects medical necessity for, the services provided on any given day.
- Client’s arrival or departure from the home setting is documented with the time of arrival, departure, mode of transportation, and who accompanied the client.

2.12.5 Claims Information

TMHP processes CFC claims. CFC providers must submit claims for services in an approved electronic claims format or on the appropriate claim form based on their provider type. CFC providers, other than home health agencies, that are enrolled as PAS-only providers, FMSAs, or SRO providers must file CFC claims using a CMS-1500 paper claim form. Home health agencies, including those enrolled as an FMSA, or an SRO provider, must file PCS claims using the UB-04 CMS-1450 paper claim form.

TMHP does not supply the forms. Home health agencies and consumer-directed agencies that bill for CFC using procedure code T1019 must include the prior authorization number on claims submitted for reimbursement. Additionally, providers utilizing paper, TexMedConnect, or billing through EDI must include the prior authorization number with all claims submissions.
2.12.5.1 Managed Care Clients

CFC services are carved-out of the Medicaid Managed Care Program for State of Texas Access Reform (STAR) clients and must be billed to TMHP for payment consideration. Carved-out services are those that are rendered to Medicaid Managed Care clients but are administered by TMHP and not the client’s MCO. Claims for STAR Health and STAR+PLUS are not carved out and must be submitted to the client’s MCO for payment consideration.

2.13 Private Duty Nursing (PDN)(CCP)

Refer to: The Home Health Nursing and Private Duty Nursing Services Handbook (Vol. 2, Provider Handbooks) for information about private duty nursing (PDN) (CCP) services.

2.14 Prescribed Pediatric Extended Care Centers (PPECC) (CCP)

PPECC services may be a benefit of the Texas Health Steps (THSteps) Comprehensive Care Program (CCP) for Medicaid clients who are:

- 20 years of age and younger;
- THSteps - CCP eligible;
- Medically or technologically dependent;

Note: The term "medically dependent or technologically dependent client" does not include a minor or occasional medical condition that does not require continuous nursing care, including asthma or diabetes, or a condition that requires an epinephrine injection.

- Have an acute or chronic condition;
- Require ongoing skilled nursing care beyond the level of Skilling Nursing (SN) visits normally authorized under Texas Medicaid Home Health Skilled Nursing (HHSN) and Home Health Aide (HHA) Services;
- Meet the medical necessity criteria for admission to a PPECC detailed in the authorization and medical necessity requirements, including a prescription from the client’s ordering physician, and;
- Have chosen to receive PPECC services.

A PPECC does not provide emergency services. PPECCs must follow the safety provisions in state PPECC licensure requirements, including the adoption and enforcement of policies and procedures for a client’s medical emergency. PPECCs must call for emergency transport to the nearest hospital when emergency services are needed by a PPECC client.

2.14.1 Services, Benefits, and Limitations

PPECC services are provided in a non-residential facility licensed by HHSC. PPECCs serve four or more medically dependent or technologically dependent clients who are 20 years of age or younger and who require ongoing skilled nursing prescribed by the client’s physician to avert death or further disability or require the routine use of a medical device to compensate for a deficit in life-sustaining body function.

Services must be included in a PPECC plan of care (POC) and are limited to no more than 12 hours in a 24-hour period. PPECC services may not be provided overnight. PPECC services are intended as an alternative to private duty nursing (PDN). When the services duplicate, PPECC services must be a one-to-one replacement of private duty nursing (PDN) hours, unless additional hours are medically necessary.

PPECCs must comply with:

- Medicaid program rules, as well as PPECC licensing statute and rules;
- Mandatory reporting of suspected abuse and neglect of children;
• Texas Medicaid provider participation requirements; and
• The requirements of the TMPPM.

Clients who receive PPECC services through THSteps-CCP require ongoing medical supervision by the ordering physician who has a therapeutic relationship with and ongoing clinical knowledge of the client. A face-to-face evaluation must be performed each year by the ordering physician for each client. A physician order is required for each authorization period including initial, revisions, and recertification. A physician in a relationship with a PPECC (employed by or contracted with a PPECC) cannot provide the physician’s order, unless the physician is the client’s treating physician and has examined the client outside of the PPECC setting.

The following services may be rendered at a PPECC, but are not considered part of the PPECC services covered by Texas Medicaid, and must be billed separately by Medicaid-enrolled service providers:

• Speech, physical, and occupational therapies
• Certified respiratory care practitioner services
• Early intervention services provided through the Early Childhood Intervention (ECI) program, which are subject to ECI policies.

When the client’s plan of care indicates that therapy services are required while the client is at the PPECC, clients must be provided a choice in speech, occupational, and physical therapy providers, as well as certified respiratory care providers. PPECC providers must coordinate care with the therapy providers to ensure the client receives therapy services as required in the PPECC setting.

Clients from birth through 36 months of age must be given the option of receiving ECI services in addition to their PPECC services. The PPECC providers must coordinate care with the client’s ECI service coordinator.

ECI services rendered in a PPECC are provided by entities that are contracted with the state to provide early intervention services.

When therapy services (occupational, speech, and/or physical therapy), or certified respiratory care services are rendered in a PPECC, they may be provided by:

• Medicaid-enrolled providers contracted with or employed by the PPECC or Medicaid-enrolled providers not employed by or contracted with the PPECC.
• Independent therapists
• Home health therapists
• Certified respiratory care providers

Refer to: The Physical Therapy, Occupational Therapy, and Speech Therapy Services Handbook (Vol. 2, Provider Handbooks) for additional information about CCP therapy services.

The Certified Respiratory Care Practitioner (CRCP) Services Handbook (Vol. 2, Provider Handbooks) for information about CRCP (CCP) services.

An admission authorized under this section is not intended to supplant the right of a client to access private duty nursing (PDN), personal care services (PCS), home health skilled nursing (HHSN), home health aide (HHA), and therapies (PT, OT, ST), as well as certified respiratory care practitioner services and early childhood intervention (ECI) services rendered in the client’s residence when medically necessary.
PPECC providers must collaborate and coordinate care with the client’s existing service providers, including physicians, therapists, certified respiratory care practitioners, and home health agencies rendering services such as private duty nursing and/or home health skilled nursing, home health aide services, personal care services, hospice, and other providers who render medically necessary services. The PPECC must ensure the provision of the following basic services:

- The development, implementation, and monitoring of a comprehensive POC in collaboration with the client or the client’s responsible adult that addresses the client’s medical, nursing, psychosocial, therapeutic, and developmental services, including the following prescribed services:
  - Skilled nursing
  - Personal care services to assist with activities of daily living or instrumental activities of daily living while in the PPECC
  - Functional developmental services
  - Nutritional and dietary services, including nutritional counseling

  Note: Nutritional services must comply with standards in HHSC licensure rules related to nutritional counseling and dietary services.

- Occupational, physical and speech therapy
- Respiratory care
- Psychosocial services
- Physician’s oversight of services

- The POC must also include the following, as applicable:
  - Training for the client’s responsible adult associated with caring for a medically or technologically dependent client.
  - Transportation services needed by a client to access PPECC services.
    - Transportation must be provided by a PPECC when a client has a stated need or a prescription for transportation to the PPECC.
    - When a PPECC provides transportation to a PPECC client, Registered Nurse (RN) or Licensed Vocational Nurse (LVN) employed by the PPECC must be on board the transport vehicle.
    - The client does not need to be accompanied by the client’s responsible adult when a PPECC provides transportation.
    - When a client has a stated need or prescription for transportation, the client must be able to utilize transportation services offered by the PPECC with the assistance of a PPECC nurse to and from the PPECC, rather than a non-emergency ambulance.
    - A non-emergency ambulance may not be utilized for transport to and from a PPECC.

  Note: A client may decline a PPECC’s transportation, and choose to be transported by other means, including his or her responsible adult.

- Direct care staff, defined in licensure regulations, provides assistance with personal care services.
- PPECC services must be:
  - Individualized, specific, and consistent with symptoms or confirmed diagnosis of the condition, illness or injury under treatment, not in excess of the client’s needs;
  - Consistent with generally accepted professional medical standards as determined by the Medicaid program and may not be experimental or investigational;
• Reflective of the level of service that can be safely and effectively furnished;
• Furnished in a manner not primarily intended for the convenience of the client, the client’s responsible adult, or the provider.

Note: The fact that a client’s ordering physician has prescribed, recommended, or approved medical care, goods or services does not, in itself, make such care or services medically necessary or a covered service.

2.14.1.1 Prior Authorization and Documentation Requirements
Prior authorization is required for PPECC services, excluding PPECC transportation. All requests for PPECC services must be based on the client’s current medical needs. Texas Medicaid defines medically necessary THSteps services as health care, diagnostic services, treatments, and other measures necessary to correct or ameliorate any disability, physical or mental illness, or chronic conditions.

Documentation of medical necessity is required for PPECC services. PPECC services are considered medically necessary when a client meets all of the following admission criteria:
• Eligible for THSteps-CCP;
• 20 years of age or younger;
• Requires ongoing skilled nursing care and supervision, skillful observations, judgments and therapeutic interventions all or part of the day to correct or ameliorate health status;
• Considered to be a medically dependent or technologically dependent client in accordance with Texas Health and Safety Code chapter 248A;
• Stable for outpatient medical services, and does not present significant risk to other clients or personnel at the PPECC;
• Requires ongoing and frequent skilled interventions to maintain or ameliorate health status, and delayed skilled intervention is expected to result in:
  • Deterioration of a chronic condition;
  • Loss of function;
  • Imminent risk to health status due to medical fragility; or
  • Risk of death.
• Has a prescription for PPECC services signed and dated by an ordering physician who has personally examined the client within 30 calendar days prior to admission and reviewed all appropriate medical records;
• Has consent for the client’s admission to the PPECC signed and dated by the client or the client’s responsible adult. Admission must be voluntary and based on the preference for PPECC services in place of PDN by the client or client’s responsible adult in both managed care and non-managed care service delivery systems.
• Resides with the responsible adult and does not reside in any 24-hour inpatient facility, including the following:
  • General acute hospital
  • Skilled nursing facility
  • Intermediate care facility
  • Special care facility, including sub-acute units or facilities for the treatment of AIDS.
• The PPECC will hold interdisciplinary conferences when PPECC services are initiated, recertified, or revised, and at least every 90 calendar days. Interdisciplinary conferences should include the client’s responsible adult and the following, as applicable:

• The client’s Department of Family and Protective Services case worker.

• The client’s therapy provider(s) and

• Hospice provider.

Note: For clients who receive their PPECC services through a Medicaid managed care organization, the MCO service coordinator and/or service manager should be included in interdisciplinary conferences.

When the sole purpose of PPECC services is to train and educate the client’s responsible adult or the client (e.g., how to administer total parenteral nutrition (TPN) or how to manage a chronic condition), PPECC services will not be approved.

Training in a home setting for certain services such as how to administer TPN may be considered through intermittent home health skilled nursing visits.

Refer to: The Home Health Nursing and Private Duty Nursing Services Handbook (Vol. 2, Provider Handbooks) for more details on training and education for the client or the client’s responsible adult on TPN administration in a home setting.

2.14.1.1.1 Initial Authorization Requests

Initial requests may be prior authorized for a maximum of 90 calendar days. Requests for the prior authorization, including all required documentation, must be submitted to the Texas Medicaid Claims Administrator by electronic portal, fax, or mail no later than 3 business days following the start of care (SOC). Requests received after the 3 business day period allowed will be denied for dates of service (DOS) that occurred before the date the request is received.

When PPECC services are authorized, the authorized period begins on the day of the week that prior authorization starts. For example, if services hours are authorized on a weekly basis, the period would begin from the day of the week the prior authorization period begins and continue for 7 calendar days. PPECC services may be authorized on a daily, weekly, or hourly basis.

Consistent with PPECC licensure requirements, an initial nursing assessment must be completed, signed and dated by the PPECC Registered Nurse (RN) no earlier than 3 business days before the SOC at the PPECC. The initial nursing assessment must be performed by a PPECC RN and cannot be delegated. The initial nursing assessment is used to establish the POC and must support medical necessity for the client to receive on-going skilled nursing care. The assessment must include, but is not limited to the following:

• Complexity and intensity of the client’s care;

• Stability and predictability of the client’s condition;

• Frequency of the client’s need for skilled nursing services;

• Identified medical, nursing, psychosocial, therapeutic, nutritional, dietary, functional, educational, and developmental needs and goals, and any training needs for the client or the client’s responsible adult;

• Description of wounds, if present;

• The client’s equipment needs and whether the setting can support the health and safety needs of the client and is adequate to accommodate the use, maintenance, and cleaning of all medical devices, equipment, and supplies required by the client;

• The comprehension level of the client’s responsible adult; and
• Receptivity to training and ability level of the responsible adult.

Note: The PPECC provider may be asked to submit additional documentation to support medical necessity as defined in this section.

Initial prior authorization requests for PPECC services must include the following documentation:

• A completed CCP Prior Authorization Request form signed and dated by the ordering physician.

• A completed Prescribed Pediatric Extended Care Center (PPECC) Plan of Care (POC) form signed and dated by the ordering physician, the PPECC RN completing the POC, and client or client’s responsible adult. A PPECC may also submit the POC on their own form, but the POC must contain the elements listed in this section. A written or verbal physician approval of the POC from the ordering physician must be in place by the SOC. If the PPECC has a verbal approval of the POC at the time the prior authorization request is submitted, the dated documentation of this POC verbal approval must be submitted with the POC, followed by the physician-signed and dated POC within 14 calendar days from receipt by the Texas Medicaid Claims Administrator.

• A completed Nursing Addendum to Plan of Care for Private Duty Nursing and/or Prescribed Pediatric Extended Care Centers form signed and dated by the ordering physician, RN completing the assessment, and client or client’s responsible adult. This completed form must include:
  • Updated problem list
  • Updated rationale and summary page
  • A contingency plan
  • A 24-hour daily care flow sheet
  • Physician and client acknowledgment
  • A written or verbal order for PPECC services from the ordering physician. A physician’s order (written or verbal) must be in place by the SOC. If the PPECC has a verbal order at the time the prior authorization request is submitted, dated documentation of this verbal order must be submitted separately, or it must be included on the POC.

• Per PPECC licensure requirements, the physician order must include:
  • Client’s name, date of birth, gender, and Medicaid ID number
  • Provider name, address, phone number, TPI number, and NPI number
  • Date the client was last seen by the physician
  • Description of current medical diagnosis or condition
  • Nursing services
  • Medication administration, if applicable
  • Dietary needs, if applicable
  • Permitted activities, if applicable
  • Therapies, if applicable
  • Transportation authorization, if applicable
  • Other services, if applicable
  • Approval of the client’s admission to the PPECC

Note: For authorization purposes, a physician signature on the PPECC plan of care serves as the physician order. However, the physician order as outlined above must be maintained in the client’s medical records.
Signed and dated consent of the client or client’s responsible adult documenting his/her choice of PPECC services. The signed consent must include an acknowledgement by the client or the client’s responsible adult that he/she has been informed that their private duty nursing might be reduced as a result of accepting PPECC services. Consent to share the client’s personal health information with the client’s other providers to ensure coordination of care must also be obtained.

- A client signature on the Nursing Addendum to Plan of Care for Private Duty Nursing and/or Prescribed Pediatric Extended Care Centers form meets the client consent requirements.

- The POC must be developed by a PPECC RN, in collaboration with an interdisciplinary team, in compliance with PPECC licensure requirements. The POC, using either the Prescribed Pediatric Extended Care (PPECC) POC form or a PPECC-developed form, must include the following components:
  - The client’s name, date of birth and Medicaid number;
  - The PPECC’s name, TPI, NPI, and hours of operation, as well as address, phone, and fax numbers
  - The ordering physician’s name, phone number, TPI and NPI
  - Date the PPECC nursing assessment was completed and name, title, and credentials of the RN who completed the POC and his/her dated signature
  - Name, title and credentials of the team member who completed the POC and his/her dated signature
  - Date the client was last seen by the ordering physician
  - The requested SOC date for PPECC services
  - All pertinent diagnoses and known allergies
  - Nursing services to be provided, including amount, duration, and frequency
  - The client’s prognosis
  - The client’s mental status
  - Rehabilitation potential
  - The equipment and/or supplies required
  - Therapies (occupational, physical, speech, and respiratory care), including how those therapies are accessed, amount, duration, and frequency. Therapies provided in the PPECC, as well as outside the PPECC (e.g., school based), must be documented.
  - Other prescribed services, including amount, duration, and frequency
  - Nutritional requirements, including type, method of administration, and frequency
  - Medications, including the dose, route, frequency and any medication-related allergies if known
  - Treatments, including amount and frequency
  - Wound care orders and measurements
  - Safety measures to protect against injury
  - Functional developmental services and psychosocial services, including amount, duration and frequency
  - Name, phone number and signature of responsible adult when the client is a minor child
  - Client emergency contact name and phone number
• Confirmation that a signed contingency plan is in place in circumstances when PPECC services are not available (e.g., fire, flood, windstorm, or electrical malfunctions), and for emergencies that occur while the client is in the care of the PPECC

• List of services the client receives in the home and school settings. (e.g., ECI, therapies, School Health and Related Services [SHARS], PCS, PDN, therapies, skilled home health, case management services, hospice, and Medicaid waiver programs such as Medically Dependent Children’s Program [MDCP], Home and Community-Based Services [HCS], Deaf-Blind Multiples Disabilities [DBMD], Texas Home Living [TxHmL], and Community Living Assistance and Support Services [CLASS]).

Note: Services provided under these programs will not prevent a client from obtaining medically necessary services.

• Client-specific measurable goals, including, if receiving PDN, the goal of ensuring coordination of ongoing skilled nursing services with the PDN provider, if receiving PDN

• Responsible adult training needs

• Prior and current functional or medical limitations

• Permitted activities

• Client’s scheduled days and hours of attendance

• Confirmation of a discharge plan, including instructions for timely discharge or referral

• Emergency contact information

• Method of transportation

• Private Duty Nursing provider name, TPI, NPI, phone, address and fax number, if known

• Ordering physician signature and date of signature

The ordering physician, PPECC RN and client or client responsible adult signatures must be current. Current is defined as signed and dated within the 30 calendar day period before the SOC. To be current, the ordering physician’s dated signature must be within the 14 calendar day period following the receipt of the authorization request by the Texas Medicaid claims administrator, when services are initiated by verbal order. All the following documentation requires the ordering physician’s signature with date, the CCP Prior Authorization Request form, the Prescribed Pediatric Extended Care Center (PPECC) Plan of Care, and the Nursing Addendum to Plan of Care for Private Duty Nursing and/or Prescribed Pediatric Extended Care Centers form.

If documentation is submitted solely with the ordering physician’s verbal order, it must be resubmitted with the ordering physician’s dated signature within 14 calendar days of the receipt of the authorization request by the Texas Medicaid Claims Administrator.

If the request is not received with a dated physician signature within 14 calendar days of the receipt of the authorization request by the Texas Medicaid Claims Administrator, the prior authorization will be considered incomplete and will be denied.

When there is documentation of a verbal order, if all required documentation is not signed and dated by the ordering physician and received by the Texas MedicaidClaims Administrator within 14 calendar days of the receipt of the authorization request, claims with dates of services prior to the receipt of the signed and dated documentation will be denied.

Requests for authorizations of PPECC services should always be commensurate with the client’s medical needs.
• The length of the authorization is determined on an individual basis and is based on the goals and timelines identified by the physician, provider, and client or responsible adult. PPECC services will not be authorized for more than 90 calendar days from the SOC for an initial authorization.

  **Note:** Clients enrolled in a Medicaid managed care health plan may receive services from a PPECC. Authorization must be received from the health plan.

### 2.14.1.1.2 Revisions to the Plan of Care

The PPECC provider may request a revision to the plan of care at any time during an authorization period. Requests for changes in the service hours during a current authorization period should be submitted if there is a change in the client’s condition, or the authorized services are not commensurate with the client’s medical needs and additional authorized hours are medically necessary.

  **Note:** Schedule changes that do not affect overall authorized ongoing skilled nursing hours do not require a revision authorization request, but must be documented in the client’s medical record.

Requests for revisions must be submitted to the Texas Medicaid Claims Administrator as soon as the PPECC identifies the need for a revision. Revision requests may be submitted by electronic portal, fax, or mail.

Requests for revisions must be submitted within 3 business days of the revised SOC date. Requests received after the 3 business days will be denied for dates of service that occurred before the request is received.

When a client’s condition changes during the course of the authorization period that impacts the amount or duration of services, a reassessment performed by a PPECC RN is required. A reassessment is not necessary if there is not a change in the client’s condition.

The PPECC provider must notify the Texas Medicaid Claims Administrator and the client’s ordering physician at any time during an authorization period if the client’s condition changes, the authorized services are not commensurate with the client’s medical needs, and the client requires additional hours of ongoing skilled nursing services. Submission of a revision authorization request, with physician signatures on required documentation, meets the notification requirement.

Revisions require all the following documentation:

- A completed CCP Prior Authorization Request form signed and dated by the ordering physician.

- An updated Prescribed Pediatric Extended Care Center (PPECC) Plan of Care form signed and dated by the ordering physician, the PPECC RN completing the POC, and client or client’s responsible adult. A PPECC may also submit the POC on its own form, but the POC must contain all required elements listed under Initial Authorizations in this section. A written or verbal physician approval of the POC from the ordering physician must be in place by the revised SOC. If the PPECC has a verbal approval of the POC at the time the prior authorization request is submitted, the dated documentation of this POC verbal approval must be submitted with the POC, followed by the physician signed and dated POC within 14 calendar days from the receipt of the authorization request by the Texas Medicaid Claims Administrator.

- A completed Nursing Addendum to Plan of Care for Private Duty Nursing and/or Prescribed Pediatric Extended Care Centers form signed and dated by the ordering physician, RN completing the assessment, and client or client’s responsible adult. This completed form must include:
  - Updated problem list
  - Updated rationale and summary page
  - A contingency plan
  - A 24-hour daily care flow sheet
• Physician and client acknowledgment

• A written or verbal order for PPECC services from the ordering physician. A physician’s order (written or verbal) must be in place by the revised SOC. If the PPECC has a verbal order at the time the prior authorization request is submitted, dated documentation of this verbal order must be submitted separately or it must be included on the POC. The signed, dated order must be received within 14 calendar days of the receipt of the authorization request by the Texas Medicaid Claims Administrator.

  **Note:** For authorization purposes, a physician signature on the PPECC plan of care serves as the physician order. However, the physician order, as detailed in “Initial Authorizations,” must be maintained in the client’s medical records.

• Signed and dated consent of the client or client’s responsible adult documenting his/her choice of PPECC services. The signed consent must include an acknowledgment by the client or the client’s responsible adult that he/she has been informed that their private duty nursing might be reduced as a result of accepting PPECC services. Consent to share the client’s personal health information with the client’s other providers to ensure coordination of care must also be obtained.

  **Note:** A client signature on the Nursing Addendum to Plan of Care for Private Duty Nursing and/or Prescribed Pediatric Extended Care Centers form meets the client consent requirements.

• The ordering physician, PPECC RN, and client or client responsible adult signatures must be current. Current is defined as signed and dated within the 30 calendar day period before the SOC. To be current, the ordering physician dated signature may be submitted within the 14 calendar day period following the receipt of the authorization request by the Texas Medicaid Claims Administrator, when services are initiated by verbal order. All the following revision documentation requires the ordering physician’s dated signature: the CCP Prior Authorization Request Form, the Prescribed Pediatric Extended Care Center (PPECC) Plan of Care form, and the Nursing Addendum to Plan of Care for Private Duty Nursing and/or Prescribed Pediatric Extended Care Centers form.

• Revisions during a current authorization period must fall within that authorization period. If the revision is requested beyond the existing authorization period, the provider must request a recertification authorization and submit all required documentation for a recertification.

When there is a revision request, and documentation is submitted solely with the ordering physician’s verbal order, it must be resubmitted with the ordering physician’s signature and date within 14 calendar days of the receipt of the authorization request by the Texas Medicaid Claims Administrator. If the request is not received with a dated physician signature within 14 calendar days of the receipt of the authorization request by the Texas Medicaid Claims Administrator, the prior authorization will be considered incomplete and will be denied.

When there is documentation of a verbal order and all of the required documentation is not signed and dated by the ordering physician and received by TMHP within 14 calendar days of the receipt of the authorization request by the Texas Medicaid Claims Administrator, claims with dates of services prior to receipt of the signed and dated documentation will be denied.

2.14.1.1.3 PPECC Provider Change During an Existing Authorization Period

If a provider or client discontinues PPECC services during an existing prior authorized period and the client requests services through a new PPECC provider, the new PPECC provider must follow all of the processes and submit documentation required for an initial request, as well as the following:

A change of provider letter signed and dated by the client or the client’s responsible adult documenting the date the client ended PPECC services (effective date of the change) with the previous provider, the names of the previous and new providers, and an explanation of why providers were changed.
When the new provider submits an authorization request, including all required documentation for an initial request, it will be authorized for no more than 90 calendar days. Regardless of the number of provider changes, clients may not receive PPECC services beyond the limitations outlined in this section.

2.14.1.1.4 Recertification

A recertification is a new authorization period that may be approved for up to a maximum of 180 calendar days when the client meets medical necessity criteria. Revision requests may be submitted by electronic portal, fax, or mail. The client or the client’s responsible adult, physician, and PPECC provider must agree in writing that the recertification is appropriate each certification period.

An updated nursing assessment must be performed by the PPECC RN no more than 30 calendar days before the current authorization period expires. If there is no change in the client’s condition, the POC must document medical necessity to support continued PPECC services.

A recertification request must be submitted no more than 30 calendar days and no fewer than 7 calendar days before a current authorization period will expire. Requests received after the current authorization expires will be denied for dates of service that occurred before the date the request is received. The following documentation is required for a recertification request:

- A completed CCP Prior Authorization Request form signed and dated by the ordering physician within 30 calendar days prior to the SOC date.
- A completed Prescribed Pediatric Extended Care Center (PPECC) Plan of Care form, signed and dated by the ordering physician, the PPECC RN completing the POC, and client or client’s responsible adult within 30 calendar days prior to the SOC date. A PPECC may also submit the POC on their own form, but the POC must contain the elements listed under “Initial Authorization Request” requirements in this section.
- The PPECC provider is responsible for ensuring that the ordering physician reviews and signs the POC within 30 calendar days of the expiration of the authorization period and this documentation must be maintained in the client’s record.
- A completed Nursing Addendum to Plan of Care for Private Duty Nursing and/or Prescribed Pediatric Extended Care Centers form signed and dated by the ordering physician, RN completing the assessment, and client or client’s responsible adult within 30 calendar days prior to the SOC date. The addendum must include an updated 24-hour nursing services flow sheet and if there are changes, an updated problem list, and updated rationale summary page, a contingency plan, and a signed physician and client acknowledgment.
- A written order for PPECC services signed and dated by the client’s ordering physician. A physician’s order must be in place by the SOC.

Note: For authorization purposes, a physician signature on the PPECC plan of care serves as the physician order. However, the physician order, with elements outlined in “Initial Authorization Requests,” must be maintained in the client’s medical record.

- Signed, dated consent of the client or client’s responsible adult documenting their choice of PPECC services. The signed consent must include an acknowledgment by the client or the client’s responsible adult that he/she has been informed that other services such as private duty nursing might be reduced as a result of accepting PPECC services. Signed and dated consent to share the client’s personal health information with the client’s other providers, as needed to ensure coordination of care, must also be obtained.

Note: A client signature on the Nursing Addendum to Plan of Care for Private Duty Nursing and/or Prescribed Pediatric Extended Care Centers form meets the client consent requirements.
The provider may request a revision of a recertification at any time during the recertification period. Revisions must follow the instructions outlined under Revisions in this section. The provider must notify the claims administrator at any time during a recertification period if the client’s condition changes and the authorized services are not commensurate with the client’s medical needs.

All authorization timelines apply to recertification.

2.14.1.1.5 Termination of Authorizations

Authorization for PPECC services will be terminated when:

- The client is no longer eligible for THSteps-CCP.
- The client no longer meets the medical necessity criteria for PPECC services.
- The place of service cannot ensure the health and safety of the client.
- The client or the client’s responsible adult refuses to comply with the service plan and compliance is necessary to assure the health and safety of the client.
- The client changes providers, and the change of notification is submitted to the claims administrator in writing with a PA request from the new provider.
- After receiving PPECC services, the client opts to decline PPECC services and receive his or her services at home. The home health agency or independent provider offering ongoing skilled nursing (e.g., PDN) must submit or update all required authorization documentation to the claims administrator.

2.14.1.1.6 Appeal of Authorization Decisions

Providers may appeal denials or modifications of requested PPECC services with documentation to support the medical necessity of the requested PPECC services.

Appeals must be submitted to the Medicaid Claims Administrator’s CCP department with complete documentation and any additional information within two weeks of the date on the decision letter. If changes are made to the authorization based on this documentation, CCP claims administrators will go back no more than 3 business days for initial, or revision requests; and no more than seven calendar days for recertification requests when additional documentation is submitted.

The client or the client’s responsible adult will be notified of any denial or modification of requested services and will be given information about how to appeal the claims administrator’s decision or request a fair hearing.

PPECC services may be denied when:

- The client does not meet medical necessity criteria for admission.
- The client does not have an ordering physician.
- The client is not 20 years of age or younger.
- The client’s needs are not beyond the scope of services available through Medicaid Title XIX Home Health SN and/or HHA Services because the needs can be met on a part-time or intermittent basis through a visiting nurse.
- The services are primarily intended to provide respite care or child care.
- The services are provided for the sole purpose of responsible adult training.
- The signed and dated POC is not received by the claims administrator within fourteen business days from the SOC.
- The request is incomplete.
- The information in the request is inconsistent.
The requested services are not ongoing skilled nursing services.

There is a duplication of services.

Prior authorization requests must be submitted for processing to the Texas Medicaid Claims Administrator Prior Authorization Department (fee-for-service clients).

**Note:** Clients enrolled in a managed care health plan may receive services from a PPECC. Prior authorization requests for these clients must be submitted solely to the client’s managed care organization.

### 2.14.1.1.7 Documentation Requirements

In addition to documentation requirements outlined in the “Authorization Requirements” section, the following documentation requirements apply. Services not supported by documentation are subject to recoupment.

All services outlined in this section are subject to retrospective review to ensure that the documentation in the client’s medical record supports the medical necessity of the service(s) provided.

PPECCs must maintain documentation in the client’s medical record, including but not limited to the following:

- Evidence that the client’s condition will allow safe delivery of PPECC services as described in the POC.
- The PPECC nursing assessment.
- The client’s individualized PPECC plan of care and documentation of medical necessity.
- The physician’s specific, written, signed and dated orders for PPECC services. Documentation of verbal orders must also be maintained.
- All prior authorization request forms for Medicaid.
- The signed, dated consent of the client or the client’s responsible adult.
- The PPECC must provide documentation that the client or the client’s responsible adult has been informed about how care will be coordinated between the client’s providers (e.g., client signature on the Nursing Addendum to Plan of Care for Private Duty Nursing and/or Prescribed Pediatric Extended Care Centers).
- The PPECC must maintain evidence in the medical record that client or the client’s responsible adult has been involved in the development of the POC. (e.g., client signature on the Prescribed Pediatric Extended Care Center [PPECC] Plan of Care).
- Evidence of PDN provider notification when a child receives PDN, and the date notification was provided.
- Notes from interdisciplinary team meetings.
- Documentation of all discrepancies between the weekly service hours scheduled and the service hours provided. Examples include but are not limited to, doctor’s appointments; the PPECC was closed one day for unforeseen reasons; the child was hospitalized; or the client’s responsible adult was ill and could not provide services that he or she would normally provide.
- For each day that PPECC services are provided, the client’s medical record must identify:
  - The names of the specific person (e.g. nursing, direct care staff, therapist) providing services,
  - Date of service,
  - Type of services performed, and
  - The start and end times of services performed.
• The PPECC must be able to calculate the cost by practitioner and type of service provided as requested by HHSC.

To complete a prior authorization process by paper, the provider must complete and submit the prior authorization documentation through fax or mail and must maintain a copy of the prior authorization request and all submitted documentation in the client’s medical record at the PPECC’s place of business.

To complete a prior authorization process electronically, the provider must complete and submit the prior authorization documentation through any approved electronic method, and must maintain a copy of the prior authorization request and all submitted documentation in the client’s medical record at the PPECC’s place of business.

The ordering physician must also maintain a copy of the signed and dated physician order and signed and dated POC in the client’s medical record.

PPECC service providers must provide written notice to clients of their intent to voluntarily terminate PPECC services at least fifteen (15) calendar days prior to terminating services, except in situations of a potential threat to the provider’s personal safety.

The PPECC must sign, date, and indicate the time the client is boarded on PPECC transportation, and the time when the client arrives at the PPECC. The PPECC must also sign, date, and indicate the time when the client is boarded for a return trip from PPECC services, as well as the arrival time at the client’s destination. The PPECC provider may use any reliable method to record times, dates, and signatures provided that it is accurate and allows for an auditable review of the records, including electronic census, time-stamp, scanning, and signature records.

For any Medicaid client that is in transport for longer than one hour, the PPECC must document the reason for the extended time in transport.

A responsible adult must sign and confirm the time that the client is boarded on PPECC transportation, as well as when a client returns from the PPECC. If a responsible adult provides the transportation, the responsible adult must sign and indicate the date and time that the client is dropped off and picked up from a PPECC. The PPECC provider must keep these records in case of an audit or monitoring.

A responsible adult must be provided daily a written, one-page summary of services provided to the client for each day that the client is in the PPECC’s care.

The PPECC must maintain documentation in the client’s medical record of the notification provided to the client and/or the client’s responsible adult of an intent to transfer or discharge the client as follows:

• A copy of the written notification provided,
• Personal contact with the client and/or the client’s responsible adult, and
• The client’s ordering physician was notified of the date of transfer or discharge.

The PPECC and the therapy provider must have a written agreement for each client regarding the provision of therapy services when therapy services (occupational, speech, physical, and respiratory care) are provided at the PPECC. The written agreement must address responsibilities of both parties, and how the parties will coordinate related to the client’s plan of care. The written agreement must be kept in the client’s medical record.

The PPECC and hospice provider must have a written agreement for each client regarding the provision of hospice services when hospice is provided at the PPECC. The written agreement must address responsibilities of both parties, and how the parties will coordinate related to the client’s plan of care. The written agreement must be kept in the client’s medical record.

2.14.1.1.8 Exclusions
The services that are not covered by the PPECC benefit include the following:

• Baby food or formula.
• PPECC services to clients related to the PPECC owner by blood, marriage or adoption.

• Services that are intended to provide mainly respite care or child care and do not directly relate to the client’s medical needs or disability.

• PPECC services rendered to a client who does not meet the definition of a medically or technologically dependent minor.

• Services covered separately by Texas Medicaid, such as:
  • Speech, occupational, physical, respiratory therapy services, and early childhood intervention services.
  • Durable medical equipment (DME), medical supplies, nutritional products provided to the client by Medicaid’s DME and medical supply service providers.
  • Private duty nursing, skilled nursing and home health aid services provided in the home setting when medically needed in addition to the PPECC services authorized.
  • Services that are the legal responsibility of a local school district.
  • Individualized comprehensive case management beyond required service coordination.

2.14.1.1.9 Claims Filing and Reimbursement

PPECC services may be reimbursed when billed with procedure codes T1025, T1026, or T2002.

Services begin when the PPECC assumes responsibility for the care of the client (i.e., the point the client boards the PPECC transportation, or when the client is brought to the PPECC by a responsible adult) and ends when the care is relinquished to the client’s responsible adult.

Providers must use appropriate procedure codes for the PPECC services performed. Procedure codes T1025 and T2002 are limited to once per day.

The PPECC per diem code (T1025) and hourly procedure code (T1026) may not be billed on the same day.

Procedure code T1026 is allowed on an hourly basis, up to four hours. Services beyond four hours must be billed using T1025. At a minimum, four hours and fifteen minutes of services must be provided before T1025 may be billed.

Procedure code T2002 is not allowed without a PPECC service on the same day, same provider.

For procedure code T1026, a minimum of 15 minutes of service is required to round up to a full hour after the first hour.

Therapy services are billed separately by Medicaid-enrolled licensed therapists, including ECI providers, and are subject to prior authorization and policies governing Physical, Occupational, and Speech Therapy - Children (Acute and Chronic), or ECI services, as applicable.

If hospice services are rendered in a PPECC setting, they must be billed separately by Medicaid-enrolled hospice providers, and are subject to prior authorization and policies governing hospice reimbursement.

The following services may be billed on the same day as PPECC services, but they may not be billed simultaneously with PPECC services. These services may be billed before or after PPECC services:

• Private Duty Nursing
• Home Health Skilled Nursing
• Home Health Aide services

PCS services provided in a PPECC are considered part of the PPECC billable rate. PCS services rendered in a client’s home may be billed before or after PPECC services on the same day.
PPECC services may be reimbursed only to a licensed PPECC.

Note: Texas Medicaid will not reimburse PPECC services that duplicate services that are the legal responsibility of the school districts. The school district, through the SHARS program, is required to meet the client’s skilled nursing needs while the client is at school. However, if those needs cannot be met by SHARS or the school district, documentation supporting medical necessity may be submitted to the Texas Medicaid Claims Administrator.

Parental accompaniment is not required for PPECC reimbursement.

Non-emergency ambulance service providers will not be reimbursed for transportation to and from a PPECC.

PPECC services are subject to retrospective review and possible recoupment when the medical record does not document the provision of PPECC services is medically necessary based on the client’s situation and needs. The PPECC provider must explain all discrepancies between the service hours approved and the service hours provided. For example: The parents withdrew their client from a PPECC and released the provider from all responsibility for the service hours; the PPECC was closed one day for unforeseen reasons; the client was hospitalized; or the responsible adult was ill and could not provide services that he or she would normally provide.

Payment will not be rendered for services that are not prior authorized.

2.15 Therapy Services (CCP)

Refer to: Section 5, “Children’s Therapy Services Clients birth through 20 years of age” in the Physical Therapy, Occupational Therapy, and Speech Therapy Services Handbook (Vol. 2, Provider Handbooks) for information about CCP therapy services.

2.16 Inpatient Psychiatric Hospital or Facility (Freestanding) (CCP)

Inpatient psychiatric treatment in a nationally accredited freestanding psychiatric facility or a nationally accredited state psychiatric hospital is a benefit of Texas Medicaid for clients who are birth through 20 years of age at the time of the service request and service delivery, if the client meets certain conditions.


2.17 Inpatient Rehabilitation Facility (Freestanding) (CCP)

2.17.1 Enrollment

Note: Rehabilitation provided at an acute care facility is covered through Texas Medicaid fee-for-service.

To be eligible to participate in CCP, a freestanding inpatient rehabilitation facility must be certified by Medicare, have a valid provider agreement with HHSC, and have completed the TMHP enrollment process. Texas Medicaid enrolls and reimburses freestanding inpatient rehabilitation facilities for CCP services and Medicare deductibles or coinsurance according to current payment guidelines. The information in this section is applicable to CCP services only.

Refer to: Subsection 2.1.2, “Enrollment” in this handbook for more information about CCP enrollment procedures.
2.17.1.1 Continuity of Hospital Eligibility Through Change of Ownership

Under procedures set forth by the CMS and HHSC, a change in ownership of a hospital does not terminate Medicare eligibility; therefore, Medicaid participation may be continued subject to the following requirements:

- The provider must obtain recertification as a Title XVIII (Medicare) hospital.
- The hospital under new ownership must submit a new signed and dated HHSC Medicaid Provider Agreement between the hospital and HHSC.

Providers can download the HHSC Medicaid Provider Agreement from the TMHP website at www.tmhp.com.

2.17.2 Services, Benefits, and Limitations

Inpatient rehabilitation services include medically necessary items and services ordinarily furnished by a Medicaid hospital or by an approved out-of-state hospital under the direction of a physician for the care and treatment of inpatient clients. Inpatient rehabilitation services will be considered for an acute problem or an acute exacerbation of a chronic problem resulting in a significant decrease in functional ability that will benefit from inpatient rehabilitation services. A condition is considered to be acute or an acute exacerbation of a chronic condition only during the six months from the onset date of the acute condition or the acute exacerbation of the chronic condition.

When a client is admitted to an inpatient facility for acute care physical, occupational, or speech therapy services, the therapy services are reimbursed as part of the inpatient hospital reimbursement methodology (Diagnosis-Related Group [DRG] or Tax Equity and Fiscal Responsibility Act [TEFRA]) and not reimbursed separately to the individual therapist. The hospital must include the physician’s written treatment plan that supports the medical necessity of the hospitalization and services.

2.17.2.1 Comprehensive Treatment

The intensity of necessary rehabilitative service cannot be provided in the outpatient setting.

Comprehensive rehabilitation treatment must be under the leadership of a physician. Comprehensive rehabilitation treatment must be an active interdisciplinary team, defined as at least two types of therapies.

Comprehensive treatment must consist of at least two appropriate physical modalities designed to resolve or improve the client’s condition (OT, PT, and ST), and must be provided for a minimum of three hours per day for five days per week.

2.17.3 Prior Authorization and Documentation Requirements

All inpatient rehabilitation services provided to clients who are birth through 20 years of age in a freestanding inpatient rehabilitation facility require prior authorization.

Prior authorization will be considered when the client has met all of the following criteria:

- The client has an acute problem or an acute exacerbation of a chronic problem resulting in a significant decrease in functional ability that will benefit from inpatient rehabilitation services.
- The intensity of necessary rehabilitative service cannot be provided in the outpatient setting.
- The client requires and will receive multidisciplinary team care defined as at least two therapies (OT, PT, or ST).
- This therapy will be provided for a minimum of three hours per day, five days per week.

The physician and the provider must maintain all documentation in the client’s medical record.
Inpatient rehabilitation may be prior authorized for up to two months when the attending physician submits documentation of medical necessity. The treatment plan must indicate that the client is expected to improve within a 60-day period and be restored to a more functional lifestyle for an acute condition or the previous level of function for an acute exacerbation of a chronic condition.

Requests for subsequent services for increments up to 60 days may be prior authorized based on medical necessity. Requests for prior authorization of subsequent services must be received before the end-date of the preceding prior authorization.

A prior authorization request for an additional 60 days of therapy will be considered with documentation supporting medical necessity.

Supporting documentation for an initial request must include the following:

- The request for inpatient rehabilitation and the treatment plan must be signed and dated by the physician. The physician’s signature is valid for no more than 60 days prior to the requested start of care date.
- A CCP Prior Authorization Request Form signed and dated by the physician.
- A current therapy evaluation with the documented age of the client at the time of evaluation.
- Therapy goals related to the client’s individual needs; goals may include improving or maintaining function, or slowing of deterioration of function.
- An updated written comprehensive treatment plan established by the attending physician or by the therapist to be followed during the inpatient rehabilitation admission that:
  - Is under the leadership of a physician and includes a description of the specific therapy being prescribed, diagnosis, treatment goals related to the client’s individual needs, and duration and frequency of therapy.
  - Includes the date of onset of the illness or injury requiring the freestanding inpatient rehabilitation facility admission.
  - Includes the requested dates of service.
  - Incorporates an active interdisciplinary team.
  - Consists of at least two appropriate physical modalities (OT, PT, and ST) designed to resolve or improve the client’s condition.
  - Includes a minimum of three hours of team interaction with the client every day, five days per week.
- In addition to the documentation for an initial request, supporting documentation for a request for subsequent services must include the following:
  - A brief synopsis of the outcomes of the previous treatment relative to the debilitating condition.
  - The expected results to be achieved by an extension of the active treatment plan, and the time interval at which this extension outcome should be achieved.
  - Discussion why the initial two months of inpatient rehabilitation has not met the client’s needs and why the client cannot be treated in an outpatient setting.

After receiving the documentation establishing the medical necessity and plan of medical care by the treating physician, prior authorization is considered by CCP for the initial service and an extension of service as applicable. A request for prior authorization must include documentation from the provider to support the medical necessity of the service.
2.17.4 Claims Information

Providers must submit inpatient rehabilitation services to TMHP in an approved electronic claims format or on a UB-04 CMS-1450 paper claim form. Providers must purchase the UB-04 CMS-1450 paper claim forms from the vendor of their choice. TMHP does not supply the forms.

For OT, PT, and ST services, freestanding inpatient rehabilitation facilities and acute care hospitals can use revenue codes 128, 420, 424, 430, 434, 440, and 444.

TMHP must receive claims for payment consideration according to filing deadlines for inpatient claims. Claims for services that have been prior authorized must reflect the PAN in Block 63 of the UB-04 CMS-1450 paper claim form or its electronic equivalent.

Refer to: “Section 3: TMHP Electronic Data Interchange (EDI)” (Vol. 1, General Information) for information on electronic claims submissions.

“Section 6: Claims Filing” (Vol. 1, General Information) for general information about claims filing.

Subsection 6.6, “UB-04 CMS-1450 Paper Claim Filing Instructions” in “Section 6: Claims Filing” (Vol. 1, General Information) for paper claims completion instructions.

Inpatient Rehabilitation Facility (Freestanding) (CCP Only) on the TMHP website at www.tmhp.com for a claim form example.

2.17.5 Reimbursement

Reimbursement for care provided in the freestanding inpatient rehabilitation facility is made under the Texas Diagnosis-Related Group (DRG) Payment System.

Refer to: Subsection 3.7.4.4.1, “Day Outliers” in the Inpatient and Outpatient Hospital Services Handbook (Vol. 2, Provider Handbooks) for reimbursement information.

2.17.5.1 Client Transfers

When more than one hospital provides care for the same case, the hospital furnishing the most significant amount of care receives consideration for a full DRG payment.

The other hospital(s) is/are paid a per diem rate based on the lesser of the mean length of stay for the DRG or eligible days in the facility. The DRG modifier PT on the R&S Report indicates per diem pricing related to a client transfer.

Client transfers within the same facility are considered one continuous stay and receive only one DRG payment. The facility must bill only one claim.

After all hospital claims have been submitted, HHSC performs a post-payment review to determine whether the hospital furnishing the most significant amount of care received the full DRG. If the review reveals that the hospital furnishing the most significant amount of care did not receive the full DRG, an adjustment is initiated.
3 School Health and Related Services (SHARS)

3.1 Overview

Medicaid services provided by school districts in Texas to Medicaid-eligible students are known as SHARS. The oversight of SHARS is a cooperative effort between the Texas Education Agency (TEA) and HHSC. SHARS allows local school districts, including public charter schools, to obtain Medicaid reimbursement for certain health-related services provided to students in special education under IDEA that are documented in a student’s Individualized Education Program (IEP).

**Important:** CMS requires school districts to be enrolled as a SHARS Medicaid provider, participate in the Random Moment Time Study (RMTS), claim on an interim basis, and submit an annual SHARS Cost Report.

SHARS reimbursement is provided for students who meet all of the following requirements:

- Are 20 years of age and younger and eligible for Medicaid
- Meet eligibility requirements for special education described in IDEA
- Have IEPs that prescribe the needed services

Services covered by SHARS includes:

- Audiology services
- Counseling
- Nursing services
- Occupational therapy (OT)
- Personal care services (PCS)
- Physical therapy (PT)
- Physician services
- Psychological services, including assessments
- Speech therapy (ST)
- Transportation in a school setting

These services must be provided by qualified personnel who are under contract with or employed by the school district.

3.1.1 Random Moment Time Study (RMTS)

CMS requires SHARS providers to participate in the RMTS to be eligible to submit claims and receive reimbursement for SHARS services. SHARS providers must comply with the Texas Time Study Guide, which includes, but is not limited to, Mandatory Annual RMTS Contact training certification of RMTS participants for all three annual RMTS quarters, and compliance with participation requirements for selected sampled moments. The three annual RMTS quarters are October through December, January through March, and April through June. A July through September RMTS is not conducted.

An existing school district can only become a SHARS provider effective October 1, each year and they must participate in all three RMTS quarters for that annual period. SHARS providers that do not participate in all three required RMTS quarters, or are RMTS non-compliant, cannot be a SHARS provider for that entire annual period (October 1 through September 30) and will be required to return any Medicaid payments received for SHARS services delivered during that annual cost report period. The school district can return to participating in the SHARS program the following federal fiscal year beginning on October 1.
A new school district (i.e., a newly formed district that began operations after October 1) can become a SHARS provider effective with the first day of the federal quarter in which it participates in the RMTS. New SHARS providers may not submit claims or be reimbursed for SHARS services provided prior to the RMTS quarter in which they begin to participate and they must participate in all remaining RMTS quarters for that annual period.

School districts can access the Texas Time Study Guide, on the HHSC website at https://rad.hhs.texas.gov/time-study/time-study-independent-school-districts-isd and refer to the link titled Guides/Manuals.

SHARS providers can contact the HHSC Time Study Unit by email at TimeStudy@hhsc.state.tx.us or by telephone at 1-512-491-1715.

### 3.1.2 Eligibility Verification

The following are means to verify Medicaid eligibility of students:

- Verify electronically through third party software or TexMedConnect.
- School districts may inquire about the eligibility of a student by submitting the student’s Medicaid number or two of the following: name, date of birth, or Social Security number (SSN). A search can be narrowed further by entering the county code or sex of the student. Verifications may be submitted in batches without limitations on the number of students.
- Contact AIS at 1-800-925-9126.

### 3.2 Enrollment

#### 3.2.1 SHARS Enrollment

To enroll in Texas Medicaid as a SHARS provider, school districts, including public charter schools, must employ or contract with individuals or entities that meet certification and licensing requirements in accordance with the Texas Medicaid State Plan for SHARS to provide program services. Since public school districts are government entities, they should select “public entity” on the enrollment application.

SHARS providers are required to notify parents or guardians of their rights to a “freedom of choice of providers” (42 CFR §431.51) under Texas Medicaid. Most SHARS providers currently provide this notification during the initial Admission, Review, and Dismissal (ARD) process. If a parent requests that someone other than the employees or currently contracted staff of the SHARS provider (school district) provide a required service listed in the student’s IEP, the SHARS provider must make a good faith effort to comply with the parent’s request. The SHARS provider can negotiate with the requested provider to provide the services under contract. The requested provider must meet, comply with, and provide all of the employment criteria and documentation that the SHARS provider normally requires of its employees and currently contracted staff. The SHARS provider can negotiate the contracted fee with the requested provider and is not required to pay the same fee that the requested provider might receive from Medicaid for similar services.

**Refer to:** Subsection 2.2, “Fee-for-Service Reimbursement Methodology” in “Section 2: Texas Medicaid Fee-for-Service Reimbursement” (Vol. 1, General Information) for more information.

#### 3.2.2 Private School Enrollment

A private school may not participate in the SHARS program as a SHARS provider.
3.3 Services, Benefits, Limitations, and Prior Authorization

All of the SHARS procedures listed in the following sections require a valid diagnosis code. SHARS includes audiology services, counseling, physician services, nursing services, psychological services, OT, PT, or ST services, personal care services, and transportation.

Reminder: SHARS are the services determined by the ARD committee to be medically necessary and reasonable to ensure that children with disabilities who are eligible for Medicaid and who are 20 years of age and younger receive the benefits accorded to them by federal and state law in order to participate in the educational program.

3.3.1 Audiology

Audiology evaluation services include:

- Identification of children with hearing loss
- Determination of the range, nature, and degree of hearing loss, including the referral for medical or other professional attention for the habilitation of hearing
- Determination of the child’s need for group and individual amplification

Audiology therapy services include the provision of habilitation activities, such as language habilitation, auditory training, audiological maintenance, speech reading (lip reading), and speech conversation.

Audiology services must be provided by a professional who holds a valid state license as an audiologist or by an audiology assistant who is licensed by the state when the assistant is acting under the supervision of a qualified audiologist. State licensure requirements are equal to American Speech-Language-Hearing Association (ASHA) certification requirements.

Audiology evaluation is billable on an individual (procedure code 92620) basis only. Audiology evaluation (procedure code 92620) is limited to a combined maximum total of twelve units in a 30-day period.

Audiology therapy is billable on an individual (procedure code 92507) and group (procedure code 92508) basis.

Only the time spent with the student present is billable; time spent without the student present is not billable.

Session notes for evaluations are not required; however, documentation must include the billable start time, billable stop time, and total billable minutes with a notation of the activity performed (e.g., audiology evaluation).

Session notes are required for therapy. Session notes must include the billable start time, billable stop time, total billable minutes, activity performed during the session, student observation, and the related IEP objective.

3.3.1.1 Audiology Billing Table

<table>
<thead>
<tr>
<th>POS*</th>
<th>Procedure Code</th>
<th>Individual or Group</th>
<th>Therapist or Assistant</th>
</tr>
</thead>
<tbody>
<tr>
<td>1, 2, or 9</td>
<td>92507 with modifier U9</td>
<td>Individual</td>
<td>Licensed audiologist</td>
</tr>
<tr>
<td>1, 2, or 9</td>
<td>92507 with modifier U1</td>
<td>Individual</td>
<td>Licensed assistant</td>
</tr>
<tr>
<td>1, 2, or 9</td>
<td>92508 with modifier U9</td>
<td>Group</td>
<td>Licensed audiologist</td>
</tr>
<tr>
<td>1, 2, or 9</td>
<td>92508 with modifier U1</td>
<td>Group</td>
<td>Licensed assistant</td>
</tr>
<tr>
<td>1, 2, or 9</td>
<td>92620</td>
<td>Individual</td>
<td>Licensed audiologist</td>
</tr>
</tbody>
</table>

*Place of Service: 1=office; 2=home; 9=other locations

Providers must use a 15-minute unit of service for billing.

Refer to: Subsection 3.5.1.2, “Billing Units Based on 15 Minutes” in this handbook.
The recommended maximum billable time for audiology evaluation is three hours, which may be billed over several days. The recommended maximum billable time for direct audiology therapy (individual or group) is one hour per day. Providers must submit documentation of the reasons for the additional time, if more than the recommended maximum time is billed.

### 3.3.2 Counseling Services

Counseling services are provided to help a child with a disability benefit from special education and must be listed in the IEP. Counseling services include, but are not limited to, the following:

- Assisting the child or parents in understanding the nature of the child’s disability
- Assisting the child or parents in understanding the special needs of the child
- Assisting the child or parents in understanding the child’s development
- Health and behavior interventions to identify the psychological, behavioral, emotional, cognitive, and social factors that are important to the prevention, treatment, or management of physical health problems
- Assessing the need for specific counseling services

Counseling services must be provided by a professional who has one of the following certifications or licensures: a licensed professional counselor (LPC), a licensed clinical social worker (LCSW), or a licensed marriage and family therapist (LMFT).

Counseling services are billable on an individual (procedure codes 96158 and 96159) or group (procedure codes 96164 and 96165) basis. Session notes are required and documentation must include the billable start time, billable stop time, total billable minutes, activity performed during the session, student observation, and the related IEP objective.

School districts may receive reimbursement for emergency counseling services as long as the student’s IEP includes a behavior improvement plan that documents the need for emergency services.

#### 3.3.2.1 Counseling Services Billing Table

<table>
<thead>
<tr>
<th>Procedure Code</th>
<th>Individual or Group</th>
<th>Unit of Service</th>
<th>POS*</th>
</tr>
</thead>
<tbody>
<tr>
<td>96158 with modifier UB</td>
<td>Individual</td>
<td>30 minutes</td>
<td>1, 2, or 9</td>
</tr>
<tr>
<td>96159 with modifier UB (add-on procedure code)</td>
<td>Individual</td>
<td>15 minutes</td>
<td>1, 2, or 9</td>
</tr>
<tr>
<td>96164 with modifier UB</td>
<td>Group</td>
<td>30 minutes</td>
<td>1, 2, or 9</td>
</tr>
<tr>
<td>96165 with modifier UB (add-on procedure code)</td>
<td>Group</td>
<td>15 minutes</td>
<td>1, 2, or 9</td>
</tr>
</tbody>
</table>

*Place of Service: 1 = Office; 2 = Home; 9 = Other Locations

Services are limited to a total of three units (one hour) per day, any provider, as follows:

- The initial 30-minute unit (procedure codes 96158 and 96164) is limited to one unit per day.
- Each additional 15-minute unit (procedure codes 96159 and 96165) is limited to two units per day.

The recommended maximum billable time (individual or group) is one hour per day. Providers must submit documentation of the reasons for the additional time, if more than the recommended maximum time is billed.
3.3.3 Psychological Testing and Services

3.3.3.1 Psychological Testing

Evaluations or assessments include activities related to the evaluation of the functioning of a student for the purpose of determining eligibility, the needs for specific SHARS services, and the development or revision of IEP goals and objectives. An evaluation or assessment is billable if it leads to the creation of an IEP for a student with disabilities who is eligible for Medicaid and who is 20 years of age or younger, whether or not the IEP includes SHARS.

Evaluations or assessments (procedure codes 96130 and 96131) must be provided by a professional who is a licensed specialist in school psychology (LSSP), a licensed psychologist, or a licensed psychiatrist in accordance with 19 TAC §89.1040(b)(1) and 42 CFR §440.60(a).

Evaluation or assessment billable time includes the following:

- Psychological, educational, or intellectual testing time spent with the student present
- Necessary observation of the student associated with testing
- A parent/teacher consultation with the student present that is required during the assessment because a student is unable to communicate or perform certain activities
- Time spent without the student present for the interpretation of testing results
- Report writing

Time spent gathering information without the student present or observing a student is not billable evaluation or assessment time.

Session notes are not required; however, documentation must include the billable start time, billable stop time, total billable minutes, and must note which assessment activity was performed (e.g., testing, interpretation, or report writing).

3.3.3.1.1 Evaluation or Assessment Billing Table

<table>
<thead>
<tr>
<th>POS*</th>
<th>Procedure Code</th>
<th>Unit of Service</th>
</tr>
</thead>
<tbody>
<tr>
<td>1, 2, or 9</td>
<td>96130</td>
<td>Initial (1 hour)</td>
</tr>
<tr>
<td>1, 2, or 9</td>
<td>96131</td>
<td>Each additional hour</td>
</tr>
</tbody>
</table>

*Place of Service: 1=office; 2=home; 9=other locations

Important: One unit (1.0) is equivalent to one hour or 60 minutes. Providers may bill in partial hours, expressed as 1/10th of an hour (six-minute segments). For example, express 30 minutes as a billed quantity of 0.5.

Refer to: Subsection 3.5.1.2, “Billing Units Based on 15 Minutes” in this handbook.

When billing, minutes of Evaluations or Assessments are not accumulated over multiple days. Minutes of Evaluations or Assessments can only be billed per calendar day.

The recommended maximum billable time for psychological testing is eight hours (8.0 units) over a 30-day period. Time spent for the interpretation of testing results without the student present is billable time. Providers must submit documentation of the reasons for the additional time, if more than the recommended maximum time is billed.

3.3.3.2 Psychological Services

Psychological services are counseling services provided to help a child with a disability benefit from special education and must be listed in the IEP.
Psychological services must be provided by a licensed psychiatrist, a licensed psychologist, or an LSSP. Nothing in this rule prohibits public schools from contracting with licensed psychologists, licensed psychological associates, and provisionally licensed psychologists who are not LSSPs to provide psychological services, other than school psychology, in their areas of competency. School districts may contract for specific types of psychological services, such as clinical psychology, counseling psychology, neuropsychology, and family therapy, that are not readily available from the LSSP who is employed by the school district. Such contracting must be on a short-term or part-time basis and cannot involve the broad range of school psychological services listed in 22 TAC §465.38(1)(B).

All psychological services are billable on an individual (procedure codes 96158 and 96159) or group (procedure codes 96164 and 96165) basis.

Session notes are required. Session notes must include the billable start time, billable stop time, total billable minutes, activity performed during the session, student observation, and the related IEP objective.

School districts may receive reimbursement for emergency psychological services as long as the student’s IEP includes a behavior improvement plan that documents the need for the emergency services.

### 3.3.3.2.1 Psychological Services Billing Table

<table>
<thead>
<tr>
<th>Procedure Code</th>
<th>Individual or Group</th>
<th>Time Limitations</th>
<th>POS*</th>
</tr>
</thead>
<tbody>
<tr>
<td>96158 with modifier AH</td>
<td>Individual</td>
<td>30 minutes</td>
<td>1, 2, or 9</td>
</tr>
<tr>
<td>96159 with modifier AH</td>
<td>Individual</td>
<td>15 minutes</td>
<td>1, 2, or 9</td>
</tr>
<tr>
<td>96164 with modifier AH</td>
<td>Group</td>
<td>30 minutes</td>
<td>1, 2, or 9</td>
</tr>
<tr>
<td>96165 with modifier AH</td>
<td>Group</td>
<td>15 minutes</td>
<td>1, 2, or 9</td>
</tr>
</tbody>
</table>

*Place of Service: 1 = Office; 2 = Home; 9 = Other Locations

Services are limited to a total of three units (one hour) per day, any provider, as follows:

- The initial 30-minute unit (procedure codes 96158 and 96164) is limited to one unit per day.
- Each additional 15-minute unit (procedure codes 96159 and 96165) is limited to two units per day.

The recommended maximum billable time for direct psychological therapy (individual or group) is a total of one hour per day for nonemergency situations. Providers must maintain documentation of the reasons for the additional time, if more than the recommended maximum time is billed.

### 3.3.4 Nursing Services

Nursing services are SN tasks, as defined by the Texas BON, that are included in the student’s IEP. Nursing services may be direct nursing care or medication administration. Examples of reimbursable nursing services include, but are not limited to, the following:

- Inhalation therapy
- Ventilator monitoring
- Nonroutine medication administration
- Tracheostomy care
- Gastrostomy care
- Ileostomy care
- Catheterization
- Tube feeding
• Suctioning
• Client training
• Assessment of a student’s nursing and personal care services needs

Direct nursing care services are billed in 15-minute increments and medication administration is reimbursed on a per-visit increment. The RN or APRN determines whether these services must be billed as direct nursing care or medication administration.

Nursing services must be provided by an RN, an APRN (including NPs and CNSs), LVN, LPN, or a school health aide or other trained, unlicensed assistive person delegated by an RN or APRN.

Nursing services are billable on an individual or group basis. Only the time spent with the student present is billable. Time spent without the student present is not billable. Session notes are not required for nursing services; however, documentation must include the billable start time, billable stop time, total billable minutes, and must note the type of nursing service that was performed.

### 3.3.4.1 Nursing Services Billing Table

<table>
<thead>
<tr>
<th>POS*</th>
<th>Procedure Code</th>
<th>Individual or Group</th>
<th>Unit of Service</th>
</tr>
</thead>
<tbody>
<tr>
<td>1, 2, or 9</td>
<td>T1002 with modifier TD</td>
<td>Individual</td>
<td>15 minutes</td>
</tr>
<tr>
<td>1, 2, or 9</td>
<td>T1002 with modifier TD and UD</td>
<td>Group</td>
<td>15 minutes</td>
</tr>
<tr>
<td>1, 2, or 9</td>
<td>T1502 with modifier TD</td>
<td>Medication administration, per visit</td>
<td></td>
</tr>
<tr>
<td>1, 2, or 9</td>
<td>T1002 with modifier U7</td>
<td>Delegation, Individual</td>
<td>15 minutes</td>
</tr>
<tr>
<td>1, 2, or 9</td>
<td>T1002 with modifier U7 and UD</td>
<td>Delegation, group</td>
<td>15 minutes</td>
</tr>
<tr>
<td>1, 2, or 9</td>
<td>T1502 with modifier U7</td>
<td>Delegation, medication administration, per visit</td>
<td></td>
</tr>
<tr>
<td>1, 2, or 9</td>
<td>T1003 with modifier TE</td>
<td>Individual</td>
<td>15 minutes</td>
</tr>
<tr>
<td>1, 2, or 9</td>
<td>T1003 with modifier TE and UD</td>
<td>Group</td>
<td>15 minutes</td>
</tr>
<tr>
<td>1, 2, or 9</td>
<td>T1502 with modifier TE</td>
<td>Medication, administration per visit</td>
<td></td>
</tr>
</tbody>
</table>

*Place of Service: 1=office; 2=home; 9=other locations
Modifiers:
- TD = nursing services provided by an RN or APRN
- U7 = nursing services delivered through delegation
- TE = nursing services delivered by an LVN/LPN
- UD = nursing services delivered on a group basis

The Medicaid-allowable fee is determined based on 15-minute increments. Providers must use a 15-minute unit of service for billing.

All of the nursing services minutes that are delivered to a student during a calendar day must be added together before they are converted to units of service. Do not convert minutes of nursing services separately for each nursing task that was performed.

Minutes of nursing services cannot be accumulated over multiple days. Minutes of nursing services can only be billed per calendar day. If the total number of minutes of nursing services is less than eight minutes for a calendar day, then no unit of service can be billed for that day, and that day’s minutes cannot be added to minutes of nursing services from any previous or subsequent days for billing purposes.
**Refer to:** Subsection 3.5.1.2, “Billing Units Based on 15 Minutes” in this handbook.

The recommended maximum billable time for direct nursing services is four hours per day. The recommended maximum billable units for procedure code T1502 with modifier TD, T1502 with modifier U7, or T1502 with modifier TE is a total of four medication administration visits per day. Providers must submit documentation of the reasons for the additional time, if more than the recommended maximum time is billed.

### 3.3.5 Occupational Therapy (OT)

#### 3.3.5.1 Referral

In order for a student to receive OT through SHARS, the name and complete address or the provider identifier of the licensed physician who prescribed the OT must be provided.

#### 3.3.5.2 Description of Services

OT evaluation services include determining what services, assistive technology, and environmental modifications a student requires for participation in the special education program.

OT includes:

- Improving, developing, maintaining, or restoring functions impaired or lost through illness, injury, or deprivation.
- Improving the ability to perform tasks for independent functioning when functions are impaired or lost.
- Preventing, through early intervention, initial or further impairment or loss of function.

OT must be provided by a professional who is licensed by the Texas Board of Occupational Therapy Examiners or an occupational therapy assistant (OTA) acting under the supervision of a qualified occupational therapist.

OT evaluation is billable on an individual (procedure code 97165, 97166, or 97167) basis only. Procedure codes 97165, 97166, and 97167 may be submitted for initial evaluations and reevaluations. OT is billable on an individual (procedure code 97530) or group (procedure code 97150) basis.

If an evaluation is performed over several days, the provider must submit the same evaluation procedure code for each evaluation session. The procedure code submitted must reflect the complexity level of the entire evaluation.

The therapist who performs the evaluation should use professional clinical judgment to decide which evaluation code to use. The selection of low (procedure code 97165), moderate (procedure code 97166), or high complexity (procedure code 97167) evaluation codes must be based on professional clinical judgment and may not be made by staff other than the rendering therapist.

The occupational therapist or COTA can only bill for time spent with the student present, including time spent assisting the student with learning to use adaptive equipment and assistive technology.

Time spent without the student present, such as training teachers or aides to work with the student (unless the student is present during the training time), report writing, and time spent manipulating or modifying the adaptive equipment is not billable.

Session notes are not required for procedure codes 97165, 97166, and 97167; however, documentation must include the billable start time, billable stop time, total billable minutes, and must note the activity that was performed (e.g., OT evaluation).

Session notes are required for procedure codes 97530 and 97150. Session notes must include the billable start time, billable stop time, total billable minutes, activity performed during the session, student observation, and the related IEP objective.
3.3.5.3 Occupational Therapy Billing Table

<table>
<thead>
<tr>
<th>POS*</th>
<th>Procedure Code</th>
<th>Individual or Group</th>
<th>Therapist or Assistant</th>
</tr>
</thead>
<tbody>
<tr>
<td>1, 2, or 9</td>
<td>97165, 97166, and 97167</td>
<td>Individual</td>
<td>Licensed therapist</td>
</tr>
<tr>
<td>1, 2, or 9</td>
<td>97150 with modifier GO</td>
<td>Group</td>
<td>Licensed therapist</td>
</tr>
<tr>
<td>1, 2, or 9</td>
<td>97150 with modifier GO and U1</td>
<td>Group</td>
<td>Licensed therapy assistant</td>
</tr>
<tr>
<td>1, 2, or 9</td>
<td>97530 with modifier GO</td>
<td>Individual</td>
<td>Licensed therapist</td>
</tr>
<tr>
<td>1, 2, or 9</td>
<td>97530 with modifier GO and U1</td>
<td>Individual</td>
<td>Licensed therapy assistant</td>
</tr>
</tbody>
</table>

*Place of Service: 1=office; 2=home; 9=other locations

Providers must use a 15-minute unit of service for billing.

Refer to: Subsection 3.5.1.2, “Billing Units Based on 15 Minutes” in this handbook.

The recommended maximum billable time for OT evaluation is three hours, which may be billed over several days within a 30 day period. The recommended maximum billable time for direct therapy (individual or group) is a total of one hour per day. Providers must submit documentation of the reasons for the additional time, if more than the recommended maximum time is billed.

3.3.6 Personal Care Services

Personal care services are provided to help a child with a disability or chronic condition benefit from special education. Personal care services include a range of human assistance provided to persons with disabilities or chronic conditions which enables them to accomplish tasks that they would normally do for themselves if they did not have a disability. An individual may be physically capable of performing ADLs and IADLs but may have limitations in performing these activities because of a functional, cognitive, or behavioral impairment.

Refer to: Subsection 2.11, “Personal Care Services (PCS) (CCP)” in this handbook for a list of ADLs and IADLs.

For personal care services to be billable, they must be listed in the student’s IEP. Personal care services are billable on an individual (procedure code T1019 with modifier U5 or U6) or group (procedure code T1019 with modifier U5 and UD or U6 and UD) basis.

Session notes are not required for procedure codes T1019 with modifier U5 or T1019 with modifier U5 and UD; however, documentation must include the billable start time, billable stop time, total billable minutes, and must note the type of personal care service that was performed.

Procedure codes T1019 with modifier U6 and T1019 with modifier U6 and UD are billed using a one-way trip unit of service.

3.3.6.1 Personal Care Services Billing Table

<table>
<thead>
<tr>
<th>POS*</th>
<th>Procedure Code</th>
<th>Individual or Group</th>
<th>Unit of Service</th>
</tr>
</thead>
<tbody>
<tr>
<td>1, 2, or 9</td>
<td>T1019 with modifier U5</td>
<td>Individual, school</td>
<td>15 minutes</td>
</tr>
<tr>
<td>1, 2, or 9</td>
<td>T1019 with modifier U5 and UD</td>
<td>Group, school</td>
<td>15 minutes</td>
</tr>
<tr>
<td>1, 2, or 9</td>
<td>T1019 with modifier U6 and UD</td>
<td>Individual, bus</td>
<td>Per one-way trip</td>
</tr>
<tr>
<td>1, 2, or 9</td>
<td>T1019 with modifier U6 and UD</td>
<td>Group, bus</td>
<td>Per one-way trip</td>
</tr>
</tbody>
</table>

*Place of Service: 1=office; 2=home; 9=other locations

Refer to: Subsection 3.5.1.2, “Billing Units Based on 15 Minutes” in this handbook.
The recommended maximum billable units for T1019 with modifier U6 or T1019 with modifier U6 and UD is a total of four one-way trips per day. Providers must submit documentation of the reasons for the additional time, if more than the recommended units of service are billed.

3.3.7 Physical Therapy (PT)

3.3.7.1 Referral

In order for a student to receive PT through SHARS, the name and complete address or the provider identifier of the licensed physician who prescribes the PT must be provided.

3.3.7.2 Description of Services

PT evaluation includes evaluating the student’s ability to move throughout the school and to participate in classroom activities and the identification of movement dysfunction and related functional problems.

PT is provided for the purpose of preventing or alleviating movement dysfunction and related functional problems.

PT must be provided by a professional who is licensed by the Texas Board of Physical Therapy Examiners or a licensed physical therapist assistant (LPTA) acting under the supervision of a qualified physical therapist.

PT evaluation is billable on an individual (procedure code 97161, 97162, or 97163) basis only. Procedure codes 97161, 97162, and 97163 may be submitted for initial evaluations and reevaluations. PT is billable on an individual (procedure code 97110) or group (procedure code 97150) basis.

If an evaluation is performed over several days, the provider must submit the same evaluation procedure code for each evaluation session. The procedure code submitted must reflect the complexity level of the entire evaluation.

The therapist who performs the evaluation should use professional clinical judgment to decide which evaluation code to use. The selection of low (procedure code 97161), moderate (procedure code 97162), or high complexity (procedure code 97163) evaluation codes must be based on professional clinical judgment and may not be made by staff other than the rendering therapist.

The physical therapist can only bill time spent with the student present, including time spent helping the student to use adaptive equipment and assistive technology.

Time spent without the student present, such as training teachers or aides to work with the student (unless the student is present during the training time) and report writing, is not billable.

Session notes are not required for procedure codes 97161, 97162, and 97163; however, documentation must include the billable start time, billable stop time, total billable minutes, and must note the activity that was performed (e.g., PT evaluation). Session notes are required for procedure codes 97110 and 97150.

Session notes must include the billable start time, billable stop time, total billable minutes, activity performed during the session, student observation, and the related IEP objective.

3.3.7.3 Physical Therapy Billing Table

<table>
<thead>
<tr>
<th>POS*</th>
<th>Procedure Code</th>
<th>Individual or Group</th>
<th>Therapist or Assistant</th>
</tr>
</thead>
<tbody>
<tr>
<td>1, 2, or 9</td>
<td>97161, 97162, and 97163</td>
<td>Individual</td>
<td>Licensed therapist</td>
</tr>
<tr>
<td>1, 2, or 9</td>
<td>97110 with modifier GP</td>
<td>Individual</td>
<td>Licensed therapist</td>
</tr>
<tr>
<td>1, 2, or 9</td>
<td>97110 with modifier GP and U1</td>
<td>Individual</td>
<td>Licensed therapy assistant</td>
</tr>
<tr>
<td>1, 2, or 9</td>
<td>97150 with modifier GP</td>
<td>Group</td>
<td>Licensed therapist</td>
</tr>
</tbody>
</table>

*Place of Service: 1=office; 2=home; 9=other locations
3.3.7.4 Providers must use a 15-minute unit of service for billing.

Refer to: Subsection 3.5.1.2, “Billing Units Based on 15 Minutes” in this handbook.

The recommended maximum billable time for PT evaluation is three hours, which may be billed over several days within a 30 day period. The recommended maximum billable time for direct therapy (individual or group) is a total of one hour per day. Providers must submit documentation of the reasons for the additional time, if more than the recommended maximum time is billed.

3.3.8 Physician Services

Diagnostic and evaluation services are reimbursable under SHARS physician services. Physician services must be provided by a licensed physician (M.D. or D.O.). A physician prescription is required before PT or OT services may be reimbursed under SHARS. ST services require either a physician prescription or a referral from a licensed SLP before the ST services may be reimbursed under the SHARS program. The school district must maintain the prescription or referral. The prescription or referral must relate directly to specific services listed in the IEP. If a change is made to a service on the IEP that requires a prescription or referral, the prescription or referral must be revised accordingly.

The expiration date for the physician prescription is the earlier of either the physician’s designated expiration date on the prescription or three years, in accordance with the IDEA three-year re-evaluation requirement.

SHARS physician services are billable only when they are provided on an individual basis. The determination as to whether not the provider needs to see the student while reviewing the student’s records is left up to the professional judgment of the provider. Therefore, billable time includes the following:

- The diagnosis or evaluation time spent with the student present
- The time spent without the student present reviewing the student’s records for the purpose of writing a prescription or referral for specific SHARS services
- The diagnosis or evaluation time spent with the student present, or the time spent without the student present reviewing the student’s records for the evaluation of the sufficiency of an ongoing SHARS service to see whether any changes are needed in the current prescription or referral for that service

Session notes are not required for procedure code 99499; however, documentation must include the billable start time, billable stop time, total billable minutes, and must note the medical activity that was performed.

3.3.8.1 Physician Services Billing Table

<table>
<thead>
<tr>
<th>POS*</th>
<th>Procedure Code</th>
<th>Individual or Group</th>
<th>Therapist or Assistant</th>
</tr>
</thead>
<tbody>
<tr>
<td>1, 2, or 9</td>
<td>97150 with modifier GP and U1</td>
<td>Group</td>
<td>Licensed therapy assistant</td>
</tr>
</tbody>
</table>

*Place of Service: 1=office; 2=home; 9=other locations

Refer to: Subsection 3.5.1.2, “Billing Units Based on 15 Minutes” in this handbook.

The recommended maximum billable time is one hour per day. Providers must submit documentation of the reasons for the additional time, if more than the recommended maximum time is billed.
3.3.9  Speech Therapy (ST)

3.3.9.1  Referral
The name and complete address or the provider identifier or license number of the referring licensed physician or licensed SLP is required before ST services can be billed under SHARS. A licensed SLP’s evaluation and recommendation for the frequency, location, and duration of ST serves as the speech referral.

3.3.9.2  Description of Services
ST evaluation services include the identification of children with speech or language disorders and the diagnosis and appraisal of specific speech and language disorders. ST services include the provision of speech and language services for the habilitation or prevention of communicative disorders.

ST evaluation is billable on an individual (procedure codes 92521, 92522, 92523, and 92524) basis only. ST is billable on an individual (procedure code 92507) or group (procedure code 92508) basis.

Procedure codes 92521, 92522, 92523, and 92524 are limited to a total of 12 units and may be reimbursed for each client per provider in a 30-day period.

Procedure code 92522 will be denied if it is submitted with the same date of service as procedure code 92523.

Procedure code 92523 will be denied if it is submitted with the same date of service as procedure code 92522.

Providers can only bill time spent with the student present, including assisting the student with learning to use adaptive equipment and assistive technology.

Time spent without the student present, such as report writing and training teachers or aides to work with the student (unless the student is present during training), is not billable. Session notes are not required for procedure codes 92521, 92522, 92523, and 92524; however, documentation must include the billable start time, billable stop time, total billable minutes, and must note the activity that was performed (e.g., speech evaluation).

Session notes are required for procedure codes 92507 and 92508. Session notes must include the billable start time, billable stop time, total billable minutes, activity performed during the session, student observation, and the related IEP objective.

3.3.9.3  Provider and Supervision Requirements
ST services are eligible for reimbursement when they are provided by a qualified SLP, who holds a Texas license or an ASHA-equivalent SLP (has a master’s degree in the field of speech-language pathology and a Texas license). ST services are also eligible for reimbursement when provided by an SLP with a state education agency certification, a licensed SLP intern, a grandfathered SLP when acting under the supervision or direction of an SLP, or a licensed assistant in speech-language pathology acting under the supervision or direction of an SLP.

The supervision must meet the following provisions:

- The supervising SLP must provide supervision that is sufficient to ensure the appropriate completion of the responsibilities that were assigned.
- The direct involvement of the supervising SLP in overseeing the services that were provided must be documented.
- The SLP who provides the direction must ensure that the personnel who carry out the directives meet the minimum qualifications set forth in the rules of the TDLR that relate to Licensed Interns or Assistants in Speech-Language Pathology.
CMS interprets “under the direction of a speech-language pathologist,” as an SLP who:

- Is directly involved with the individual under his direction.
- Accepts professional responsibility for the actions of the personnel he agrees to direct.
- Sees each student at least once.
- Has input about the type of care provided.
- Reviews the student’s speech records after the therapy begins.
- Assumes professional responsibility for the services provided.

### 3.3.9.4 Speech Therapy Billing Table

<table>
<thead>
<tr>
<th>POS*</th>
<th>Procedure Code</th>
<th>Individual or Group</th>
<th>Therapist or Assistant</th>
</tr>
</thead>
<tbody>
<tr>
<td>1, 2, or 9</td>
<td>92521, 92522, 92523, or 92524 with modifier GN</td>
<td>Individual</td>
<td>Licensed therapist</td>
</tr>
<tr>
<td>1, 2, or 9</td>
<td>92507 with modifier GN and U8</td>
<td>Individual</td>
<td>Licensed therapist</td>
</tr>
<tr>
<td>1, 2, or 9</td>
<td>92507 with modifier GN and U1</td>
<td>Individual</td>
<td>Licensed assistant</td>
</tr>
<tr>
<td>1, 2, or 9</td>
<td>92508 with modifier GN and U8</td>
<td>Group</td>
<td>Licensed therapist</td>
</tr>
<tr>
<td>1, 2, or 9</td>
<td>92508 with modifier GN and U1</td>
<td>Group</td>
<td>Licensed assistant</td>
</tr>
</tbody>
</table>

*Place of Service: 1=office; 2=home; 9=other locations

Providers must use a 15-minute unit of service for billing.

Refer to: Subsection 3.5.1.2, “Billing Units Based on 15 Minutes” in this handbook.

The recommended maximum billable time for evaluation is three hours, which may be billed over several days. The recommended maximum billable time for direct therapy (individual or group) is a total of one hour per day. Providers must submit documentation of the reasons for the additional time, if more than the recommended maximum time is billed.

### 3.3.10 Transportation Services in a School Setting

Transportation services in a school setting may be reimbursed when they are provided on a specially adapted vehicle and if the following criteria are met:

- Provided to or from a Medicaid-covered service on the day for which the claim is made
- A child requires transportation in a specially adapted vehicle to serve the needs of the disabled
- A child resides in an area that does not have school bus transportation, such as those in close proximity to a school
- The Medicaid services covered by SHARS are included in the student’s IEP
- The special transportation service is included in the student’s IEP

A specially adapted vehicle is one that has been physically modified (e.g., addition of a wheelchair lift, addition of seatbelts or harnesses, addition of child protective seating, or addition of air conditioning). A bus monitor or other personnel accompanying children on the bus is not considered an allowable
special adaptive enhancement for Medicaid reimbursement under SHARS specialized transportation. Specialized transportation services reimbursable under SHARS requires the Medicaid-eligible special education student has the following documented in his or her IEP:

- The student requires a specific physical adaptation or adaptations of a vehicle in order to be transported
- The reason the student needs the specialized transportation

Children with special education needs who ride the regular school bus to school with other nondisabled children are not required to have the transportation services in a school setting listed in their IEP. Also, the cost of the regular school bus ride cannot be billed to SHARS. Therefore, the fact that a child may receive a service through SHARS does not necessarily mean that the transportation services in a school setting may be reimbursed for them.

Reimbursement for covered transportation services is on a student one-way trip basis. If the student receives a billable SHARS service (including personal care services on the bus) and is transported on the school’s specially adapted vehicle, the following one-way trips may be billed:

- From the student’s residence to school
- From the school to the student’s residence
- From the student’s residence to a provider’s office that is contracted with the district
- From a provider’s office that is contracted with the district to the student’s residence
- From the school to a provider’s office that is contracted with the district
- From a provider’s office that is contracted with the district to the student’s school
- From the school to another campus to receive a billable SHARS service
- From the campus where the student received a billable SHARS service back to the student’s school

Covered transportation services from a child’s residence to school and return are not reimbursable if, on the day the child is transported, the child does not receive Medicaid services covered by SHARS (other than transportation). Documentation of each one-way trip provided must be maintained by the school district (e.g., trip log). This service must not be billed by default simply because the student is transported on a specially adapted bus.

### 3.3.10.1 Transportation Services in a School Setting Billing Table

<table>
<thead>
<tr>
<th>POS*</th>
<th>Procedure Code</th>
<th>Unit of Service</th>
</tr>
</thead>
<tbody>
<tr>
<td>1, 2, or 9</td>
<td>T2003</td>
<td>Per one-way trip</td>
</tr>
</tbody>
</table>

*Place of Service: 1=office; 2=home; 9=other locations

The recommended maximum billable units for procedure code T2003 is a total of four one-way trips per day.

### 3.3.11 Prior Authorization

Prior authorization is not required for SHARS services.
3.4  Documentation Requirements

3.4.1  Record Retention
Student-specific records that are required for SHARS become part of the student’s educational records and must be maintained for seven years. All records that are pertinent to SHARS billings must be maintained by the school district until all audit questions, appeal hearings, investigations, or court cases are resolved. Records must be stored in a readily accessible location and format and must be available for state or federal audits.

The following is a checklist of the minimum documents to collect and maintain:

- Signed consent to bill Medicaid by parent or guardian
- IEP
- Current provider qualifications (licenses)
- Attendance records
- Prescriptions and referrals
- Medical necessity documentation (e.g., diagnoses and history of chronic conditions or disability)
- Session notes or service logs, including provider signatures
- Supervision logs
- Special transportation logs
- Claims submittal and payment histories

All services require documentation to support the medical necessity of the service rendered, including SHARS services. SHARS services are subject to retrospective review and recoupment if documentation does not support the service billed.

3.5  Claims Filing and Reimbursement
During the cost report period, school districts participating in SHARS are reimbursed on an interim claiming basis using SHARS interim rates. It is important that SHARS providers understand that SHARS interim payments are provisional in nature. The total allowable costs for providing services for SHARS must be documented by submitting the required annual cost report.

3.5.1  Claims Information
Claims for SHARS must be submitted to TMHP in an approved electronic claims format or on a CMS-1500 claim form. Providers may purchase CMS-1500 paper claim forms from the vendor of their choice. TMHP does not supply the forms.

Claims must be submitted within 365 days from the date of service, or no later than 95 days after the end of the Federal Fiscal Year (i.e., January 3), whichever comes first.

Refer to: “Section 3: TMHP Electronic Data Interchange (EDI)” (Vol. 1, General Information) for information on electronic claims submissions.

“Section 6: Claims Filing” (Vol. 1, General Information) for general information about claims filing.

Subsection 6.5, “CMS-1500 Paper Claim Filing Instructions” in “Section 6: Claims Filing” (Vol. 1, General Information) for instructions on completing paper claims.
3.5.1.1 Appealing Denied SHARS Claims

SHARS providers that appeal claims denied for exceeding benefit limitations must submit documentation of medical necessity with the appeal. Documentation submitted with an appeal must include the pages from the IEP and ARD documents that show the authorization of the services, including the specified frequency and duration and the details of the need for additional time or the reasons for exceeding the benefit limitations.

Each page of the documentation must have the client’s name and Medicaid number.

3.5.1.2 Billing Units Based on 15 Minutes

All claims for reimbursement are based on the actual amount of billable time associated with the SHARS service. For those services for which the unit of service is 15 minutes (1 unit = 15 minutes), partial units must be rounded up or down to the nearest quarter hour.

Reminder: Enter the number of billing units in Block 24G of the CMS-1500 paper claim form. Claims without this information may be reimbursed as a unit of 1.

To calculate billing units, count the total number of billable minutes for the calendar day for the SHARS student, and divide by 15 to convert to billable units of service. If the total billable minutes are not divisible by 15, the minutes are converted to one unit of service if they are greater than seven and converted to 0 units of service if they are seven or fewer minutes.

For example, 68 total billable minutes/15 = 4 units + 8 minutes. Since the 8 minutes are more than 7 minutes, those 8 minutes are converted to one unit. Therefore, 68 total billable minutes = 5 units of service.

Examples:

<table>
<thead>
<tr>
<th>Minutes</th>
<th>Units</th>
</tr>
</thead>
<tbody>
<tr>
<td>0 min–7 mins</td>
<td>0 units</td>
</tr>
<tr>
<td>8 mins–22 mins</td>
<td>1 unit</td>
</tr>
<tr>
<td>23 mins–37 mins</td>
<td>2 units</td>
</tr>
<tr>
<td>38 mins–52 mins</td>
<td>3 units</td>
</tr>
<tr>
<td>53 mins–67 mins</td>
<td>4 units</td>
</tr>
<tr>
<td>68 mins–82 mins</td>
<td>5 units</td>
</tr>
</tbody>
</table>

3.5.1.3 Billing Units Based on an Hour

All claims for reimbursement are based on the actual amount of billable time associated with the SHARS service. For those services for which the unit of service is an hour (1 unit = 60 minutes = one hour), partial units must be billed in tenths of an hour and rounded up or down to the nearest six-minute increment.

Enter the number of billing units in Block 24G of the CMS-1500 paper claim form. Claims without this information may be reimbursed as a unit of 1.

To calculate billing units, count the total number of billable minutes for the calendar day for the SHARS student and divide by 60 to convert to billable units of service. If the total billable minutes are not divisible by 60, the minutes are converted to partial units of service as follows:

<table>
<thead>
<tr>
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<th>Units</th>
</tr>
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<tbody>
<tr>
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<td>0 units</td>
</tr>
<tr>
<td>4 mins–9 mins</td>
<td>0.1 unit</td>
</tr>
<tr>
<td>10 mins–15 mins</td>
<td>0.2 unit</td>
</tr>
</tbody>
</table>
### Other examples:

<table>
<thead>
<tr>
<th>Minutes</th>
<th>Units</th>
</tr>
</thead>
<tbody>
<tr>
<td>16 mins–21 mins</td>
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</tr>
<tr>
<td>22 mins–27 mins</td>
<td>0.4 unit</td>
</tr>
<tr>
<td>28 mins–33 mins</td>
<td>0.5 unit</td>
</tr>
<tr>
<td>34 mins–39 mins</td>
<td>0.6 unit</td>
</tr>
<tr>
<td>40 mins–45 mins</td>
<td>0.7 unit</td>
</tr>
<tr>
<td>46 mins–51 mins</td>
<td>0.8 unit</td>
</tr>
<tr>
<td>52 mins–57 mins</td>
<td>0.9 unit</td>
</tr>
</tbody>
</table>

#### 3.5.2 Managed Care Clients

SHARS services are carved-out of the Medicaid Managed Care Program and must be billed to TMHP for payment consideration. Carved-out services are those that are rendered to Medicaid Managed Care clients, but are administered by TMHP and not the client’s MCO.

#### 3.5.3 Reimbursement

Providers are reimbursed for medical and transportation services provided under the SHARS Program on a cost basis using federally mandated allocation methodologies in accordance with 1 TAC §355.8443.

In order to accommodate participating SHARS districts that require interim cash flow to offset the financial burden of providing for students, an interim fee-for-service claiming system still exists for SHARS. The interim claims are based on SHARS interim rates but are provisional in nature.

The provider’s final reimbursement amount is arrived at by a cost report, cost reconciliation, and cost settlement process. The provider’s total costs for both direct medical and transportation services as reported in the cost report are adjusted using the federally mandated allocation methodologies.

If a provider’s interim payments exceed 99 percent of the provider’s federal portion of the total certified Medicaid allowable costs, the provider must repay the over payments or HHSC will offset all of the provider’s future claims payments until the amount is recovered.

If 99 percent of the provider’s federal portion of the total certified Medicaid allowable costs exceeds the interim Medicaid payments, HHSC will pay the difference to the provider in accordance with the final actual certification agreement.

Submittal of a SHARS cost report is mandatory for each provider that requests and receives interim payments. Failure to file a SHARS cost report will result in sanctions, which includes recoupment of all interim payments for the cost report period in which the default occurs.


For additional information SHARS providers can contact a SHARS Rate Analyst by email at ra_shars@hhsc.state.tx.us or by telephone at 1-512-730-4300.
Refer to: Subsection 2.2, “Fee-for-Service Reimbursement Methodology” in “Section 2: Texas Medicaid Fee-for-Service Reimbursement” (Vol. 1, General Information) for more information. Subsection 2.9, “Federal Medical Assistance Percentage (FMAP)” in “Section 2: Texas Medicaid Fee-for-Service Reimbursement” (Vol. 1, General Information).

3.5.3.1 Quarterly Certification of Funds

SHARS providers are required to certify on a quarterly basis the amount reimbursed during the previous federal fiscal quarter. TMHP Provider Enrollment mails the quarterly Certification of Funds statement to SHARS providers after the end of each quarter of the federal fiscal year (October 1 through September 30). The purpose of the statement is to verify that the school district incurred costs on the dates of service that were funded from state or local funds in an amount equal to, or greater than, the combined total of its interim rates times the paid units of service. While the payments were received the previous federal fiscal quarter, the actual dates of service could have been many months prior. Therefore, the certification of public expenditures is for the date of service and not the date of payment.

In order to balance amounts in the Certification of Funds, providers will receive, or have access to, the Certification of Funds Claims Information Report. For help balancing the amounts in the statement, providers can contact the TMHP Contact Center at 1-800-925-9126.

Refer to: “Subsection A.12.4, "TMHP Provider Relations” in “Appendix A: State, Federal, and TMHP Contact Information” (Vol. 1, General Information)” for more information about provider relations representatives.

The Certification of Funds statement must be:

- Signed by the business officer or other financial representative who is responsible for signing other documents that are subject to audit.
- Notarized.
- Returned to TMHP within 25 calendar days of the date printed on the letter.

Failure to do so may result in recoupment of funds or the placement of a vendor hold on the provider’s payments until the signed Certification of Funds statement is received by TMHP. Providers must contact the TMHP Contact Center at 1-800-925-9126 if they do not receive their Certification of Funds statement.

On an annual basis, SHARS providers are required to certify through their cost reports their total, actual, incurred costs, including the federal share and the nonfederal share. Refer to subsection 3.6, “Cost Reporting, Cost Reconciliation, and Cost Settlement” in this handbook for additional information about cost reporting.

3.6 Cost Reporting, Cost Reconciliation, and Cost Settlement

CMS requires annual cost reporting, cost reconciliation, and cost settlement processes for all Medicaid SHARS services delivered by school districts. CMS requires that school districts, as public entities, not be paid in excess of their Medicaid-allowable costs and that any overpayments be recouped through the cost reconciliation and cost settlement processes. In an effort to minimize any potential recoupments, HHSC has assigned SHARS interim rates that are as close as possible to each district’s Medicaid-allowable costs for providing each SHARS service.

3.6.1 Cost Reporting

Each SHARS provider is required to complete an annual cost report for all SHARS that were delivered during the previous federal fiscal year (October 1 through September 30). The cost report is due on or before April 1 of the year following the reporting period.

The following certification forms must be submitted and received by HHSC for the cost report. The annual cost report includes two certification forms which must be completed to certify the provider’s incurred actual costs:

- Cost report certification
- Claimed expenditures

The certification forms received by HHSC for the cost report must be:

- The original certification pages.
- Signed by the business officer or other financial representative who is responsible for legally binding the district.
- Notarized.

The primary purpose of the cost report is to document the provider’s costs for delivering SHARS, including direct costs and indirect costs, and to reconcile the provider’s interim payments for SHARS with its actual total Medicaid-allowable costs. All annual SHARS cost reports that are filed are subject to desk review by HHSC or its designee.

For additional information, SHARS providers can contact a SHARS Rate Analyst by email at ra_shars@hhsc.state.tx.us or by telephone at 1-512-730-7400.

### 3.6.2 Cost Reconciliation and Cost Settlement

The cost reconciliation process must be completed within 24 months of the end of the reporting period covered by the annual SHARS cost report. The total Medicaid-allowable costs are compared to the provider’s interim payments for SHARS delivered during the reporting period, which results in a cost reconciliation.

If a provider has not complied with all cost report requirements or a provider’s interim payments exceed the actual certified Medicaid-allowable costs of the provider for SHARS to Medicaid clients, HHSC will recoup the federal share of the overpayment by one of the following methods:

- Offset all future claims payments to the provider until the amount of the federal share of the overpayment is recovered
- Recoup an agreed-upon percentage from future claims payments to the provider to ensure recovery of the overpayments within one year
- Recoup an agreed-upon dollar amount from future claims payments to ensure recovery of the overpayment within one year

If the actual certified Medicaid-allowable costs of a provider for SHARS exceed the provider’s interim payments, HHSC will pay the federal share of the difference to the provider in accordance with the final, actual certification agreement and submit claims to CMS for reimbursement of that payment in the federal fiscal quarter following payment to the provider.

HHSC issues a notice of settlement that denotes the amount due to or from the provider.

### 3.6.3 Informal Review of Cost Reports Settlement

An ISD or the Superintendent, Chief Financial Officer, Business Officer, or other ISD Official with legal authority who disagrees with the adjustments made during the cost reconciliation process has the right to request an informal review of the adjustments. Requests for informal reviews must be sent by certified mail and received by HHSC within the time frame designated on the settlement notice. Furthermore, the request for informal review must include a concise statement of the specific actions or determinations
the district disputes, the ISD’s recommended resolution, and any supporting documentation deemed relevant to the dispute. Failure to follow these instructions will result in the denial of the request for an informal review.

School districts can access published cost report guidance documents, on the HHSC website at https://rad.hhs.texas.gov/acute-care/school-health-and-related-services/shars-cost-report-information. For additional information, SHARS providers can contact a SHARS Rate Analyst by email at ra_shars@hhsc.state.tx.us or by telephone at 1-512-730-7400.

4 * Texas Health Steps (THSteps) Dental

Medicaid dental services rules are described under Title 25 Texas Administrative Code (TAC) Part 1, Chapter 33. The online version of TAC is available at the Secretary of State’s website at www.sos.state.tx.us/tac/index.shtml. All dental providers must comply with the rules and regulations of the Texas State Board of Dental Examiners (TSBDE), including standards for documentation and record maintenance as stated in 22 TAC §108.7, Minimum Standard of Care, General, and §108.8, Records of the Dentist.

Note: THSteps dental benefits are administered as Children’s Medicaid Dental Services by dental managed care organizations for most Medicaid fee-for-service and managed care clients who are 20 years of age and younger.

Referto: [Revised] The Medicaid Managed Care Handbook (Vol. 2, Provider Handbooks) or to the HHSC website at hhs.texas.gov/services/health/medicaid-chip, for additional information about children’s Medicaid dental Services.

Under the Early Periodic Screening Diagnostic, and Treatment (EPSDT) regulation, known in Texas as Texas Health Steps (THSteps), Section 1905(r) of the Social Security Act mandates that all Medicaid eligible beneficiaries who are birth through 20 years of age receive medically necessary services to treat, correct, and ameliorate illnesses and conditions identified if the service is covered in the state’s Medicaid plan or is an optional Medicaid service. It is the responsibility of the state to determine medical necessity on a case-specific basis. No arbitrary limitations on services are allowed (e.g., one pair of eyeglasses or 10 therapy sessions per year) if determined to be medically necessary.

Services not covered under this section include:

- Experimental or investigational treatment.
- Services or items not generally accepted as effective and/or not within the normal course and duration of treatment.
- Services for the caregiver or provider convenience.

All EPSDT requirements must be adhered to for beneficiaries who receive services under managed care arrangements.

4.1 Enrollment

To become a provider of THSteps or intermediate care facility for persons with intellectual disability (ICF-IID) dental services, a dentist must:

- Practice within the scope of the provider’s professional licensure.
- Complete the Dental Provider Enrollment Application and return it to TMHP.

Dental providers are required to maintain an active license status with the TSBDE. TMHP receives a monthly automated board feed from TSBDE to update licensure information. If licensure cannot be verified with the automated board feed, it is the providers’ responsibility to provide a copy of the active TSBDE license to TMHP. If TSBDE has a delay in processing license applications and renewals, the
The provider must request a letter from TSBDE for their individual provider information and send the letter of verification of current licensure to TMHP. The letter must contain the provider’s specific identification information, license number, and licensure period.

If TMHP cannot verify a valid license at the time of enrollment, it is the providers’ responsibility to provide a copy of the active TSBDE license to TMHP.

A dental provider cannot be enrolled if his or her dental license is due to expire within 30 days; a current license must be submitted. Dental licensure for owners of a dental practice is a requirement of the Occupations Code, Vernon’s Texas Codes Annotated (VTCA), Subtitle D, Chapters 251-267 (the Texas Dental Practice Act).

Providers can download and print dental provider enrollment application forms from the TMHP website at www.tmhp.com or call the TMHP Contact Center at 1-800-925-9126 to request them.

All owners of a dental practice must maintain an active license status with the TSBDE to receive reimbursement from Texas Medicaid. Any change in ownership or licensure status for any enrolled dentist must be immediately reported in writing to TMHP Provider Enrollment and will affect reimbursement by Texas Medicaid.

A dentist must complete the Dental Provider Enrollment Application for each separate practice location and will receive a unique provider identifier for each practice location if the application is approved.

The application form includes a written agreement with HHSC.

Dental providers may enroll in the THSteps Dental program and ICF-IID Dental Programs or as a Doctor of Dentistry Practicing as a Limited Physician, or both. The enrollment requirements are different with respect to the category of enrollment.

- All dental providers must declare one or more of the following categories:
  - General practice
  - Pediatric dentist
  - Periodontist
  - Endodontist
  - Oral and maxillofacial surgeon
  - Orthodontist
  - Other (prosthodontist, public health, and others)

Dentists (D.D.S., D.M.D.) who want to provide orthodontic services must be enrolled as a dentist or orthodontist provider for THSteps and must have at least one of the following qualifications.

THSteps dental providers may perform and be reimbursed for orthodontic services if they have attested to at least one of the following requirements:

- Completion of a dental pediatric specialty residency
- Completion of a minimum of 200 hours of continuing education in orthodontics within the last 10 years (8 hours can be online or self instruction) (Proof of the completion of continuing education hours is not required to be submitted with a request for prior authorization of orthodontic services; however, documentation must be produced by the dentist during retrospective review.)

Orthodontist providers are eligible to provide orthodontic services. In order to comply with the TSBDE rules and regulations, this designation can only be associated with dentists who are board-eligible or board-certified by an American Dental Association (ADA) recognized orthodontic specialty board.

Referto: “Section 1: Provider Enrollment and Responsibilities” (Vol. 1, General Information).
Dental residents may provide dental services in a teaching facility under the guidance of the attending staff/faculty member(s) as long as the facility’s dental staff by-laws and standards by the Commission on Dental Accreditation (CODA) are met, and the attending dentist/faculty member has determined the resident to be competent to perform the dental services. THSteps does not require the supervising dentist to examine the client as long as these conditions are met.

In a clinic, an attending dentist/faculty member must be present in the dental clinic for consultation, supervision, and active teaching when residents are treating patients in scheduled clinic sessions. This does not preclude occasional situations where a faculty member cannot be available. A dentist must assume responsibility for the clinic’s operation.

4.1.1 THSteps Dental Eligibility
The client must be Medicaid- and THSteps-eligible (birth through 20 years of age) at the time of the service request and service delivery. However, Medicaid-approved orthodontic services already in progress may be continued even after the client loses Medicaid eligibility if the orthodontic treatment:

- Began before the loss of Medicaid eligibility.
- Began before the day of the client’s 21st birthday.
- Was completed within 36 months of the beginning date.

The client is not eligible for a THSteps medical checkup or THSteps dental benefits if the client’s Your Texas Benefits Medicaid card or Medicaid Eligibility Verification Form (Forms H1027 and H1027-A-C) states any of the following:

- Emergency
- Presumptive eligibility (PE)
- Qualified Medicare beneficiary (QMB)
- Healthy Texas Women (HTW) program

4.1.2 THSteps Dental and ICF-IID Dental Services
A provider may enroll as an individual dentist, a group practice, or both. Regardless of the category of practice designation under THSteps Dental, providers can only submit claims for THSteps and ICF-IID Dental Services.


4.1.3 THSteps Dental Checkup and Treatment Facilities
All THSteps dental checkup and treatment policies apply to examinations and treatment completed in a dentist’s office, a health department, clinic setting, hospital operating room, or in a mobile/satellite unit. Enrollment of a mobile/satellite unit must be under a dentist or clinic name. Mobile units can be a van or any temporary site away from the primary office and are considered extensions of that office and are not separate entities. The physical setting must be appropriate so that all elements of the checkup or treatment can be completed. The checkup must meet the requirements detailed in subsection D.5, “Parental Accompaniment” in this handbook. The provider with a mobile unit or who uses portable dental equipment must obtain a permit for the mobile unit from the TSBDE.

4.1.4 Doctor of Dentistry Practicing as a Limited Physician
Dentists who serve clients and submit claims using medical (CPT) procedure codes, such as oral-maxillofacial surgeons, may enroll as a doctor of dentistry practicing as a limited physician. Providers may enroll as an individual dentist or as a dental group. To enroll as a doctor of dentistry practicing as a limited physician, a dentist must:

- Be currently licensed by the TSBDE or currently licensed in the state where the service was performed.
• Have a Medicare provider identification number before applying for a Medicaid provider identifier.

• Enroll as a Medicaid provider with a limited physician provider identifier.

4.1.5 Client Rights

Dental providers enrolled in Texas Medicaid enter into a written contract with HHSC to uphold the following rights of the Medicaid client:

• To receive dental services that meet or exceed the standards of care established by the laws relating to the practice of dentistry and the rules and regulations of the TSBDE.

• To receive information following a dental examination about the dental diagnosis; scope of proposed treatment, including alternatives and risks; anticipated results; and the need and risks for administration of sedation or anesthesia.

• To have full participation in the development of the treatment plan and the process of giving informed consent.

• To have freedom from physical, mental, emotional, sexual, or verbal abuse, or harm from the provider or staff.

• To have freedom from overly aggressive treatment in excess of that required to address documented medical necessity.

A provider’s failure to ensure any of the client rights may result in termination of the provider agreement or contract and other civil or criminal remedies.

4.1.6 Complaints and Resolution

Complaints about dental services are typically received through the TMHP Contact Center, although a complaint is accepted from any source. A complaint is researched by TMHP and resolved or escalated as appropriate. Examples of complaints from clients about providers include:

• The provider did not consult with the client, explain what services were necessary, or obtain parent or guardian informed consent.

• The treating provider refused to make the child’s record available to the new provider.

• The provider did not give the child the appropriate local anesthesia or pain medication.

• The provider did not use sterile procedures; the facility or equipment were not clean.

• The provider or his staff were verbally abusive.

• The client did not receive a service, but the provider submitted a claim to Texas Medicaid.

• The provider charged a Medicaid client for benefits covered by Medicaid.

4.2 Services, Benefits, Limitations, and Prior Authorization

4.2.1 THSteps Dental Services

THSteps is the Texas version of the Medicaid program known as Early and Periodic Screening, Diagnosis, and Treatment (EPSDT).

THSteps dental services are mandated by Medicaid to provide for the early detection and treatment of dental health problems for Medicaid-eligible clients who are birth through 20 years of age. THSteps dental service standards are designed to meet federal regulations and incorporate the recommendations of representatives of national and state dental professional organizations.

THSteps’ designated staff (DSHS, HHSC, or contractor), through outreach and informing, encourage eligible children to use THSteps dental checkups and services when children first become eligible for Medicaid, and each time children are periodically due for their next dental checkup.
Children within Medicaid have free choice of Medicaid-enrolled providers and are given names of enrolled providers. A list of THSteps dental providers in a specific area can be obtained using the Online Provider Lookup on the TMHP website at www.tmhp.com, or by calling 1-877-847-8377.

Upon a provider’s request, DSHS (or its contractor) will assist eligible children with the scheduling of free transportation to their dental appointment or clients can call the Medical Transportation Program at 1-877-633-8747.

Refer to: The Medical Transportation Program Handbook (Vol. 2, Provider Handbooks) for information about transportation arrangements.

4.2.1.1 Eligibility for THSteps Dental Services

A client is eligible for THSteps dental services from birth through 20 years of age. The eligibility period is determined by the client’s age on the first of the month. If a client’s birthday is not on the first of a month, the new eligibility period begins on the first day of the following month. When the client turns 21 years of age during a month, the client is eligible for THSteps dental non-CCP services through the end of that month.

A client is eligible for Comprehensive Care Program (CCP) dental services until their 21st birthday. The eligibility period ends on their 21st birthday and does not continue through the end of the month in which the birthday falls.

4.2.1.2 Parental Accompaniment

Clients who are 14 years of age and younger must be accompanied to Texas Health Steps dental checkups and visits by the client’s parent, legal guardian, or another adult who is authorized by the parent or legal guardian. The authorized adult can be the client’s relative. The individual accompanying the client must wait for the client while the appointment takes place. For additional information and exceptions, see “D.5 Parental Accompaniment” in this handbook.

4.2.2 Substitute Dentist

In accordance with TAC §§354.1121 and 354.1221, related to Medicaid billing for the services of substitute dentists, dentists who are temporarily absent from their practice are allowed to submit claims for reimbursement of Medicaid services rendered to their Medicaid clients by a substitute dentist.

Dentists may bill for the services of a substitute dentist pursuant to 42 CFR §447.10.

The following are conditions for reimbursement of services rendered by a substitute dentist:

- Dentists who take a leave of absence for no more than 90 days may bill for the services of a substitute dentist who renders services on an occasional basis when the primary dentist is unavailable to provide services. Services must be rendered at the practice location of the dentist who has taken the leave of absence. A locum tenens arrangement is not allowed for dentists.

- This arrangement is limited to no more than 90 consecutive days. Under this temporary basis, the primary dentist (who is the billing agent dentist) may not submit a claim for services furnished by a substitute dentist to address long-term vacancies in a dental practice. The billing agent dentist may submit claims for the services of a substitute dentist for longer than 90 consecutive days if the dentist has been called or ordered to active duty as a member of a reserve component of the Armed Forces. Medicaid and CSHCN accepts claims from the billing agent dentist for services provided by the substitute dentist for the duration of the billing agent dentist’s active duty as a member of a reserve component of the Armed Forces.

- Providers billing for services provided by a substitute dentist must bill with modifier U5 in Block 19 of the American Dental Association (ADA) claim form.

- The billing agent dentist may recover no more than the actual administrative cost of submitting the claim on behalf of the substitute dentist. This cost is not reimbursable by Medicaid or CSHCN.
• The billing agent dentist must bill substitute dentist services on a different claim form from his or her own services. The billing agent dentist services cannot be billed on the same claim form as substitute dentist services.

• The substitute dentist must be licensed to practice in the state of Texas, must be enrolled in Texas Medicaid, and must not be on the Texas Medicaid provider exclusion list.

• The dentist who is temporarily absent from the practice must be indicated on the claim as the billing agent dentist, and his or her name, address, and National Provider Identifier (NPI) must appear in Blocks 53, 54, and 56 of the ADA claim form.

• The substitute dentist’s NPI number must be documented in Block 35 of the ADA claim form. Electronic submissions do not require a provider signature.

Dentists must familiarize themselves with these requirements and document accordingly. Those services not supported by the required documentation, as detailed above, will be subject to recoupment.

4.2.3 Texas Health Steps Dental Checkups

Texas Health Steps dental checkups include an oral evaluation, prophylaxis, topical fluoride, and appropriate radiographs.

The Texas Health Steps dental periodicity schedule for preventative and diagnostic procedures does not apply to clients living in an intermediate care facility for individuals with intellectual or developmental disabilities (ICF-IID) who are 21 years of age and older.

4.2.3.1 Exception-to-Periodicity Oral Evaluation, Dental Checkup, and Emergency or Trauma Related Services

Oral evaluations and dental checkups allow for the early diagnosis and treatment of dental problems. Oral evaluations and dental checkups might be needed at more frequent intervals than noted in the periodicity schedule.

If needed, a dental checkup or oral evaluation can still be reimbursed when the service falls outside the periodicity schedule. The rules for such exceptions are outlined below.

4.2.3.2 Exception-to-Periodicity Oral Evaluation

A Texas Health Steps exception-to-periodicity oral evaluation is limited to dental procedure code D0120.

An exception-to-periodicity oral evaluation is allowed when the service is:

• Medically necessary and based on risk factors and health needs for clients birth through 6 months of age.

• Mandated service required to meet federal or state exam requirements for Head Start, daycare, foster care or preadoption.

Providers must include all appropriate procedure codes on the dental claim submission form. Additionally, dental providers must include modifier SC or 32 to identify the reason for the exception.

4.2.3.3 Exception-to-Periodicity Dental Checkup

A Texas Health Steps exception-to-periodicity dental checkup is allowed when the client will not be available for the next periodically due dental checkup. This includes clients whose parents are migrant or seasonal workers.

Providers must include all appropriate procedure codes on the dental claim submission form. Additionally, dental providers must include modifier SC to identify the reason for the exception.
4.2.3.4 Exception-to-Periodicity Emergency or Trauma Related Oral Evaluation

A Texas Health Steps exception-to-periodicity emergency or trauma related oral evaluation is limited to dental procedure code D0140.

A Texas Health Steps exception-to-periodicity emergency or trauma related dental service is allowed when the service is:

- Required for immediate treatment and any follow-up treatment.
- Required for therapeutic services needed to complete a case for clients who are 5 months of age and younger, when initiated as emergency services, trauma, or early childhood caries.

When submitting a claim for emergency or trauma related dental services, the provider must include:

- “Trauma” or “Emergency” in Block 30 “Description” field.
- The original date of treatment or incident in Block 35, “Remark” field.
- Completion of Block 45, “Treatment Resulting from” field, if applicable.

Providers must include all appropriate procedure codes on the dental claim submission form. Additionally, dental providers must include modifier ET to identify the reason for the exception.


4.2.3.5 Documentation

The client’s dental medical record must include documentation for the exception-to-periodicity Texas Health Steps oral evaluation, dental checkup or exception-to periodicity emergency or trauma related service for medical necessity.

Dental services are subject to retrospective review and recoupment if documentation does not support the services submitted for payment.

4.2.3.6 Reimbursement

Providers must include the appropriate procedure code and one of the modifiers to identify the reason for the exception must be included on the ADA dental claim submission form. Procedure codes must be included in Block 29. Modifiers must be included in Block 19:

<table>
<thead>
<tr>
<th>Procedure Code</th>
<th>Modifier to Identify Exemption</th>
<th>Modifier Description</th>
</tr>
</thead>
<tbody>
<tr>
<td>D0120</td>
<td>32</td>
<td>Mandated Service</td>
</tr>
<tr>
<td>D0120</td>
<td>SC</td>
<td>Medically Necessary Service</td>
</tr>
<tr>
<td>D0140, D9110</td>
<td>ET</td>
<td>Emergency - Trauma Related Services</td>
</tr>
</tbody>
</table>

4.2.3.7 Diagnostic Services

Diagnostic services should be performed for all clients, starting within the first six months of the eruption of the first primary tooth, but no later than one year of age.
The provider must document medical necessity and the specific tooth or area of the mouth on the claim for procedure codes D0140, D0160, and D0170.

Documentation supporting medical necessity for procedure codes D0140, D0160, and D0170 must also be maintained by the provider in the client’s medical record and must include the following:

- The client’s complaint supporting medical necessity for the examination
- The specific area of the mouth that was examined or the tooth involved
- A description of what was done during the visit
- Supporting documentation of medical necessity which may include, but is not limited to, radiographs or photographs

Documentation supporting medical necessity for procedure code D0180 must be maintained by the provider in the client’s medical record and must include the following:

- The client’s complaint supporting medical necessity for the examination

### Procedure Code | Limitations
<table>
<thead>
<tr>
<th><strong>Clinical Oral Evaluations</strong></th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>Procedure codes D0140, D0160, D0170, and D0180 are limited dental codes and may be paid in addition to a comprehensive oral exam (procedure code D0150) or periodic oral exam (procedure code D0120), when submitted within a six-month period. When submitting a claim for procedure code D0140, D0160, D0170, or D0180, the provider must indicate documentation of medical necessity on the claim. These claims are subject to retrospective review. If no comments are indicated on the claim form, the payment may be recouped.</strong></td>
</tr>
<tr>
<td><strong>D0120</strong></td>
</tr>
<tr>
<td><strong>D0140</strong></td>
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<tr>
<td><strong>D0145</strong></td>
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<tr>
<td><strong>D0150</strong></td>
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<tr>
<td><strong>D0160</strong></td>
</tr>
<tr>
<td><strong>D0170</strong></td>
</tr>
<tr>
<td><strong>D0180</strong></td>
</tr>
</tbody>
</table>

A=Age range limitations, N=Narrative required, FMX=Full-mouth radiographs (nonpanoramic), MTID=Missing tooth ID(s), PPXR=Preoperative and postoperative radiographs required, PXR=Preoperative and postoperative photographs required, PC=Periodontal charting required, PATH=Pathology report required and must be retained in the client’s record, CCP=Comprehensive Care Program, NC=No charge to Medicaid and may not bill the client, and *= Services payable to an FQHC for a client encounter
• A description of what was done during the treatment
• Supporting documentation of medical necessity which may include, but is not limited to, radiographs or photographs

A caries risk assessment procedure code (D0601, D0602, or D0603) is required on the same claim when dental examination procedure code D0120, D0145, or D0150 are submitted for reimbursement on the same claim, for the same date of service, by the same provider.

Procedure codes D0601, D0602, and D0603 are informational only, and are not payable. Information-only procedure codes must be billed in the amount of at least $0.01 in the cost column on the claim form.

The client’s dental condition(s) that justifies the risk assessment classification submitted with the claim must be maintained by the provider in the client’s medical record, and it must be clearly documented using a caries risk assessment tool or in narrative charting. The client’s medical record is subject to retrospective review.

Professionally developed caries risk assessment tools are available at:
• American Dental Association (ADA)
• American Academy of Pediatric Dentistry (AAPD)
• Texas Health Steps (THSteps)

<table>
<thead>
<tr>
<th>Procedure Code</th>
<th>Limitations</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>Radiographs/Diagnostic Imaging (Including Interpretation)</strong></td>
<td></td>
</tr>
<tr>
<td>Number of films required is dependent on the age of the client. A minimum of eight films is required to be considered a full-mouth series. Adults and children who are 12 years of age and older require 12–20 films, as is appropriate. The panoramic radiographic image (D0330) with four bitewing radiographic images (D0274) may be considered equivalent to the complete or full-mouth series of radiographic images (D0210), and the submitted amount for either combination is equivalent to the maximum fee.</td>
<td></td>
</tr>
<tr>
<td>D0210</td>
<td>Limited to one service every three years by the same provider. Will be denied when submitted on an emergency claim. A 2–20</td>
</tr>
<tr>
<td>D0220</td>
<td>Limited to one service per day by the same provider. A 1–20</td>
</tr>
<tr>
<td>D0230</td>
<td>The total cost of periapicals and other radiographs cannot exceed the payment for a complete intraoral series. A 1–20</td>
</tr>
<tr>
<td>D0240</td>
<td>Limited to two services per day by the same provider. Periapical films taken at an occlusal angle must be submitted as periapical radiograph, procedure code D0230. May be submitted as an emergency service. A Birth–20</td>
</tr>
<tr>
<td>D0250</td>
<td>Limited to one service per day by the same provider. A 1–20, N, CCP</td>
</tr>
<tr>
<td>D0270</td>
<td>Limited to one service per day by the same provider. A 1–20</td>
</tr>
<tr>
<td>D0272</td>
<td>Limited to one service per day by the same provider. Denied when submitted for the same DOS as D0210 by any provider. A 1–20</td>
</tr>
<tr>
<td>D0273</td>
<td>Limited to one service per day by the same provider. Denied when submitted for the same DOS as D0210 by any provider. A 1–20</td>
</tr>
<tr>
<td>D0274</td>
<td>Limited to one service per day by the same provider. Denied when submitted for the same DOS as D0210 by any provider. A 2–20</td>
</tr>
</tbody>
</table>

A=Age range limitations, N=Narrative required, FMX=Full-mouth radiographs (nonpanoramic), MTID=Missing tooth ID(s), PPXR=Preoperative and postoperative radiographs required, PXR=Preoperative radiographs required, PHO=preoperative and postoperative photographs required, PC=Periodontal charting required, PATH=Pathology report required and must be retained in the client’s record, CCP=Comprehensive Care Program, NC=No charge to Medicaid and may not bill the client, and *= Services payable to an FQHC for a client encounter

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Procedure code D0350 must be used to submit claims for photographs, and will be accepted only when diagnostic-quality radiographs cannot be taken. Supporting documentation and photographs must be maintained in the client’s medical record when medical necessity is not evident on radiographs for dental caries or the following procedure codes. Medical necessity must be documented on the electronic or paper claim.

<table>
<thead>
<tr>
<th>Procedure Code</th>
<th>Limitations</th>
</tr>
</thead>
<tbody>
<tr>
<td>D0277</td>
<td>Limited to one service per day by the same provider. Not to be submitted within 36 months of D0210 or D0330. Denied when submitted for the same DOS as D0330 by the same provider. Denied when submitted for the same DOS as D0210 by any provider. A 2–20</td>
</tr>
<tr>
<td>D0310</td>
<td>A 1–20, N, CCP</td>
</tr>
<tr>
<td>D0320</td>
<td>A 1–20, N, CCP</td>
</tr>
<tr>
<td>D0321</td>
<td>A 1–20, N, CCP</td>
</tr>
<tr>
<td>D0322</td>
<td>A 1–20, N, CCP</td>
</tr>
<tr>
<td>D0330*</td>
<td>Limited to one service per day by the same provider. Not allowed on emergency claims unless third molars or a traumatic condition is involved. For clients who are 2 years of age and younger, must document the necessity of a panoramic film. The panoramic radiographic image (D0330) with four bitewing radiographic images (D0274) may be considered equivalent to the complete or full-mouth series of radiographic images (D0210), and the submitted amount for either combination is equivalent to the maximum fee. A 3–20</td>
</tr>
<tr>
<td>D0340*</td>
<td>Limited to one service per day by the same provider. Not reimbursable separately when a comprehensive orthodontic or crossbite therapy workup is performed. A 1–20, N, CCP</td>
</tr>
<tr>
<td>D0350*</td>
<td>Limited to one service per day by the same provider. Not reimbursable separately when a comprehensive orthodontic or crossbite therapy workup is performed. A Birth–20</td>
</tr>
<tr>
<td>D0367</td>
<td>Prior authorization is required. Limited to a combined maximum of three services per year, any provider. Additional services may be considered with documentation of medical necessity. A Birth-20</td>
</tr>
</tbody>
</table>

**Note:** Radiograph codes do not include the exam. If an exam is also performed, providers must submit the appropriate ADA procedure code.

**Procedure Codes**

<table>
<thead>
<tr>
<th>Procedure Code</th>
<th>Limitations</th>
</tr>
</thead>
<tbody>
<tr>
<td>D4210</td>
<td>D4211</td>
</tr>
<tr>
<td>D4240</td>
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<td>D4283</td>
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</tr>
<tr>
<td>D4355</td>
<td>D4910</td>
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</tbody>
</table>

**Procedure Code**

**Limitations**

**Tests and Examinations**

A=Age range limitations, N=Narrative required, FMX=Full-mouth radiographs (nonpanoramic), MTID=Missing tooth ID(s), PPXR=Preoperative and postoperative radiographs required, PXR=Preoperative radiographs required, PHO=Preoperative and postoperative photographs required, PC=Periodontal charting required, PATH=Pathology report required and must be retained in the client’s record, CCP=Comprehensive Care Program, NC=No charge to Medicaid and may not bill the client, and *= Services payable to an FQHC for a client encounter.
### Preventive Services

<table>
<thead>
<tr>
<th>Procedure Code</th>
<th>Limitations</th>
</tr>
</thead>
<tbody>
<tr>
<td>D0415</td>
<td>A 1–20, N, CCP</td>
</tr>
<tr>
<td>D0425</td>
<td>Not reimbursable separately. Considered part of another dental procedure.</td>
</tr>
<tr>
<td>D0460</td>
<td>Limited to one service per day by the same provider. Not payable for primary teeth. Will deny when submitted for the same DOS as any endodontic procedure. A 1-20, N, CCP</td>
</tr>
</tbody>
</table>

#### Tests and Examinations continued

<table>
<thead>
<tr>
<th>Procedure Code</th>
<th>Limitations</th>
</tr>
</thead>
<tbody>
<tr>
<td>D0470*</td>
<td>Not reimbursable separately when crown, fixed prosthodontics, diagnostic workup, or crossbite therapy workup is performed. A 1-20, N, CCP</td>
</tr>
</tbody>
</table>

### Oral Pathology Laboratory

<table>
<thead>
<tr>
<th>Procedure Code</th>
<th>Limitations</th>
</tr>
</thead>
<tbody>
<tr>
<td>D0472</td>
<td>By pathology laboratories only. (refer to CPT codes)</td>
</tr>
<tr>
<td>D0473</td>
<td>By pathology laboratories only. (refer to CPT codes)</td>
</tr>
<tr>
<td>D0474</td>
<td>By pathology laboratories only. (refer to CPT codes)</td>
</tr>
<tr>
<td>D0476</td>
<td>By pathology laboratories only. (refer to CPT codes)</td>
</tr>
<tr>
<td>D0502</td>
<td>A 1–20, N, CCP</td>
</tr>
<tr>
<td>D0999</td>
<td>A 1–20, N, CCP</td>
</tr>
</tbody>
</table>

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A=Age range limitations, N=Narrative required, FMX=Full-mouth radiographs (nonpanoramic), MTID=Missing tooth ID(s), PPXR=Preoperative and postoperative radiographs required, PXR=Preoperative radiographs required, PHO=Preoperative and postoperative photographs required, PC=Periodontal charting required, PATH=Pathology report required and must be retained in the client’s record, CCP=Comprehensive Care Program, NC=No charge to Medicaid and may not bill the client, and * = Services payable to an FQHC for a client encounter
<table>
<thead>
<tr>
<th>Procedure Code</th>
<th>Limitations</th>
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<tbody>
<tr>
<td><strong>Other Preventive Services</strong></td>
<td></td>
</tr>
<tr>
<td>D1310</td>
<td>Denied as part of all preventative, therapeutic and diagnostic dental procedures. A client requiring more involved nutrition counseling may be referred to a THSteps primary care physician.</td>
</tr>
<tr>
<td>D1320</td>
<td>A client requiring tobacco counseling may be referred to a THSteps primary care provider.</td>
</tr>
<tr>
<td>D1330</td>
<td>Requires documentation of the type of instructions, number of appointments, and content of instructions. This procedure refers to services above and beyond routine brushing and flossing instruction and requires that additional time and expertise have been directed toward the client’s care. Denied when billed for the same DOS as dental prophylaxis (D1110 or D1120) or topical fluoride treatments (D1206 or D1208) by any provider. Limited to once per client, per year, by any provider. A 1–20, N, CCP</td>
</tr>
<tr>
<td>D1351*</td>
<td>Sealants may be applied to the occlusal, buccal, and lingual pits and fissures of any tooth that is at risk for dental decay and is free of proximal caries and free of restorations on the surface to be sealed. Sealants are a benefit when applied to deciduous (baby or primary) teeth or permanent teeth. Indicate the tooth numbers and surfaces on the claim form. Reimbursement will be considered on a per-tooth basis, regardless of the number of surfaces sealed. Denied when billed for the same DOS as any D4000 series periodontal procedure code. Sealants and replacement sealants are limited to one every 3 years per tooth by the same provider or provider group. Dental sealants performed more frequently than once every three years by a different provider are also a benefit if the different provider is not associated with the provider or provider group that initially placed the sealant on the tooth. If submitted on emergency claim, procedure code will be denied. A Birth–20</td>
</tr>
<tr>
<td>D1352</td>
<td>Denied if a caries risk assessment (procedure code D0602 or D0603) has not been submitted, by any provider, within 180 days prior. Denied when submitted for the same DOS as any D4000 series periodontal procedure code. A 5–20</td>
</tr>
<tr>
<td><strong>Space Maintenance (Passive Appliances)</strong></td>
<td></td>
</tr>
<tr>
<td>Space maintainers are a benefit of Texas Medicaid after premature loss of primary or secondary molars (TID A, B, I, J, K, L, S, and T for clients who are 1 through 12 years of age, and after loss of permanent molars (TID 3, 14, 19, and 30) for clients who are 3 through 20 years of age. Limited to 1 space maintainer per TID, per lifetime, per client.</td>
<td></td>
</tr>
<tr>
<td>When procedure code D1510, D1516, or D1517 have been previously reimbursed, the recementation of space maintainers (procedure code D1551, D1552, or D1553) may be considered for reimbursement to either the same or different THSteps dental provider. Replacement space maintainers may be considered upon appeal with documentation supporting medical necessity. Removal of a fixed space maintainer (procedure code D1556, D1557, or D1558) is not payable to the provider or dental group practice that originally placed the device. Procedure codes D1553 and D1556 are limited to once per quadrant, per day, same provider.</td>
<td></td>
</tr>
<tr>
<td><strong>Space Maintenance (Passive Appliances) continued</strong></td>
<td></td>
</tr>
<tr>
<td>D1516*</td>
<td>A 1-20 (TIDs #A, B, I, J), MTID</td>
</tr>
</tbody>
</table>

A=Age range limitations, N=Narrative required, FMX=Full-mouth radiographs (nonpanoramic), MTID=Missing tooth ID(s), PPXR=Preoperative and postoperative radiographs required, PXR=Preoperative radiographs required, PHO=Preoperative and postoperative photographs required, PC=Periodontal charting required, PATH=Pathology report required and must be retained in the client’s record, CCP=Comprehensive Care Program, NC=No charge to Medicaid and may not bill the client, and *= Services payable to an FQHC for a client encounter.
4.2.3.9 Therapeutic Services

Medicaid reimbursement is contingent on compliance with the following limitations:

- Documentation requirements
  
  *Refer to:* Subsection 4.3, “Documentation Requirements” in this handbook.

- More than one restoration on a single surface is considered a single restoration.

- Multiple surface restorations must show definite crossing of the plane of each surface listed for each primary and permanent tooth completed.

- A multiple surface restoration cannot be submitted as two or more separate one-surface restorations.

- Restorations and therapeutic care are provided as a Medicaid service based on medical necessity and reimbursed only for therapeutic reasons and not preventive purposes (refer to CDT).

All dental restorations and prosthetic appliances that require lab fabrication may be submitted for reimbursement using the date the final impression was made as the DOS. If the client did not return for final seating of the restoration or appliance, a narrative must be included on the claim form and in the client’s chart in lieu of a postoperative radiograph. The 95-day filing deadline is in effect from the date of the final impression. If the client returns to the office after the claim has been filed, the dentist is
obligated to attempt to seat the restoration or appliance at no cost to the client or Texas Medicaid. For records retention requirements, refer to subsection 4.3, “Documentation Requirements” in this handbook.

4.2.4 Comprehensive Care Program (CCP)

The Omnibus Budget Reconciliation Act (OBRA) of 1989 mandated the expansion of the federal EPSDT program to include any service that is medically necessary and for which federal financial participation (FFP) is available, regardless of the limitations of Texas Medicaid. This expansion is referred to as the Comprehensive Care Program (CCP).

CCP services are provided only for those clients who are birth through 20 years of age who are eligible to receive THSteps services. When the client becomes 21 years of age, all CCP benefits stop. Dental services that are a benefit through CCP are designated in the Limitations column of the tables with the notation “CCP” beginning in subsection 4.2.3.7, “Diagnostic Services” in this handbook.

4.2.5 Children’s Medicaid Dental Plan Choices

Children’s Medicaid dental services benefits are administered by three dental managed care organizations (i.e., dental plans) across the state of Texas.

<table>
<thead>
<tr>
<th>Medicaid Managed Care Dental Plan</th>
<th>Dental Plan Provider Services</th>
</tr>
</thead>
<tbody>
<tr>
<td>DentaQuest</td>
<td>1-800-685-9971</td>
</tr>
<tr>
<td>MCNA Dental</td>
<td>1-855-776-6262</td>
</tr>
<tr>
<td>United Healthcare Dental</td>
<td>1-800-527-1764</td>
</tr>
</tbody>
</table>

Note: Services provided to Medicaid managed care clients must be provided by their main dentist.

4.2.6 Authorization Transfers for Medicaid Managed Care Dental Orthodontic Services

If a client transitions to a managed care dental plan after their orthodontic services were initially authorized by TMHP, the claims for the orthodontic services will be processed and reimbursed by the managed care dental plan. Providers should check client eligibility to identify the managed care dental plan to which the client transitions.

Claims for orthodontic services remain the responsibility of the dental managed care plan until the authorized services are completed, even if the client loses dental managed care or Medicaid eligibility.

4.2.7 ICF-IID Dental Services

ICF-IID dental services are mandated by Medicaid. Reimbursement is provided for treatment of dental problems for Medicaid-eligible residents of ICF-IID facilities who are 21 years of age and older. Residents of ICF-IID facilities who are 20 years of age and younger receive services through the regular THSteps Program. Eligibility for ICF-IID services is determined by HHSC.

Procedure codes that do not have a CCP designation in the “Limitations” column of the dental fee schedule may be submitted in a routine manner for ICF-IID clients. These procedures must be documented as medically necessary and appropriate. ICF-IID clients are not subject to periodicity for preventive care. For procedure codes that have a CCP designation, a provider may request authorization with documentation or provide documentation on the submitted claim.

Refer to: Subsection 4.2.14, “Medicaid Dental Benefits, Limitations, and Fee Schedule” in this handbook.
4.2.7.1 THSteps and ICF-IID Provision of Dental Services

All THSteps and ICF-IID dental services must be performed by the Medicaid-enrolled dental provider except for permissible work that is delegated to a licensed dental hygienist, dental assistant, or dental technician in a dental laboratory on the premises where the dentist practices, or in a commercial laboratory registered with the TSBDE. The Texas Dental Practice Act and the rules and regulations of the TSBDE (22 TAC, Part 5) define the scope of work that dental auxiliary personnel may perform. Any deviations from these practice limitations shall be reported to the TSBDE and HHSC, and could result in sanctions or other actions imposed against the provider.

THSteps and ICF-IID clients must receive:

- Dental services specified in the treatment plan that meet the standards of care established by the laws relating to the practice of dentistry and the rules and regulations of the TSBDE.
- Dental services free from abuse or harm from the provider or the provider’s staff.
- Only the treatment required to address documented medical necessity that meets professionally recognized standards of health care.

4.2.7.2 Children in Foster Care

Clients in foster care receive services from Superior HealthPlan’s dental contractor. Providers may contact DentaQuest at 1-888-308-9345 for more information.

Paper claims and requests for prior authorization must be mailed to:

DentaQuest
12121 North Corporate Parkway
Mequon, WI 53092
Fax: 1-262-241-7150 or 1-888-313-2883

4.2.8 Written Informed Consent and Standards of Care

As outlined in 22 TAC §108.7, the dental provider must maintain written informed consent signed by the patient, or a parent or legal guardian of the patient if the patient is a minor, or a legal guardian of the patient if the patient has been adjudicated incompetent to manage the patient’s personal affairs.

Additionally, as required in 25 TAC §33.6 and §33.20, THSteps providers must obtain legally effective, written informed consent before providing THSteps dental checkups and treatment services. Such consent is required for all oral evaluations; dental diagnostic, preventative, and therapeutic services; and treatment plans. The written informed consent must identify the tooth and surface IDs associated with the proposed treatment and should disclose risks or hazards that could influence a reasonable person in making a decision to give or withhold consent.

THSteps clients or their parents or legal guardians who can give written informed consent must receive information following a dental examination about the dental diagnosis, scope of proposed treatment, including alternatives and risks, anticipated results, and need for and risks of the administration of sedation or anesthesia. Additionally, they must receive a full explanation of the treatment plan and give written informed consent before treatment is initiated. The parent or guardian being present at the time of the dental visit facilitates the provider obtaining written informed consent. Dentists must comply with TSBDE Rule 22 TAC §108.2, “Fair Dealing.”

4.2.9 First Dental Home

Based on the American Academy of Pediatric Dentistry’s (AAPD) definition, Texas Medicaid defines a dental home as the dental provider who supports an ongoing relationship with the client that includes all aspects of oral health care delivered in a comprehensive, continuously accessible, coordinated, and family-centered way. Establishment of a client’s dental home begins no later than 6 months of age and includes referrals to dental specialists when appropriate.
In providing a dental home for a client, the dental provider enhances the ability to assist clients and their parents in obtaining optimum oral health care. The first dental home visit can be initiated as early as 6 months of age and must include, but is not limited to, the following:

- Comprehensive oral examination
- Oral hygiene instruction with primary caregiver
- Dental prophylaxis, if appropriate
- Topical fluoride varnish application when teeth are present
- Caries risk assessment
- Dental anticipatory guidance

Clients who are from 6 through 35 months of age may be seen for dental checkups by a certified First Dental Home provider.

First Dental Home services are submitted using procedure code D0145. The dental home provider must retain supporting documentation for procedure code D0145 in the client’s record. The supporting documentation must include, but is not limited to, the following:

- Oral and physical health history review
- Dental history review
- Primary caregiver’s oral health
- Oral evaluation
- Caries risk assessment
- Dental prophylaxis, which may include a toothbrush prophylaxis
- Oral hygiene instruction with parent or caregiver
- Fluoride varnish application
- An appropriate preventive oral health regimen (recall schedule)
- Anticipatory guidance communicated to the client’s parent, legal guardian, or primary caregiver to include the following:
  - Oral health and home care
  - Oral health of primary caregiver/other family members
  - Development of mouth and teeth
  - Oral habits
  - Diet, nutrition, and food choices
  - Fluoride needs
  - Injury prevention
  - Medications and oral health
  - Any referrals, including dental specialist’s name

Procedure code D0145 is limited to individual dentists certified by Texas Health Steps to perform this service. Training for certification as a First Dental Home provider is available as a free continuing education course on the THSteps website at www.txhealthsteps.com.
Procedure codes D0120, D0150, D0160, D0170, D0180, D1120, D1206, D1208, and D8660 are denied if procedure code D0145 is submitted for the same DOS by any provider. A First Dental Home examination is limited to ten services per client lifetime with at least 60 days between visits by any provider to prevent denials of the service.

4.2.10 Dental Referrals by THSteps Primary Care Providers

Dental providers may receive referrals for clients who are 6 months of age and older from THSteps primary care providers. The primary care provider must provide information about the initiation of routine dental services with the recommendation to the client’s parent or guardian that an appointment be scheduled with a dental provider in order to establish a dental home. If a THSteps dental checkup reveals a dental health condition that requires follow-up diagnosis or treatment, the provider performing the dental checkup should assist the client in planning follow-up care within their practice or in making a referral to another qualified dental provider.

Note: For clients who are 20 years of age and younger, the client’s guardian may refer the client for dental services or a client of legal age may refer themselves for dental services.

4.2.11 Change of Provider

A provider may refer a client to another dental provider for treatment for any of the following reasons:

- Treatment by a dental specialist such as a pediatric dentist, periodontist, oral surgeon, endodontist, or orthodontist is indicated and is in the best interests of the THSteps client.
- The services needed are outside the skills or scope of practice of the initial provider.

A provider may discontinue treatment if there is documented failure to keep appointments by the client, noncompliance with the treatment plan, or conflicts with the client or other family members. In any such action to discontinue treatment, providers must comply with 22 TAC §108.5, "Patient Abandonment."

The client also may select another provider, if desired. HHSC may refer the client to another provider as a result of adverse information obtained during a utilization review or resolution of a complaint from either provider or client.

4.2.11.1 Interrupted or Incomplete Orthodontic Treatment Plans

Authorizations for orthodontic or extensive restorative treatment plans that have been prior authorized for a provider are not transferable to another provider. If a client’s treatment plan is interrupted and the services are not completed, the original or new provider must request a new prior authorization to complete the interrupted, incomplete, and prior authorized treatment plan.

To complete the treatment plan, the client must be eligible for Medicaid. It is the provider’s responsibility to verify the client’s eligibility through TexMedConnect, the Medicaid Client Portal for Providers, or the TMHP Contact Center.

If the client does not return for the completion of services and there is a documented failure to keep appointments by the client, the dental provider who initiated the services may submit a claim for reimbursement in compliance with the 95-day filing deadline.

Refer to: Subsection 4.2.24.4, “Premature Termination of Comprehensive Orthodontic Treatment” in this section.

4.2.12 * Periodicity for THSteps Dental Services

[Revised] For clients who are 6 months through 20 years of age, dental checkups may occur at 6-month (181-day) intervals. Texas Medicaid has adopted the AAPD’s “Guideline on Periodicity of Examination, Preventive Dental Services, Anticipatory Guidance/Counseling, and Oral Treatment for Infants, Children, and Adolescents” to serve as a guide and reference for dentists when scheduling and providing services to THSteps clients.
In November 2004, the ADA, in conjunction with the FDA, established “Guidelines for Prescribing Dental Radiographs.” The guidelines include type of encounters relevant to the client’s age and dental developmental stage. Texas Medicaid has adopted the ADA guidelines to serve as a guide and reference for dentists who treat THSteps clients.


THSteps dental providers may provide any medically necessary dental services such as emergency, diagnostic, preventive, therapeutic, and orthodontic services that are within the Texas Medicaid guidelines and limitations specified for each area as long as the client’s Medicaid eligibility is current for the date that dental services are being provided.

4.2.12.1 Exceptions to Periodicity

If a periodic dental checkup has been conducted within the last six months, the client still may be able to receive another periodic dental checkup in the same six-month period by any provider. For THSteps clients, exceptions to the six-month periodicity schedule for dental checkup services may be approved for one of the following reasons:

- Medically necessary service, based on risk factors and health needs (includes clients who are birth through 6 months of age).
- Required to meet federal or state exam requirements for Head Start, daycare, foster care, preadoption, or to provide a checkup prior to the next periodically-due checkup if the client will not be available when due. This includes clients whose parents are migrant or seasonal workers.
- Clients’ choice to request a second opinion or change service providers (not applicable to referrals).
- Subsequent therapeutic services necessary to complete a case for clients who are 5 months of age and younger when initiated as emergency services, for trauma, or early childhood caries.
- Medical checkup prior to a dental procedure requiring general anesthesia.
- A First Dental Home client can be seen up to ten times within the age of 6 through 35 months.

It is the provider’s responsibility to verify that the client is eligible for the date that dental services are to be provided. Eligibility may be verified through TexMedConnect, the Medicaid Client Portal for Providers, or the TMHP Contact Center.

When the need for an exception to periodicity is established, a narrative explaining the reason for the exception to periodicity limitations must be documented in the client’s file and on the claim submission. For claims filed electronically, check “yes” when prompted. For claims filed on paper, place comments in Block 35.

For ICF-IID clients who are 21 years of age and older, the periodicity schedule for preventive dental procedures (exams, prophylaxis, fluoride, and radiographs) does not apply.
4.2.13 Tooth Identification (TID) and Surface Identification (SID) Systems

Claims are denied if the procedure code is not compatible with TID or SID. Use the alpha characters to describe tooth surfaces or any combination of surfaces. For SID designation on anterior teeth, use facial (F) and incisal (I). For SID purposes, use buccal (B) and occlusal (O) designations for posterior teeth.

4.2.13.1 Supernumerary Tooth Identification

Each identified permanent tooth and each identified primary tooth has its own identifiable supernumerary number. This developed system can be found in the Current Dental Terminology (CDT) published by the ADA.

The TID for each identified supernumerary tooth will be used for paper and electronic claims and can only be submitted for payment with the following procedure codes:

- For primary teeth only: D7111.
- For both primary and permanent teeth the following codes can be submitted: D7140, D7210, D7220, D7230, D7240, D7241, D7250, D7285, D7286, and D7510.

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<th>Tooth #</th>
<th>T</th>
<th>S</th>
<th>R</th>
<th>Q</th>
<th>P</th>
<th>O</th>
<th>N</th>
<th>M</th>
<th>L</th>
<th>K</th>
</tr>
</thead>
<tbody>
<tr>
<td>Super #</td>
<td>TS</td>
<td>SS</td>
<td>RS</td>
<td>QS</td>
<td>PS</td>
<td>OS</td>
<td>NS</td>
<td>MS</td>
<td>LS</td>
<td>KS</td>
</tr>
</tbody>
</table>
4.2.14  Medicaid Dental Benefits, Limitations, and Fee Schedule

For THSteps clients, dental procedure limitations may be waived when all the following have been met. The dental procedure is:

- Medically necessary and FFP is available for it.
- Prior authorized by the TMHP Dental Director.
- Properly documented in the client’s record.

Refer to: Subsection 4.3, “Documentation Requirements” in this handbook.

For ICF-IID clients, services designated as CCP-type are available. In the “Limitations” column of the fee schedule, abbreviations indicate the age range limitations and documentation requirements. The following abbreviations also appear in a table at the bottom of each page of the fee schedule:

<table>
<thead>
<tr>
<th>Acronym</th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td>A</td>
<td>Age range limitations</td>
</tr>
<tr>
<td>CCP</td>
<td>Payable under CCP for clients who are 20 years of age and younger when THSteps benefits or limits are exceeded</td>
</tr>
<tr>
<td>DOS</td>
<td>Date of service</td>
</tr>
<tr>
<td>FMX</td>
<td>Intraoral radiographs—complete series</td>
</tr>
<tr>
<td>MTID</td>
<td>Missing tooth ID(s)</td>
</tr>
<tr>
<td>N</td>
<td>Narrative of medical necessity for the procedure must be retained in the client’s record</td>
</tr>
<tr>
<td>NC</td>
<td>Not reimbursed by Medicaid. Services may not be charged to the client.</td>
</tr>
<tr>
<td>PATH</td>
<td>Pathology report must accompany the claim and must be retained in the client’s record</td>
</tr>
<tr>
<td>PC</td>
<td>Periodontal charting must be retained in the client’s record</td>
</tr>
<tr>
<td>PHO</td>
<td>Preoperative and postoperative photographs required and must be maintained in the client’s medical record</td>
</tr>
<tr>
<td>PPXR</td>
<td>Preoperative and postoperative radiographs required when the procedure is performed and must be retained in the client’s record; do not send with initial claims</td>
</tr>
<tr>
<td>PXR</td>
<td>Preoperative radiographs are required when the procedure is performed and must be retained in the client’s record; do not send with initial claims</td>
</tr>
</tbody>
</table>

4.2.15  Restorative Services

<table>
<thead>
<tr>
<th>Procedure Code</th>
<th>Limitations</th>
</tr>
</thead>
<tbody>
<tr>
<td>D2140*</td>
<td>A Birth–20, PXR</td>
</tr>
<tr>
<td>D2150*</td>
<td>A Birth–20, PXR</td>
</tr>
<tr>
<td>D2160*</td>
<td>A 1–20, PXR</td>
</tr>
<tr>
<td>D2161*</td>
<td>A 1–20, PXR</td>
</tr>
</tbody>
</table>

Procedure codes D2140, D2150, D2160, and D2161 are benefits for clients who are 21 years of age or older residing in an ICF-IID facility.

A=Age range limitations, N=Narrative required, FMX=Full-mouth radiographs (nonpanoramic), MTID=Missing tooth ID(s), PPXR=Preoperative and postoperative radiographs required, PXR=Preoperative radiographs required, PHO=Preoperative and postoperative photographs required, PC=Periodontal charting required, PATH=Pathology report required and must be retained in the client’s record, CCP=Comprehensive Care Program, NC=No charge to Medicaid and may not bill the client, *= Services payable to an FQHC for a client encounter
## Resin-Based Composite Restorations—Direct

<table>
<thead>
<tr>
<th>Procedure Code</th>
<th>Limitations</th>
</tr>
</thead>
<tbody>
<tr>
<td>D2390*</td>
<td>A Birth–20, PXR</td>
</tr>
<tr>
<td>D2391*</td>
<td>A Birth–20, PXR</td>
</tr>
<tr>
<td>D2392*</td>
<td>A Birth–20, PXR</td>
</tr>
<tr>
<td>D2393*</td>
<td>A 1–20, PXR</td>
</tr>
<tr>
<td>D2394*</td>
<td>A 1–20, PXR</td>
</tr>
</tbody>
</table>

## Inlay/Onlay Restorations (Permanent Teeth only)

For procedure codes D2510 through D2664, inlay/onlay (permanent teeth only), porcelain is allowed on all teeth. Prior authorization is required for all inlays/onlays or permanent crowns. Procedure codes D2542, D2543, D2544, and D2662 through D2664 are payable once per client, per tooth every ten years.

<table>
<thead>
<tr>
<th>Procedure Code</th>
<th>Limitations</th>
</tr>
</thead>
<tbody>
<tr>
<td>D2510</td>
<td>A 13–20, N, PPXR, CCP</td>
</tr>
<tr>
<td>D2520</td>
<td>A 13–20, N, PPXR, CCP</td>
</tr>
<tr>
<td>D2530</td>
<td>A 13–20, N, PPXR, CCP</td>
</tr>
<tr>
<td>D2542</td>
<td>Same as D2520. A 13–20, N, PPXR, CCP</td>
</tr>
<tr>
<td>D2543</td>
<td>All materials accepted. A 13–20, N, PPXR, CCP</td>
</tr>
<tr>
<td>D2544</td>
<td>All materials accepted. A 13–20, N, PPXR, CCP</td>
</tr>
<tr>
<td>D2650</td>
<td>All materials accepted. A 13–20, N, PPXR, CCP</td>
</tr>
<tr>
<td>D2651</td>
<td>All materials accepted. A 13–20, N, PPXR, CCP</td>
</tr>
<tr>
<td>D2652</td>
<td>All materials accepted. A 13–20, N, PPXR, CCP</td>
</tr>
<tr>
<td>D2662</td>
<td>All materials accepted. A 13–20, N, PPXR, CCP</td>
</tr>
<tr>
<td>D2663</td>
<td>All materials accepted. A 13–20, N, PPXR, CCP</td>
</tr>
<tr>
<td>D2664</td>
<td>All materials accepted. A 13–20, N, PPXR, CCP</td>
</tr>
</tbody>
</table>

A=Age range limitations, N=Narrative required, FMX=Full-mouth radiographs (nonpanoramic), MTID=Missing tooth ID(s), PPXR=Preoperative and postoperative radiographs required, PXR=Preoperative radiographs required, PHO=Preoperative and postoperative photographs required, PC=Periodontal charting required, PATH=Pathology report required and must be retained in the client's record, CCP=Comprehensive Care Program, NC=No charge to Medicaid and may not bill the client, and *= Services payable to an FQHC for a client encounter.
## Crowns—Single Restorations Only

For procedure codes D2710 through D2794, single crown restorations (permanent teeth only), the following limitations apply:

- Prior authorization is required for codes D2710 through D2794.
- Reimbursement for crowns and onlay restorations require submission of post-operative bitewing radiograph(s) (for posterior teeth); post-operative periapical radiograph(s) (for anterior teeth) will need to be submitted with the claim to verify that the restoration meets the standard of care.
- Radiographs are reviewed to verify that the restoration meets both medical necessity and standard of care to approve reimbursement.
- Reimbursement for crowns and onlay restorations are payable once per client, per tooth every ten years.

<table>
<thead>
<tr>
<th>Procedure Code</th>
<th>Limitations</th>
</tr>
</thead>
<tbody>
<tr>
<td>D2710</td>
<td>All materials accepted. A 13–20, N, PPXR, CCP</td>
</tr>
<tr>
<td>D2720</td>
<td>All materials accepted. A 13–20, N, PPXR, CCP</td>
</tr>
<tr>
<td>D2721</td>
<td>All materials accepted. A 13–20, N, PPXR, CCP</td>
</tr>
<tr>
<td>D2722</td>
<td>All materials accepted. A 13–20, N, PPXR, CCP</td>
</tr>
<tr>
<td>D2740</td>
<td>All materials accepted. A 13–20, N, PPXR, CCP Limited to TID #4–13 and 20–29 only.</td>
</tr>
<tr>
<td>D2750*</td>
<td>All materials accepted. A 13–20, N, PPXR, CCP Limited to TID #4–13 and 20–29 only.</td>
</tr>
<tr>
<td>D2751*</td>
<td>All materials accepted. A 13–20, N, PPXR, CCP Limited to TID #4–13 and 20–29 only.</td>
</tr>
<tr>
<td>D2752</td>
<td>All materials accepted. A 13–20, N, PPXR, CCP Limited to TID #4–13 and 20–29 only.</td>
</tr>
<tr>
<td>D2780</td>
<td>A 13–20, N, PPXR, CCP</td>
</tr>
<tr>
<td>D2781</td>
<td>A 13–20, N, PPXR, CCP</td>
</tr>
<tr>
<td>D2782</td>
<td>A 13–20, N, PPXR, CCP</td>
</tr>
<tr>
<td>D2783</td>
<td>Anterior teeth only (#6–11 and 22–27). A 13–20, N, PPXR, CCP</td>
</tr>
<tr>
<td>D2790</td>
<td>Posterior teeth only (#1–5, 12–21, and 28–32). All materials accepted. A 13–20, N, PPXR, CCP</td>
</tr>
<tr>
<td>D2791*</td>
<td>Posterior teeth only (#1–5, 12–21, and 28–32). All materials accepted. A 13–20, N, PPXR</td>
</tr>
<tr>
<td>D2792*</td>
<td>Posterior teeth only (#1–5, 12–21, and 28–32). All materials accepted. A 13–20, N, PPXR, CCP</td>
</tr>
<tr>
<td>D2794</td>
<td>A 13–20, N, PPXR, CCP</td>
</tr>
</tbody>
</table>

### Other Restorative Services

<table>
<thead>
<tr>
<th>Procedure Code</th>
<th>Limitations</th>
</tr>
</thead>
<tbody>
<tr>
<td>D2910</td>
<td>A 13–20, PXR</td>
</tr>
<tr>
<td>D2915</td>
<td>A 4–20</td>
</tr>
<tr>
<td>D2920</td>
<td>A 1–20, PXR</td>
</tr>
<tr>
<td>D2930*</td>
<td>A Birth–20, PXR</td>
</tr>
</tbody>
</table>

A=Age range limitations, N=Narrative required, FMX=Full-mouth radiographs (nonpanoramic), MTID=Missing tooth ID(s), PPXR=Preoperative and postoperative radiographs required, PXR=Preoperative radiographs required, PHO=Preoperative and postoperative photographs required, PC=Periodontal charting required, PATH=Pathology report required and must be retained in the client’s record, CCP=Comprehensive Care Program, NC=No charge to Medicaid and may not bill the client, and *= Services payable to an FQHC for a client encounter.
<table>
<thead>
<tr>
<th>Procedure Code</th>
<th>Limitations</th>
</tr>
</thead>
<tbody>
<tr>
<td>D2931*</td>
<td>A 1–20, PXR. A benefit for clients who are 21 years of age and older residing in an ICF–IID facility.</td>
</tr>
<tr>
<td>D2932*</td>
<td>A 1–20, PXR (TID #C–H, M–R, and 1-32), all permanent teeth. A benefit for clients who are 21 years of age and older residing in an ICF–IID facility.</td>
</tr>
<tr>
<td>D2933*</td>
<td>Limited to anterior primary teeth only (TID #C–H, M–R). A Birth–20, N, CCP, PXR</td>
</tr>
<tr>
<td>D2934*</td>
<td>Limited to anterior primary teeth only (TID #C–H, M–R). A Birth–20, N, CCP, PXR</td>
</tr>
<tr>
<td>D2940*</td>
<td>Not allowed on the same date as permanent restoration. A Birth–20, PXR</td>
</tr>
<tr>
<td>D2950*</td>
<td>Provider payments received in excess of $45.00 for restorative work performed within six months of a crown procedure on the same tooth will be deducted from the subsequent crown procedure reimbursement. Not allowed on primary teeth. A 4–20, N, CCP, PXR</td>
</tr>
<tr>
<td>D2951</td>
<td>Not allowed on primary teeth. A 4–20, PXR</td>
</tr>
<tr>
<td>D2952</td>
<td>Not payable with D2950. Not allowed on primary teeth. A 13–20, CCP, PXR</td>
</tr>
<tr>
<td>D2953</td>
<td>Must be used with D2952. Not allowed on primary teeth. A 13–20</td>
</tr>
<tr>
<td>D2954*</td>
<td>Not payable with D2952 or D3950 on the same TID by the same provider. Not allowed on primary teeth. A 13–20, N, CCP, PXR</td>
</tr>
<tr>
<td>D2955</td>
<td>For removal of posts (for example, fractured posts) not to be used in conjunction with endodontic retreatment (D3346, D3347, D3348). Not allowed on primary teeth. A 4–20, CCP, PXR</td>
</tr>
<tr>
<td>D2957</td>
<td>Must be used with D2954. Not allowed on primary teeth. A 13–20, PXR, CCP</td>
</tr>
<tr>
<td>D2960</td>
<td>A 13–20, N, PPXR, CCP</td>
</tr>
<tr>
<td>D2961</td>
<td>A 13–20, N, PPXR, CCP</td>
</tr>
<tr>
<td>D2962</td>
<td>A 13–20, N, PPXR, CCP</td>
</tr>
<tr>
<td>D2971*</td>
<td>May be reimbursed up to four services per lifetime for each tooth. Payable to any THSteps dental provider who performed the original cementation of the crown. A 13–20</td>
</tr>
<tr>
<td>D2980</td>
<td>A 1–20, PXR (permanent teeth only)</td>
</tr>
<tr>
<td>D2999</td>
<td>A 1–20, N, CCP, PXR</td>
</tr>
</tbody>
</table>

A=Age range limitations, N=Narrative required, FMX=Full-mouth radiographs (nonpanoramic), MTID=Missing tooth ID(s), PPXR=Preoperative and postoperative radiographs required, PXR=Preoperative radiographs required, PHO=Preoperative and postoperative photographs required, PC=Periodontal charting required, PATH=Pathology report required and must be retained in the client’s record, CCP=Comprehensive Care Program, NC=No charge to Medicaid and may not bill the client, and *= Services payable to an FQHC for a client encounter

The following dental restoration procedure codes will be limited to once per rolling year, for the same TID, by the same provider:

<table>
<thead>
<tr>
<th>Procedure Codes</th>
</tr>
</thead>
<tbody>
<tr>
<td>D2140</td>
</tr>
<tr>
<td>D2394</td>
</tr>
</tbody>
</table>
Procedure codes D2335 and D2390 when provided to primary teeth will be limited to once per lifetime, same TID, any provider, and will be denied if any of the following anterior restoration procedure codes have been paid within a rolling year, for the same TID, by the same provider:

<table>
<thead>
<tr>
<th>Procedure Codes</th>
</tr>
</thead>
<tbody>
<tr>
<td>D2140</td>
</tr>
<tr>
<td>D2932</td>
</tr>
</tbody>
</table>

Total reimbursement for direct restorations on primary teeth cannot exceed the total dollar amount for a prefabricated stainless steel crown on a primary tooth (procedure code D2930) per TID, per date of service.

Total reimbursement for direct restorations on permanent teeth cannot exceed the total dollar amount for a prefabricated stainless steel crown on a permanent tooth (procedure code D2931) per TID, per date of service. This limitation does not apply to procedure code D2335 for primary or permanent teeth.

4.2.15.1 Direct Restorations and Other Restorative Services

Direct restoration of a primary tooth with the use of a prefabricated crown will be considered as a once in a lifetime restoration, same TID, any provider. Exceptions may be considered when pre-treatment X-ray images, intra-oral photos, and narrative documentation clearly support the medical necessity for the replacement of the prefabricated crown (procedure codes D2930, D2932, D2933, and D2934) during pre-payment review.

Procedure code D2930 will be denied if the following procedure codes have been billed within a rolling year, for the same TID, by the same provider:

<table>
<thead>
<tr>
<th>Procedure Codes</th>
</tr>
</thead>
<tbody>
<tr>
<td>D2140</td>
</tr>
<tr>
<td>D2932</td>
</tr>
</tbody>
</table>

Procedure codes D2933 and D2934 will be denied if the following procedure codes have been billed within a rolling year, for the same TID, by the same provider:

<table>
<thead>
<tr>
<th>Procedure Codes</th>
</tr>
</thead>
<tbody>
<tr>
<td>D2140</td>
</tr>
</tbody>
</table>

Procedure codes D2931 and D2932 will be denied if the following procedure codes have been billed within a rolling year, for the same TID, by the same provider:

<table>
<thead>
<tr>
<th>Procedure Codes</th>
</tr>
</thead>
<tbody>
<tr>
<td>D2140</td>
</tr>
<tr>
<td>D2932</td>
</tr>
</tbody>
</table>

4.2.16 Endodontics Services

Therapeutic pulpotomy (procedure code D3220) and apexification and recalcification procedures (procedure codes D3351, D3352, and D3353) are considered part of the root canal (procedure codes D3310, D3320, and D3330) or retreatment of a previous root canal (procedure codes D3346, D3347, and D3348). When therapeutic pulpotomy or apexification and recalcification procedures are submitted with root canal codes, the reimbursement rate is adjusted to ensure that the total amount reimbursed does not exceed the total dollar amount allowed for the root canal procedure.
Reimbursement for a root canal includes all appointments necessary to complete the treatment. Pulpotomy and radiographs performed pre, intra, and postoperatively are included in the root canal reimbursement.

Root canal therapy that has only been initiated, or taken to some degree of completion, but not carried to completion with a final filling, may not be submitted as a root canal therapy code. It must be submitted using code D3999 with a narrative description of what procedures were completed in the root canal therapy.

Documentation supporting medical necessity must be kept in the client’s record and include the following: the medical necessity as documented through periapical radiographs of tooth treated showing pre-treatment, during treatment, and post-treatment status; the final size of the file to which the canal was enlarged; and the type of filling material used. Any reason that the root canal may appear radiographically unacceptable must be documented in the client’s record.

If the client is pregnant and does not want radiographs, use alternative treatment (temporary) until after delivery.

Direct pulp caps (procedure code D3110) and indirect pulp caps (procedure code D3120) are a benefit for permanent teeth only (TID 1-32).

Direct pulp caps may be reimbursed when billed with the following procedure codes for the same tooth ID on the same date of service by the same provider.

<table>
<thead>
<tr>
<th>Procedure Codes</th>
</tr>
</thead>
<tbody>
<tr>
<td>D2140</td>
</tr>
<tr>
<td>D2391</td>
</tr>
<tr>
<td>D2650</td>
</tr>
<tr>
<td>D2722</td>
</tr>
<tr>
<td>D2790</td>
</tr>
</tbody>
</table>

Indirect pulp caps (procedure code D3120) may be reimbursed when billed with procedure code D2940 for the same tooth ID, on the same date of service by the same provider.

<table>
<thead>
<tr>
<th>Procedure Code</th>
<th>Limitations</th>
</tr>
</thead>
<tbody>
<tr>
<td>Pulp Capping</td>
<td></td>
</tr>
<tr>
<td>D3110</td>
<td>A 1–20, N, PXR, CCP</td>
</tr>
<tr>
<td>D3120</td>
<td>A 1–20, N, PXR, CCP</td>
</tr>
</tbody>
</table>

A=Age range limitations, N=Narrative required, FMX=Full-mouth radiographs (nonpanoramic), MTID=Missing tooth ID(s), PPXR=Preoperative and postoperative radiographs required, PXR=Preoperative radiographs required, PHO=Preoperative and postoperative photographs required, PC=Periodontal charting required, PATH=Pathology report required and must be retained in the client’s record, CCP=Comprehensive Care Program, NC=No charge to Medicaid and may not bill the client, and *= Services payable to an FQHC for a client encounter.
### Pulpotomy

<table>
<thead>
<tr>
<th>Procedure Code</th>
<th>Limitations</th>
</tr>
</thead>
<tbody>
<tr>
<td>D3220*</td>
<td>Denied when performed within six months of D3230, D3240, D3310, D3320, or D3330 for the same primary TID, same provider. Denied when performed within six months of D3310, D3320, or D3330 on the same permanent TID, same provider. A Birth–20, PXR. Limited to once per lifetime, per primary tooth (TIDs A-T). Re-treatment claims for an incomplete pulpotomy performed by a dentist not associated with the original treating dentist or dental group will be considered for reimbursement upon appeal. Documentation of medical necessity and the incomplete initial pulpotomy must be submitted with the appeal. The appeal must include a written narrative and pre- and post-treatment X-rays, which will be reviewed by a Texas licensed dentist. <strong>Note:</strong> The identified, original treating dentist or dental group will not be considered for reimbursement.</td>
</tr>
</tbody>
</table>

### Endodontic Therapy on Primary Teeth

<table>
<thead>
<tr>
<th>Procedure Code</th>
<th>Limitations</th>
</tr>
</thead>
<tbody>
<tr>
<td>D3230*</td>
<td>Anterior primary incisors and cuspids. TIDs #C–H, M–R. A 1–20, PXR</td>
</tr>
</tbody>
</table>

### Endodontic Therapy (including Treatment Plan, Clinical Procedures, and Follow-up Care)

<table>
<thead>
<tr>
<th>Procedure Code</th>
<th>Limitations</th>
</tr>
</thead>
<tbody>
<tr>
<td>D3310*</td>
<td>A 6–20, PPXR</td>
</tr>
<tr>
<td>D3320*</td>
<td>A 6–20, PPXR</td>
</tr>
<tr>
<td>D3330*</td>
<td>A 6–20, PPXR</td>
</tr>
</tbody>
</table>

### Endodontic Retreatment

<table>
<thead>
<tr>
<th>Procedure Code</th>
<th>Limitations</th>
</tr>
</thead>
<tbody>
<tr>
<td>D3346*</td>
<td>A 6–20, PPXR</td>
</tr>
<tr>
<td>D3347*</td>
<td>A 6–20, PPXR</td>
</tr>
<tr>
<td>D3348*</td>
<td>A 6–20, PPXR</td>
</tr>
</tbody>
</table>

### Apexification/Recalcification Procedures

<table>
<thead>
<tr>
<th>Procedure Code</th>
<th>Limitations</th>
</tr>
</thead>
<tbody>
<tr>
<td>D3351*</td>
<td>A 6–20, N, PXR, CCP</td>
</tr>
<tr>
<td>D3352*</td>
<td>A 6–20, N, PXR, CCP</td>
</tr>
<tr>
<td>D3353*</td>
<td>A 6–20, PXR, CCP</td>
</tr>
</tbody>
</table>

### Apicoectomy/Periradicular Services

<table>
<thead>
<tr>
<th>Procedure Code</th>
<th>Limitations</th>
</tr>
</thead>
<tbody>
<tr>
<td>D3410</td>
<td>A 6–20, N, PPXR, CCP</td>
</tr>
<tr>
<td>D3421</td>
<td>A 6–20, N, PPXR, CCP</td>
</tr>
<tr>
<td>D3425</td>
<td>A 6–20, N, PPXR, CCP</td>
</tr>
<tr>
<td>D3426</td>
<td>A 6–20, N, PPXR, CCP</td>
</tr>
<tr>
<td>D3430</td>
<td>A 6–20, N, PPXR, CCP</td>
</tr>
<tr>
<td>D3450</td>
<td>A 6–20, N, PXR, CCP</td>
</tr>
<tr>
<td>D3460</td>
<td>Prior authorization required. Submit request with periapical radiographs, for each tooth involved. A 16–20, N, PPXR, CCP</td>
</tr>
</tbody>
</table>

### Procedure Code Limitations

A=Age range limitations, N=Narrative required, FMX=Full-mouth radiographs (nonpanoramic), MTID=Missing tooth ID(s), PPXR=Preoperative and postoperative radiographs required, PXR=Preoperative radiographs required, PHO=Preoperative and postoperative photographs required, PC=Periodontal charting required, PATH=Pathology report required and must be retained in the client’s record, CCP=Comprehensive Care Program, NC=No charge to Medicaid and may not bill the client, and *= Services payable to an FQHC for a client encounter
4.2.17 Periodontal Services

Procedure codes D4210 and D4211, when submitted for clients who are 12 years of age and younger, will be initially denied, but may be appealed with documentation of medical necessity. Preoperative and postoperative photographs are required for the following procedure codes: D4210, D4211, D4270, D4273, D4275, D4276, D4277, D4278, D4283, D4285, D4355, and D4910.

Procedure codes D4283 and D4285 are limited to three teeth per site, same day same provider. Procedure code D4283 must be billed along with procedure code D4273 and procedure code D4285 must be billed along with procedure code D4275 on the same claim, for the same date of service, by the same provider.

Preoperative and postoperative photographs are required when medical necessity is not evident on radiographs for the following procedure codes: D4240, D4241, D4245, D4266, and D4267. Documentation is required when medical necessity is not evident on radiographs for the following procedure codes: D4210, D4211, D4240, D4241, D4245, D4266, D4267, D4270, D4273, D4275, D4276, D4277, D4278, D4283, D4285, D4355, and D4910.

Procedure code D4278 must be billed on the same date of service as procedure code D4277 or the service will be denied.

Full mouth debridement (procedure code D4355) will be denied when submitted for the same date of service as the following procedure codes by any provider:

<table>
<thead>
<tr>
<th>Procedure Code</th>
<th>Limitations</th>
</tr>
</thead>
<tbody>
<tr>
<td>D3470</td>
<td>A 6–20, N, PXR, CCP</td>
</tr>
<tr>
<td>D3910</td>
<td>A 1–20, N, CCP</td>
</tr>
<tr>
<td>D3920</td>
<td>A 6–20, N, PXR, CCP</td>
</tr>
<tr>
<td>D3950</td>
<td>A 6–20, N, PXR, CCP</td>
</tr>
<tr>
<td>D3999</td>
<td>A 1–20, N, PXR, CCP</td>
</tr>
</tbody>
</table>

Other Endodontic Procedures

<table>
<thead>
<tr>
<th>Procedure Code</th>
<th>Limitations</th>
</tr>
</thead>
<tbody>
<tr>
<td>D3910</td>
<td>A 1–20, N, CCP</td>
</tr>
<tr>
<td>D3920</td>
<td>A 6–20, N, PXR, CCP</td>
</tr>
<tr>
<td>D3950</td>
<td>A 6–20, N, PXR, CCP</td>
</tr>
<tr>
<td>D3999</td>
<td>A 1–20, N, PXR, CCP</td>
</tr>
</tbody>
</table>

Limitations:
A=Age range limitations, N=Narrative required, FMX=Full-mouth radiographs (nonpanoramic), MTID=Missing tooth ID(s), PPXR=Preoperative and postoperative radiographs required, PXR=Preoperative radiographs required, PHO=Preoperative and postoperative photographs required, PC=Periodontal charting required, PATH=Pathology report required and must be retained in the client’s record, CCP=Comprehensive Care Program, NC=No charge to Medicaid and may not bill the client, and *= Services payable to an FQHC for a client encounter

Claims for preventive dental procedure codes D1110, D1120, D1206, D1208, D1351, and D1351 will be denied when submitted for the same DOS as any D4000 series periodontal procedure codes.

Procedure codes D4266 and D4267 may be appealed with documentation of medical necessity. Medical necessity for third molar sites are:
- Medical or dental history documenting need due to inadequate healing of bone following third molar extraction, including the date of third molar extraction.
- Secondary procedure several months postextraction.
- Position of the third molar preoperatively.
- Postextraction probing depth to document continuing bony defect.
• Postextraction radiographs documenting continuing bony defect.
• Bone graft and barrier material used.

Medical necessity for other than third molar sites are:
• Medical or dental history documenting comorbid condition (e.g., juvenile diabetes, cleft palate, avulsed tooth or teeth, traumatic oral injuries).
• Intra- or extra-oral radiographs of treatment site(s).
• If not radiographically evident, intraoral photographs are optional unless requested preoperatively by HHSC or its agent.
• Periodontal probing depths.
• Number of intact walls associated with an angular bony defect.
• Bone graft and barrier material used.

<table>
<thead>
<tr>
<th>Procedure Code</th>
<th>Limitations</th>
</tr>
</thead>
<tbody>
<tr>
<td>D4210</td>
<td>A 13–20, N, PPXR, PHO, CCP</td>
</tr>
<tr>
<td>D4211</td>
<td>A 13–20, N, PHO, CCP</td>
</tr>
<tr>
<td>D4230</td>
<td>A 13–20, N, PHO, PXR, CCP</td>
</tr>
<tr>
<td>D4231</td>
<td>A 13–20, N, PHO, PXR, CCP</td>
</tr>
<tr>
<td>D4240</td>
<td>A 13–20, N, FMX, PXR, PHO when medical necessity is not evident on radiographs, PC, CCP</td>
</tr>
<tr>
<td>D4241</td>
<td>Limited to once per year. A 13–20, N, FMX, PXR, PHO when medical necessity is not evident on radiographs, PC</td>
</tr>
<tr>
<td>D4245</td>
<td>Per quadrant. A 13–20, N, PXR, PHO when medical necessity is not evident on radiographs, CCP</td>
</tr>
<tr>
<td>D4249</td>
<td>A six- to eight-week healing period following crown lengthening before final tooth preparation, impression making, and fabrication of a final restoration is required for claims submission of this code. A 13–20, N, PPXR, CCP</td>
</tr>
<tr>
<td>D4260</td>
<td>Limited to once per quadrant, per day, same provider. A 13–20, N, FMX, PXR, PC, CCP</td>
</tr>
<tr>
<td>D4261</td>
<td>Limited to once per quadrant, per day, same provider. A 13–20, N, FMX, PXR, PC</td>
</tr>
<tr>
<td>D4266</td>
<td>A 13–20, N, PXR, PHO when medical necessity is not evident on radiographs, CCP</td>
</tr>
<tr>
<td>D4267</td>
<td>A 13–20, N, PXR, PHO when medical necessity is not evident on radiographs, CCP</td>
</tr>
<tr>
<td>D4270</td>
<td>A 13–20, N, PXR, PHO, CCP</td>
</tr>
<tr>
<td>D4273</td>
<td>This procedure is performed to create or augment gingiva, to obtain root coverage or to eliminate frenum pull, or to extend the vestibular fornix. A 13–20, N, PXR, PHO, CCP</td>
</tr>
</tbody>
</table>

A=Age range limitations, N=Narrative required, FMX=Full-mouth radiographs (nonpanoramic), MTID=Missing tooth ID(s), PPXR=Preoperative and postoperative radiographs required, PXR=Preoperative radiographs required, PHO=Preoperative and postoperative photographs required, PC=Periodontal charting required, PATH=Pathology report required and must be retained in the client’s record, CCP=Comprehensive Care Program, NC= No charge to Medicaid and may not bill the client, and *= Services payable to an FQHC for a client encounter
<table>
<thead>
<tr>
<th>Procedure Code</th>
<th>Limitations</th>
</tr>
</thead>
<tbody>
<tr>
<td>D4274</td>
<td>This procedure is performed in an edentulous area adjacent to a periodontally involved tooth. Gingival incisions are used to allow removal of a tissue wedge to gain access and correct the underlying osseous defect and to permit close flap adaptation. A 13–20, N, PXR, CCP</td>
</tr>
<tr>
<td>D4275</td>
<td>Limited to once per day, same provider. A 13–20, PXR, PHO</td>
</tr>
<tr>
<td>D4276</td>
<td>Prior authorization is required. Not payable in addition to D4273 or D4275 for the same DOS. A 13–20, PXR, PHO</td>
</tr>
<tr>
<td>D4277</td>
<td>A 13–20, N, PXR, PHO, CCP</td>
</tr>
<tr>
<td>D4278</td>
<td>A 13–20, N, PXR, PHO, CCP</td>
</tr>
<tr>
<td>D4283</td>
<td>A 13–20, N, PHO</td>
</tr>
<tr>
<td>D4285</td>
<td>A 13–20, N, PHO</td>
</tr>
<tr>
<td><strong>Nonsurgical Periodontal Services</strong></td>
<td></td>
</tr>
<tr>
<td>D4320</td>
<td>A 1–20, PXR</td>
</tr>
<tr>
<td>D4321</td>
<td>A 1–20, PXR</td>
</tr>
<tr>
<td>D4341*</td>
<td>Prior authorization is required. Denied when submitted on the same date of service as D4355. Denied when submitted for the same DOS as other D4000 series codes. A 13–20, FMX, PC, N, CCP</td>
</tr>
<tr>
<td>D4342</td>
<td>Prior authorization is required. Denied when submitted on the same date of service as D4355. Denied when submitted for the same DOS as other D4000 series codes. A 13–20, PC, FMX, N</td>
</tr>
<tr>
<td>D4355*</td>
<td>Denied when submitted for the same DOS as other D4000 series codes. A 13–20, N, PXR, PHO, CCP</td>
</tr>
<tr>
<td>D4381</td>
<td>This procedure does not replace conventional or surgical therapy required for debridement, respective procedures, or regenerative therapy. The use of controlled-release chemotherapeutic agents is an adjunctive therapy or for cases in which systemic disease or other factors preclude conventional or surgical therapy. A 13–20, N, PXR, CCP</td>
</tr>
<tr>
<td><strong>Other Periodontal Services</strong></td>
<td></td>
</tr>
<tr>
<td>D4910</td>
<td>Payable only following active periodontal therapy by any provider as evidenced either by a submitted claim for procedure code D4240, D4241, D4260, or D4261 or by evidence through client records of periodontal therapy while not Medicaid-eligible. Not payable within 90 days after D4355, not payable for the same DOS as any other evaluation procedure. Limited to once per 12 calendar months by the same provider. A 13–20, N, PXR, PHO, CCP</td>
</tr>
<tr>
<td>D4920</td>
<td>A 13–20, N, PXR, CCP</td>
</tr>
<tr>
<td>D4999</td>
<td>A 13–20, N, PXR, CCP</td>
</tr>
</tbody>
</table>

* A=Age range limitations, N=Narrative required, FMX=Full-mouth radiographs (nonpanoramic), MTID=Missing tooth ID(s), PPXR=Preoperative and postoperative radiographs required, PXR=Preoperative radiographs required, PHO=Preoperative and postoperative photographs required, PC=Periodontal charting required, PATH=Pathology report required and must be retained in the client’s record, CCP=Comprehensive Care Program, NC=No charge to Medicaid and may not bill the client, and *= Services payable to an FQHC for a client encounter
## 4.2.18 Prosthodontic (Removable) Services

<table>
<thead>
<tr>
<th>Procedure Code</th>
<th>Limitations</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>Complete Dentures (Including Routine Post Delivery Care)</strong></td>
<td></td>
</tr>
<tr>
<td>D5110</td>
<td>A 3–20, PXR</td>
</tr>
<tr>
<td>D5120</td>
<td>A 3–20, PXR</td>
</tr>
<tr>
<td>D5130</td>
<td>A 13–20, N, PXR, CCP</td>
</tr>
<tr>
<td>D5140</td>
<td>A 13–20, N, PXR, CCP</td>
</tr>
<tr>
<td><strong>Partial Dentures (Including Routine Post Delivery Care)</strong></td>
<td></td>
</tr>
<tr>
<td>D5211*</td>
<td>A 6–20, PXR, MTID</td>
</tr>
<tr>
<td>D5212*</td>
<td>A 6–20, PXR, MTID</td>
</tr>
<tr>
<td>D5213</td>
<td>A 9–20, N, PXR, MTID, CCP</td>
</tr>
<tr>
<td>D5214</td>
<td>A 9–20, N, PXR, MTID, CCP</td>
</tr>
<tr>
<td><strong>Adjustments to Dentures</strong></td>
<td></td>
</tr>
<tr>
<td>D5410</td>
<td>A 3–20, PXR</td>
</tr>
<tr>
<td>D5411</td>
<td>A 3–20, PXR</td>
</tr>
<tr>
<td>D5421</td>
<td>A 6–20, PXR</td>
</tr>
<tr>
<td>D5422</td>
<td>A 6–20, PXR</td>
</tr>
<tr>
<td><strong>Repairs to Complete Dentures</strong></td>
<td></td>
</tr>
<tr>
<td>D5511</td>
<td>Cost of repairs cannot exceed replacement costs. A 3–20, PXR</td>
</tr>
<tr>
<td>D5512</td>
<td>Cost of repairs cannot exceed replacement costs. A 3–20, PXR</td>
</tr>
<tr>
<td>D5520</td>
<td>Cost of repairs cannot exceed replacement costs. A 3–20, PXR</td>
</tr>
<tr>
<td><strong>Repairs to Partial Dentures</strong></td>
<td></td>
</tr>
<tr>
<td>D5611*</td>
<td>A 3–20, PXR</td>
</tr>
<tr>
<td>D5612*</td>
<td>A 3–20, PXR</td>
</tr>
<tr>
<td>D5630*</td>
<td>A 6–20, PXR</td>
</tr>
<tr>
<td>D5640*</td>
<td>A 6–20, PXR</td>
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<tr>
<td>D5650*</td>
<td>A 6–20, PXR</td>
</tr>
<tr>
<td>D5660*</td>
<td>A 6–20, PXR</td>
</tr>
<tr>
<td>D5670*</td>
<td>Will be denied as part of procedure codes D5211, D5213, and D5640. A 6–20</td>
</tr>
<tr>
<td>D5671*</td>
<td>Will be denied as part of procedure codes D5212, D5214, and D5640. A 6–20</td>
</tr>
<tr>
<td><strong>Denture Rebase Procedures</strong></td>
<td></td>
</tr>
<tr>
<td>D5710</td>
<td>A 4–20, PXR</td>
</tr>
<tr>
<td>D5711</td>
<td>A 4–20, PXR</td>
</tr>
<tr>
<td>D5720*</td>
<td>A 7–20, PXR</td>
</tr>
<tr>
<td>D5721*</td>
<td>A 7–20, PXR</td>
</tr>
<tr>
<td><strong>Denture Reline Procedures</strong></td>
<td></td>
</tr>
</tbody>
</table>

A=Age range limitations, N=Narrative required, FMX=Full-mouth radiographs (nonpanoramic), MTID=Missing tooth ID(s), PPXR=Preoperative and postoperative radiographs required, PXR=Preoperative radiographs required, PHO=Preoperative and postoperative photographs required, PC=Periodontal charting required, PATH=Pathology report required and must be retained in the client’s record, CCP=Comprehensive Care Program, NC=No charge to Medicaid and may not bill the client, and *= Services payable to an FQHC for a client encounter.
<table>
<thead>
<tr>
<th>Procedure Code</th>
<th>Limitations</th>
</tr>
</thead>
<tbody>
<tr>
<td>D5730</td>
<td>A 4–20, N, PXR, CCP</td>
</tr>
<tr>
<td>D5731</td>
<td>A 4–20, N, PXR, CCP</td>
</tr>
<tr>
<td>D5740*</td>
<td>A 7–20, N, PXR, CCP</td>
</tr>
<tr>
<td>D5741*</td>
<td>A 7–20, N, PXR, CCP</td>
</tr>
<tr>
<td>D5750</td>
<td>A 4–20, PXR</td>
</tr>
<tr>
<td>D5751</td>
<td>A 4–20, PXR</td>
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<tr>
<td>D5760*</td>
<td>A 7–20, PXR</td>
</tr>
<tr>
<td>D5761*</td>
<td>A 7–20, PXR</td>
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</tbody>
</table>

**Interim Prosthesis**

<table>
<thead>
<tr>
<th>Procedure Code</th>
<th>Limitations</th>
</tr>
</thead>
<tbody>
<tr>
<td>D5810</td>
<td>A 3–20, N, PXR, CCP</td>
</tr>
<tr>
<td>D5811</td>
<td>A 3–20, N, PXR, CCP</td>
</tr>
<tr>
<td>D5820</td>
<td>A 3–20, N, PXR, CCP</td>
</tr>
<tr>
<td>D5821</td>
<td>A 3–20, N, PXR, CCP</td>
</tr>
</tbody>
</table>

**Other Removable Prosthetic Services**

<table>
<thead>
<tr>
<th>Procedure Code</th>
<th>Limitations</th>
</tr>
</thead>
<tbody>
<tr>
<td>D5850</td>
<td>A 3–20, N, PXR, CCP</td>
</tr>
<tr>
<td>D5851</td>
<td>A 3–20, N, PXR, CCP</td>
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<tr>
<td>D5862</td>
<td>A 4–20, N, PXR, CCP</td>
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<td>D5866</td>
<td>A 4–20, N, PXR, CCP</td>
</tr>
<tr>
<td>D5899</td>
<td>A 1–20, N, PXR, CCP</td>
</tr>
</tbody>
</table>

**Maxillofacial Prosthetics**

<table>
<thead>
<tr>
<th>Procedure Code</th>
<th>Limitations</th>
</tr>
</thead>
<tbody>
<tr>
<td>D5911</td>
<td>A 1–20, N, PXR, CCP</td>
</tr>
<tr>
<td>D5912</td>
<td>A 1–20, N, PXR, CCP</td>
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<tr>
<td>D5913</td>
<td>A 1–20, N, PXR, CCP</td>
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<tr>
<td>D5914</td>
<td>A 1–20, N, PXR, CCP</td>
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<td>D5915</td>
<td>A 1–20, N, PXR, CCP</td>
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<tr>
<td>D5916</td>
<td>A 1–20, N, PXR, CCP</td>
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<td>D5919</td>
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<td>D5922</td>
<td>A 1–20, N, PXR, CCP</td>
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<td>D5923</td>
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<tr>
<td>D5924</td>
<td>A 1–20, N, PXR, CCP</td>
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<tr>
<td>D5925</td>
<td>A 1–20, N, PXR, CCP</td>
</tr>
<tr>
<td>D5926</td>
<td>A 1–20, N, PXR, CCP</td>
</tr>
</tbody>
</table>

A=Age range limitations, N=Narrative required, FMX=Full-mouth radiographs (nonpanoramic), MTID=Missing tooth ID(s), PPXR=Preoperative and postoperative radiographs required, PXR=Preoperative radiographs required, PHO=Preoperative and postoperative photographs required, PC=Periodontal charting required, PATH=Pathology report required and must be retained in the client’s record, CCP=Comprehensive Care Program, NC= No charge to Medicaid and may not bill the client, and *= Services payable to an FQHC for a client encounter
### 4.2.19 Prosthodontic (Fixed) Services

Prosthodontic procedure codes require prior authorization.

**Refer to:** Subsection 4.2.29, “Mandatory Prior Authorization” in this handbook for documentation requirements.

<table>
<thead>
<tr>
<th>Procedure Code</th>
<th>Limitations</th>
</tr>
</thead>
<tbody>
<tr>
<td>D5927</td>
<td>A 1–20, N, PXR, CCP</td>
</tr>
<tr>
<td>D5928</td>
<td>A 1–20, N, PXR, CCP</td>
</tr>
<tr>
<td>D5929</td>
<td>A 1–20, N, PXR, CCP</td>
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<tr>
<td>D5931</td>
<td>A 1–20, N, PXR, CCP</td>
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<td>D5932</td>
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<tr>
<td>D5935</td>
<td>A 1–20, N, PXR, CCP</td>
</tr>
<tr>
<td>D5936</td>
<td>A 1–20, N, PXR, CCP</td>
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<tr>
<td>D5937</td>
<td>Not for temporo-mandibular dysfunction (TMD) treatment. A 1–20, N, PXR, CCP</td>
</tr>
<tr>
<td>D5951</td>
<td>Prior authorization. A Birth–20, N, PXR</td>
</tr>
<tr>
<td>D5952</td>
<td>Prior authorization. A Birth–20, N, PXR</td>
</tr>
<tr>
<td>D5953</td>
<td>Prior authorization. A 13–20, N, PXR</td>
</tr>
<tr>
<td>D5954</td>
<td>Prior authorization. A Birth–20, N, PXR</td>
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<tr>
<td>D5955</td>
<td>Prior authorization. A Birth–20, N, PXR</td>
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<tr>
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<td>Prior authorization. A Birth–20, N, PXR</td>
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</tr>
<tr>
<td>D5999*</td>
<td>A 1–20, N, PXR, CCP</td>
</tr>
</tbody>
</table>

A=Age range limitations, N=Narrative required, FMX=Full-mouth radiographs (nonpanoramic), MTID=Missing tooth ID(s), PPXR=Preoperative and postoperative radiographs required, PXR=Preoperative radiographs required, PHO=Preoperative and postoperative photographs required, PC=Periodontal charting required, PATH=Pathology report required and must be retained in the client’s record, CCP=Comprehensive Care Program, NC= No charge to Medicaid and may not bill the client, and *= Services payable to an FQHC for a client encounter
Periapical radiographs are required for each tooth involved in the authorization request. The criteria used by the TMHP Dental Director are:

- At least one abutment tooth requires a crown (based on traditional requirements of medical necessity and dental disease).
- The space cannot be filled with a removable partial denture.
- The purpose is to prevent the drifting of teeth in all dimensions (anterior, posterior, lateral, and the opposing arch).
- Each abutment or each pontic constitutes a unit in a bridge.
- Porcelain is allowed on all teeth.

<table>
<thead>
<tr>
<th>Procedure Code</th>
<th>Limitations</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>Fixed Partial Dental Pontics</strong></td>
<td></td>
</tr>
<tr>
<td>D6210</td>
<td>A 16–20, PPXR, MTID, CCP</td>
</tr>
<tr>
<td>D6211</td>
<td>A 16–20, PPXR, MTID, CCP</td>
</tr>
<tr>
<td>D6212</td>
<td>A 16–20, PPXR, MTID, CCP</td>
</tr>
<tr>
<td>D6240</td>
<td>A 16–20, PPXR, MTID, CCP</td>
</tr>
<tr>
<td>D6241</td>
<td>A 16–20, PPXR, MTID, CCP</td>
</tr>
<tr>
<td>D6242</td>
<td>A 16–20, PPXR, MTID, CCP</td>
</tr>
<tr>
<td>D6245</td>
<td>A 16–20, PPXR, MTID, CCP</td>
</tr>
<tr>
<td>D6250</td>
<td>A 16–20, PPXR, MTID, CCP</td>
</tr>
<tr>
<td>D6251</td>
<td>A 16–20, PPXR, MTID, CCP</td>
</tr>
<tr>
<td>D6252</td>
<td>A 16–20, PPXR, MTID, CCP</td>
</tr>
<tr>
<td><strong>Fixed Partial Dental Retainers—Inlays/Onlays</strong></td>
<td></td>
</tr>
<tr>
<td>D6545</td>
<td>A 16–20, PPXR, CCP</td>
</tr>
<tr>
<td>D6548</td>
<td>A 16–20, PPXR, CCP</td>
</tr>
<tr>
<td>D6549</td>
<td>A 16-20, PPXR, CCP</td>
</tr>
<tr>
<td><strong>Fixed Partial Dental Retainers—Crowns</strong></td>
<td></td>
</tr>
<tr>
<td>D6720</td>
<td>A 16–20, PPXR, CCP</td>
</tr>
<tr>
<td>D6721</td>
<td>A 16–20, PPXR, CCP</td>
</tr>
<tr>
<td>D6722</td>
<td>A 16–20, PPXR, CCP</td>
</tr>
<tr>
<td>D6740</td>
<td>A 16–20, PPXR, CCP</td>
</tr>
<tr>
<td>D6750</td>
<td>A 16–20, PPXR, CCP</td>
</tr>
<tr>
<td>D6751</td>
<td>A 16–20, PPXR, CCP</td>
</tr>
<tr>
<td>D6752</td>
<td>A 16–20, PPXR, CCP</td>
</tr>
<tr>
<td>D6780</td>
<td>A 16–20, PPXR, CCP</td>
</tr>
<tr>
<td>D6781</td>
<td>A 16–20, PPXR, CCP</td>
</tr>
<tr>
<td>D6782</td>
<td>A 16–20, PPXR, CCP</td>
</tr>
<tr>
<td>D6783</td>
<td>A 16–20, PPXR, CCP</td>
</tr>
</tbody>
</table>

A=Age range limitations, N=Narrative required, FMX=Full-mouth radiographs (nonpanoramic), MTID=Missing tooth ID(s), PPXR=Preoperative and postoperative radiographs required, PXR=Preoperative radiographs required, PHO=Preoperative and postoperative photographs required, PC=Periodontal charting required, PATH=Pathology report required and must be retained in the client’s record, CCP=Comprehensive Care Program, NC=No charge to Medicaid and may not bill the client, and *= Services payable to an FQHC for a client encounter.
### 4.2.20 Oral and Maxillofacial Surgery Services

All oral surgery procedures include local anesthesia, suturing, if needed, and visits for routine postoperative care.

<table>
<thead>
<tr>
<th>Procedure Code</th>
<th>Limitations</th>
</tr>
</thead>
<tbody>
<tr>
<td>D6790</td>
<td>Permanent posterior teeth only. A 16–20, PPXR, CCP</td>
</tr>
<tr>
<td>D6791</td>
<td>Permanent posterior teeth only. A 16–20, PPXR, CCP</td>
</tr>
<tr>
<td>D6792</td>
<td>Permanent posterior teeth only. A 16–20, PPXR, CCP</td>
</tr>
</tbody>
</table>

**Other Fixed Partial Dental**

<table>
<thead>
<tr>
<th>Procedure Code</th>
<th>Limitations</th>
</tr>
</thead>
<tbody>
<tr>
<td>D6920</td>
<td>A 16–20, PXR, CCP</td>
</tr>
<tr>
<td>D6930</td>
<td>A 16–20, PXR, CCP</td>
</tr>
<tr>
<td>D6940</td>
<td>A 16–20, N, PXR, CCP</td>
</tr>
<tr>
<td>D6950</td>
<td>A 16–20, N, PXR, CCP</td>
</tr>
<tr>
<td>D6980</td>
<td>A 16–20, N, PXR, CCP</td>
</tr>
<tr>
<td>D6999</td>
<td>A 16–20, N, PXR, CCP</td>
</tr>
</tbody>
</table>

**Surgical Extractions**

- **D7210**
  - Includes removal of the roots of a previously erupted tooth missing its clinical crown. A 1–20, PXR
- **D7220**
  - A 1–20, PXR
- **D7230**
  - A 1–20, PXR
- **D7240**
  - A 1–20, PXR
- **D7241**
  - Document unusual circumstance. A 1–20, N, PXR
- **D7250**
  - Involves tissue incision and removal of bone to remove a permanent or primary tooth root left in the bone from a previous extraction, caries, or trauma. Usually some degree of soft or hard tissue healing has occurred. A 1–20, N, PXR

**Other Surgical Procedures**

- **D7260**
  - Requires prior authorization. A 1–20, N, PXR; TIDs #1–16 only.
- **D7261**
  - May not be paid for the same DOS as D7260; TIDs #1–16 only. A 1–20
- **D7270**
  - A 1–20, N, PXR, CCP
- **D7272**
  - Requires prior authorization. A 1–20, N, PXR, CCP

---

<table>
<thead>
<tr>
<th>Procedure Code</th>
<th>Limitations</th>
</tr>
</thead>
<tbody>
<tr>
<td>D7280</td>
<td>A 1–20, N, PXR. Procedure code D7280 will be denied unless billed with an authorized procedure code D7283 for the same tooth, on the same day, by the same provider.</td>
</tr>
<tr>
<td>D7282</td>
<td>Permanent TIDs #1–32 only; may not be paid for the same DOS as D7280. A 4–20</td>
</tr>
<tr>
<td>D7283</td>
<td>Requires prior authorization. A 1–20; TIDs #2-15 and 18-31. To obtain prior authorization, the following items must be submitted: a prior authorization request form, a copy of an authorized Medicaid orthodontic treatment plan, and a current panoramic radiograph to determine medical necessity.</td>
</tr>
<tr>
<td>D7285</td>
<td>A 1–20, PXR, PATH, CCP</td>
</tr>
<tr>
<td>D7286*</td>
<td>A 1–20, PXR, PATH, CCP</td>
</tr>
<tr>
<td>D7290</td>
<td>A 1–20, N, PXR, CCP</td>
</tr>
<tr>
<td>D7291</td>
<td>A 4–20, N, PXR, CCP</td>
</tr>
</tbody>
</table>

**Alveoplasty—Surgical Preparation of Ridge for Dentures**

<table>
<thead>
<tr>
<th>Procedure Code</th>
<th>Limitations</th>
</tr>
</thead>
<tbody>
<tr>
<td>D7310</td>
<td>A 1–20, N, PXR, CCP</td>
</tr>
<tr>
<td>D7320</td>
<td>A 1–20, N, PXR, CCP</td>
</tr>
</tbody>
</table>

**Vestibuloplasty**

<table>
<thead>
<tr>
<th>Procedure Code</th>
<th>Limitations</th>
</tr>
</thead>
<tbody>
<tr>
<td>D7340</td>
<td>A 1–20, N, PXR, CCP</td>
</tr>
<tr>
<td>D7350</td>
<td>A 1–20, N, PXR, CCP</td>
</tr>
</tbody>
</table>

**Surgical Excision of Soft Tissue Lesions**

<table>
<thead>
<tr>
<th>Procedure Code</th>
<th>Limitations</th>
</tr>
</thead>
<tbody>
<tr>
<td>D7410</td>
<td>A 1–20, PXR, PATH</td>
</tr>
<tr>
<td>D7411</td>
<td>A 1–20, PXR, PATH</td>
</tr>
<tr>
<td>D7413</td>
<td>The incidental removal of cysts/lesions attached to the root(s) of a simple extraction is considered part of the extraction or surgical fee. A 1–20, N, PXR, PATH, CCP</td>
</tr>
<tr>
<td>D7414</td>
<td>The incidental removal of cysts/lesions attached to the root(s) of an extracted tooth is considered part of the extraction or surgical fee. A 1–20, N, PXR, PATH, CCP</td>
</tr>
</tbody>
</table>

**Surgical Excision of Intraosseous Lesions**

<table>
<thead>
<tr>
<th>Procedure Code</th>
<th>Limitations</th>
</tr>
</thead>
<tbody>
<tr>
<td>D7440</td>
<td>The incidental removal of cysts/lesions attached to the root(s) of an extracted tooth is considered part of the extraction or surgical fee. A 1–20, N, PXR, PATH, CCP</td>
</tr>
<tr>
<td>D7441</td>
<td>The incidental removal of cysts/lesions attached to the root(s) of an extracted tooth is considered part of the extraction or surgical fee. A 1–20, N, PXR, PATH, CCP</td>
</tr>
<tr>
<td>D7450</td>
<td>The incidental removal of cysts/lesions attached to the root(s) of an extracted tooth is considered part of the extraction or surgical fee. A 1–20, N, PXR, PATH, CCP</td>
</tr>
<tr>
<td>D7451</td>
<td>The incidental removal of cysts/lesions attached to the root(s) of an extracted tooth is considered part of the extraction or surgical fee. A 1–20, N, PXR, PATH, CCP</td>
</tr>
</tbody>
</table>

A=Age range limitations, N=Narrative required, FMX=Full-mouth radiographs (nonpanoramic), MTID=Missing tooth ID(s), PPXR=Preoperative and postoperative radiographs required, PXR=Preoperative radiographs required, PHO=Preoperative and postoperative photographs required, PC=Periodontal charting required, PATH=Pathology report required and must be retained in the client’s record, CCP=Comprehensive Care Program, NC=No charge to Medicaid and may not bill the client, and *= Services payable to an FQHC for a client encounter.
<table>
<thead>
<tr>
<th>Procedure Code</th>
<th>Limitations</th>
</tr>
</thead>
<tbody>
<tr>
<td>D7460</td>
<td>The incidental removal of cysts/lesions attached to the root(s) of a simple extraction is considered part of the extraction or surgical fee. A Birth–20, N, PXR, PATH, CCP</td>
</tr>
<tr>
<td>D7461</td>
<td>The incidental removal of cysts/lesions attached to the root(s) of a simple extraction is considered part of the extraction or surgical fee. A Birth–20, N, PXR, PATH, CCP</td>
</tr>
<tr>
<td>D7465</td>
<td>The incidental removal of cysts/lesions attached to the root(s) of a simple extraction is considered part of the extraction or surgical fee. A 1–20, N, PXR, PATH, CCP</td>
</tr>
</tbody>
</table>

**Excision of Bone Tissue**

D7472 Prior authorization is required. A 1–20

**Surgical Incision**

<table>
<thead>
<tr>
<th>Procedure Code</th>
<th>Limitations</th>
</tr>
</thead>
<tbody>
<tr>
<td>D7510*</td>
<td>TID required. A 1–20, PXR</td>
</tr>
<tr>
<td>D7520</td>
<td>A 1–20, N, PXR, CCP</td>
</tr>
<tr>
<td>D7530</td>
<td>A 1–20, N, PXR</td>
</tr>
<tr>
<td>D7540</td>
<td>A 1–20, N, PXR</td>
</tr>
<tr>
<td>D7550*</td>
<td>A 1–20, N, PXR</td>
</tr>
<tr>
<td>D7560</td>
<td>A 1–20, N, PXR, CCP</td>
</tr>
<tr>
<td>D7670</td>
<td>A 1–20, N, PXR, CCP</td>
</tr>
</tbody>
</table>

**Reduction of Dislocation and Management of Other Temporomandibular Joint Dysfunctions**

<table>
<thead>
<tr>
<th>Procedure Code</th>
<th>Limitations</th>
</tr>
</thead>
<tbody>
<tr>
<td>D7820</td>
<td>A 1–20, N, PXR</td>
</tr>
<tr>
<td>D7880</td>
<td>A 1–20, N, PXR, CCP</td>
</tr>
<tr>
<td>D7899</td>
<td>A 1–20, N, PXR, CCP</td>
</tr>
</tbody>
</table>

**Repair of Traumatic Wounds**

<table>
<thead>
<tr>
<th>Procedure Code</th>
<th>Limitations</th>
</tr>
</thead>
<tbody>
<tr>
<td>D7910*</td>
<td>A 1–20, N, PXR, CCP</td>
</tr>
</tbody>
</table>

**Complicated Suturing**

<table>
<thead>
<tr>
<th>Procedure Code</th>
<th>Limitations</th>
</tr>
</thead>
<tbody>
<tr>
<td>D7911</td>
<td>A 1–20, N, PXR, CCP</td>
</tr>
<tr>
<td>D7912</td>
<td>A 1–20, N, PXR, CCP</td>
</tr>
</tbody>
</table>

**Other Repair Procedures**

<table>
<thead>
<tr>
<th>Procedure Code</th>
<th>Limitations</th>
</tr>
</thead>
<tbody>
<tr>
<td>D7955</td>
<td>A 1–20</td>
</tr>
<tr>
<td>D7961</td>
<td>A 12–20. N, PXR, Prior authorization is required.</td>
</tr>
<tr>
<td>D7970*</td>
<td>A 1–20, N, PXR, CCP</td>
</tr>
<tr>
<td>D7971*</td>
<td>A 1–20, N, PXR, CCP</td>
</tr>
<tr>
<td>D7972</td>
<td>TIDs #1, 16, 17, and 32 only; may not be paid in addition to D7971 for the same DOS. A 13–20</td>
</tr>
<tr>
<td>D7980</td>
<td>A 1–20, N, PXR, CCP</td>
</tr>
<tr>
<td>D7983</td>
<td>A 1–20, N, PXR, CCP</td>
</tr>
</tbody>
</table>

A=Age range limitations, N=Narrative required, FMX=Full-mouth radiographs (nonpanoramic), MTID=Missing tooth ID(s), PPXR=Preoperative and postoperative radiographs required, PXR=Preoperative radiographs required, PHO=Preoperative and postoperative photographs required, PC=Periodontal charting required, PATH=Pathology report required and must be retained in the client’s record, CCP=Comprehensive Care Program, NC=No charge to Medicaid and may not bill the client, and *= Services payable to an FQHC for a client encounter.
### 4.2.21 Adjunctive General Services

<table>
<thead>
<tr>
<th>Procedure Code</th>
<th>Limitations</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>D7997</strong></td>
<td>Per arch, appliance removal (not by the dentist who placed the appliance). Includes removal of arch bar. Prior authorization is required. A 1–20, N, PXR, CCP</td>
</tr>
<tr>
<td><strong>D7999</strong></td>
<td>A 1–20, N, PXR, CCP</td>
</tr>
</tbody>
</table>

Refer to: Subsection 4.2.21.1, "Benefit Limitations for Adjunctive General Services" in this handbook for benefit limitations.

Emergency service only. The type of treatment rendered and TID must be indicated. It must be a service other than a prescription or topical medication. Each claim submitted for payment must be marked as “Emergency” in the Description field, Block 30, and the original date of treatment or incident must be referenced in the “Remarks” field, Block 35. The appropriate box must be checked in the “Treatment Resulting From” field, Block 45, if applicable, and modifier ET must be used to indicate an emergency.

Documentation to support the emergency and the treatment performed must be maintained in the client’s dental medical record.

Refer to: Subsection 4.2.27, “Emergency or Trauma Related Services for All THSteps Clients and Clients Who Are 5 Months of Age and Younger” in this handbook

**A=Age range limitations, N=Narrative required, FMX=Full-mouth radiographs (nonpanoramic), MTID=Missing tooth ID(s), PPXR=Preoperative and postoperative radiographs required, PXR=Preoperative radiographs required, PHO=Preoperative and postoperative photographs required, PC=Periodontal charting required, PATH=Pathology report required and must be retained in the client’s record, CCP=Comprehensive Care Program, NC=No charge to Medicaid and may not bill the client, and *= Services payable to an FQHC for a client encounter**
<table>
<thead>
<tr>
<th>Procedure Code</th>
<th>Limitations</th>
</tr>
</thead>
<tbody>
<tr>
<td>D9230*</td>
<td>May not be submitted more than one per client, per day. Denied if submitted with D9248. A 1–20.</td>
</tr>
<tr>
<td>D9239</td>
<td>Primary procedure code indicating first 15 minutes of intravenous moderate (conscious) sedation. Limited to 15 minutes per day, same provider. Denied if submitted with D9222 or D9248. A 1–20</td>
</tr>
<tr>
<td>D9243</td>
<td>Add-on procedure code indicating additional 15 minute increments of intravenous moderate (conscious) sedation. Limited to one hour and fifteen minutes per day, same provider. Must be billed with procedure code D9239. Denied if submitted with D9248. A 1–20</td>
</tr>
<tr>
<td>D9248*</td>
<td>May be submitted twice within a 12-month period. Denied if submitted with D9420, any provider. A 1–20</td>
</tr>
</tbody>
</table>

**Professional Consultation**

D9310 An oral evaluation by a specialist of any type who is also providing restorative or surgical services must be submitted as D0160. A 1–20, N, CCP

**Professional Visits**

D9410 Narrative required on claim form. A 1–20, N

D9420 Limited to twice per rolling year, per client, any provider. Documentation supporting the medical necessity of a dental hospital call, including any medical, physical, (e.g., traumatic event), mental or behavioral disability, and a description of the service performed that requires a hospital call must be retained in the client’s dental record and will be subject to retrospective review. Charts are subject to retrospective review. A 1–20, N

D9430 During regularly scheduled hours, no other services performed. Visits for routine postoperative care are included in all therapeutic and oral surgery fees. A 1–20, N

D9440 Visits for routine postoperative care are included in all therapeutic and oral surgery fees. A 1–20, N

**Drugs**

Procedure code D9630 is not payable for take home fluorides or drugs. Prescriptions should be given to clients to be filled by the pharmacy for these medications as the pharmacy is reimbursed by the Medicaid Vendor Drug Program. Procedure code D9630 is payable for medications (antibiotics, analgesics, etc.) administered to a client in the provider’s office. Documentation of dosage and route of administration must be provided in the Remarks section of the claim.

Refer to: The Outpatient Drug Services Handbook (Vol. 2, Provider Handbooks).

D9610 May not be submitted with code D9248. A 1–20, N

D9612 A 13–20, N, PXR

D9630 Includes, but is not limited to, oral antibiotics, oral analgesic, and oral sedatives administered in the office. May not be submitted with codes D9230, D9241, D9248, D9610, and D9920. A 1–20, N

**Miscellaneous Services**

D9910 Per whole mouth application, does not include fluoride. Not to be used for bases, liners, or adhesives under or with restorations. Limited to once per year. A 18–20, N, CCP

A=Age range limitations, N=Narrative required, FMX=Full-mouth radiographs (nonpanoramic), MTID=Missing tooth ID(s), PXR=Preoperative and postoperative radiographs required, PHO=Preoperative and postoperative photographs required, PC=Periodontal charting required, PATH=Pathology report required and must be retained in the client’s record, CCP=Comprehensive Care Program, NC=No charge to Medicaid and may not bill the client, *= Services payable to an FQHC for a client encounter
4.2.21.1 Benefit Limitations for Adjunctive General Services

Procedure code D9110 is a benefit for the following:

- Sedative or periodontal dressing
- Starting root canal procedure; i.e., open and drain tooth or re-medication of previously opened tooth
- Smoothing fractured tooth that is cutting lips or cheek
- Debridement or curettage of wound
- Excision of operculum over an erupting tooth
- Limited gingivectomy
- Suture removal by dentist other than the dentist who placed suture(s)
• Placement of a temporary crown by other than the patient’s regular dentist and one who is not in the process, has not previously, or does not in the future intend to perform an acrylic, polycarbonate, stainless steel or cast crown on this same tooth

• Tissue conditioning of a full or partial denture
• Removal of spontaneously or post-surgically sequestered bone spicule
• Spot or limited scaling and root planing
• Procedures necessary to treat a dry socket
• Procedures necessary to control bleeding
• Non-surgical reduction of TMJ dislocation
• Procedures necessary to relieve pain associated with pericoronitis, particularly third molars

Procedure code D9110 is not a benefit for the following:

• A written prescription
• Medication given or administered
• Application of topical medication to teeth or gums
• Occlusal adjustments
• Oral hygiene instructions

4.2.22 Dental Anesthesia

Dental providers must have the following information on file with TMHP to be eligible for reimbursement for dental anesthesia:

• A current anesthesia permit level issued by the TSBDE.
• Providers must have a Level 4 permit and an anesthesiology residency recognized by the American Dental Board of Anesthesiology to bill the enhanced rate for procedure codes D9222 and D9223.

All dental providers must comply with the American Academy of Pediatric Dentistry (AAPD) guidelines and TSBDE rules and regulations, including the standards for documentation and record maintenance for dental anesthesia.

4.2.22.1 Anesthesia Permit Levels

The following table shows the levels of anesthesia permits that are issued by the TSBDE:

<table>
<thead>
<tr>
<th>Permit Level</th>
<th>Description of Level</th>
<th>Permit Privileges</th>
</tr>
</thead>
<tbody>
<tr>
<td>Nitrous oxide/oxygen inhalation conscious sedation</td>
<td>Stand-alone permit</td>
<td></td>
</tr>
<tr>
<td>Level 1</td>
<td>Minimal sedation</td>
<td>Stand-alone permit</td>
</tr>
<tr>
<td>Level 2</td>
<td>Moderate enteral</td>
<td>Automatically qualifies for Level 1 and Level 2 permit privileges</td>
</tr>
<tr>
<td>Level 3</td>
<td>Moderate parenteral</td>
<td>Automatically qualifies for Level 1, Level 2, and Level 3 permit privileges</td>
</tr>
<tr>
<td>Level 4</td>
<td>Deep sedation/general anesthesia</td>
<td>Automatically qualifies for Level 1, Level 2, Level 3, and Level 4 permit privileges</td>
</tr>
</tbody>
</table>
All providers must have the appropriate anesthesia permit when proceeding with the procedure codes in the table below. The following table indicates the anesthesia procedure codes and the minimum anesthesia permit level to be reimbursed for the procedure codes:

<table>
<thead>
<tr>
<th>Procedure Codes</th>
<th>Minimum Anesthesia Permit Level</th>
</tr>
</thead>
<tbody>
<tr>
<td>D9211</td>
<td>Level 3</td>
</tr>
<tr>
<td>D9212</td>
<td>Level 3</td>
</tr>
<tr>
<td>D9222</td>
<td>Level 4</td>
</tr>
<tr>
<td>D9223</td>
<td>Level 4</td>
</tr>
<tr>
<td>D9230</td>
<td>Stand-alone permit for nitrous oxide/oxygen inhalation conscious sedation or Level 1</td>
</tr>
<tr>
<td>D9239</td>
<td>Level 3</td>
</tr>
<tr>
<td>D9243</td>
<td>Level 3</td>
</tr>
<tr>
<td>D9248</td>
<td>Level 2</td>
</tr>
</tbody>
</table>

Local anesthesia in conjunction with operative or surgical services (procedure code D9215) is all inclusive with any other dental service and is not reimbursed separately.

**4.2.22.2 Method for Counting Minutes for Timed Procedure Codes**

All claims for reimbursement of procedure codes paid in 15-minute increments are based on the actual amount of billable time associated with the service. For those services for which the unit of service is 15 minutes (1 unit = 15 minutes), partial units should be rounded up or down to the nearest quarter hour.

Time intervals for 1 through 12 units are as follows:

<table>
<thead>
<tr>
<th>Units</th>
<th>Number of Minutes</th>
</tr>
</thead>
<tbody>
<tr>
<td>0 units</td>
<td>0 minutes through 7 minutes</td>
</tr>
<tr>
<td>1 unit</td>
<td>8 minutes through 22 minutes</td>
</tr>
<tr>
<td>2 units</td>
<td>23 minutes through 37 minutes</td>
</tr>
<tr>
<td>3 units</td>
<td>38 minutes through 52 minutes</td>
</tr>
<tr>
<td>4 units</td>
<td>53 minutes through 67 minutes</td>
</tr>
<tr>
<td>5 units</td>
<td>68 minutes through 82 minutes</td>
</tr>
<tr>
<td>6 units</td>
<td>83 minutes through 97 minutes</td>
</tr>
<tr>
<td>7 units</td>
<td>98 minutes through 112 minutes</td>
</tr>
<tr>
<td>8 units</td>
<td>113 minutes through 127 minutes</td>
</tr>
<tr>
<td>9 units</td>
<td>128 minutes through 142 minutes</td>
</tr>
<tr>
<td>10 units</td>
<td>143 minutes through 157 minutes</td>
</tr>
<tr>
<td>11 units</td>
<td>158 minutes through 172 minutes</td>
</tr>
<tr>
<td>12 units</td>
<td>173 minutes through 187 minutes</td>
</tr>
</tbody>
</table>

All levels of sedation must have clinical documentation and a narrative in the client’s dental record to support the necessity of the service. Documentation must include the sedation record that indicates sedation start and end times in accordance with the American Academy of Pediatric Dentistry (AAPD) guidelines. The client’s dental record must be available for review by representatives of HHSC or its designee.
4.2.23 Hospitalization and ASC/HASC

Dental services performed in an ASC, HASC, or a hospital (either as an inpatient or an outpatient) may be benefits of THSteps based on the medical or behavioral justification provided, or if one of the following conditions exist:

- The procedures cannot be performed in the dental office.
- The client is severely disabled.

To satisfy the preadmission history and physical examination requirements of the hospital, ASC, or HASC, a THSteps medical checkup for dental rehabilitation or restoration may be performed by the child’s primary care provider. Physicians who are not enrolled as THSteps medical providers must submit claims for the examination of a client before the procedure with the appropriate evaluation and management procedure code from the following table:

<table>
<thead>
<tr>
<th>Procedure Code</th>
<th>Place of Service (POS)</th>
</tr>
</thead>
<tbody>
<tr>
<td>99202</td>
<td>POS 1 (office)</td>
</tr>
<tr>
<td>99222</td>
<td>POS 3 (inpatient hospital)</td>
</tr>
<tr>
<td>99282</td>
<td>POS 5 (outpatient hospital)</td>
</tr>
</tbody>
</table>

Refer to: Subsection 4.2.12.1, “Exceptions to Periodicity” in this handbook.

Note: The dental provider must submit claims to TMHP using the ADA Dental Claim Form to be considered for reimbursement through THSteps Dental Services.

The dental provider is responsible for obtaining prior authorization for the services performed under general anesthesia. Hospitals, ASC’s, and anesthesiologists must obtain the prior authorization number from the dental provider.

Contact the individual HMO for precertification requirements related to the hospital procedure. If services are precertified, the provider receives a precertification number effective for 90 days.

In those areas of the state with Medicaid managed care, the provider should contact the managed care plan for specific requirements or limitations. It is the dental provider’s responsibility to obtain precertification from the client’s HMO or managed care plan for facility and general anesthesia services if precertification is required.

To be reimbursed by the HMO, the provider must use the HMO’s contracted facility and anesthesia provider. These services are included in the capitation rates paid to HMOs, and the facility or anesthesiologist risk nonpayment from the HMO without such approval. Coordination of all specialty care is the responsibility of the client’s primary care provider. The primary care provider must be notified by the dentist or the HMO of the planned services.

Dentists providing sedation or anesthesia services must have the appropriate current permit from the TSBDE for the level of sedation or anesthesia provided.

The dental provider must be in compliance with the guidelines detailed in General Information.

Note: Post-treatment authorization will not be approved for codes that require mandatory prior authorization.

4.2.24 Orthodontic Services (THSteps)

Orthodontic services are a benefit for THSteps clients who are 13 years of age and older who have either permanent dentition and a severe handicapping malocclusion or one of the following special medical conditions:

- Cleft palate
- Head-trauma injury involving the oral cavity
• Skeletal anomalies involving the oral cavity

A severe handicapping malocclusion is defined by Texas Medicaid as dysfunctional masticatory (chewing) capacity as a result of the existing relationship between the maxillary (upper) and mandibular (lower) dental arches or teeth that without correction will result in damage to the temporomandibular joint (s) (TMJ) or other supporting oral structures (e.g., bone, tissues, intra- or extra-oral muscles, etc.).

Exception to the age restriction may be considered for clients who are 12 years of age and younger if medical necessity has been verified by the dental director for one of the following:

• Interceptive orthodontic treatment services
• Crossbite therapy
• Limited orthodontic treatment and minor treatment to control harmful habits
• Special medical conditions

Dental services that are not covered by THSteps Dental Services but are medically necessary and allowable may be a benefit under CCP according to federal Medicaid guidelines and TAC.

As required by the Texas Human Resources Code, clients who are 14 years of age and younger must be accompanied to Texas Health Steps dental checkups and visits by the client’s parent, legal guardian, or another adult who is authorized by the parent or legal guardian. The authorized adult can be the client’s relative. The individual accompanying the client must wait for the client while the appointment takes place.

Exempt entities (school health clinics, Head Start program, or childcare facilities) that provide services must as a condition of reimbursement:

• Obtain written, unrevoked consent for the services from the client’s parent or legal guardian within a one-year period before the date of service.
• Encourage parental involvement in and management of the health care of the clients who receive services from the clinic, program, or facility.

The following definitions of dentition established by the ADA’s Current Dental Terminology (CDT) manual are recognized by Texas Medicaid:

• Primary Dentition: Teeth developed and erupted first in order of time.
• Transitional Dentition: The final phase of the transition from primary to adult teeth, in which the deciduous molars and canines are in the process of shedding and the permanent successors are emerging.
• Adolescent Dentition: The dentition that is present after the normal loss of primary teeth and prior to cessation of growth that would affect orthodontic treatment.
• Adult Dentition: The dentition that is present after the cessation of growth that would affect orthodontic treatment.

The American Association of Orthodontists classification of occlusion or malocclusion is as follows:

• Class I: A Class I occlusion exists with the teeth in a normal relationship when the mesialbuccal cusp of the maxillary first permanent molar coincides with the buccal groove of the mandibular first molar.
• Class II: A Class II malocclusion occurs when the mandibular teeth are distal or behind the normal relationship with the maxillary teeth. This can be due to a deficiency of the lower jaw or an excess of the upper jaw and therefore, presents two types:
  • Division I is when the mandibular arch is behind the upper jaw with a consequential protrusion of the upper front teeth.
• Division II exists when the mandibular teeth are behind the upper teeth, with a retrusion of the maxillary front teeth. Both of these malocclusions have a tendency toward a deep bite because of the uncontrolled migration of the lower front teeth upwards.

• *Class III:* A Class III malocclusion occurs when the lower dental arch is in front of (mesial to) the upper dental arch. People with this type of occlusion usually have a strong or protrusive chin, which can be due to either horizontal mandibular excess or horizontal maxillary deficiency. Commonly referred to as an underbite.

### 4.2.24.1 Benefits and Limitations for Orthodontic Services

Comprehensive orthodontic services must be provided by a board-eligible or board-certified orthodontist.

**Note:** Exceptions to a board-eligible or board-certified orthodontist may be considered for clients in a rural or frontier area or where access to care is an issue.

The diagnostic workup is considered part of the pre-orthodontic treatment visit (procedure code D8660). The following procedure codes are used to submit claims for the diagnostic workup:

<table>
<thead>
<tr>
<th>Diagnostic Workup Procedure Codes</th>
</tr>
</thead>
<tbody>
<tr>
<td>D0330</td>
</tr>
</tbody>
</table>

Comprehensive orthodontic services include all of the following:

• Diagnostic workups
• Banding
• Initial brackets
• Replacement brackets
• Monthly visits
• Initial retainers
• Special orthodontic treatment appliance(s)

The following procedure codes are used to submit claims for orthodontic services:

<table>
<thead>
<tr>
<th>Orthodontic Services Procedure Codes</th>
</tr>
</thead>
<tbody>
<tr>
<td>D8080</td>
</tr>
</tbody>
</table>

Full banding is allowed on permanent dentition only, and treatment should be accomplished in one stage and is limited to once per lifetime.

**Exception:** Cases of mixed dentition may be considered when the treatment plan includes extractions of remaining primary teeth or in the case of cleft palate.

### 4.2.24.2 Crossbite Therapy

Crossbites (anterior and posterior) are defined by the American Academy of Pediatric Dentistry (AAPD) as malocclusions involving one or more teeth in which the maxillary teeth occlude lingually with the mandibular antagonistic (opposing) teeth. A crossbite can be of a dental or skeletal origin or a combination of both.

The intent of crossbite therapy is to prevent the need for comprehensive orthodontic treatment. This treatment may lessen the severity or future effects of a malformation, eliminate its cause, or may include localized tooth movement.
Crossbite therapy (limited orthodontics) is allowed for primary or transitional dentition. Crossbite therapy will not be considered for transitional dentition when there is a need for full banding of the adult teeth.

Crossbite therapy must be submitted with procedure code D8050 or D8060. Clients with special medical conditions may be considered for interceptive orthodontic services of the primary dentition if the services are medically necessary and submitted with procedure code D8050.

Crossbite therapy is an inclusive charge for treating the crossbite to completion. Adjustments, maintenance, diagnostic models, and diagnostic workup procedures are not reimbursed separately.

4.2.24.3 **Minor Treatment to Control Harmful Habits**

Special orthodontic appliances are a benefit for minor treatment to control harmful habits.

Orthodontic appliances for minor treatment to control harmful habits must be submitted with procedure codes D8210, D8220, and D8670.

Monthly adjustments (procedure code D8670) for minor treatment to control harmful habits are limited up to 10 visits.

Claims for panoramic films (procedure code D0330), cephalometric films (procedure code D0340), oral/facial photographic images obtained intraorally or extraorally (procedure code D0350) and diagnostic models (procedure code D0470) will be denied when they are submitted with procedure code D8210 or D8220.

Each orthodontic appliance (procedure code D8210 and D8220) are limited to once per arch, per lifetime.

4.2.24.4 **Premature Termination of Comprehensive Orthodontic Treatment**

Premature termination of comprehensive orthodontic treatment includes the following:

- Removal of the brackets and arch wires
- Removal of appliances with the fabrication of retainers
- Delivery of orthodontic retainers

Documentation of one of the following must be retained for premature termination of comprehensive orthodontic treatment:

- Documentation of a lack of cooperation from the client.
- Documentation that the client requested premature removal and a release of liability form has been signed by the parent, guardian, or client if he or she is at least 18 years of age.

Premature termination of comprehensive orthodontic treatment must be submitted with procedure code D8680.

Removal of the appliance (procedure code D8680) will be denied if the claim is submitted by any provider on the same date of service as orthodontic treatment (procedure codes D8050, D8060, and D8080).

Providers must keep a copy of the release of liability form on file and are responsible for this documentation during a review process.

If premature removal of the appliances is requested before completion of treatment, future orthodontic services may not be considered. The provider must document why the premature removal was necessary.
4.2.24.5 Other Orthodontic Services
Replacement brackets (procedure code D8690) are a benefit when the client transfers from one provider to another or when trauma is involved.

Providers are responsible for any replacement brackets that are required as part of the comprehensive orthodontic treatment. Additional reimbursement for replacement brackets (procedure code D8690) is limited to a combined total amount of $100.00, same provider.

Only one retainer per arch per lifetime (procedure code D8680) is allowed; however, each retainer may be replaced with prior authorization once per lifetime due to loss or breakage. Retainer adjustments are not reimbursed separately.

Appliances required as part of the cleft palate treatment plan may be reimbursed separately.

Special orthodontic appliances may be used with full banding and crossbite therapy when approved by the TMHP Dental Director or Associate Dental Director.

4.2.24.6 Non-covered Services
Single arch comprehensive orthodontic treatment is not a benefit of Texas Medicaid.

Orthodontic services that are performed solely for cosmetic purposes are not a benefit of Texas Medicaid. Although aesthetics is an important part of self-esteem, services primarily for self-worth are not within the scope of this Texas Medicaid benefit.

Orthodontic services for a client who initiated orthodontic treatment through a private arrangement while Medicaid-eligible are not a benefit of Texas Medicaid.

An initial orthodontic or pre-orthodontic treatment visit (procedure code D8660) is considered part of the exam in an oral evaluation (procedure codes D0120 or D0150).

4.2.24.7 Comprehensive Orthodontic Treatment
Comprehensive orthodontic services (procedure code D8080) are restricted to clients who are 13 years of age and older or clients who have exfoliated all primary dentition.

National procedure codes do not allow for any work-in-progress or partial submission of a claim by separating the three orthodontic components: diagnostic workup, orthodontic appliance (upper), or orthodontic appliance (lower).

When submitting claims for comprehensive orthodontic treatment procedure code D8080, three local codes must be submitted as remarks codes along with procedure code D8080. Local codes (procedure codes Z2009, Diagnostic workup approved; Z2011, Orthodontic appliance, upper; or Z2012, Orthodontic appliance, lower) must be placed in the Remarks Code field on electronic claims or Block 35 on paper claims.

Note: If the remarks code and procedure code D8080 are not submitted, the claim will be denied.

Each remarks code pays the correct reimbursement rate which, when combined, totals the maximum payment of $775. Procedure code D8080 must be submitted on three separate details, with the appropriate remarks code, even if the claim submission is for the workup and full banding. Submission of only one detail for a total of $775 will not be accepted.

Example 1: A client is approved for full banding, but after the initial workup, the client discontinues treatment. This provider would submit the national procedure code D8080 and place the local code Z2009, Diagnostic workup approved, in the Remarks/comment field. The claim would pay $175.

Example 2: A client is approved for full banding. The provider continues treatment and places the maxillary bands. The provider would submit the national procedure code D8080 and place the local procedure code Z2009, Diagnostic workup approved, and Z2011, Maxillary bands, in the Remarks/comment field. The claim would pay $475.
All electronic claims for procedure code D8080 must have the appropriate remarks code associated with the procedure code.

Providers must adhere to the following guidelines for electronic claim submission so TMHP can accurately apply the correct remarks code to the appropriate claim detail.

A Diagnostic Procedure Code (DPC) remarks code must be submitted, only once, in the first three bytes of the NTE02 at the 2400 loop.

**Example 1:** For a claim with one detail, submitted with procedure code D8080 and remarks code Z2009, enter the information as follows: DPCZ2009. The total submitted would be $175.

**Example 2:** For a claim with two details, where details one and two are procedure code D8080 and the remarks codes are Z2009 and Z2011, enter the information as follows: DPCZ2009Z2011. The total submitted would be $475.

**Example 3:** For a claim with three details, where all three details are submitted separately with procedure code D8080, enter the remarks code based on the order of the claim detail as follows: DPCZ2009Z2011Z2012. The total submitted would be $775.

This method ensures accurate and appropriate payment for services rendered and addresses the need for submission of a partial claim.

### 4.2.24.8 Orthodontic Procedure Codes and Fee Schedule

When submitting claims for orthodontic procedures, use the following procedure codes:

<table>
<thead>
<tr>
<th>Procedure Code</th>
<th>Limitations</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>Orthodontic Services</strong></td>
<td></td>
</tr>
<tr>
<td>D0330*, D0340*, D0350*, and D0470*</td>
<td></td>
</tr>
<tr>
<td>D7280</td>
<td>A 1-20</td>
</tr>
<tr>
<td><strong>Interceptive Orthodontic Treatment</strong></td>
<td></td>
</tr>
<tr>
<td>D8050*</td>
<td>Replaces Z2018 and 8110D. Limited to one per lifetime.</td>
</tr>
<tr>
<td>D8060*</td>
<td>Replaces Z2018 and 8120D. Limited to one per lifetime.</td>
</tr>
<tr>
<td><strong>Comprehensive Orthodontic Treatment</strong></td>
<td></td>
</tr>
<tr>
<td><strong>Minor Treatment to Control Harmful Habits</strong></td>
<td></td>
</tr>
<tr>
<td>D8210*</td>
<td>Refer to subsection 4.2.25, “Special Orthodontic Appliances” in this handbook for associated remarks field code.</td>
</tr>
<tr>
<td>D8220*</td>
<td>Refer to subsection 4.2.25, “Special Orthodontic Appliances” in this handbook for associated remarks field code.</td>
</tr>
<tr>
<td><strong>Other Orthodontic Services</strong></td>
<td></td>
</tr>
<tr>
<td>D8660*</td>
<td>Replaces Z2008. Denied when submitted for the same DOS as D0145 by any provider. Denied when submitted for the same DOS as D0120 or D0150 by the same provider.</td>
</tr>
<tr>
<td>D8670*</td>
<td>Replaces Z2013.</td>
</tr>
<tr>
<td>D8680*</td>
<td>Replaces Z2014 and Z2015; one retainer per arch per lifetime; may be replaced once because of loss or breakage (prior authorization is required).</td>
</tr>
</tbody>
</table>

* = Services payable to an FQHC for a client encounter.
4.2.25 Special Orthodontic Appliances

All removable or fixed special orthodontic appliances must be prior authorized. The prior authorization request must include both the national code and remarks code. However, prior authorization requests may omit the DPC prefix to the eight-digit remarks code.

All removable or fixed special orthodontic appliances must be submitted with national procedure code D8210 or D8220. To ensure appropriate claims processing, the DPC remarks code (local procedure code) reflecting the specific service is also required. The appropriate remarks codes must be entered on the prior authorization request form. Failure to follow the following steps will cause the claims to deny. Failure to enter the DPC remarks code and the appropriate procedure code will not result in claim denial; however, manual intervention is required to process the claim, which may result in a delay of payment.

For paper claim submissions, providers must enter the local procedure code in Block 35 (Remarks) of the ADA claim form.

For electronic submissions, providers enter the DPC remarks code in the Comments field to ensure correct authorization, accurate records, and reimbursement.

For electronic submissions other than TexMedConnect submissions, providers must use the following instructions to ensure that TMHHP accurately applies the correct local procedure code to the appropriate claim detail:

- The DPC prefix must be submitted, only once, in the first three bytes of the NTE02 at the 2400 loop.
- In bytes 4–8, providers must submit the remark code (local procedure code) based on the order of the claim detail. Do not enter any spaces or punctuation between remark codes, unless to designate the detail is not submitted with D8210 or D8220.

**Example:** For a claim with three details, where details one and three are submitted with procedure code D8210 and detail two is not, enter the following information in the NTE02 at the 2400 loop: DPC1014D 1046D. (The space shows that detail two needs no local code.) If all details require a local code, enter DPC, no spaces, and the appropriate local codes.

To submit using TexMedConnect, providers must enter the local code into the Remarks Code field, located under the details header. The Remarks Code field is the field directly after the Procedure Code field. TexMedConnect submitters are not required to manually enter the DPC prefix as it is placed in the appropriate field on the TexMedConnect electronic claim.

The following table identifies the appropriate DPC remarks codes to use when requesting prior authorization or submitting a claim for procedure code D8210 or D8220:

<table>
<thead>
<tr>
<th>Procedure Code</th>
<th>Remarks Code</th>
<th>Remarks Code Description</th>
</tr>
</thead>
<tbody>
<tr>
<td>D8690*</td>
<td></td>
<td>Bracket replacement.</td>
</tr>
<tr>
<td>D8999</td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

* = Services payable to an FQHC for a client encounter.
<table>
<thead>
<tr>
<th>Procedure Code</th>
<th>Remarks Code</th>
<th>Remarks Code Description</th>
</tr>
</thead>
<tbody>
<tr>
<td>D8210*</td>
<td>DPC1004D</td>
<td>Bite plate/bite plane</td>
</tr>
<tr>
<td>D8210*</td>
<td>DPC1005D</td>
<td>Bionator</td>
</tr>
<tr>
<td>D8210*</td>
<td>DPC1006D</td>
<td>Bite block</td>
</tr>
<tr>
<td>D8210*</td>
<td>DPC1007D</td>
<td>Bite-plate with push springs</td>
</tr>
<tr>
<td>D8220*</td>
<td>DPC1008D</td>
<td>Bonded expansion device</td>
</tr>
<tr>
<td>D8210*</td>
<td>DPC1010D</td>
<td>Chateau appliance (face mask, palatal exp and hawley)</td>
</tr>
<tr>
<td>D8210*</td>
<td>DPC1011D</td>
<td>Coffin spring appliance</td>
</tr>
<tr>
<td>D8220*</td>
<td>DPC1012D</td>
<td>Crib</td>
</tr>
<tr>
<td>D8210*</td>
<td>DPC1013D</td>
<td>Dental obturator, definitive (obturator)</td>
</tr>
<tr>
<td>D8210*</td>
<td>DPC1014D</td>
<td>Dental obturator, surgical (obturator, surgical stayplate, immediate temporary obturator)</td>
</tr>
<tr>
<td>D8220*</td>
<td>DPC1015D</td>
<td>Distalizing appliance with springs</td>
</tr>
<tr>
<td>D8220*</td>
<td>DPC1016D</td>
<td>Expansion device</td>
</tr>
<tr>
<td>D8210*</td>
<td>DPC1017D</td>
<td>Face mask (protraction mask)</td>
</tr>
<tr>
<td>D8220*</td>
<td>DPC1018D</td>
<td>Fixed expansion appliance</td>
</tr>
<tr>
<td>D8220*</td>
<td>DPC1019D</td>
<td>Fixed lingual arch</td>
</tr>
<tr>
<td>D8220*</td>
<td>DPC1020D</td>
<td>Fixed mandibular holding arch</td>
</tr>
<tr>
<td>D8210*</td>
<td>DPC1021D</td>
<td>Fixed rapid palatal expander</td>
</tr>
<tr>
<td>D8210*</td>
<td>DPC1022D</td>
<td>Frankel appliance</td>
</tr>
<tr>
<td>D8210*</td>
<td>DPC1023D</td>
<td>Functional appliance for reduction of anterior openbite and crossbite</td>
</tr>
<tr>
<td>D8210*</td>
<td>DPC1024D</td>
<td>Headgear (face bow)</td>
</tr>
<tr>
<td>D8220*</td>
<td>DPC1025D</td>
<td>Herbst appliance (fixed or removable)</td>
</tr>
<tr>
<td>D8220*</td>
<td>DPC1026D</td>
<td>Inter-occlusal cast cap surgical splints</td>
</tr>
<tr>
<td>D8210*</td>
<td>DPC1027D</td>
<td>Intrusion arch</td>
</tr>
<tr>
<td>D8220*</td>
<td>DPC1028D</td>
<td>Jasper jumpers</td>
</tr>
<tr>
<td>D8220*</td>
<td>DPC1029D</td>
<td>Lingual appliance with hooks</td>
</tr>
<tr>
<td>D8220*</td>
<td>DPC1030D</td>
<td>Mandibular anterior bridge</td>
</tr>
<tr>
<td>D8220*</td>
<td>DPC1031D</td>
<td>Mandibular bihelix (similar to a quad helix for mandibular expansion to attempt nonextraction treatment)</td>
</tr>
<tr>
<td>D8210*</td>
<td>DPC1032D</td>
<td>Mandibular lip bumper</td>
</tr>
<tr>
<td>D8220*</td>
<td>DPC1036D</td>
<td>Mandibular lingual 6x6 arch wire</td>
</tr>
<tr>
<td>D8210*</td>
<td>DPC1037D</td>
<td>Mandibular removable expander with bite plane (crozat)</td>
</tr>
<tr>
<td>D8210*</td>
<td>DPC1038D</td>
<td>Mandibular ricketts rest position splint</td>
</tr>
<tr>
<td>D8210*</td>
<td>DPC1039D</td>
<td>Mandibular splint</td>
</tr>
<tr>
<td>D8210*</td>
<td>DPC1040D</td>
<td>Maxillary anterior bridge</td>
</tr>
<tr>
<td>D8210*</td>
<td>DPC1041D</td>
<td>Maxillary bite-opening appliance with anterior springs</td>
</tr>
<tr>
<td>D8220*</td>
<td>DPC1042D</td>
<td>Maxillary lingual arch with spurs</td>
</tr>
<tr>
<td>D8220*</td>
<td>DPC1043D</td>
<td>Maxillary and mandibular distalizing appliance</td>
</tr>
<tr>
<td>D8220*</td>
<td>DPC1044D</td>
<td>Maxillary quad helix with finger springs</td>
</tr>
</tbody>
</table>

* = Services payable to an FQHC for a client encounter.
4.2.26 Handicapping Labio-lingual Deviation (HLD) Index

The orthodontic provider must complete and sign the HLD Index (Angle classification).

The HLD index requires the use of an HLD score sheet and a Boley gauge for measuring.

<table>
<thead>
<tr>
<th>Procedure Code</th>
<th>Remarks Code</th>
<th>Remarks Code Description</th>
</tr>
</thead>
<tbody>
<tr>
<td>D8220*</td>
<td>DPC1045D</td>
<td>Maxillary and mandibular retainer with pontics</td>
</tr>
<tr>
<td>D8210*</td>
<td>DPC1046D</td>
<td>Maxillary Schwarz</td>
</tr>
<tr>
<td>D8210*</td>
<td>DPC1047D</td>
<td>Maxillary splint</td>
</tr>
<tr>
<td>D8210*</td>
<td>DPC1048D</td>
<td>Mobile intraoral Arch-Mia (similar to a BiHelix for nonextraction treatment)</td>
</tr>
<tr>
<td>D8220*</td>
<td>DPC1049D</td>
<td>Modified quad helix appliance</td>
</tr>
<tr>
<td>D8220*</td>
<td>DPC1050D</td>
<td>Modified quad helix appliance (with appliance)</td>
</tr>
<tr>
<td>D8220*</td>
<td>DPC1051D</td>
<td>Nance appliance</td>
</tr>
<tr>
<td>D8220*</td>
<td>DPC1052D</td>
<td>Nasal stent</td>
</tr>
<tr>
<td>D8210*</td>
<td>DPC1053D</td>
<td>Occlusal orthotic device</td>
</tr>
<tr>
<td>D8210*</td>
<td>DPC1054D</td>
<td>Orthopedic appliance</td>
</tr>
<tr>
<td>D8210*</td>
<td>DPC1055D</td>
<td>Other mandibular utilities</td>
</tr>
<tr>
<td>D8210*</td>
<td>DPC1056D</td>
<td>Other maxillary utilities</td>
</tr>
<tr>
<td>D8220*</td>
<td>DPC1057D</td>
<td>Palatal bar</td>
</tr>
<tr>
<td>D8220*</td>
<td>DPC1058D</td>
<td>Post-surgical retainer</td>
</tr>
<tr>
<td>D8220*</td>
<td>DPC1059D</td>
<td>Quad helix appliance held with transpalatal arch horizontal projections</td>
</tr>
<tr>
<td>D8220*</td>
<td>DPC1060D</td>
<td>Quad helix maintainer</td>
</tr>
<tr>
<td>D8220*</td>
<td>DPC1061D</td>
<td>Rapid palatal expander (RPE), such as quad Helix, Haas, or Menne</td>
</tr>
<tr>
<td>D8210*</td>
<td>DPC1062D</td>
<td>Removable bite plate</td>
</tr>
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<td>D8210*</td>
<td>DPC1063D</td>
<td>Removable mandibular retainer</td>
</tr>
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<td>D8210*</td>
<td>DPC1064D</td>
<td>Removable maxillary retainer</td>
</tr>
<tr>
<td>D8210*</td>
<td>DPC1065D</td>
<td>Removable prosthesis</td>
</tr>
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<td>D8210*</td>
<td>DPC1066D</td>
<td>Sagittal appliance 2 way</td>
</tr>
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<td>D8210*</td>
<td>DPC1067D</td>
<td>Sagittal appliance 3 way</td>
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<td>D8220*</td>
<td>DPC1068D</td>
<td>Stapled palatal expansion appliance</td>
</tr>
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<td>D8210*</td>
<td>DPC1069D</td>
<td>Surgical arch wires</td>
</tr>
<tr>
<td>D8210*</td>
<td>DPC1070D</td>
<td>Surgical splints (surgical stent/wafer)</td>
</tr>
<tr>
<td>D8210*</td>
<td>DPC1071D</td>
<td>Surgical stabilizing appliance</td>
</tr>
<tr>
<td>D8220*</td>
<td>DPC1072D</td>
<td>Thumbsucking appliance, requires submission of models</td>
</tr>
<tr>
<td>D8210*</td>
<td>DPC1073D</td>
<td>Tongue thrust appliance, requires submission of models</td>
</tr>
<tr>
<td>D8210*</td>
<td>DPC1074D</td>
<td>Tooth positioner (full maxillary and mandibular)</td>
</tr>
<tr>
<td>D8210*</td>
<td>DPC1075D</td>
<td>Tooth positioner with arch</td>
</tr>
<tr>
<td>D8220*</td>
<td>DPC1076D</td>
<td>Transpalatal arch</td>
</tr>
<tr>
<td>D8220*</td>
<td>DPC1077D</td>
<td>Two bands with transpalatal arch and horizontal projections forward</td>
</tr>
<tr>
<td>D8220*</td>
<td>DPC1078D</td>
<td>Appliance</td>
</tr>
</tbody>
</table>

* = Services payable to an FQHC for a client encounter.
Refer to: The Texas Medicaid Handicapping Labio-Lingual Deviation (HLD) Index Score Sheet on the TMHP website at www.tmhp.com.

Providers should be conservative in scoring. The client must be considered severe handicapping malocclusion with dysfunctional masticatory (chewing) capacity as a result of the existing relationship between the maxillary (upper) and mandibular (lower) dental arches and/or teeth that, without correction, will result in damage to the temporomandibular joint(s) (TMJ) and/or other supporting oral structures (e.g., bone, tissues, intra and/or extra oral muscles, etc.) and have a minimum of 26 points on the HLD index to be considered for any orthodontic care other than crossbite correction. “Half-mouth” treatment cannot be approved.

With the client or models in the centric position, the HLD index is to be scored as follows. Record all measurements rounded-off to the nearest millimeter (mm). Enter a score of “0” if the condition is absent.

Cleft Palate

A cleft palate case request for mixed dentition will be considered only if narrative justification supports treatment before the client reaches full dentition.

Note: Intermittent treatment requests may exceed the allowable 26 reimbursable treatment visits.

Severe Traumatic Deviations

Refers to facial accidents only. Points cannot be awarded for congenital deformity. Severe traumatic deviations do not include traumatic occlusions for crossbites.

Overjet in Millimeters

Score the case exactly as measured. The measurement must be recorded from the most protrusive incisor, then subtract 2 mm (considered the norm), and enter the difference as the score.

Overbite in Millimeters

Score the case exactly as measured. The measurement must be recorded from the labio-incisal edge of the overlapped anterior tooth or teeth to the point of maximum coverage, then subtract 3 mm (considered the norm), and enter the difference as the score.

Mandibular Protrusion in Millimeters

Score the client exactly as measured. The measurement must be recorded from the “line of occlusion” of the permanent teeth, not from the ectopically erupted teeth in the anterior segment.

Open Bite in Millimeters

Score the case exactly as measured. Measurement must be recorded from the “line of occlusion” of the permanent teeth, not from the ectopically erupted teeth in the anterior segment. Caution is advised in undertaking treatment of open bites in older teenagers, because of the frequency of relapse.

Ectopic Eruption

An unusual pattern of eruption, such as high labial cuspids or teeth that are grossly out of the long axis of the alveolar ridge.

Ectopic eruption does not include teeth that are rotated or teeth that are leaning or slanted especially when the enamel-gingival junction is within the long axis of the alveolar ridge.

Note: Record the more serious condition. Do not include (score) teeth from an arch if that arch is to be counted in the category of Anterior Crowding. For each arch, either the ectopic eruption or anterior crowding may be scored, but not both.
Anterior Crowding
Arch length insufficiency must exceed 3.5 mm to be considered as crowding in either arch. Mild rotations that may react favorably to stripping or moderate expansion procedures are not to be scored as crowded.

Excessive Anterior Spacing in Millimeters
The score for this category must be the total, in millimeters, of the anterior spaces.

Providers should be conservative in scoring. Liberal scoring will not be helpful in the evaluation and approval of the case. The case must be considered dysfunctional and have a minimum of 26 points on the HLD index to qualify for any orthodontic care other than crossbite correction. Half-mouth cases cannot be approved.

The intent of the program is to provide orthodontic care to clients with handicapping malocclusion to improve function. Although aesthetics is an important part of self-esteem, services that are primarily for aesthetics are not within the scope of benefits of this program.

The proposals for treatment services should incorporate only the minimal number of appliances required to properly treat the case. Requests for multiple appliances to treat an individual arch will be reviewed for duplication of purpose.

If attaining a qualifying score of 26 points is uncertain, providers must include a brief narrative when submitting the case. The narrative may reduce the time necessary to gain final approval and reduce shipping costs incurred to resubmit records.

Providers must properly label and protect all records (especially plaster diagnostic models) when shipping. If plaster diagnostic models are requested by and shipped to TMHP, the provider should assure that the models are adequately protected from breakage during shipping. TMHP will return intact models to the provider.

4.2.27 Emergency or Trauma Related Services for All THSteps Clients and Clients Who Are 5 Months of Age and Younger
THSteps clients who are birth through 5 months of age are not eligible for routine dental checkups; however:

- They can be seen for emergency dental services by the dentist at any time for trauma, early childhood caries, or other oral health problems.
- They may be referred to a dentist by their primary care provider when a medical checkup identifies the medical necessity for dental services.

Prior authorization is not required for emergency or trauma-related dental services. Claims for these dental services must be filed separately from nonemergency dental services. Only one emergency or trauma-related dental claim per client, per day, may be considered for reimbursement. Routine therapeutic procedures are not considered emergency or trauma-related procedures.

When submitting a claim for emergency or trauma-related dental services, the provider must:

- Enter the word “Emergency” or “Trauma” in the description field (Block 30) of the claim form (also enter a brief description of the CDT procedure code used). Claims are subject to retrospective review. If no comments are indicated on the claim form, the payment may be recouped.
- If checking the Other Accident box, briefly describe in the Remarks field, Block 35 of the claim form, what caused the emergency or trauma.
- Check the appropriate box in Block 45, Treatment Resulting From, of the claim form (the options to check are Occupational Illness/Injury, Auto Accident, or Other Accident).
Documentation to support the diagnosis and treatment of trauma must be retained in the client’s record.

**Note:** Indicating Trauma in the description field allows the provider to be reimbursed for treatment on an emergency, continuing, and long-term basis without regard to periodicity, subject to the client’s eligibility and program limitations. An exception to periodicity for THSteps dental services is granted automatically for immediate treatment and any future follow-up treatment, as long as each claim submitted for payment is marked “Trauma” in the Description field, Block 30, and the original date of treatment or incident is referenced in the Remarks field, Block 35.

**Refer to:**
Subsection 4.1, “General Medicaid Eligibility” in “Section 4: Client Eligibility” (Vol. 1, General Information).
Subsection 4.2.14, “Medicaid Dental Benefits, Limitations, and Fee Schedule” in this handbook.

**4.2.28 Emergency Services for Medicaid Clients Who Are 21 Years of Age and Older**

Limited dental services are available for clients who are 21 years of age and older (not residing in an ICF-IID facility) whose dental diagnosis is secondary to and causally related to a life-threatening medical condition.

**Refer to:**

**4.2.28.1 Long Term Care (LTC) Emergency Dental Services**

HHSC provides a limited range of dental services for Medicaid-eligible residents of LTC facilities. All claims for dental services provided to LTC residents are submitted to HHSC. For information, providers should contact the appropriate LTC facility or HHSC at 1-800-252-9240.

**4.2.28.2 Laboratory Requirements**

Dental laboratories must be registered with TSBDE laboratories, and technicians must not be under restrictions imposed by TSBDE or a court.

**4.2.29 Mandatory Prior Authorization**

Mandatory prior authorization is required for consideration of reimbursement to dental providers who render the following services:

- Orthodontia
- Endodontic endosseous implants
- Fixed prosthetic services
- Removable prosthodontics
- Dental general anesthesia
- All inlays/onlays or permanent crowns
- Procedure code D4276
- Nonsurgical periodontal service (procedure codes D4341 and D4342)
• Procedure code D7272
• Procedure code D7283
• Procedure code D7472

• Limited dental services for clients who are 21 years of age and older (not residing in an ICF-IID facility) whose dental diagnosis is secondary to and causally related to a life-threatening medical condition
• Cone beam imaging

Approved orthodontic treatment plans must be initiated before the client’s loss of Medicaid eligibility and before the 21st birthday, and must be completed within 36 months of the authorization date. Authorization for other procedures is valid for up to 90 days.

To obtain prior authorization for crowns, onlays, endodontic endosseous implants, and fixed prosthodontics, a prior authorization form together with documentation supporting medical necessity and appropriateness must be submitted. Required documentation includes, but is not limited to:

• The THSteps Dental Mandatory Prior Authorization Request Form.
• Current, dated, pre-operative periapical radiographs completely showing the apex of the tooth to be treated.
• Current, dated, pre-operative full arch radiographs are required for fixed prosthodontics.
• Documentation supporting that the mouth is free of disease; no untreated periodontal or endodontic disease, or rampant caries.
• Documentation supporting only one virgin abutment tooth; at least one tooth must require a crown unless a Maryland Bridge is being considered.
• Provider documentation supporting the medical necessity and appropriateness of the recommended treatment.
• Tooth Identification (TID) System noting only permanent teeth.
• Documentation supporting that a removable partial is not a viable option to fill the space between the teeth.

Prior authorization will not be given when films show two abutment teeth (virgin teeth do not require a crown, except for Maryland Bridge) or there is untreated periodontal or endodontic disease, or rampant caries which would contraindicate the treatment.


Removable prosthodontics (procedure codes D5951, D5952, D5953, D5954, D5955, D5958, D5959, and D5960) for clients with cleft lip or cleft palate requires prior authorization with a completed THSteps Dental Mandatory Prior Authorization Request Form and narrative documenting the medical need for these appliances. Additional information may be requested by the TMHP Dental Director if necessary before making a determination.

The prior authorization number is required on claims for processing. If the client is not eligible for Medicaid on the DOS or the claim is incomplete, it will affect reimbursement. Prior authorization is a condition for reimbursement; it is not a guarantee of payment.

Note: Post-treatment authorization will not be approved for codes that require mandatory prior authorization.
**4.2.29.1 Cone Beam Imaging**

Prior authorization is required for procedure code D0367.

Cone beam imaging is used to determine the best course of treatment for cleft palate repair, skeletal anomalies, post-trauma care, implanted or fixed prosthodontics, and orthodontic or orthognathic procedures. Cone beam imaging is limited to initial treatment planning, surgery, and postsurgical follow up.

To obtain prior authorization, a THSteps Dental Mandatory Prior Authorization Request Form must be submitted with documentation supporting medical necessity and appropriateness. Required documentation includes, but is not limited to, the following:

- Presenting conditions
- Medical necessity
- Status of the client’s treatment

**4.2.29.2 General Anesthesia for Dental Treatment**

Prior authorization is required for the use of general anesthesia while rendering treatment (to include the dental service fee, the anesthesia fee, and facility fee) regardless of place of service. A client must meet the minimum requirement of 22 total points on the Criteria for Dental Therapy Under General Anesthesia form.

**Refer to:** [Criteria for Dental Therapy Under General Anesthesia](http://www.tmhp.com) on the TMHP website.

In those areas of the state with Medicaid Managed Care, precertification or approval is required from the client’s health maintenance organization (HMO) for anesthesia and facility charges. It is the dental provider’s responsibility to obtain precertification from the client’s HMO or managed care plan for facility and general anesthesia services. A medical checkup prior to a dental procedure requiring general anesthesia is considered an exception to THSteps periodicity. A referral to the client’s primary care physician is not required. Prior authorization is available for exceptions to periodicity. Provider must include all appropriate supporting documentation with the submittal. The criteria for general anesthesia applies only to treatment of clients who are 20 years of age and younger or ICF-IID program clients.

**4.2.29.2.1 Dental Therapy Under General Anesthesia**

Providers must comply with TSBDE Rules and Regulations, Title 22 TAC, Part 5, Chapter 110, §§110.6 –110.10. Any anesthesia type services are paid only to the provider. The dental provider is responsible for determining whether a client meets the minimum criteria necessary for receiving general anesthesia. A local anesthesia fee is not paid in addition to other restorative, operative, or surgical procedure fees.

For clients who are six years of age or younger, the following will apply:

- All Level 4 sedation/general anesthesia services provided by a dentist (procedure codes D9222 and D9223), and any anesthesia services provided by an anesthesiologist (M.D./D.O.) or certified registered nurse anesthetist (CRNA) to be provided in conjunction with dental therapeutic services (procedure code 00170 with modifier U3) must be prior authorized.
- The dentist performing the therapeutic dental procedure is responsible for obtaining prior authorization from TMHP and is responsible for providing the anesthesia prior authorization information to the anesthesiology provider.
- Prior authorization for both dental services and Level 4 sedation/general anesthesia service is mandatory for the reimbursement of either service.
Dental general anesthesia using procedure code D9222, D9223, or 00170 with modifier U3 is limited to once per six calendar months per client, by any provider. Add-on procedure code D9223 must be billed in conjunction with primary procedure code D9222, same provider.

Each distinct dental procedure code to be performed that requires prior authorization must be listed on the Texas Health Steps Dental Mandatory Prior Authorization Request Form. Repetitive dental procedure codes must be listed to indicate the total quantity to be performed. Claims submitted with unauthorized procedure codes will be denied, but may be appealed with documentation of medical necessity.

Requests for prior authorization must include, but is not limited to, the following client-specific documents and information:

- A completed Criteria for Dental Therapy Under General Anesthesia form
- A completed THSteps Dental Mandatory Prior Authorization Request Form
- The location of where the procedure(s) will be performed (office, inpatient hospital, or outpatient hospital)
- A narrative unique to the client, detailing the reasons for the proposed level of sedation (indicate procedure code D9222, D9223, or 00170). The narrative must include a history of prior treatment, information about failed attempts at other levels of sedation, behavior in the dental chair, proposed restorative treatment (tooth ID and surfaces), urgent need to provide comprehensive dental treatment based on extent of diagnosed dental caries, and any relevant medical condition(s).
- Diagnostic quality radiographs or photographs

**Note:** When appropriate radiographs or photographs cannot be taken prior to general anesthesia, the narrative must support the reasons for an inability to perform diagnostic services. For special cases that receive authorization, diagnostic quality radiographs or photographs will be required for payment and will be reviewed by the TMHP dental director.

The current process of scoring 22 points on the Criteria for Dental Therapy Under General Anesthesia form does not guarantee authorization or reimbursement for clients who are six years of age and younger.

**Note:** In cases of an emergency medical condition, accident, or trauma, prior authorization is not necessary. However, a narrative and appropriate pre- and post-treatment radiographs or photographs must be submitted with the claim, which will be reviewed by the TMHP dental director.

A copy of the Criteria for Dental Therapy Under General Anesthesia form must be maintained in the client’s dental record. The client’s dental record must be available for review by representatives of the Health and Human Services Commission (HHSC) or its designee.

Prior authorization is required for medically necessary dental general anesthesia that exceeds once per six months, per client, any provider. The dental provider is responsible for obtaining prior authorization for the services performed under general anesthesia. Hospitals, ASCs, and anesthesiologists must obtain the prior authorization number from the dental provider.

**Refer to:** Criteria for Dental Therapy Under General Anesthesia on the TMHP website at www.tmhp.com. Dental rehabilitation or restoration services requiring general anesthesia may be performed in an office, inpatient, or outpatient facility.

Surgical services related to THSteps dental services requiring general anesthesia must be coded as follows:

- Procedure code 00170 with modifier U3 is for the anesthesiologist or certified registered nurse anesthetist (CRNA) to use on the claim form.
• Procedure code 41899 with modifier U3 is for the facility to use on the claim form. Procedure code 41899 does not require prior authorization for ASCs and Hospital-based Ambulatory Surgical Centers (HASCs).

• An appropriate diagnosis code must be used on the claim form.

• Modifier U3 identifies that the service is associated with THSteps.

The claim forms used are the CMS-1500 or the UB-04 CMS-1450 paper claim forms. The examining physician, anesthesiologist, hospital, ASC, or HASC must submit claims to TMHP separately for the medical and facility components of their services.

Refer to: THSteps Dental Mandatory Prior Authorization Request Form on the TMHP website at www.tmhp.com.

The dental provider must include dental procedure codes and modifiers on the Texas Health Steps Dental Mandatory Prior Authorization Request Form for prior authorization to be considered for any Texas Health Steps dental services.

4.2.29.3 Orthodontic Services

Prior authorization is required for all orthodontic services.

Orthodontic services do not include any related services outside those listed in this section (e.g., extractions or surgeries); however, all services must be included in the orthodontic treatment plan.

Approved orthodontic treatment plans must be initiated before clients lose Medicaid eligibility or reach 21 years of age, and all active orthodontic treatments must be completed within 36 months of the authorization date. Services cannot be added or approved after eligibility has expired.

Note: If a client reaches 21 years of age or loses Medicaid eligibility before the authorized orthodontic services are completed, reimbursement is provided to complete the orthodontic treatment plan that was authorized and initiated while the client was 20 years of age or younger and eligible for Texas Medicaid as long as the orthodontic treatment plan is completed within the appropriate time frames.

Any non-orthodontic service that is included as part of the treatment plan (extractions or surgeries) must be completed before the client loses eligibility or reaches 21 years of age in order to be reimbursed through Texas Medicaid. Services cannot be added or approved after Texas Medicaid eligibility has expired.

Once prior authorization is obtained, the provider is obligated to advise the client that he or she is able to receive the approved orthodontic service (including monthly orthodontic adjustment visits and retainers) even if the client loses eligibility or reaches his or her 21st birthday.

All requests must be reviewed by the TMHP Dental Director or other state dental contractor’s board-eligible or board-certified orthodontist employee or consultant who is licensed in Texas.

To avoid unnecessary denials, providers must submit correct and complete information, including documentation for medical necessity for the services requested. Providers must maintain documentation of medical necessity in the client’s medical record. Requesting providers may be asked for additional information to clarify or complete a request.

A completed Texas Health Steps (THSteps) Dental Mandatory Prior Authorization Request Form must be signed and dated by the performing dental provider. The completed authorization form must include the procedure codes for all services requested along with a written statement of medical necessity for the proposed orthodontic treatment.

All prior authorization requests for orthodontic services must be accompanied by an attestation from the requesting provider that the provider is either a pediatric dentist or orthodontist.
General dentists who are requesting prior authorization for orthodontic services must attest and maintain documentation of a minimum of 200 hours of continuing dental education specifically in orthodontics within the last 10 years; 8 hours can be online or self-instruction.

Proof of the completion of continuing education hours is not required to be submitted with a request for prior authorization of orthodontic services; however, documentation must be produced by the dentist during retrospective review. All attestations are subject to compliance review and orthodontic services may be subject to recoupment.

4.2.29.3.1 Initial Orthodontic Services Request

The prior authorization form must include all of the procedures that are required to complete the requested treatment including, but not limited to, the following:

- Diagnostic workup
- Medically necessary extractions (Tooth ID must be included)
- Orthognathic surgery
- Upper and lower appliance
- Monthly adjustments
- Special orthodontic treatment appliances
- Placement of banding and brackets
- Replacement of brackets
- Removal of the brackets and arch wires
- Other special orthodontic appliances
- Fabrication of special orthodontic appliances
- Delivery of orthodontic retainers
- Appliance removal (if indicated)

A completed and scored Handicapping Labio-Lingual Deviations (HLD) Index with a diagnosis of Angle class (a minimum of 26 points are required for approval of non-cleft palate cases). If attaining a qualifying score of 26 points is uncertain, a brief narrative should be provided.

**Note:** A score of a minimum 26 points on the HLD index does not indicate an automatic approval for comprehensive orthodontics. Approval will be based on the diagnostic workup supporting the HLD index. Documentation provided must be reviewed by a qualified board eligible or board certified orthodontist.

When requesting prior authorization, providers must include diagnostic models, radiographs (X-rays), cephalometric X-ray with tracings, photographs, and other supporting documentation with the THSteps Dental Mandatory Prior Authorization Request Form.

All required documents must be submitted together in one package per prior authorization request. Prior authorization requests that are not submitted in one package per request will be considered incomplete.

**Note:** All documentation submitted with an incomplete request will be sent back to the provider with a letter that indicates the prior authorization request was incomplete. Providers must resubmit prior authorization requests with all the required documentation within 14 business days of the request receipt date, or the request will be denied as “incomplete.”
4.2.29.3.2 Diagnostic Tools

Prior authorization requests must include the date of service the diagnostic tools were obtained (the date of service the dental records were produced). All diagnostic tools must be properly labeled and protected when shipped by the provider. If any diagnostic tool is damaged during shipment, the provider may be required to reproduce the documentation for consideration of the case for prior authorization.

Note: If medical necessity cannot be determined from the diagnostic tools that are submitted with the request, the prior authorization request may be denied.

TMHP will be responsible for retaining an image of each diagnostic tool that is submitted for every complete orthodontic prior authorization request.

Copies of diagnostic models, X-rays, and any other paper diagnostic tools will be accepted and are preferred. Copies will not be returned, but providers will be required to maintain the dental records for retrospective review. Originals will be returned to the submitting provider only when the document is clearly marked “original.”

Diagnostic models in the form of plaster casts are preferred; however, providers may choose the positions in which the casts are made. E-models must be in the centric occlusion position.

Radiographs that are submitted must include, but are not limited to, the following:

- Panoramic or a full mouth series
- Cephalometric with tracings

Photographic images must be submitted with the request and must be in a 1:1 ratio format (actual size), including, but not limited to, the following:

- Full face, smiling
- Left and right profiles
- Full maxillary arch (open mouth view)
- Full mandibular arch (open mouth view)
- Right side occluded in centric occlusion
- Left side occluded in centric occlusion
- Anterior occluded in centric occlusion

X-rays must be of diagnostic quality and do not have to be submitted on photographic quality paper.

Submitting providers must attest that radiographs, photographs, and other documentation are unaltered.

4.2.29.3.3 Authorization Extensions

Extensions on allowed time frames may be considered no sooner than 60 days before the authorization expires. Extra monthly adjustments (procedure code D8670) will not be prior authorized, but the time frame may be considered for extension not to exceed 36 months of actual treatment. Providers must submit the following:

- Diagnostic workup.
  
  Note: Photographs may be substituted for models.
- The reason the treatment was not completed in the original time frame.
- An explanation of the treatment plan status.
4.2.29.3.4 Crossbite Therapy

Requests for crossbite therapy (procedure codes D8050 or D8060) require the submission of diagnostic models to receive authorization. An HLD score sheet is not required for crossbite therapy.

Providers that submit requests for crossbite therapy must maintain documentation in the client’s record that demonstrates the following criteria:

- Posterior teeth—Are not end-to-end, but the buccal cusp of the upper teeth is lingual to the buccal cusp of the lower teeth.
- Anterior teeth—The incisal edge of the upper teeth are lingual to the incisal edge of the opposing arch.

4.2.29.3.5 Minor Treatment to Control Harmful Habits

A THSteps Dental Mandatory Prior Authorization Form must be completed when requesting prior authorization for orthodontic appliances for minor treatment to control harmful habits. Documentation must support medical necessity of any appliance requested.

Providers must submit diagnostic models when requesting prior authorization for a removable appliance or fixed appliance.

Procedure codes D8210 or D8220 may only be approved for control of harmful habits including, but not limited to, thumb sucking or tongue thrusting and may not be prior authorized for services that are related to comprehensive orthodontic services.

4.2.29.3.6 Premature Termination of Orthodontic Services

Prior authorization for the premature termination of orthodontic services (procedure code D8680) is required.

Premature termination of orthodontic services includes all of the following:

- Removal of the brackets and arch wires.
- Other special orthodontic appliances.
- Fabrication of special orthodontic appliances.
- Delivery of orthodontic retainers.

The prior authorization must include all of the following for consideration:

- Panoramic radiograph (copies are preferred).
- Cephalometric radiograph with tracing (copies are preferred).
- Six intra-oral photographs (copies are preferred).
- Three extra-oral photographs (copies are preferred).
- A narrative documenting why the provider is terminating the orthodontic services early.
- Documentation that the parent, legal guardian, or the client, if he or she is 18 years of age or older or an emancipated minor, understands that the provider is terminating the orthodontic services, and the client is no longer eligible for orthodontic services by Texas Medicaid/THSteps.

In addition to the final record, the provider requesting premature termination of orthodontic services must submit a copy of the signed release form that includes the following:

A signature by one of the following:

- The parent
- Legal guardian
• The client, if he or she is 18 years of age or older or an emancipated minor

• One of the following statements:
  • The client is uncooperative or non-compliant with the treating dentist’s directions and does not intend to complete orthodontic treatment.
  • The client requested the premature removal of orthodontic appliances and does not intend to complete orthodontic treatment.

  **Note:** *A client for whom removal of an appliance has occurred due to the client’s request, or is uncooperative or non-compliant will not be eligible for any additional Medicaid orthodontic services.*

• The client has requested the premature removal of orthodontic appliances due to extenuating circumstances including, but not limited to, the following:
  • Incarceration.
  • Mental health complications with a recommendation from the treating physician.
  • Foster care placement.
  • Child of a migrant farm worker. With the intent to complete orthodontic treatment at a later date if Medicaid eligibility for orthodontic services continues.
  • Special medical conditions.

  **Note:** *If comprehensive orthodontic services are terminated due to extenuating circumstances, clients will be eligible for completion of their Medicaid orthodontic services if the services are re-initiated while the client is eligible for Medicaid.*

The requesting provider will be responsible for removal of the orthodontic appliances, final records, and fabrication and delivery of orthodontic retainers at the time of premature removal or at any future time should the client present to the treating provider’s office.

**4.2.29.3.7 Transfer of Services**

Prior authorization that is issued to a provider for orthodontic services is not transferable to another provider. The new provider must request a new prior authorization to complete the orthodontic treatment that was initiated by the original provider. The original prior authorization will be end-dated when services are transferred to another provider.

The new provider must obtain his or her own records, and the new request for orthodontic services must include the date of service on which the documentation was obtained (the date of service on which the records were produced) and the following supporting documentation:

• All of the documentation that is required for the original request

  **Note:** *Photographs may be substituted for models.*

• The reason the client left the previous provider

• An explanation of the treatment status

The authorization request for clients who are undergoing orthodontic treatment services and subsequently become eligible for Medicaid are subject to the same requirements.

**4.2.29.3.8 Orthodontic Cases Initiated Through a Private Arrangement**

Authorization may be given for continuation of orthodontic cases for clients who initiated orthodontic treatment through a private arrangement before becoming eligible for Medicaid.

Authorization will not be given for continuation of orthodontic cases for clients who initiated orthodontic treatment through a private arrangement and were eligible for Medicaid at the start of service.
4.2.30 THSteps and ICF-IID Dental Prior Authorization

Submit claims, dental correspondence, and THSteps and ICF-IID prior authorization requests to the appropriate address listed in the following table:

<table>
<thead>
<tr>
<th>Correspondence</th>
<th>Address</th>
</tr>
</thead>
<tbody>
<tr>
<td>ADA dental claim forms</td>
<td>Texas Medicaid &amp; Healthcare Partnership</td>
</tr>
<tr>
<td></td>
<td>PO Box 200555</td>
</tr>
<tr>
<td></td>
<td>Austin, TX 78720-0555</td>
</tr>
<tr>
<td>All dental correspondence</td>
<td>Texas Medicaid &amp; Healthcare Partnership</td>
</tr>
<tr>
<td>Prior authorization requests</td>
<td>Fee-for-Service and ICF-IID Dental Authorizations</td>
</tr>
<tr>
<td></td>
<td>PO Box 204206</td>
</tr>
<tr>
<td></td>
<td>Austin, TX 78720-4206</td>
</tr>
</tbody>
</table>

4.3 Documentation Requirements

All services require documentation to support the medical necessity of the service rendered, including dental services. Dental services are subject to retrospective review and recoupment if documentation does not support the service submitted for payment.

The provider must educate all staff members, including dentists, about the following documentation requirements and charting procedures:

- For THSteps and ICF-IID dental claims, providers are not required to submit preoperative and postoperative radiographs unless these are specifically requested by HHSC, the TMHP Dental Director, or are needed for prior authorization or pre-payment review.

- Documentation of all restorative, operative, crown and bridge, and fixed and removable prosthetics procedures must support the services that were performed and must demonstrate medical necessity that meets the professional standards of health care that are recognized by TSBDE. Documentation must include appropriate pretreatment, precementation and postcementation radiographs, study models and working casts, laboratory prescriptions, and invoices. Documentation must include the correct DOS. A panoramic radiograph without additional bitewing radiographs is considered inadequate as a diagnostic tool for caries detection. OIG may retrospectively recoup payment if the documentation does not support the services submitted for payment.

- All documentation must be maintained in the client’s record for a period of five years to support the medical necessity at the time of any post-payment utilization review. All documentation, including radiographs, must be of diagnostic and appropriate quality.

- In any situation where radiographs are required but cannot be obtained, intraoral photographs must be in the chart.

- Any complications, unusual circumstances encountered, morbidity, and mortality must be entered as a complete narrative in the client’s record.

- A provider must maintain a minimum standard of care through appropriate and adequate records, including a current history, limited physical examination, diagnosis, treatment plan, and written informed consent as a reasonable and prudent dentist would maintain. These records, as well as the actual treatment, must be in compliance with all state statutes, the Dental Practice Act, and the TSBDE Rules.

- Documentation for endodontic therapy must include the following: the medical necessity, pretreatment, during treatment, and post-treatment periapical radiographs, the final size of the file to which the canal was enlarged, and the type of filling material used. Any reason that the root canal may appear radiographically unacceptable must be entered in the chart. Endodontic therapy must be in compliance with the American Association of Endodontists quality assurance guidelines.
• Documentation for most periodontal services requires a six-point per tooth depth of pocket charting, a complete mouth series of periapical and bitewing radiographs, and any other narratives or supporting documentation consistent with the nationally accepted standards of care of the specialty of periodontics, and which conform to the minimum standard of care for periodontal treatment required of Texas dentists. A panoramic radiograph without additional bitewing or periapical radiographs is considered inadequate for diagnosis of periodontal problems.

• Documentation for surgical procedures requiring a definitive diagnosis for submitting a claim for a specific CDT code necessitates that a pathology report and a written record of clinical observations be present in the chart, together with any appropriate radiographs, operative reports, and appropriate supporting documentation. All impactions, surgical extractions, and residual tooth root extractions require appropriate preoperative periapical or panoramic radiographs (subject to limitations) be present in the chart.

• Any documentation requirements or limitations not mentioned in this manual that are present in the CDT are applicable. The written documentation requirements or limitations in this manual supersede those in the CDT.

4.3.1 General Anesthesia

When proceeding with Level 4 sedation/general anesthesia the dental provider is required to maintain the following documentation in the client’s dental record:

• The medical evaluation justifying the need for anesthesia
• Description of relevant behavior and reference scale
• Other relevant narratives justifying the need for general anesthesia
• Client’s demographics, including date of birth
• Relevant dental and medical history
• Dental radiographs, intraoral/perioral photography, or diagram of dental pathology
• Proposed dental plan of care
• Consent signed by parent or guardian giving permission for the proposed dental treatment and acknowledging that the reason for the use of IV sedation or general anesthesia for dental care has been explained
• Completed Criteria for Dental Therapy Under General Anesthesia form
• The parent or guardian dated signature on the Criteria for Dental Therapy Under General Anesthesia form attesting that they understand and agree with the dentist’s assessment of their child’s behavior
• Dentist’s attestation statement and signature, which may be put on the bottom of the Criteria for Dental Therapy Under General Anesthesia form or included in the client’s dental record as a stand alone form

4.3.2 Orthodontic Services

Requests for orthodontic services must be accompanied by all of the following documentation:

• An orthodontic treatment plan. The treatment plan must include all procedures required to complete full treatment (e.g., extractions, orthognathic surgery, upper and lower appliance, monthly adjustments, anticipated bracket replacements, appliance removal if indicated, special orthodontic appliances). The treatment plan should incorporate only the minimal number of appliances required to properly treat the case. Requests for multiple appliances to treat an individual arch are reviewed for duplication of purpose.
• Diagnostic models.
• Cephalometric radiograph with tracings.
• Completed and scored HLD sheet with diagnosis of Angle class (a minimum of 26 points is required for consideration of approval of non cleft palate cases).
• Facial photographs.
• Full series of radiographs or a panoramic radiograph; diagnostic-quality films are required (copies are preferred and will not be returned to the provider).
• Any additional pertinent information as determined by the dentist or requested by TMHP’s Dental Director. Requests for crossbite therapy require the submission of diagnostic models to receive authorization. Providers must maintain documentation in the client’s record that demonstrates the following criteria:
  • Posterior teeth. Not end-to-end, but buccal cusp of upper teeth should be lingual to buccal cusp of lower teeth.
  • Anterior teeth. The incisal edge of upper should be lingual to the incisal of the opposing arch.

The dentist should be certain that radiographs, photographs, and other information are properly packaged to avoid damage. TMHP is not responsible for lost or damaged materials.

Refer to: THSteps Dental Mandatory Prior Authorization Request Form on the TMHP website at www.tmhp.com.

4.4 Utilization Review

HHSC or a designated entity may conduct utilization reviews through automated analysis of a provider’s pattern(s) of practice, including peer group analysis. Such analysis may result in a subsequent on-site utilization review. HHSC or its claims processing contractor may conduct utilization reviews at the direction of the Office of Inspector General (OIG), according to HHSC rules.

DSHS may also conduct dental utilization reviews of randomly selected THSteps dental providers. These reviews compare Medicaid dental services that have been reimbursed to a dental provider to the results of an oral examination of the client as conducted by DSHS regional dentists.

Refer to: 25 TAC, §33.72 for more information about utilization review.

4.5 Claims Filing and Reimbursement

4.5.1 Reimbursement

The Medicaid rates for dentists are calculated as access-based fees in accordance with 1 TAC §§355.455(b), 355.8085, and 355.8441(11). Providers can refer to the online fee lookup (OFL) or the applicable fee schedule on the TMHP website at www.tmhp.com.

Texas Medicaid implemented mandated rate reductions for certain services. The OFL and static fee schedules include a column titled “Adjusted Fee” to display the individual fees with all mandated percentage reductions applied. Additional information about rate changes is available on the TMHP website at www.tmhp.com/resources/rate-and-code-updates.

4.5.2 Claim Submission After Loss of Eligibility

The Texas Medicaid 95-day filing deadline applies to all THSteps and ICF-IID dental services. If a client has lost Medicaid eligibility or turned 21 years of age, continue to file claims for services provided on the DOS the client was eligible. Indicate the actual DOS on the claim form, and enter the authorization number in the appropriate block on each claim filed.

4.5.3 Third Party Liability

Refer to: “Section 8: Third Party Liability (TPL)” (Vol. 1, General Information).

4.5.4 Claims Information

Dental services must be submitted to TMHP in an approved electronic format or on the ADA Dental Claim Form. Providers may purchase ADA Dental claim forms from the vendor of their choice. TMHP does not supply the forms. A sample of the ADA Dental Claim form can be found on the ADA website at www.ada.org.

When completing an ADA Dental claim form, all required information must be included on the claim, as TMHP does not key information from attachments. Superbills or itemized statements are not accepted as claim supplements.

All THSteps and ICF-IID claims must be received by TMHP within 95 days from each DOS and submitted to the following address:

Texas Medicaid & Healthcare Partnership
PO Box 200555
Austin, TX 78720-0555

Claims for emergency, orthodontic, or routine dental services must each be filed on separate forms. A claim submitted for either emergency or orthodontic services must be identified as such in Block 35 (Remarks) of the claim form.

A THSteps and ICF-IID dental provider cannot submit claims to Texas Medicaid under his individual performing provider identifier for the services provided by one or more associate dentists practicing in his office as employees or independent contractors with specific employer-employee or contractual relationships. All dentists providing services to Medicaid clients must enroll as THSteps dental providers regardless of employer relationships. The individual provider submitting claims may be reimbursed into a single accounting office to maintain these described relationships.

Claims submitted by newly-enrolled providers must be received within 95 days of the date the new provider identifier is issued, and within 365 days of the DOS.

Providers should submit claims to Texas Medicaid for their usual and customary fees.

Claims for dental services provided to children in foster care must be filed with DentaQuest, the dental claims processor for Superior HealthPlan.

Refer to: Subsection 4.2.7.2, “Children in Foster Care” in this handbook.

Claims must not be submitted to Texas Medicaid for appointments missed by clients. A client with Medicaid cannot be billed for failure to keep an appointment. Only claims for actual services rendered are considered for payment.

Refer to: “Section 3: TMHP Electronic Data Interchange (EDI)” (Vol. 1, General Information) for information on electronic claims submissions.

Section: Claims Filing” (Vol. 1, General Information).

Subsection 1.7.11, “Billing Clients” in “Section 1: Provider Enrollment and Responsibilities” (Vol. 1, General Information).

4.5.5 Claim Appeals

A claim denied because of age restrictions or other limitations listed in the Medicaid dental fee schedule may be considered for reimbursement on appeal when client medical necessity is provided to the TMHP Dental Director.

All denied claim appeals must be submitted to TMHP with the exception of a request to waive late filing deadlines. TMHP does not have the authority to waive state or federal mandates regarding claim filing deadlines.

If, after all appeal processes at TMHP have been exhausted, the provider remains dissatisfied with TMHP’s decision concerning the appeal, the provider may file a complaint with the HHSC Claims Administrator Operations Management Unit.

Refer to: Subsection 7.1.5, “Paper Appeals” in “Section 7: Appeals” (Vol. 1, General Information).

Subsection 7.3.1, “Administrative Claim Appeals” in “Section 7: Appeals” (Vol. 1, General Information).

Note: Providers must exhaust the appeals process with TMHP before filing a complaint to the HHSC Claims Administrator Operations Management Unit.

Providers may use one of three methods to appeal Medicaid claims to TMHP: telephone (AIS), paper, or electronic.

All appeals of denied claims or requests for adjustments on paid claims must be received by TMHP within 120 days of the date of disposition of the R&S Report on which the claim appears. If the 120-day appeal deadline falls on a weekend or TMHP-recognized holiday, the deadline will be extended to the next business day.

Certain claims must be appealed on paper; they cannot be appealed either electronically or by telephone.

Refer to: Subsection 7.1.5, “Paper Appeals” in “Section 7: Appeals” (Vol. 1, General Information) for information about appeals that may not be appealed electronically and claims that may not be appealed through AIS.

To appeal in writing:

If a claim cannot be appealed electronically or by telephone, appeal the claim on paper by completing the following steps:

1) Provide a copy of the R&S Report page where the claim is reported.
2) Circle one claim per R&S Report page.
3) Identify the information that was incorrectly provided and note the correct information that should be used to appeal the claim. If necessary, specify the reason for appealing the claim.
4) Attach radiographs or other necessary supporting documentation.
5) If available, attach a copy of the original claim. Claim copies are helpful when the appeal involves dental policy or procedure coding issues.
6) Do not copy supporting documentation on the opposite side of the R&S Report.
7) It is strongly recommended that providers submitting paper appeals retain a copy of the documentation being sent. It is also recommended that paper documentation be sent by certified mail with a return receipt requested to establish TMHP’s receipt of the claim and the date the claim was received. The provider is urged to retain copies of multiple claim submissions if the Medicaid provider identifier is pending.

Note: Claims submitted by newly-enrolled providers must be received within 95 days of the date the new provider identifier is issued, and within 365 days of the DOS.

8) Submit the paper appeal with supporting documentation and any radiographs and adjustment requests to the following address:

Texas Medicaid & Healthcare Partnership
Inquiry Control Unit
12357-B Riata Trace Parkway, Suite 100
Austin, TX 78727
To appeal by telephone:

1) Contact the Dental Line at 1-800-568-2460.
2) For each claim in question, have the R&S Report listing the claim and any supporting documents readily available.
3) Identify the claim submitted for appeal. The internal control number (ICN) will be requested.
4) Supply the information necessary to correct the claim, such as the missing tooth number or letter, the corrected procedure code, surface ID, or Medicaid number.

The appeal will appear as finalized or pending on the following week’s R&S Report.

Providers may also appeal electronically.

Electronic appeal submission is a method of submitting Texas Medicaid appeals using a personal computer. The electronic appeals feature can be accessed directly through the TMHP EDI Gateway or by using TexMedConnect. For additional information, contact the TMHP EDI Help Desk at 1-888-863-3638.

Electronic appeals can increase accuracy of claims processing, resulting in a more efficient case flow to the provider:

- Download and printout capabilities help maintain audit trails for the provider.
- Appeal submission windows can be automatically filled in with electronic R&S Report information, thereby reducing data entry time.

4.5.6 Frequently Asked Questions About Dental Claims

Q Why is routine dental treatment not a benefit when performed at the same visit as an emergency visit?

A The following are reasons routine dental treatment is not a benefit when performed at the same visit as an emergency visit:

- The purpose of an emergency claim is to allow the provider to treat a true emergency without the concern that routine dental procedures may be denied.
- Medicaid program policy guidelines do not allow payment for both emergency and routine services to the same provider at the same visit. True emergency claims process through the audit system correctly when “emergency” is checked on either the paper or electronic claim and the Remarks or Narrative section of the claim form describes the nature of the emergency.

Q Why are some claims for oral exams and emergency exams on the same date for the same client denied?

A Medicaid program policy does not allow claims for an initial oral exam and an emergency exam to be submitted for the same DOS for the same client. An emergency exam performed by the same provider in the same six-month time period as an initial exam may be considered for reimbursement only when the claim for the emergency exam indicates it is an emergency and the emergency block is marked and the Remarks or Narrative section is completed. If the claim is not marked as an emergency, the claim will be denied.

Q How are orthodontic bracket replacements reimbursed? Can the client be charged for bracket replacements?

A The provider must use orthodontic procedure code D8690 to claim reimbursement for bracket replacement. Medical necessity must be documented in the client record. Payment is subject to retrospective review. The client with current Medicaid eligibility must not be charged for bracket replacement. If the provider charges the client erroneously, the provider must refund any amount paid by the client.

Q Why could an appeal of a denied claim take a long time?
A An appeal can take a long time if TMHP is required to research the denied claim and determine the reason the claim did not go through the system. For faster results, providers should submit appeals as soon as possible and not use the entire 120 days allowed to submit the appeal. The following are guidelines on filing claims efficiently:

- Use R&S Report dates to track filed claims.

- File claims electronically through TMHP EDI. Electronic claims submission does not allow a claim with an incorrect date to be accepted and processed, which saves time for the provider submitting claims and TMHP in processing claims. Call 1-888-863-3638, for more information about TMHP EDI.

- File claims with the correct information included. Most denied claims result from the omission of dates, signature, or narrative, or incorrect ID numbers such as client Medicaid numbers or provider identifiers.

Q Why are only ten appeals allowed per call?
A There is a limit on appeals per call to allow all providers equal access.

Q Why do reimbursement checks sometimes take a long time to arrive?
A Reimbursement may be delayed if a provider fails to submit claims in a timely manner.

Q Does electronic claims submission result in delayed payment?
A No. Providers who submit claims electronically report faster results than when submitting claims on paper. Providers are encouraged to use TMHP EDI for claims submission.

The following are helpful hints to a more efficiently processed claim:

- Ensure the provider identifier is on all claims.

- Include the performing provider’s signature on all paper claims.

- Verify client eligibility for procedures.

- Verify if the procedure code requires a narrative on the claim; the narrative is for medical necessity.

- Include the required client information, including name, birth date, and client number.

- Dental auxiliary staff (i.e., the hygienist or the chairside assistant) cannot enroll in Texas Medicaid; therefore, they cannot submit claims to Texas Medicaid. Any procedure performed by the auxiliary must be submitted by the supervising dentist, using the dentist’s provider identifier.

Claim Submission Reminders:

- Procedure code D8660 is allowed at different age levels, per provider. If a claim for procedure code D8660 is submitted within six months of procedure code D8080, procedure code D8080 will be reduced by the amount that was paid for procedure code D8660.

- Prior authorization is required with documentation of medical necessity when replacing lost or broken orthodontic retainers (procedure code D8680). Clients may not be billed for covered services.

- Prior authorization of orthodontic services is nontransferable. If a client changes an orthodontic provider for any reason, or a provider ceases to be a Medicaid provider, the new orthodontic services provider must submit a separate request for prior authorization. The provider requesting and receiving authorization for the service also must perform the service and submit the claim. Codes listed on the authorization letters are the only codes considered for payment. All other codes submitted for payment are denied. Providing the authorization number on the submitted claim results in more efficient claims processing.
• Prior authorization is required for clients who are 7 through 20 years of age that are in need of general anesthesia and do not meet the Criteria for Dental Therapy Under General Anesthesia requirements (22 point threshold). Prior authorization is required for medically necessary dental general anesthesia that exceeds once per six months, per client, any provider. The dentist providing therapeutic services under general anesthesia is responsible for obtaining prior authorization for both services.

• General anesthesia (provided in the dentist office, ambulatory service clinic, and inpatient/outpatient hospital settings) does not require prior authorization for clients who are 7 through 20 years of age, unless the client does not meet the minimum required points for general anesthesia in Criteria for Dental Therapy Under General Anesthesia on the TMHP website at www.tmhp.com. All THSteps dental charts for dental general anesthesia are subject to retrospective, random review for compliance with the Criteria for Dental Therapy Under General Anesthesia and requirements for chart documentation.

• Providers must not bill a client unless a formal denial for the requested item or service has been issued by TMHP stating the service is not a benefit of Texas Medicaid and the client has signed the Client Acknowledgment Statement in advance of the service being provided for that specific item or service. A provider must not bill Medicaid clients if the provided service is a benefit of Texas Medicaid.

Refer to: Subsection 1.7.11.1, “Client Acknowledgment Statement” in “Section 1: Provider Enrollment and Responsibilities” (Vol. 1, General Information).

THSteps clients must receive:

• Dental services specified in the treatment plan that meet the standards of care established by the laws relating to the practice of dentistry and the rules and regulations of the TSBDE.

• Dental services that are free from abuse or harm from the provider or the provider’s staff.

• Only the treatment required to address documented medical necessity that meets professionally recognized standards of health care.

5 THSteps Medical

5.1 THSteps Medical and Dental Administrative Information

5.1.1 Overview

This section describes the administrative requirements for THSteps, including provider requirements, client eligibility requirements, and billing and claims processing information. Providers that need additional information may call 1-800-757-5691 or refer to “Appendix F. Texas Health Steps Quick Reference Guide” in this handbook for a more specific list of resources and telephone numbers. Providers may also contact the Texas Department of State Health Services (DSHS) THSteps Provider Relations staff located in DSHS regional offices by calling the appropriate regional office as listed in “Appendix A: State, Federal, and TMHP Contact Information” (Vol. 1, General Information). THSteps Provider Relations contact information is also available on the DSHS website at www.dshs.texas.gov/thsteps/regions.shtm.

In addition, THSteps has developed online educational modules to provide additional information about the program, components of the medical checkup, and other information. These modules provide free continuing education hours for a variety of providers. Providers do not have to be enrolled in THSteps. These courses may be accessed at www.txhealthsteps.com.

The Early and Periodic Screening, Diagnosis, and Treatment (EPSDT) service is Medicaid’s comprehensive preventive child health service for clients who are birth through 20 years of age. In Texas, EPSDT is known as THSteps and includes periodic screening, vision, hearing, and dental preventive and
treatment services. EPSDT was created by the 1967 amendments to the federal Social Security Act and defined by the Omnibus Budget Reconciliation Act (OBRA) of 1989. The periodic screening for a checkup consists of five federally required components as noted on the THSteps Periodicity Schedule. In addition, Section 1905(r)(5) of the Social Security Act (SSA) requires that any medically necessary health-care service listed in the Act be provided to EPSDT clients even if the service is not available under the state’s Medicaid plan to the rest of the Medicaid population. A service is medically necessary when it corrects or ameliorates the client’s disability, physical or mental illness, or chronic condition. These additional services are available through CCP. For questions about coverage, providers can call CCP at 1-800-846-7470.

5.1.2 Statutory Requirements

Several specific legislative requirements affect THSteps and the providers participating in the program. These include, but are not limited to, the following:

- Newborn Screening, Health and Safety Code, Chapter 33, Section §33.011 Newborn Screening Test Requirement.
- Subsection D.5, “Parental Accompaniment” in this handbook.
- Requirements for Reporting Abuse or Neglect, as outlined in subsection 1.7.1, “Compliance with Texas Family Code” in “Section 1: Provider Enrollment and Responsibilities” (Vol. I, General Information).
- Early Childhood Intervention (ECI), 34 Code of Federal Regulations (CFR) Part 303; Chapter 73, Texas Human Resources Code, and Title 40 TAC, Chapter 108.
- Newborn Hearing Screening, Health and Safety Code, Chapter 47.
- Teen Confidentiality Issues. There are many state statutes that may affect consent to medical care for a minor, depending on the facts of the situation. Among the relevant statutes are Chapters 32, 33, 153, and 266 of the Texas Family Code. Providers may want to consult an attorney, their licensing board, or professional organization if guidance is needed or questions arise on matters of medical consent.

Refer to: “Appendix D. Texas Health Steps Statutory State Requirements” in this handbook for more information.

5.1.3 Texas Vaccines for Children (TVFC) Program

The TVFC program provides vaccines at no cost to the provider. The vaccines are recommended according to the Recommended Childhood and Adolescent Immunization Schedule (Advisory Committee on Immunization Practices [ACIP], AAP, and the American Academy of Family Physicians [AAFP]). Medicaid does not reimburse for vaccines/toxoids that are available from TVFC. THSteps providers are strongly encouraged to enroll in TVFC at DSHS and must do so in order to obtain free vaccines for clients who are birth through 18 years of age. Local and public health departments that are not otherwise enrolled as a provider that is authorized to receive reimbursement for vaccine administration fees should enroll as a Comprehensive Care Program (CCP) provider. Providers may not charge Texas Medicaid for the cost of the vaccines obtained from TVFC; however the administration fee, not to exceed $14.85, is considered for reimbursement.

When single antigen vaccine(s)/toxoid(s) or comparable antigen vaccine(s)/toxoid(s) are available for distribution through TVFC, but the provider chooses to use an ACIP-recommended product that is not distributed through TVFC, the vaccine/toxoid will not be covered; however, the administration fee will be considered.

Note: Administered vaccines/toxoids must be reported to DSHS. DSHS submits all vaccines/toxoids reported with parental consent to a centralized repository of immunization histories for clients younger than 18 years of age. This repository is known in Texas as ImmTrac2.
For additional information about immunizations, providers can refer to the THSteps online educational module “Immunization” at www.txhealthsteps.com.

Refer to: “Appendix B. Immunizations” in this handbook.

5.1.4 Vaccine Adverse Event Reporting System (VAERS)

The National Childhood Vaccine Injury Act (NCVIA) of 1986 requires health-care providers to report:

- Any reaction listed by the vaccine manufacturer as a contraindication to subsequent doses of the vaccine.
- Any reaction listed in the Reportable Events Table that occurs within the specified time period after vaccination.

NCVIA requires health-care providers to report certain adverse events that occur following vaccination. As a result, VAERS was established by CDC and FDA in 1990. VAERS provides a mechanism for the collection and analysis of adverse events (side effects) associated with vaccines currently licensed in the United States. Adverse events are defined as health effects that occur after immunization that may or may not be related to the vaccine. VAERS data are monitored continually to detect unknown adverse events or increases in known side effects.

A copy of the Reportable Events Table can be obtained by calling VAERS at 1-800-822-7967 or by downloading it from www.dshs.texas.gov/immunize/forms/vaers_table.pdf.

Clinically significant adverse events should be reported even if it is unclear whether a vaccine caused the event. For additional information about NCVIA, providers can refer to www.dshs.texas.gov/immunize/forms/11-11246.

5.1.5 Referrals for Medicaid-Covered Services

When a provider performing a checkup determines that a referral for diagnosis or treatment is necessary for a condition found during the medical checkup, that information must be discussed with the parents or guardians. A referral must be made to a provider who is qualified to perform the necessary diagnosis or treatment services. If the performing provider is competent to treat the condition found, a referral elsewhere is not necessary, unless it is to the primary care provider to assure continuity of care.

Providers that need assistance finding a specialist who accepts clients with Medicaid coverage can call the THSteps toll-free helpline at 1-877-847-8377, or they can find one using the Online Provider Lookup on the TMHP website at www.tmhp.com.

Continuity of care is an important aspect of providing services and follow-up. Efforts should be made to determine that the appointment was kept and that the provider who received the referral has provided a diagnosis and recommendations for further care to the referring provider.

In addition to referrals for conditions discovered during a checkup or for specialized care, the following referrals may be used:


- **Hearing Services referrals.** If the hearing screening returns abnormal results, clients who are birth through 20 years of age must be referred to a Texas Medicaid provider who is an audiologist or physician who is experienced with the pediatric population and who offers auditory services.
• **Routine Dental Referrals.** The provider must refer clients to establish a dental home beginning at 6 months of age or earlier if trauma or early childhood caries are identified. For established clients after the 6-month medical checkup, the provider must confirm if a dental home has been established and is ongoing; if not, additional referrals must be made at subsequent medical checkups until the parent or caregiver confirms that a dental home has been established for the client. Clients who are birth through 5 months of age are not eligible for routine dental checkups but should be referred to a dentist if any dental issues are identified during a THSteps medical checkup or acute care visit. When possible, clients should be referred to a provider who has completed the required benefit education and is certified by Texas Health Steps to perform First Dental Home services. The First Dental Home provider may be located through the advanced search function in the Online Provider Look Up or by calling 1-877-847-8377.

• **Referrals for Dental Treatment.** If a THSteps medical provider identifies the medical necessity of dental services, the provider must refer the client to a THSteps dental provider. The THSteps medical provider can accomplish this by providing the parent or guardian a listing of THSteps dentists from the Online Provider Lookup. The parent or guardian can receive assistance in locating a THSteps dentist and assistance with scheduling of dental appointments by contacting the THSteps toll-free helpline at 1-877-847-8377. Clients who are birth through 5 months of age also can be seen for emergency dental services by the dentist at any time for trauma, early childhood caries, or other oral health problems. Clients who are birth through 20 years of age may self-refer for dental care.

• **Emergency Dental Referrals.** If a medical checkup provider identifies an emergency need for dental services, such as bleeding, infection, or excessive pain, the client may be referred directly to a participating dental provider. Emergency dental services are covered at any time for all Medicaid clients who are birth through 20 years of age.

  **Note:** Assistance in coordinating dental referrals can be obtained from the THSteps toll-free helpline at 1-877-847-8377 or the DSHS Regional THSteps Coordinator for the respective region (lists are provided in “Appendix A: State, Federal, and TMHP Contact Information” (Vol. 1, General Information). In cases of both emergency and nonemergency dental services, clients are able to make a choice when selecting a dental provider who is participating in the THSteps Dental Program.

• **Family Planning and Genetic Services Referrals.** For clients eligible for Medicaid who need genetic services or family planning services, a referral should be made. Information about Medicaid-covered genetic services is available in the Medical and Nursing Specialists, Physicians, and Physician Assistants Handbook (Vol. 2, Provider Handbooks) and information about family planning services is available in Section 2, “Medicaid Title XIX Family Planning Services” in the Gynecological, Obstetrics, and Family Planning Title XIX Services Handbook (Vol. 2, Provider Handbooks). If a THSteps medical provider also provides family planning, the provider may inform clients that these services are available.

• **ECI Referrals.** Federal and state law requires providers to refer children as soon as possible, but no longer than 7 days after identification of a suspected developmental delay or disability to the local ECI program for children who are birth through 35 months of age regardless if a referral was made to another qualified provider. The provider may call the local ECI program or the Health and Human Services Office of the Ombudsman at 1-877-787-8999 to make referrals. Children who are 3 years of age and older with a suspected developmental delay or disability should be referred to the local school district.

• **WIC Referrals.** Clients who are birth through 5 years of age or who are pregnant are eligible for WIC and should be referred to WIC for nutrition education and counseling, and food benefits.

Refer to: Section 2, “Table of Contents” in the Medicaid Managed Care Handbook (Vol. 2, Provider Handbooks) for more information about referrals.
5.1.6 THSteps Medical Checkup Facilities

All THSteps medical checkup policies apply to checkups completed in a physician’s office, a health department, clinic setting, or in a mobile/satellite unit. Enrollment of a mobile/satellite unit must be under a physician or clinic name. Mobile units can be a van or any area away from the primary office and are considered extensions of that office and are not separate entities.

The physical setting must be appropriate so that all elements of the checkup can be completed.

Refer to:
- Subsection 5.3.10, “THSteps Medical Checkups Periodicity Schedule” in this handbook on the THSteps Periodicity Schedule.
- Subsection 5.3.11, “Mandated Components” in this handbook for additional information on checkup components.

5.1.7 THSteps Dental Services

Access to THSteps dental services is mandated by Texas Medicaid and provides reimbursement for the early detection and treatment of dental health problems, including oral health preventive services, for Medicaid clients who are birth through 20 years of age. THSteps dental service standards are designed to meet federal regulations and to incorporate the recommendations of representatives of national and state dental professional groups.

OBRA 1989 mandated the expansion of the federal EPSDT program to include any service that is medically necessary and for which FFP is available, regardless of the limitations of Texas Medicaid. This expansion is referred to as CCP.

Refer to:
- Section 2, “Medicaid Children’s Services Comprehensive Care Program (CCP)” in this handbook for more information.

THSteps-designated staff (HHSC, DSHS, or its designee), through outreach and education, encourage the parents or caregivers of eligible clients to use THSteps dental checkups and preventive care when clients first become eligible for Medicaid and each time clients are due for their next periodic dental checkup.

Upon request, THSteps-designated staff (HHSC, DSHS, or its designee) assist the parents or caregivers of eligible clients with scheduling appointments and transportation. Medicaid clients have freedom of choice of providers and are given names of enrolled providers. Call the THSteps toll-free helpline at 1-877-847-8377 for a list of THSteps dental providers in a specific area.

For additional information about dental health, providers can refer to the THSteps online educational modules "Oral Health For Primary Care Providers" and "Oral Health Examinations for Dental Professionals" at www.txhealthsteps.com.

5.2 Enrollment

5.2.1 THSteps Medical Provider Enrollment

Providers cannot be enrolled if their professional license is due to expire within 30 days of application. Facility providers must submit a current copy of the supervising practitioner’s license. To provide Medicaid services, each NP or CNS must be licensed as an RN and be recognized as an APRN by Texas BON.

Refer to: Subsection 1.1, “Provider Enrollment” in “Section 1: Provider Enrollment and Responsibilities” (Vol. 1, General Information) for information about enrollment procedures.

The following provider types may provide THSteps preventive services within his or her scope of practice, must be enrolled in Texas Medicaid, but do not have to be enrolled as a THSteps provider:

- A federal qualified health center (FQHC)
- A rural health clinic (RHC)
The following provider types may provide THSteps preventive services within his or her scope of practice and must be enrolled in Texas Medicaid as a THSteps provider:

- A physician (doctor of medicine [M.D.] or doctor of osteopathy [D.O.]) or physician group
- A physician assistant (PA)
- A clinical nurse specialist (CNS)
- A nurse practitioner (NP)
- A certified nurse midwife (CNM)
- A health-care provider or facility with physician supervision including, but not limited to:
  - Community-based hospital and clinic
  - Family planning clinic
  - Home health agency
  - Local or regional health department
  - Maternity clinic
  - Migrant health center
  - School-based health center

**Medical Residents**

Medical residents may provide medical checkups in a teaching facility under the guidance of the attending staff as long as the facility’s medical staff by-laws and requirements of the Graduate Medical Education (GME) Program are met, and the attending physician has determined the intern or resident to be competent to perform checkups. THSteps does not require the supervising physician to examine the client as long as these conditions are met.

**Clinics**

In a clinic, a physician is not required to be present at all times during the hours of operation unless otherwise required by federal regulations. A physician must assume responsibility for the clinic’s operation.

5.2.1.1 **Requirements for Registered Nurses Who Provide Medical Checkups**

RNs without a CNS, NP, or CNM recognition as an APRN by the Texas BON may provide medical checkups only under direct physician supervision, meaning the physician is either on site during the checkup or immediately available to furnish assistance and direction to the RN during the checkup.

Required online education modules developed by THSteps must be completed prior to providing checkup services. All modules are approved for continuing education units (CEUs) for RN’s as well as other medical disciplines. Required THSteps online education modules are available on the [RN Information page](#) of the THSteps website. The RN or the RN’s employer must maintain documentation that the required modules were completed.

Online modules are updated regularly to include new content. RNs that have completed the required modules previously are encouraged, but not required to retake online modules.

Before a physician delegates a THSteps checkup to an RN, the physician must establish the RN’s competency to perform the service as required by the physician’s scope of practice. The delegating physician is responsible for supervising the RN who performs the services. The delegating physician remains responsible for any service provided to a client.

**Refer to:** Subsection 5.2.1, “THSteps Medical Provider Enrollment” in this handbook for more information about enrollment procedures.
5.3  Services, Benefits, Limitations, and Prior Authorization

5.3.1  Eligibility for THSteps Services and Checkup Due Dates
Through outreach, THSteps staff (DSHS, HHSC, or contractors) encourage clients to use THSteps preventive medical checkup services when they first become eligible for Medicaid and each time thereafter when they are periodically due for their next medical checkup. THSteps will send clients a letter when they are due for a medical checkup.

A client is eligible for THSteps services, including medical checkups, from birth through 20 years of age.

Although the Medicaid Eligibility Verification Letter (Form H1027) identifies eligible clients when the client’s Your Texas Benefits Medicaid card is lost or has not yet been issued, Form H1027 does not indicate whether the client is due for medical checkup services. Providers can verify the client’s eligibility through TexMedConnect, the Medicaid Client Portal for Providers, or the TMHP Contact Center.

A client is due for a THSteps medical checkup based on his or her date of birth and the ages indicated on the periodicity schedule. Children younger than three years of age are due at frequent intervals. Children and youth three years of age and older are considered due for a checkup on their birthday and are encouraged to have a yearly checkup as soon as practical. In addition, for children enrolled in Medicaid managed care, a new member is due for a THSteps medical checkup as soon as practicable, but in no case later than 14 days of enrollment for newborns, and no later than 90 days of enrollment for all other eligible child members.

Providers should schedule checkups based on the ages in the periodicity schedule, but circumstances may support the need for a checkup prior to the client’s birthday (for example, a 4-year checkup could be performed prior to the child’s 4th birthday if the child is a member of a migrant family that is leaving the area). THSteps fee-for-service policy creates this flexibility by allowing a total number of checkups at each age range.

Refer to: “Subsection 5.3.6, “THSteps Medical Checkups” in this handbook for additional details.

Providers are encouraged to notify the client when they are due for the next checkup according to the THSteps periodicity schedule.

A checkup that is necessary more frequently than indicated on the periodicity schedule is considered an exception-to-periodicity.

Refer to: Subsection 5.3.7, “Exception-to-Periodicity Checkups” in this handbook for additional details about billing for a checkup performed as an exception-to-periodicity checkup.

5.3.2  Prior Authorization
Prior authorization is not required for preventive care medical checkups.

5.3.3  Additional Consent Requirements
Additional parental or guardian consent may be required if online or web-based screening tools are used that could result in client data being stored electronically in an outside database other than the provider’s electronic medical record system, or if the data is used for purposes other than THSteps screening. The provider should seek legal advice regarding the need for this consent.

5.3.4  Verification of Medical Checkups
The first source of verification that a THSteps medical checkup has occurred is a paid claim or encounter. THSteps encourages providers to file a claim either electronically or on a CMS-1500 paper claim form as soon as possible after the date of service, as the paid claim updates client information. The provider may contact TMHP through the TMHP website at www.tmhp.com or AIS at 1-800-925-9126 to verify that the client is due for a checkup.
A second source of acceptable verification is a physician’s written statement that the checkup occurred. If the provider chooses to give the client written verification, it must include the client’s name, Medicaid ID number, date of the medical checkup, and a notation that a complete THSteps medical checkup was performed.

**Note:** Verification of medical checkups must not be sent to THSteps but must be maintained by the client to be provided as needed by an HHSC eligibility caseworker.

If neither the first nor the secondary source of verification is available, a THSteps outreach worker may contact the provider’s office for verification.

### 5.3.5 Medical Home

HHSC and DSHS encourage the provision of the THSteps medical checkup as part of a medical home. Texas Medicaid defines a medical home as a model of delivering care that is accessible, continuous, comprehensive, family-centered, and coordinated. In providing a medical home for the client, the primary care clinician directs care coordination together with the client or youth and/or family.

Medical checkup providers with mobile units should encourage the families to establish a medical home for their child(ren) and obtain future checkups from their primary care provider.

When a checkup is provided in the home setting, mobile unit, or clinic other than the medical home, it should be in coordination with the medical home and the results must be provided to the medical home as soon as possible.

A mobile unit is an extension of the provider’s office and must be able to provide a complete checkup.

For additional information on the medical home, providers can refer to the “Introduction to the Medical Home” module provided by THSteps at [www.txhealthsteps.com](http://www.txhealthsteps.com).

### 5.3.6 THSteps Medical Checkups

THSteps medical checkups reflect the federal and state requirements for a preventive checkup. Preventive care medical checkups are a benefit of the THSteps program if they are provided by enrolled THSteps providers and all of the required components are completed. An incomplete preventive medical checkup is not a benefit. The THSteps periodicity schedule specifies screening procedures required at each stage of the client’s life to ensure that health screenings occur at age-appropriate points in a client’s life.

Components of a medical checkup that have an available CPT code are not reimbursed separately on the same day as a medical checkup, with the exception of initial point-of-care blood lead testing, mental health screening for adolescents, postpartum depression screening, tuberculin skin test (TST), developmental and autism screening, vaccine administration, and oral evaluation and fluoride varnish (OEFV).

**Note:** Initial blood lead testing, other than point-of-care, must be sent to the DSHS Laboratory for testing.

**Reminder:** Incomplete medical checkups are subject to recoupment unless there is documentation supporting why a component was not completed.

**Refer:** Subsection 5.3.11.1.3, “Mental Health Screening” in this handbook for more information about required mental health screenings.

Sports physical examinations are not a benefit of Texas Medicaid. If the client is due for a THSteps medical checkup and a comprehensive medical checkup is completed, a THSteps medical checkup may be reimbursed and the provider may complete the documentation for the sports physical.

THSteps preventive medical checkups are not a benefit under telemedicine or telehealth.

**Refer:** The THSteps Medical Checkups Periodicity Schedule which may be found at [https://hhs.texas.gov/doing-business-hhs/provider-portals/health-services-providers/texas-health-steps/medical-providers](https://hhs.texas.gov/doing-business-hhs/provider-portals/health-services-providers/texas-health-steps/medical-providers).
Checkups should be scheduled, to the extent possible, based on the ages on the periodicity schedule to accommodate the need for flexibility when scheduling checkup appointments.

The following table lists the number of visits allowed at each age range:

<table>
<thead>
<tr>
<th>Age Range</th>
<th>Number of Visits</th>
</tr>
</thead>
<tbody>
<tr>
<td>Birth through 11 months (does not include 12 month checkup)</td>
<td>6</td>
</tr>
<tr>
<td>1 through 4 years</td>
<td>7</td>
</tr>
<tr>
<td>5 through 11 years</td>
<td>7</td>
</tr>
<tr>
<td>12 through 17 years</td>
<td>6</td>
</tr>
<tr>
<td>18 through 20 years</td>
<td>3</td>
</tr>
</tbody>
</table>

All of the checkups listed on the periodicity schedule were developed according to the recommendations of the AAP and in consultation with recognized authorities in pediatric preventive health. In Texas, the THSteps periodicity schedule may differ from the AAP periodicity schedule based on the scheduling of laboratory or other tests in federal EPSDT or state regulations.

For more information about conducting a THSteps checkup, providers can refer to the THSteps online educational modules at [www.txhealthsteps.com](http://www.txhealthsteps.com).

The following table includes the procedure codes, required condition indicators, and the resulting referral status for medical checkups. Condition indicators must be used in addition to a provider type modifier at each THSteps checkup. A condition indicator must be submitted on the claim with the periodic medical checkup procedure code. Condition indicators are required whether a referral was made or not. If a referral is made, then providers must use the Y referral status. If no referral is made, then providers must use the N referral status.

<table>
<thead>
<tr>
<th>Procedure Codes</th>
<th>Condition Indicators</th>
<th>Referral Status</th>
</tr>
</thead>
<tbody>
<tr>
<td>99381, 99382, 99383, 99384, and 99385 (new client preventive visit) -or- 99391, 99392, 99393, 99394, and 99395 (Established client preventive visit)</td>
<td>NU (not used)</td>
<td>N (no referral given)</td>
</tr>
<tr>
<td>99381, 99382, 99383, 99384, and 99385 (new client preventive visit) -or- 99391, 99392, 99393, 99394, and 99395 (established client preventive visit)</td>
<td>S2 (under treatment) or ST* (new services requested)</td>
<td>Y (yes THSteps or EPSDT referral was given to the client)</td>
</tr>
</tbody>
</table>

* The ST condition indicator should only be used when a referral is made to another provider or the client must be rescheduled for another appointment with the same provider. It does not include treatment initiated at the time of the checkup.

THSteps preventive care medical checkups for clients who are 18 through 20 years of age must be submitted with procedure codes 99385 or 99395 and diagnosis code Z0000 or Z0001.

Claims for procedure codes 99381, 99382, 99383, 99384, 99391, 99392, 99393, and 99394 must be submitted with the appropriate age related diagnosis code listed in the following table:

<table>
<thead>
<tr>
<th>Client Age</th>
<th>Diagnosis Code</th>
</tr>
</thead>
<tbody>
<tr>
<td>Birth through 7 days</td>
<td>Z00110</td>
</tr>
<tr>
<td>8 through 28 days</td>
<td>Z00111</td>
</tr>
</tbody>
</table>
The age-appropriate diagnosis code for a preventive care medical checkup must be submitted on the claim. If an immunization is administered as part of the preventive care medical checkup, diagnosis code Z23 may also be included on the claim, in addition to the age-appropriate diagnosis.

If an immunization is the only service provided during an office visit, providers may submit only diagnosis code Z23 on the claim.

**Note:** *A THSteps preventive care medical checkup will not be reimbursed if the office visit is only for immunization.*

Modifier AM, SA, TD, or U7 must be submitted with the THSteps medical checkups procedure code to indicate the practitioner who performed the unclothed physical examination during the medical checkup.

<table>
<thead>
<tr>
<th>Modifier</th>
<th>Practitioner</th>
</tr>
</thead>
<tbody>
<tr>
<td>AM</td>
<td>Physician, team member service</td>
</tr>
<tr>
<td>SA</td>
<td>Nurse practitioner rendering service in collaboration with a physician</td>
</tr>
<tr>
<td>TD</td>
<td>Registered nurse</td>
</tr>
<tr>
<td>U7</td>
<td>Physician assistant</td>
</tr>
</tbody>
</table>

THSteps medical checkups performed in an FQHC or RHC setting are paid an all-inclusive rate per encounter, which includes immunizations, developmental screening, autism screening, mental health screening for adolescents, postpartum depression screening, TST, blood lead test, and OEFV. When submitting claims for THSteps checkups and services, RHC providers must use the national POS code 72, and FQHC providers must use modifier EP in addition to the modifiers used to identify who performed the medical checkup. In accordance with the federal rules for RHCs and FQHCs, an RN in an RHC or FQHC may not perform THSteps checkups independently of a physician’s interactions with the client.

**Refer to:** Section 4, “Federally Qualified Health Center (FQHC)” in the *Clinics and Other Outpatient Facility Services Handbook (Vol. 2, Provider Handbooks)* for information related to billing.

Checkups, exception-to-periodicity checkups, and follow-up visits are limited to once per day any provider.

A checkup and the associated follow-up visit may not be reimbursed on the same date of service. The follow-up visit will be denied.

An incomplete checkup is subject to recoupment unless there is documentation to support why the component was not completed as part of the checkup.

A new patient is one who has not received any professional services within the preceding three years from the provider or from another provider of the same specialty who belongs to the same group practice. As an exception, a new preventive care medical checkup (procedure code 99381, 99382, 99383, 99384, or 99385) may be billed when no prior checkups have been billed by the same provider or provider group, even if an acute care new patient E/M service was previously performed by the same provider.
An additional new checkup is allowed only when the client has not received any professional services in the preceding three years from the same provider or another provider who belongs to the same group practice, because subsequent acute care visits to the new patient THSteps checkup continues the established relationship with the provider.

If the provider that performs the medical checkup provides treatment for an identified condition on the same day, the provider may submit a separate claim for an acute care established-client office visit. The separate claim must include the established-client procedure code that is appropriate for the diagnosis and treatment of the identified problem. Treatment of minor illnesses or conditions (e.g., follow-up of a mild upper respiratory infection) during the THSteps medical checkup may not warrant additional billing.

### Acute Care Visits

When a new patient checkup is billed for the same date of service as a new patient acute care visit, both new patient services may be reimbursed when billed by the same provider or provider group if no other acute care visits or preventive care medical checkups have been billed in the past three years.

Providers must use modifier 25 to describe circumstances in which an acute care E/M visit was provided at the same time as a checkup. Providers must submit modifier 25 with the E/M procedure code when the rendered services are distinct and provided for a different diagnosis. Providers must bill an appropriate level E/M procedure code with the diagnosis that supports the acute care visit. The medical record must contain documentation that supports the medical necessity and the level of service of the E/M procedure code that is submitted for reimbursement.

An acute care E/M visit for an insignificant or trivial problem or abnormality billed on the same date of service as a checkup or exception-to-periodicity checkup is subject to recoupment.

Providers must bill an acute care visit with their provider identifier on a separate claim without benefit code EP1.

Refer to: [Acute Care Visit on the Same Day as a THSteps Preventive Visit Checkup](https://www.tmhp.com) on the TMHP website at www.tmhp.com for a claim form example.

[THSteps Preventive Visit Checkup with Immunization and Vaccine Administration](https://www.tmhp.com) on the TMHP website at www.tmhp.com for a claim form example.

### 5.3.7 Exception-to-Periodicity Checkups

Exception-to-periodicity checkups are complete medical checkups completed outside the timeframes listed in the THSteps Periodicity Schedule due to extenuating circumstances.

Exception-to-periodicity checkups are complete medical checkups, which are medically necessary and might cause the total number of checkups to exceed the number allowed for the client’s age range if the client were to have all regular scheduled checkups. An exception-to-periodicity checkup is allowed when:

- Medically necessary, for example, for a client with developmental delay, suspected abuse, or other medical concerns or a client in a high-risk environment, such as living with a sibling with elevated blood lead.
- Required to meet state or federal exam requirements for Head Start, day care, foster care, or preadoption.
- When needed before a dental procedure requiring general anesthesia.

As noted in the Periodic Checkup Age Range table, the number of checkups is set for each age range. This may avoid an exception-to-periodicity checkup and allow flexibility for the provider and family to schedule a checkup including before the child’s birthday.
If a client is due for a medical checkup, a checkup outside of the regular THSteps schedule must be billed as a regular checkup rather than an exception to periodicity.

The checkup is considered complete when all the required components are documented in the client’s medical record or supporting documentation, which details the reason a component(s) was not completed. A plan to complete the component(s) if not due to reasons of conscious or parental concerns must be included in the documentation.

**Note:** A sports physical is not a reason for an exception-to-periodicity checkup.

When billing for an exception-to-periodicity visit, provider must also include the most appropriate exception-to-periodicity modifiers. Claims for periodic THSteps medical checkups exceeding periodicity that do not include one for these modifiers will be denied as exceeding periodicity.

<table>
<thead>
<tr>
<th>Modifier</th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td>SC</td>
<td>Medically necessary service or supply</td>
</tr>
<tr>
<td>23</td>
<td>Unusual Anesthesia: Occasionally, a procedure that usually requires either no anesthesia or local anesthesia must be done under general anesthesia because of unusual circumstances. This circumstance may be reported by adding the modifier “23” to the procedure code of the basic service.</td>
</tr>
<tr>
<td>32</td>
<td>Mandated Services: Services related to mandated consultation or related services (e.g., PRO, third party payer, governmental, legislative, or regulatory requirement) may be identified by adding the modifier “32” to the basic procedure.</td>
</tr>
</tbody>
</table>

THSteps medical exception-to-periodicity services must be billed with the same procedure codes, provider type, modifier, and condition indicators as a medical checkup. Additionally, providers must use modifiers 23, 32, and SC to indicate the exception.

**5.3.8 Medical Checkup Follow-up Visit**

Use procedure code 99211 with the provider identifier and THSteps benefit code when billing for a follow-up visit.

**Note:** Reimbursement for the follow-up visit includes all elements of the visit. Reimbursement may not be allowed for the follow-up visit when submitted with certain procedure codes. For example: In accordance with CMS NCCI requirements, modifier 25 guidelines do not apply for procedure code 99211 when billed with other procedure codes that are included in the visit as related elements, including, but not limited to, administration of immunizations.

**Refer to:** Subsection 6.4.1, “National Correct Coding Initiative (NCCI) Guidelines” in “Section 6: Claims Filing” (Vol. 1, General Information) for additional information.

Medical Checkup Follow-up Visit with Immunization Administration on the TMHP website at [www.tmhp.com](http://www.tmhp.com) for a claim form example.

Medical Checkup Follow-up Visit with TB Skin Test on the TMHP website at [www.tmhp.com](http://www.tmhp.com) for a claim form example.

A follow-up visit may be required to complete necessary procedures related to a checkup or exception-to-periodicity checkup, such as:

- Reading the TST.
- Administering immunizations in cases where the client’s immunizations were not up-to-date, medically contraindicated, or unable to be given during the checkup.
- Collection of specimens for laboratory testing that were not obtained during the checkup or the original specimen could not be processed.
• Completion of sensory or developmental screening that was not completed at the time of the checkup due to the client’s condition.

A return visit to follow up on treatment initiated during a checkup or to make a referral is not a follow-up visit, but is considered an acute care visit under an appropriate E/M procedure code for an established client.

If the parent or guardian did not give consent for a component during the initial checkup, and supporting documentation is provided, no follow-up visit is necessary.

### 5.3.9 Newborn Examination

Providers do not have to be enrolled as THSteps providers to bill newborn examination procedure codes 99460, 99461, or 99463.

Newborn examinations that are billed with procedure code 99460, 99461, or 99463 may qualify as a THSteps medical checkup when all required components are completed according to the THSteps Periodicity Schedule and documented in the medical record.

Providers must use their provider identifier without benefit code EP1 when billing newborn examination services.

**Note:** In Texas, the mandated newborn hearing screening and newborn screening test is included as part of the in-hospital newborn exam.

A newborn hearing screening is included in the reimbursement to the hospital for the newborn hospital stay and is not reimbursed separately. The screening is covered as part of the newborn delivery. A newborn hearing screening must be offered to each newborn by the facility where the birth occurs, through a program mandated by the Texas Legislature and certified by the Department of State Health Services (DSHS). If a facility is not required by legislative mandate to perform newborn hearing screenings, a referral must be made to a facility that offers the screening.

If an infant is not born in a birthing facility and is not admitted to a birthing facility, the infant must be referred to a facility that provides newborn hearing screening.

State-mandated newborn screening for critical congenital heart disease (CCHD) is offered by and performed in the birth facility in accordance with Health and Safety Code (HSC) § 33.011 and 25 TAC §§37.75–37.79.

Providers billing these newborn codes are not required to be THSteps providers, but they must be enrolled as Medicaid providers. TMHP encourages THSteps enrollment for all providers that offer a medical home for clients and provide them with medical checkups and immunizations. Physicians and hospital staff are encouraged to inform parents eligible for Medicaid that the next THSteps checkup on the periodicity schedule should be scheduled from discharge to five days of age and that regular checkups should be scheduled during the first year and after.

**Refer to:** Subsection 9.2.44, “Newborn Services” in the Medical and Nursing Specialists, Physicians, and Physician Assistants Handbook (Vol. 2, Provider Handbooks) for additional information on inpatient newborn services.

The THSteps online education module “Newborn Hearing Screening” on the THSteps website at www.txhealthsteps.com for additional information about conducting a newborn hearing screen.

### 5.3.10 THSteps Medical Checkups Periodicity Schedule

The client is periodically eligible for medical checkup services based on the THSteps Medical Checkups Periodicity Schedule. All the checkups listed on the periodicity schedule have been developed based on recommendations of the AAP and recognized authorities in pediatric preventive health. In Texas, THSteps has modified the AAP periodicity schedule based on the scheduling of a laboratory or other test in federal EPSDT or state regulations.
The THSteps Medical Checkups Periodicity Schedule is available on the DSHS website at https://hhs.texas.gov/doing-business-hhs/provider-portals/health-services-providers/texas-health-steps/medical-providers.

5.3.11 Mandated Components

THSteps medical checkups must include regularly scheduled examinations and screenings of the general physical and mental health, growth, development, and nutritional status of infants, children, and youth.

The following federal and state mandated components must be documented in the client's medical record for the checkup to be considered complete:

- Comprehensive health and developmental history, including physical and mental health development
- Comprehensive unclothed physical examination
- Immunizations appropriate for age and health history
- Laboratory test appropriate to age and risk, including lead toxicity at specific federally-mandated ages
- Health education including anticipatory guidance
- Dental referral

The client's medical record must include documentation to support the rationale a component was not completed, and a plan to complete the component(s) if not due to parent or caregiver concern or reasons of conscience, including religious beliefs. THSteps provides optional clinical records to assist the provider in the documentation of the required components. These forms may be found at https://hhs.texas.gov/doing-business-hhs/provider-portals/health-services-providers/texas-health-steps/forms.

If the client has a condition that has been previously diagnosed and is currently receiving treatment, the associated standardized screening may be omitted with proper documentation.

Documented test or screening results obtained within the preceding 30 days for clients who are two years of age and younger, and the preceding 90 days for clients who are three years of age and older may be used to meet the testing or screening requirements. Results must include the dates of service and one of the following:

- A clear reference to the previous visit by the same provider
- Results obtained from another provider

5.3.11.1 Comprehensive Health and Developmental History

5.3.11.1.1 Nutritional Screening

Dietary practices must be evaluated at each checkup to identify and address nutritional issues or concerns.

5.3.11.2 Developmental Surveillance or Screening

Developmental surveillance or screening is a required component of every checkup for clients who are birth through 6 years of age. Autism screening is required at 18 months of age and again at 24 months of age. If not completed at 24 months of age, or if there is a particular concern it should be completed at 30 months of age.

As THSteps medical services, developmental screening (procedure code 96110) and autism screening (procedure code 96110 with modifier U6) are limited to once per day, per client, by the same provider or provider group. This service will be denied unless a checkup, exception-to-periodicity checkup, or follow-up visit was reimbursed for the same date of service by the same provider.
Standardized developmental screening is required at the ages listed in the “Required Screening Ages and Recommended Tools” table. Providers must use one of the validated, standardized tools listed in the table when performing a developmental or autism screening. A standardized screen is not required at other checkups up to and including the 6-year checkup; however, developmental surveillance is required at these checkups and includes a review of milestones (gross and fine motor skills, communication skills, speech-language development, self-help/care skills, and social, emotional, and cognitive development) and mental health and is not considered a separate service.

Providers may be reimbursed separately when using one of the required screening tools listed in the following table in addition to the checkup visit at specific age visits. THSteps requires one of the following required standardized tools at the following ages for a checkup to be considered complete:

<table>
<thead>
<tr>
<th>Screening Ages</th>
<th>Developmental Screening Tools</th>
<th>Autism Screening Tools</th>
</tr>
</thead>
<tbody>
<tr>
<td>9 months</td>
<td>Ages and Stages Questionnaire (ASQ) or Parents’ Evaluation of Development Status (PEDS)</td>
<td>N/A</td>
</tr>
<tr>
<td>18 months</td>
<td>ASQ or PEDS</td>
<td>Modified Checklist for Autism in Toddlers (M-CHAT) or M-CHAT Revised with Follow-Up (M-CHAT R/F)</td>
</tr>
<tr>
<td>24 months</td>
<td>ASQ or PEDS</td>
<td>M-CHAT or M-CHAT R/F</td>
</tr>
<tr>
<td>3 years</td>
<td>ASQ, Ages and Stages Questionnaire: Social-Emotional (ASQ:SE) or PEDS</td>
<td>N/A</td>
</tr>
<tr>
<td>4 years</td>
<td>ASQ, ASQ:SE or PEDS</td>
<td>N/A</td>
</tr>
</tbody>
</table>

If a developmental screening that is required in the Required Screening Ages and Recommended Tools table is not completed during a checkup or if the client is being seen for the first time, standardized developmental screening must be completed through 6 years of age.

If a provider administers a standardized and validated developmental screening at additional checkups other than those listed in the Required Screening Ages and Recommended Tools table, the provider must document the rationale for the additional screening, which may be due to provider or parental concerns.

Developmental screening that is completed without the use of one of the required standardized screening tools is not a separately payable benefit, and the checkup will be considered incomplete.

Standardized developmental screening as part of a medical checkup and for ages other than required on the periodicity schedule is not covered when completed for the sole purpose of meeting day care, Head Start, or school program requirements.

Standardized developmental screening may be performed outside a THSteps medical checkup as part of development and neurological assessment testing.


Referral for an in-depth developmental evaluation is determined by the criteria of the specific tool or at the provider’s discretion. Referral for in-depth evaluation of development should be provided when parents express concern about their child’s development, regardless of scoring on a standardized developmental screening tool. A medical diagnosis or a confirmed developmental delay is not required for referrals.
The ECI program serves clients who are birth through 35 months of age with disabilities or developmental delays. Under federal and state regulations, all health-care professionals are required to refer children to the Texas HHS ECI program as soon as possible, but no longer than 7 days after identifying a disability or a suspected delay in development, even if referred to an appropriate provider for further testing. If the client is 3 years of age or older, referral should be made to the local school district’s special education program.

5.3.11.1.3 Mental Health Screening

Mental health screening for behavioral, social, and emotional development is required at each THSteps checkup. Comparable to the American Academy of Pediatrics (AAP) Recommendations for Preventive Health Care guidelines, THSteps allows clients who are 12 through 18 years of age to receive a mental health screening using one of THSteps recognized mental health screening tools.

The following validated, standardized mental health screening tools are recognized by THSteps for mental health screening in adolescents who are 12 through 18 years of age:

- Pediatric Symptom Checklist (PSC-17)
- Pediatric Symptom Checklist (PSC-35)
- Pediatric Symptom Checklist for Youth (Y-PSC)
- Patient Health Questionnaire (PHQ-9)
- Patient Health Questionnaire (PHQ-9) Modified for Adolescents (PHQ-A [depression screen])
- Car, Relax, Alone, Forget, Family, and Trouble Checklist (CRAFFT)
- Patient Health Questionnaire (PHQ-A [anxiety, eating problems, mood problems and substance use])

A mental health screening must be submitted with procedure code 96160 for a screening tool completed by the adolescent, or procedure code 96161 for a screening tool completed by the parent or caregiver on behalf of the adolescent. When claims with procedure code 96160 or 96161 are submitted for mental health screenings, one of the validated, standardized mental health screening tools recognized by THSteps must be used.

Only one procedure code (96160 or 96161) may be reimbursed for the mental health screening per client per calendar year based on the description of the procedure code and the service rendered. Procedure codes 96160 and 96161 will not be reimbursed for the same client for any date of service.

Procedure code 96160 or 96161 must be submitted on the same date of service by the same provider as procedure code 99384, 99385, 99394, or 99395, and reimbursement is limited to once per calendar year, any provider.

The client’s medical record must include documentation identifying the tool that was used, the screening results, and any referrals that are made.

When the clinician conducting the mental health screen has the appropriate training and credentials to conduct the mental health evaluation and provide treatment, the clinician may choose to provide the mental health services or refer the client to an appropriate clinician. Clinicians who do not have these qualifications must refer clients to a qualified Medicaid-enrolled mental health specialist for such care.

For additional information about conducting a mental health screen, providers can refer to the THSteps online educational module “Mental Health Screening” at www.txhealthsteps.com.

5.3.11.1.4 Postpartum Depression Screening

Postpartum depression screening is a benefit of Texas Medicaid. Procedure codes G8431 and G8510 may be reimbursed when billing for postpartum depression screening in the office setting.
THSteps medical providers may receive separate reimbursement for postpartum depression screening, in addition to the infant’s Texas Health Steps medical checkup or follow-up visit. The reimbursement amount for procedure codes G8431 and G8510 covers all postpartum depression screenings provided during the infant’s medical checkup or follow-up visit.

Postpartum depression screening must be submitted under the infant’s Medicaid client number and will be restricted to clients who are 12 months of age and younger.

Procedure codes G8431 and G8510 must be submitted on the same claim, for the same date of service, by the same provider as one of the following THSteps medical checkup or follow-up visit procedure codes:

<table>
<thead>
<tr>
<th>Procedure Codes</th>
</tr>
</thead>
<tbody>
<tr>
<td>99211</td>
</tr>
</tbody>
</table>

Only one procedure code, either G8431 or G8510, may be reimbursed per provider in the 12 months following the infant’s birth.

**Postpartum Depression Screening and Referral Services**

The American Academy of Pediatrics (AAP) recommends the infant’s provider screen mothers for postpartum depression. Postpartum depression is the most common form of postpartum mood disturbance. Screening mothers for postpartum depression is appropriate for the general postpartum population and is recommended within the first few months following birth, up to the infant’s first birthday.

Postpartum depression meets the same clinical criteria as major depressive disorder, with the main difference being onset during pregnancy or after delivery.

While postpartum depression is the most common form of postpartum mood disturbance, providers should be aware that other mood disorders that may arise during the postpartum period include anxiety and panic disorders, obsessive-compulsive disorder, and postpartum psychosis.

Postpartum psychosis is a much more severe form of postpartum depression accompanied by psychotic features. Postpartum psychosis is rare, typically develops in the first few days to weeks after delivery, and is a psychiatric emergency requiring immediate medical attention.

In addition to postpartum psychosis, immediate or emergent medical attention may be necessary when the risk of imminent harm or danger is present.

**Screening Guidelines**

Screening using a validated tool is required. At a minimum, screening should occur at least once during the postpartum period. Validated tools may include the following:

- Edinburgh Postnatal Depression Scale
- Postpartum Depression Screening Scale
- Patient Health Questionnaire 9

Screening alone is inadequate for improving clinical outcomes. A positive screening for postpartum depression requires the THSteps provider to develop a referral plan with the mother.

**Positive Screenings: Referrals and Follow-Up**

THSteps providers must discuss the screening results with the mother, discuss the possibility of depression, and the impact depression may have on the mother, family, and health of the infant.
The THSteps provider and mother should discuss the mother’s options so the provider can refer her to an appropriate provider. Screening and referral is not contingent upon the mother’s Medicaid eligibility. When needed, referrals should be made regardless of the funding source, including referral to local mental health authorities and local behavioral health authorities.

THSteps providers should refer the mother to a provider who can perform further evaluation and determine an appropriate course of treatment. Appropriate providers include, but are not limited to:

- Mental health clinicians
- The mother’s primary care provider
- Obstetricians and gynecologists
- Family physicians
- Community resources such as Local Mental Health Authorities (LMHAs)

**Note:** Referral to an emergency center may be necessary when the risk for imminent harm or danger is present, such as mothers who report suicidal thoughts or thoughts of harming herself or the baby.

Resources should be provided for support in the interim until the mother is able to access care.

Scheduling a return visit for the infant sooner than the next scheduled visit may be appropriate in some cases.

**Prior Authorization Requirements**

Screening for postpartum mood disorders at the checkup or follow up visit does not require prior authorization.

**Note:** While recommended, screening for postpartum depression at the THSteps visit is not a compulsory requirement of the infant visit.

**Documentation Requirements**

Documentation in the infant’s record must include the name of the screening tool used and date the screening was completed.

If the mother screens positive for depression, at a minimum, the provider must note that a referral plan was discussed with the mother and a referral to a provider was made.

Providers may give the mother a copy of the completed screening tool to take with her to referral appointments.

Documentation should also include any health education or anticipatory guidance provided, along with the time period recommended for the infant’s next appointment.

**5.3.11.1.5 Tuberculosis (TB) Screening**

Administer the TB risk screening tool annually beginning at 12 months of age and thereafter at other medical checkups.

The TB risk screening tool is available on the HHSC website at [https://hhs.texas.gov/doing-business-hhs/provider-portals/health-services-providers/texas-health-steps/forms](https://hhs.texas.gov/doing-business-hhs/provider-portals/health-services-providers/texas-health-steps/forms).

A TST is to be administered when the screening tool indicates a risk for possible exposure. Providers must use procedure code 86580 when a TST is administered.

A TST may be reimbursed separately when performed as part of a THSteps medical checkup. TB screenings are part of the encounter rates for FQHCs and RHCs and are not reimbursed separately.

A follow-up visit (procedure code 99211) is required to read all TSTs. The provider may bill the follow-up visit with a provider identifier and THSteps benefit code.
If further evaluation is required to diagnose either latent TB infection or active TB disease, the provider may bill the appropriate E/M office visit code. Diagnosis and treatment are provided as a medical office visit. Providers can also call the TB program at 1-512-533-3000 for additional clinical information.


5.3.11.2 Comprehensive Unclothed Physical Examination

An age-appropriate unclothed physical examination is required at each checkup.

Recording of measurements and percentiles as appropriate to age to document growth and development including:

- Length or height and weight.
- The World Health Organization (WHO) growth charts, which are recommended for clients from birth to 2 years of age.
- The Centers for Disease Control and Prevention (CDC) growth charts, which are recommended for clients from 2 years of age and older.
- Fronto-occipital circumference (FOC) through the first 24 months of age.
- Body mass index (BMI) calculated beginning at 2 years of age.
- Blood pressure beginning at 3 years of age.

5.3.11.2.1 Oral Health Screening

Oral health screening is a part of the medical checkup physical examination.

5.3.11.2.2 Sensory Screening

Documentation of test results from a school vision or hearing screening program may replace the required audiometric or visual acuity screening if conducted within 12 months prior to the checkup.

Clients who are birth through 35 months of age with suspected or confirmed hearing or visual impairment must be referred to ECI as soon as possible, but no longer than 7 days after identification.

5.3.11.2.3 Hearing Screening

State-mandated newborn hearing screening is offered by and performed in the birth facility in accordance with Health and Safety Code (HSC), Chapter 47, §§ 47.001–47.009 and 25 TAC §§ 37.501–37.507.

A newborn hearing screening must be completed at the birthing facility. Automated auditory brainstem response (AABR) or transient evoke or distortion product otoacoustic emissions (OAE) may be performed.

Screening Results

Birthing facilities must report all newborn hearing screening results to DSHS within five business days using the web-based Texas Early Hearing Detection and Intervention (TEHDI) Management Information System (MIS) if written parental consent is obtained. Documented written consent must be maintained in the infant’s medical record.

Birthing facilities must provide written newborn hearing screening results to the parent or caregiver as well as the newborn’s primary care provider or medical home.
Primary care providers and medical homes (Texas Health Steps [THSteps] providers) must obtain a copy of the newborn hearing screening results within the TEHDI MIS if not provided by the birthing facility. The primary care provider or medical home must review all newborn hearing screening results with the parent or caregiver at the first checkup and determine if any additional follow-up is necessary.

**Note:** The primary care provider or medical home is responsible for managing and coordinating care for the child. Refer to the American Academy of Pediatrics Position Statement at [http://pediatrics.aappublications.org/content/129/5/996](http://pediatrics.aappublications.org/content/129/5/996).

Newborns who pass the newborn hearing screening must have their hearing monitored per the THSteps periodicity schedule. The primary care provider or medical home may opt to use the following tools to monitor developmental milestone benchmarks in newborns that pass their newborn hearing screening:

- TEHDI: A Roadmap for Families (English)
- Hearing Checklist for Parents (English)
- Hearing Checklist for Parents (Spanish)

Newborns who do not pass the initial screen must be rescreened a second time in the birthing facility before discharge.

**Outpatient Rescreening**

Newborns who do not pass the second screen in the birthing facility must be referred to a Medicaid-enrolled provider for an outpatient follow-up rescreen. The rescreen provider must have access to AABR or OAE screening and must be experienced with the pediatric population under age three.

Newborns who do not receive a referral from the birthing facility, after not passing the second screen in the birthing facility, must be referred by their primary care provider or medical home to a Medicaid-enrolled provider for an outpatient follow-up rescreen, unless their primary care provider or medical home is adequately equipped to provide the service. The optimal time frame for the outpatient follow-up rescreen is when the infant is between 10 and 30 days old.

Outpatient follow-up rescreens must be completed by AABR or OAE screening. Results must be reported as soon as possible to the DSHS TEHDI MIS, as well as the infant’s primary care provider or medical home.

Newborns who pass the outpatient rescreen must have their hearing monitored by their primary care provider or medical home per the THSteps periodicity schedule.

**Diagnostic Audiological Evaluation**

Newborns who do not pass the outpatient rescreen must be referred to a Medicaid-enrolled audiologist for a diagnostic audiological evaluation using the Texas Pediatric Protocol for Evaluation. Referrals should be made upon consultation with the primary care provider or medical home.

**Note:** Additional information about the Texas Pediatric Protocol for Evaluation is available at [https://www.dshs.texas.gov/tehdi/educational-materials.aspx](https://www.dshs.texas.gov/tehdi/educational-materials.aspx).

Unless the newborn or infant has been hospitalized since birth, the diagnostic audiological evaluation must be completed no later than the third month after birth, or upon referral by the primary care provider or medical home.

Diagnostic audiological evaluations completed by audiologists using the Texas Pediatric Protocol for Evaluation must include a diagnostic auditory brainstem response (ABR) and, if not previously done, a diagnostic OAE to determine cochlear involvement.

Audiologists will use equipment norms for newborns, preferably ones they have collected on their equipment.
Protocols include air and bone conduction testing using tone burst ABR, as well as click ABR, so the amplification may be appropriate to fit the individual.

**Note:** Additional information about technologies that have been evaluated by an independent investigator and DSHS, and have been found to meet the requirements for conducting newborn hearing screening is available at [www.dshs.texas.gov/tehdi/Audiologic-Evaluation-Protocol.aspx](http://www.dshs.texas.gov/tehdi/Audiologic-Evaluation-Protocol.aspx).

**Evaluation Results**

Audiologists must report all diagnostic results to DSHS TEHDI MIS and provide written hearing screening results to the primary care provider or medical home.

The newborn or infant will be fitted for hearing aids by the audiologist when appropriate and should receive continued audiological assessments and monitoring as needed.

**ECI Referrals**

Newborns or infants not passing the outpatient rescreen must also be referred by the primary care provider or medical home to ECI for provision of services. The referral should be made within the TEHDI MIS.

Newborns or infants, as required by federal law under the Individuals with Disabilities in Education Act (IDEA), may be referred to ECI twice under the following circumstances:

- Upon suspicion that the child is deaf or hard of hearing, for service coordination and possible confirmation of eligibility for ECI services
- Upon confirmation that the child is deaf or hard of hearing, for a referral to other Local Education Agency for auditory impairment services

**Late Onset Hearing Loss**

When one or more risk factors for late onset hearing loss has been identified and the newborn or infant passed their hearing screen, the outcome will not be “normal hearing” but will be “in process.” Noting the infant as “in process” allows all health-care providers in the care of the infant to be aware of the presence of risk factors to determine the frequency of risk monitoring to identify audiological issues as soon as possible. This determination depends on the type and number of the following risks identified:

- Craniofacial anomalies
- Exchange transfusion for elevated bilirubin
- Family history of deafness
- NICU > 5 days
- Apgar 0-4 at 1 minute
- Apgar 0-6 at 5 minutes
- Bacterial meningitis
- Birth weight < 1500g
- Congenital infection
- Head injury
- Neurodegenerative disorder
- Other postnatal infection
- Otitis media > 3 months (middle ear infection)
- Ototoxic medications administered
• Parental concern regarding hearing status
• Persistent pulmonary hypertension of the newborn associated with mechanical ventilation
• Syndrome

**Note:** Information about risk factors for late onset hearing loss and a risk monitoring periodicity schedule is available in Chapter 3, “Tracking, Reporting, & Follow-Up,” in The NCHAM E-Book on the National Center for Hearing Assessment and Management (NCHAM) website at [http://infanthearing.org](http://infanthearing.org).

Hearing screening must be performed at each checkup for clients who are birth through 20 years of age. Audiometric screening must be performed at specific ages indicated on the periodicity schedule. Subjective screening through provider observation or informant report is required at the other checkups.

Clients at high risk or with abnormal screening results must be referred to an appropriate Medicaid-enrolled provider who specializes in pediatric audiology services. Clients who are birth through 20 years of age enrolled with Texas Medicaid for the date(s) of service are eligible for Texas Medicaid hearing services benefits.

5.3.11.2.4 Vision Screening

Vision screening must be performed at each checkup. A visual acuity test must be performed at ages indicated on the periodicity schedule. Subjective screening through provider observation or informant report is done at the other checkups.

All clients must be screened for eye abnormalities by history, observation, and physical exam and referred to a Medicaid-enrolled optometrist or ophthalmologist experienced with the pediatric population if at high risk.

Clients with abnormal visual acuity screening results must be referred to a Medicaid-enrolled optometrist or ophthalmologist experienced with the pediatric population.

5.3.11.3 *Immunizations*

Providers must assess the immunization status at every medical checkup to ensure all age requirements have been met. The necessary vaccines and toxoids must be administered at the time of the checkup unless medically contraindicated or because of parent’s or caregiver’s reasons of conscience including religious beliefs. If an indicated vaccine or toxoid was not administered, the reason must be documented in the client’s medical record.

Vaccines and toxoids must be administered according to the current ACIP “Recommended Childhood and Adolescent Immunization Schedule - United States.” Providers must not refer clients to the local health department or other entity for immunization administration.

THSteps providers are strongly encouraged to obtain vaccines from TVFC for clients who are birth through 18 years of age. Vaccines that are identified as being distributed through TVFC are not reimbursed separately.

Vaccines and toxoids may be reimbursed through Texas Medicaid at a fee determined by HHSC when the vaccine is medically necessary for THSteps clients who are 19 through 20 years of age.

The specific diagnosis necessitating the vaccine and toxoid is required when billing with the following administration procedure codes in combination with an appropriate vaccine/toxoid procedure code:

<table>
<thead>
<tr>
<th>Procedure Codes</th>
</tr>
</thead>
<tbody>
<tr>
<td>90460</td>
</tr>
</tbody>
</table>
The age-appropriate diagnosis code for a preventive care medical checkup must be submitted on the
claim. If an immunization is administered as part of a preventive care medical checkup, diagnosis code
Z23 may also be included on the claim, in addition to the age-appropriate diagnosis.

Providers should submit only diagnosis code Z23 on the claim if an immunization is the only service
provided during an office visit.

Vaccine and toxoid administration must be billed with the following age appropriate diagnosis codes:

<table>
<thead>
<tr>
<th>Client Age</th>
<th>Diagnosis Code</th>
</tr>
</thead>
<tbody>
<tr>
<td>Birth through 7 days</td>
<td>Z00110</td>
</tr>
<tr>
<td>Eight through 28 days</td>
<td>Z00111</td>
</tr>
<tr>
<td>29 days through 17 years</td>
<td>Z00121, Z00129</td>
</tr>
<tr>
<td>18 years or older</td>
<td>Z0000, Z0001</td>
</tr>
</tbody>
</table>

Procedure codes 90460 and 90461 are benefits for services rendered to clients who are birth through
18 years of age when counseling is provided for the immunization administered.

Procedure codes 90471 and 90472 are benefits for services rendered to clients of any age when
counseling is not provided for the immunization administered.

Procedure codes 90473 and 90474 are benefits for services rendered to clients who are birth through 20
years of age when counseling is not provided for the immunization administered.

The following vaccines and toxoids are a benefit of Texas Medicaid:

<table>
<thead>
<tr>
<th>Procedure Code</th>
<th>Number of Components**</th>
</tr>
</thead>
<tbody>
<tr>
<td>90620*</td>
<td>1</td>
</tr>
<tr>
<td>90632</td>
<td>1</td>
</tr>
<tr>
<td>90644</td>
<td>2</td>
</tr>
<tr>
<td>90655*</td>
<td>1</td>
</tr>
<tr>
<td>90658*</td>
<td>1</td>
</tr>
<tr>
<td>90670*</td>
<td>1</td>
</tr>
<tr>
<td>90674</td>
<td>1</td>
</tr>
<tr>
<td>90682</td>
<td>1</td>
</tr>
<tr>
<td>90687*</td>
<td>1</td>
</tr>
<tr>
<td>90698*</td>
<td>5</td>
</tr>
<tr>
<td>90707*</td>
<td>3</td>
</tr>
<tr>
<td>90714*</td>
<td>2</td>
</tr>
<tr>
<td>90723*</td>
<td>5</td>
</tr>
<tr>
<td>90734*</td>
<td>1</td>
</tr>
<tr>
<td>90746</td>
<td>1</td>
</tr>
<tr>
<td>90756</td>
<td>1</td>
</tr>
</tbody>
</table>

* TV/FC-distributed vaccine/toxoid
** The number of components applies if counseling is provided and procedure code 90460 and 90461 are submitted.

Procedure codes 90655, 90657, 90685, and 90687 are limited to clients who are 6 through 35 months of
age.
Procedure codes 90656 and 90658 are limited to clients who are 3 years of age and older.

Procedure codes 90686 and 90688 are limited to clients who are 6 months of age and older.

Procedure code 90682 is limited to clients who are 18 years of age and older.

Procedure code 90756 is limited to clients who are 4 years of age and older.

Providers may use the state-defined modifier U1 in addition to the associated administered vaccine procedure code for clients who are birth through 18 years of age and the vaccine was unavailable through TVFC.

<table>
<thead>
<tr>
<th>Modifier</th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td>U1</td>
<td>State-defined modifier: Vaccine(s)/toxoid(s) privately purchased by provider when TVFC vaccine/toxoid is unavailable</td>
</tr>
</tbody>
</table>

Note: “Unavailable” is defined as a new vaccine approved by ACIP that has not been negotiated or added to a TVFC contract, funding for new vaccine that has not been established by TVFC, or national supply or distribution issues. Providers will be informed if a vaccine meets the definition of 'not available' from TVFC and when the provider’s privately purchased vaccine may be billed with modifier U1.

Modifier U1 may not be used for failure to enroll in TVFC, maintain sufficient TVFC vaccine/toxoid inventory, or clients who are 19 through 20 years of age.

Each vaccine or toxoid and its administration must be submitted on the claim in the following sequence: the vaccine procedure code immediately followed by the applicable immunization administration procedure code(s). All of the immunization administration procedure codes that correspond to a single vaccine or toxoid procedure code must be submitted on the same claim as the vaccine or toxoid procedure code.

Each vaccine or toxoid procedure code must be submitted with the appropriate “administration with counseling” procedure code(s) (procedure codes 90460 and 90461) or the most appropriate “administration without counseling” procedure code (procedure code 90471, 90472, 90473, or 90474). If an “administration with counseling” procedure code is submitted with an “administration without counseling” procedure code for the same vaccine or toxoid, the second administration of the vaccine or toxoid will be denied.

Administration With Counseling

Providers must submit claims for immunization administration procedure codes 90460 or 90461 based on the number of components per vaccine. Providers must specify the number of components per vaccine by billing 90460 and 90461 as defined by the procedure code descriptions:

- Procedure code 90460 is submitted for the administration of the 1st component.
- Procedure code 90461 is submitted for the administration of each additional component identified in the vaccine.

Procedure code 90461 will be denied if procedure code 90460 has not been submitted on the same claim for the same vaccine or toxoid.

The necessary counseling that is conducted by a physician or other qualified health-care professional must be documented in the client’s medical record.
The following is an example of how to submit claims for immunization administration procedure codes when counseling is provided:

<table>
<thead>
<tr>
<th>Procedure Code</th>
<th>Quantity Billed</th>
</tr>
</thead>
<tbody>
<tr>
<td>Vaccine or toxoid procedure code with 1 component</td>
<td>1</td>
</tr>
<tr>
<td>90460 (1st component)</td>
<td>1</td>
</tr>
<tr>
<td>Vaccine or toxoid procedure code with 3 components</td>
<td>1</td>
</tr>
<tr>
<td>90460 (1st component)</td>
<td>1</td>
</tr>
<tr>
<td>90461 (2nd and 3rd components)</td>
<td>2</td>
</tr>
</tbody>
</table>

**Note:** The term “components” refers to the number of antigens that prevent disease(s) caused by one organism. Combination vaccines are those that contain multiple vaccine components.

**Administration Without Counseling**

Procedure codes 90471, 90472, 90473, and 90474 may be reimbursed per vaccine based on the route of administration.

The following is an example of how to submit claims for injection administration procedure codes when counseling is not provided:

<table>
<thead>
<tr>
<th>Procedure Code</th>
<th>Quantity Billed</th>
</tr>
</thead>
<tbody>
<tr>
<td>Vaccine or toxoid procedure code</td>
<td>1</td>
</tr>
<tr>
<td>90471 (Injection administration)</td>
<td>1</td>
</tr>
<tr>
<td>Vaccine or toxoid procedure code</td>
<td>1</td>
</tr>
<tr>
<td>90472 (Injection administration)</td>
<td>1</td>
</tr>
<tr>
<td>Vaccine or toxoid procedure code</td>
<td>1</td>
</tr>
<tr>
<td>90472 (Injection administration)</td>
<td>1</td>
</tr>
</tbody>
</table>

**Vaccine Administration and Preventive E/M Visits**

For claims that are submitted with an immunization administration procedure code and a preventive E/M visit, providers may append modifier 25 to the preventive E/M visit procedure code to identify a significant, separately identifiable E/M service that was rendered by the same provider on the same day as the immunization administration. Documentation that supports the provision of a significant, separately identifiable E/M service must be maintained in the client’s medical record and made available to Texas Medicaid upon request.

**Refer to:** [Acute Care Visit on the Same Day as a THSteps Preventive Visit Checkup](http://www.tmhp.com) on the TMHP website for a claim form example.

[THSteps Preventive Visit Checkup with Immunization and Vaccine Administration](http://www.tmhp.com) on the TMHP website for a claim form example.

**5.3.11.3.1 Vaccine Information Statement (VIS)**

A VIS is required by federal mandate to inform parents and vaccine recipients of the risks and benefits of the vaccine they are about to receive. Not only is it important to explain the risks and benefits before a vaccine is administered, it is also important that providers use the most current forms available. For more about immunizations, vaccine-preventable diseases, or literature and forms, providers can call the DSHS Immunization Branch at 1-800-252-9152 or review information at [www.dshs.texas.gov/immunize](http://www.dshs.texas.gov/immunize).
5.3.11.4 **Health Education and Anticipatory Guidance**

Anticipatory guidance is a federally mandated component of the THSteps medical checkup and includes health education and counseling. Health education and counseling with parents or guardians and clients are required to assist parents in understanding what to expect in terms of the client’s development and to provide information about the benefits of healthy lifestyles and practices, as well as accident and disease prevention. Written material may also be given but does not replace counseling. The optional THSteps clinical records include age-appropriate topics on the back of each form. These forms can be found at [https://hhs.texas.gov/doing-business-hhs/provider-portals/health-services-providers/texas-health-steps/forms](https://hhs.texas.gov/doing-business-hhs/provider-portals/health-services-providers/texas-health-steps/forms).

5.3.11.5 **Dental Referral**

Based on the AAPD definition of a dental home, Texas Medicaid defines a dental home as the dental provider who supports an ongoing relationship with the client that is inclusive of all aspects of oral health care delivered in a comprehensive, continuously accessible, coordinated, and family-centered way. In Texas, establishment of a client’s dental home should begin at 6 months of age but no later than 12 months of age and includes referral to dental specialists when appropriate.

The physician must refer clients to establish a dental home beginning at 6 months of age or earlier if trauma or early childhood caries are identified. For established clients after the six-month medical checkup, the provider must confirm if a dental home has been established and is on-going; if not, additional referrals must be made at subsequent medical checkup visits until the parent or caregiver confirms that a dental home has been established for the client. The parent or caregiver of the client may self-refer for dental care at any age, including 12 months of age or younger.

5.3.11.6 **Laboratory Test**

Age-appropriate and risk-based laboratory testing as noted on the periodicity schedule is considered part of the medical checkup. The DSHS Laboratory provides supplies for specimen collection and mailing and shipping; and reporting of test results to enrolled THSteps medical providers that submit specimens to the DSHS Laboratory. These services and supplies are limited to THSteps medical checkup laboratory services provided in the course of a medical checkup to THSteps clients. Unauthorized use of services and supplies is a violation of federal regulations.

DSHS Laboratory services are available at no cost to all enrolled THSteps medical providers for THSteps medical checkups only.

*Example:* If a provider needs immediate results for the anemia screening, the specimen may be processed in the office/clinic, but will not be separately reimbursed. The test results must be documented in the client’s medical record.

*Exception:* For tests related to screening for type 2 diabetes, dyslipidemia, HIV, and syphilis, the client or specimen may be sent to the laboratory of the provider’s choice. Point-of-care testing that is performed in the provider’s office to obtain the initial blood lead specimen may be reimbursed separately.

The date of service for the laboratory testing is to be the date the specimen was obtained as part of the medical checkup, follow-up, or exception-to-periodicity checkup.
The procedure codes for any laboratory testing services other than screening for type 2 diabetes, dyslipidemia, HIV, and syphilis are informational when obtained on the same day a checkup is completed, even if an acute care visit is performed on the same date of service.

If the laboratory testing as identified on the THSteps Medical Checkup Periodicity Schedule is obtained as part of an E/M visit on a different date of service than a checkup, the services may be considered as separate services and may be sent to the laboratory of the provider’s choice.

Laboratory specimens obtained for diagnostic evaluation, rather than for screening purposes and performed on the same day as a checkup, may be considered as separate services unless the test is required as part of a checkup. If the test is required as part of the checkup, the laboratory specimens, with the exception of screening tests for dyslipidemia, type 2 diabetes, HIV, and syphilis must be submitted to the DSHS Laboratory for testing. Diagnostic specimens that are not part of the checkup can be sent to the laboratory of the provider’s choice.

Laboratory services that are related to a THSteps medical checkup are available from the DSHS Laboratory and may not be billed separately with an office visit or consultation on the same day as a THSteps medical checkup.

All of the laboratory tests that are listed on the THSteps Periodicity Schedule may be submitted to the DSHS Laboratory if the specimen submission requirements can be met. Tests that are listed in the “Laboratory Test Procedure Codes” table must be submitted to the DSHS Laboratory. Tests that must be sent to the DSHS laboratory but that are processed elsewhere are not reimbursed; however, the documentation results may be used to meet the requirements for a checkup.

The following procedure codes may not be billed separately with an office visit or consultation on the same day as a THSteps medical checkup either by a provider or laboratory. Claims for the following procedure codes submitted by a provider or a commercial laboratory for the same DOS as a THSteps medical checkup are denied and are subject to retrospective review:

<table>
<thead>
<tr>
<th>Laboratory Test Procedure Codes</th>
</tr>
</thead>
<tbody>
<tr>
<td>83655*</td>
</tr>
</tbody>
</table>

* Unless performed using point-of-care testing, the initial lead specimen must be sent to the DSHS Laboratory

For specimens sent to the DSHS Laboratory, the complete medical checkup includes the specimen collection and supplies, mailing and shipping supplies, and the review of the test results from the DSHS Laboratory.

For specimens sent to a laboratory of the provider’s choice, the checkup includes the specimen collection or ordering of the test and the review of the test results from the laboratory.

### 5.3.11.6.1 Laboratory Supplies

The DSHS Laboratory verifies enrollment of THSteps medical providers before sending laboratory supplies and the informational packet to the medical providers. Newly enrolled providers should contact the DSHS Laboratory to request laboratory supplies. Upon request, the DSHS Laboratory provides THSteps medical providers with laboratory supplies associated with specimen collection, submission, and mailing and shipping of required laboratory tests related to medical checkups. Requests for specimen submission forms are routed to the DSHS Laboratory reporting staff and mailed separately to the providers. The Child Health Laboratory Supplies Order Form lists the laboratory supplies that the DSHS Laboratory provides to THSteps medical providers.

To obtain a THSteps Child Health Laboratory Supplies Order Form, providers can call 1-512-776-7661 or 1-888-963-7111, ext. 7661, or download the form online at [www.dshs.texas.gov/lab/MRS_forms.shtm](http://www.dshs.texas.gov/lab/MRS_forms.shtm).
5.3.11.6.2 Newborn Screening Supplies

Providers that perform newborn screening (NBS) can order supplies by submitting a Newborn Screening Supplies Order Form to the DSHS Laboratory. The Newborn Screening Supplies Order Form lists the NBS supplies that the DSHS Laboratory provides to medical providers.

**Note:** For newborn screening, only the specimen collection form (NBS 3), mailing envelope and provider address labels are provided. Lancets, mailing, and shipping costs are the responsibility of the submitter.

To obtain a Newborn Screening Supplies Order Form, medical providers can call 1-512-776-7661 or 1-888-963-7111, ext. 7661, or download the form online at www.dshs.texas.gov/lab/MRS_forms.shtm.

Contact information for requesting laboratory supplies:

Container Preparation  
Laboratory Services Section, MC 1947  
Department of State Health Services  
PO Box 149347  
Austin, TX 78714-9347  
1-512-776-7661 or 1-888-963-7111, Ext. 7661  
Fax: 1-512-776-7672

5.3.11.6.3 Laboratory Submission

All required laboratory testing for THSteps clients must be performed by the Department of State Health Services (DSHS) Laboratory in Austin, TX, with the following exceptions:

- Specimens collected for type 2 diabetes, dyslipidemia, HIV, and syphilis screening may be sent to the laboratory of a provider’s choice or to the DSHS Laboratory in Austin if submission requirements can be met.

- Initial blood lead testing using point-of-care testing.

THSteps medical checkup laboratory specimens submitted to the DSHS Laboratory must be accompanied with the DSHS Laboratory Specimen Submission Form (Newborn Screening NBS 3 or G-THSTEPS as appropriate) for test(s) requested. All forms must include the client’s name and Medicaid number as they appear on the Your Texas Benefits Medicaid card. If a number is not currently available but is pending (i.e., a newborn or a newly certified client verified by a Medicaid Eligibility Verification [Form H1027] as eligible for Medicaid), providers must write “pending” in the Medicaid number space, which is located in the payor source section of the laboratory specimen submission form.

Laboratory specimens received at the DSHS Laboratory without a Medicaid number or the word “pending” written on the accompanying specimen submission form will be analyzed, and the provider will be billed.

Specimens submitted to the laboratory must also meet specific acceptance criteria. For additional information on specimen submission, providers can refer to the DSHS Laboratory web page at: www.dshs.texas.gov/lab/MRS_specimens.shtm.

**Note:** If an extreme health problem exists and telephone results are needed quickly, providers should make a request on the laboratory form. With the exception of weekends and holidays, routine specimens are analyzed and reported within three business days after receipt by the DSHS Laboratory. Critical abnormal test results (e.g., hemoglobin equal to or below 7g/dL or blood lead levels greater than or equal to 40 mcg/dL) are identified in the laboratory within 36 hours after receipt of specimens and are reported to the submitter by telephone within one hour of confirmation.
The THSteps laboratory specimens that can be mailed at ambient temperature can be sent to the DSHS Laboratory Services Section through the U.S. Postal Service at no cost using the provided business reply labels:

DSHS Laboratory Services Section
Walter Douglass
PO Box 149163
Austin, TX 78714-9803
1-512-776-7318 or 1-888-963-7111 Ext. 7318

THSteps laboratory specimens that require overnight shipping on cold packs through a courier service must be sent to the DSHS Laboratory Services Section at:

DSHS Laboratory Services Section, MC-1947
1100 West 49th Street
Austin, TX 78756-3199

Newborn Screening specimens can be sent through the U.S. Postal Service to:

Texas Department of State Health Services
Laboratory Services Section
PO Box 149341
Austin, TX 78714-9341

Gonorrhea and Chlamydia specimens for regular delivery are sent to:

Department of State Health Services
Laboratory - MC 1947
Walter Douglass, 1-512-776-7569
PO Box 149163
Austin, TX 78714-9803

Gonorrhea and Chlamydia specimens that are shipped cold overnight by courier are sent to:

Department of State Health Services
Laboratory - MC 1947
Walter Douglass, 1-512-776-7569
1100 W. 49th Street
Austin, TX 78756-3199

Collectors are available from the DSHS Austin Laboratory. To order collectors, providers must complete the Order Form for Gonorrhea/Chlamydia (GC/CT) Laboratory Supplies (G-6C) that is posted on the DSHS website at www.dshs.texas.gov/lab/MRS_forms.shtml and fax the completed form to 1-512-776-7672.

Providers can call 1-512-776-6030 or toll-free 1-888-963-7111, ext. 6030, for questions about submission requirements such as collection, supplies, and mailing of specimens for THSteps gonorrhea and chlamydia adolescent screening.
5.3.11.6.4  Send Comments

Providers with comments or feedback about THSteps specimen collection supplies should contact the DSHS Laboratory. Supplies are evaluated continually, and feedback from supply users is useful. Documented comments may support, justify, or initiate a change in a provided item. Providers can send a brief letter or fax to the following address:

Quality Assurance Unit
Laboratory Services Section, MC 1947
Department of State Health Services
PO Box 149347
Austin, TX 78714-9347
Fax: 1-512-776-7294

5.3.11.6.5  Laboratory Reporting

A computer-generated result report is mailed or faxed to the submitting THSteps medical checkup provider. A monthly statistical report card is only available online for providers documenting their total number of Total Hemoglobin and Blood Lead submissions by diagnosis and adequacy. The DSHS Laboratory has web-based services (remote order or result reporting) available for THSteps and Newborn Screening laboratory services. For more information, providers can visit the DSHS website at www.dshs.texas.gov/lab/remoteData.shtm or call 1-888-963-7111, Ext. 6030.

5.3.11.6.6  Required Laboratory Tests Related to Medical Checkups

The following laboratory screening procedures are required components of the THSteps medical checkup and are to be performed in accordance with the age and frequency specified on the THSteps medical checkup periodicity schedule. Due to changes in specimen collection, handling, and submission criteria, providers should contact the DSHS Laboratory for the most current specimen requirements by calling 1-888-963-7111, Ext. 6236, 6237, or 2628, email ClinicalChemistry@dshs.texas.gov, or visiting the DSHS website at www.dshs.texas.gov/lab/mrs_labs_toc.shtm.

Anemia Screening

Anemia screening by hemoglobin or hematocrit levels is required at ages as noted on the THSteps Periodicity Schedule and the specimen must be sent to the DSHS Laboratory. If there is an urgent need for test results, these tests may be completed in a provider’s office or clinic, but they will not be reimbursed separately. These test results must be documented in the client’s medical record.

Lead Screening and Testing

In accordance with current federal regulations, THSteps requires blood lead screening at ages notated on the THSteps Periodicity Schedule and must be performed during the medical checkup. Environmental lead risk assessments, as part of anticipatory guidance, should be completed at all checkups through age 6 when testing is not mandated, and may be performed using the Lead Risk Questionnaire, Form Pb-110, which is provided in both English and Spanish at https://hhs.texas.gov/doing-business-hhs/provider-portals/health-services-providers/texas-health-steps/forms. Providers may also opt to use an equivalent form of their choice.

The initial lead testing may be performed using a venous or capillary specimen, and must either be sent to the DSHS Laboratory or performed in the provider’s office using point-of-care testing. If the client has an elevated blood lead level of 5 mcg/dL or greater, the provider must perform a confirmatory test using a venous specimen. The confirmatory specimen may be sent to the DSHS Laboratory, or the client or specimen may be sent to a laboratory of the provider’s choice.
All blood lead levels in clients who are 14 years of age or younger must be reported to DSHS Texas Childhood Lead Poisoning Prevention Program (TXCLPPP). Reports should include all information as required on the Child Blood Lead Reporting, Form F09-11709 or the Point-of-Care Blood Lead Testing report Form Pb-111, which can be found at www.dshs.texas.gov/lead/providers.shtm or by calling 1-800-588-1248.

Information related to blood lead screening and reporting for clients who are 15 years of age or older is available on the DSHS Blood Lead Surveillance Group’s website at www.dshs.texas.gov/lead/providers.shtm.

Initial blood lead testing using point-of-care testing (procedure code 83655 with modifier QW) may be reimbursed to THSteps medical providers when performed in the provider’s office. Providers must have a Clinical Laboratory Improvement Amendments of 1988 (CLIA) certificate of waiver. For waived tests, providers must use modifier QW as indicated on the CMS website.

Blood lead testing is part of the encounter rates for FQHCs and RHCs and is not reimbursed separately.

Providers may obtain more information about the medical and environmental management of lead-poisoned children from the DSHS Childhood Lead Poisoning Prevention Program by calling 1-800-588-1248 or visiting the web page at www.dshs.texas.gov/lead.

Refer to: “Appendix C. Lead Screening” in this handbook for more information on lead screening procedures and follow-up.

**Dyslipidemia**

Screening for dyslipidemia is required once for clients who are 9 through 11 years of age and once again for clients who are 18 through 20 years of age, regardless of risk. These are in addition to the current risk-based screening for clients who are 24 months through 20 years of age. Clients or specimens may be sent to the laboratory of the provider’s choice, including the DSHS Laboratory. THSteps does not provide a formal risk assessment tool. Providers may refer to the AAP policy statement on cholesterol screening for more information.

**Diabetes**

Screening for type 2 diabetes is based on risk assessment. THSteps does not provide a formal risk assessment tool. Clients and specimens may be sent to the laboratory of the provider’s choice, including the DSHS Laboratory.

**Newborn Screening**

Each newborn delivered in Texas must be subjected to two screens to test for a number of genetic and heritable disorders. Each newborn screen is indicated on the THSteps Periodicity Schedule. A current list of screened disorders is available at www.dshs.texas.gov/newborn/screened_disorders.shtm.

Additional information about newborn screening, is available on the Newborn Screening Program website at www.dshs.texas.gov/newborn/default.shtm.

The initial newborn screen specimen must be obtained between 24 and 48 hours after birth. Newborns discharged from a hospital or birthing facility before this time criteria is met must have a newborn screen blood specimen obtained immediately prior to discharge. When the newborn is an inpatient in the hospital, the hospital shall ensure that the appropriate screens are done. When the newborn is not in the hospital, the physician or health-care practitioner who attends the newborn outside of the hospital shall be responsible for causing the appropriate screens to be done. TAC Title 25, Part 1, Chapter 37, Subchapter D, Rule §37.55.

A second screen is to be obtained between one and two weeks of age by the newborn’s physician or health-care practitioner, and is a required component of the THSteps medical checkup. Clients may not be referred to the local health department or other providers for this service. If there is any doubt that a
client younger than 12 months of age was properly tested, the provider should submit a screen on DSHS Form NBS 3 to the Texas Department of State Health Services, Laboratory Services Section, Austin, Texas.

Newborn screening tests may be performed in special circumstances, such as adoption, if there is not record of previous test results. Newborn screen results are mailed or faxed to the address that the provider indicated on DSHS Form NBS 3. Providers may sign up to receive results online through the DSHS Laboratory web-based services. For more information visit the DSHS website at www.dshs.texas.gov/lab/newbornscreening.shtm or call 1-888-963-7111, Ext. 6030.

Note: Recommendations for necessary follow-up procedures are included with the newborn screen results. Newborn Screening (NBS) Clinical Care Coordination staff will contact providers when there are significant out of range newborn screening laboratory results.

5.3.11.6.7 Additional Required Laboratory Tests Related to Medical Checkups for Adolescents

The following is a list of required and risk-based laboratory tests related to medical checkups for adolescents and guidelines for testing for sexually transmitted diseases (STDs).

Testing for Sexually Transmitted Diseases

Syphilis Testing

Syphilis testing should be performed on adolescents that are at high risk for infection. Clients and specimens may be sent to the laboratory of the provider’s choice, including the DSHS Laboratory.

Gonorrhea and Chlamydia Infection Testing

Testing for gonorrhea and Chlamydia should be performed on adolescents that are at high risk for infection. Specimens must be sent to the DSHS Laboratory in Austin.

HIV Testing

Clients should be informed that the HIV test is routinely available, confidential, and completely anonymous. It is critical to maintain confidentiality when caring for clients, as well as their specimens. Testing should be performed only after informed consent is obtained from the adolescent. Informed consent does not have to be written as long as there is documentation in the medical record that the test has been explained and consent has been obtained.

Screening for HIV is required once for clients who are 16 through 18 years of age, regardless of risk. Clients or specimens may be sent to the laboratory of the providers’ choice including the DSHS Laboratory.

Screening for HIV is also based on risk assessment for clients who are 11 through 20 years of age based on risk assessment. Clients or specimens may be sent to the laboratory of the providers’ choice, including the DSHS Laboratory.

THSteps does not provide a formal HIV risk assessment tool. Providers may refer to the AAP policy statement on HIV screening and CDC guidelines on HIV screening for more information.

HIV testing may be performed for adolescents without requirement of parental consent. Adolescents at risk for HIV infection should be offered confidential HIV screening. If the client refuses the HIV test, the provider may not perform the test and must explain the option of anonymous testing and refer the client to a testing facility that offers anonymous testing. A notation must be made in the medical record that notification of the HIV test and the right to refuse was given. Providers may call the HIV/STD InfoLine for referrals to HIV/AIDS testing sites; prevention, case management, and treatment providers; STD clinics; and other related service organizations. The HIV/STD InfoLine is 1-800-299-2437. This toll-free HIV/AIDS and STD information and referral service is available for English- and Spanish-speaking callers and for those who are hearing-impaired.
Communicable Disease Reporting

Diagnoses of STDs, including HIV, are reportable conditions under 25 TAC, Chapter 97. Providers must report confirmed diagnoses of STDs as required by 25 TAC §97.132.

5.3.11.6.8 Zika Virus Testing

Refer to: Subsection 2.2.13.1, “Zika Virus Testing” in the Radiology and Laboratory Services Handbook (Vol. 2, Provider Handbooks) for information about Zika virus testing.

5.3.12 Non-mandated Components

5.3.12.1 Oral Evaluation and Fluoride Varnish (OEFV) in the Medical Home

An OEFV (procedure code 99429) is aimed at improving oral health outcomes for clients who are 6 through 35 months of age by initiating a limited set of preventive dental services (not a dental checkup) in the medical home.

The OEFV must be billed on the same date of service as a medical checkup (procedure codes 99381, 99382, 99391, and 99392) and is limited to six services per lifetime by any provider. Procedure code 99429 must be billed with modifier U5 and diagnosis code Z00121 or Z00129 for an intermediate oral evaluation with fluoride varnish application.

An OEFV is not a required component of a THSteps medical checkup, but providers are encouraged to participate in this preventive intervention. OEFV is limited to THSteps medical checkup providers who have completed the required benefit education and are certified by Texas Health Steps to perform OEFV services.

Training for certification is available as a free continuing education course on the THSteps website at www.txhealthsteps.com.

The OEFV add-on includes the following components:

- Intermediate oral evaluation
- Inspection of teeth for signs of early childhood caries, and other caries
- Inspection of the oral soft tissues for any abnormalities
- Inspection for bleeding, swelling, or infection
- Indications of lack of cleaning of the mouth

The intermediate oral evaluation components that may be performed by a trained staff member are:

- Fluoride varnish application
- Dental anticipatory guidance to include:
  - The need for thorough daily oral hygiene practices
  - Education in potential gingival manifestations for clients with diabetes and clients under long-term medication therapy
  - THSteps eligibility qualifies the client for dental services
  - Diet, nutrition, and food choices
  - Fluoride needs
  - Injury prevention
  - Antimicrobials, medications, and oral health
If the client has no erupted teeth, additional dental anticipatory guidance is expected.

*Note:* The physician must complete the intermediate oral evaluation but can delegate all other components.

### 5.4 Documentation Requirements

All THSteps services require documentation to support the medical necessity of the services rendered including THSteps medical services. THSteps services are subject to retrospective review and recoupment if documentation does not support the services billed.

The following federal and state mandated components must be documented in the client’s medical record for the checkup to be considered complete:

- Comprehensive health and developmental history, including physical and mental health development
- Comprehensive unclothed physical examination
- Immunizations appropriate for age and health history
- Laboratory test appropriate to age and risk, including lead toxicity at specific federally mandated ages
- Health education including anticipatory guidance
- Dental referral

The client’s medical record must include documentation to support the rationale a component was not completed, and a plan to complete the component(s) if not due to parent or caregiver concern or reasons of conscience, including religious beliefs.

#### 5.4.1 Separate Identifiable Acute Care Evaluation and Management Visit

If an acute or chronic condition that requires E/M beyond the required components for a medical checkup is discovered, a separate E/M procedure code may be considered for reimbursement for the same date of service as a checkup or the client can be referred for further diagnosis and treatment.

- The client’s medical record must contain documentation that the separate identifiable service(s) were medically necessary and include a diagnosis in addition to Z0000, Z0001, Z00110, Z00111, Z00121 or Z00129 and treatment. Documentation must be made available to Texas Medicaid upon request.
- An insignificant or trivial problem or abnormality that is encountered in the process of performing a checkup and does not require additional work and performance of the key components of a problem-oriented E/M service cannot be considered a separate established patient E/M acute care visit.
- Modifier 25 must be used to identify a significant, separately identifiable E/M service rendered by the same provider on the same day of the procedure or other service. Documentation that supports the provision of a significant, separately identifiable E/M service must be maintained in the client’s medical record and made available to Texas Medicaid upon request.

Refer to: [Acute Care Visit on the Same Day as a THSteps Preventive Visit Checkup](https://www.tmhp.com) on the TMHP website for a claim form example.

#### 5.5 Claims Filing and Reimbursement

Providers may refer to Volume 1 for general information about claims filing and reimbursement.
Referto: “Section 3: TMHP Electronic Data Interchange (EDI)” (Vol. 1, General Information) for information on electronic claims submissions.

“Section 6: Claims Filing” (Vol. 1, General Information) for general information about claims filing.

Subsection 6.6, “UB-04 CMS-1450 Paper Claim Filing Instructions” in “Section 6: Claims Filing” (Vol. 1, General Information) for instructions on completing paper claims.

“Section 6: Claims Filing” (Vol. 1, General Information) for paper claims completion instructions.

“Section 2: Texas Medicaid Fee-for-Service Reimbursement” (Vol. 1, General Information) for more information about reimbursement.

Subsection 2.2, “Fee-for-Service Reimbursement Methodology” in “Section 2: Texas Medicaid Fee-for-Service Reimbursement” (Vol. 1, General Information) for more information.

5.5.1 Claims Information

THSteps Medical providers are not required to bill other insurance before billing Medicaid. If a provider is aware of other insurance, the provider must choose whether or not to bill the other insurance. The provider has the following options:

- If the provider chooses to bill the other insurance, the provider must submit the claim to the client’s other insurance before submitting the claim to Medicaid.

- If the provider chooses to bill Medicaid and not the client’s other insurance, the provider is indicating that he or she accepts the Medicaid payment as payment in full. Medicaid then has the right to recovery from the other insurance. The provider does not have the right to recovery and cannot seek reimbursement from the other insurance after Medicaid has made payment.

- If the provider learns that a client has other insurance coverage after Medicaid has paid a claim, the provider must refund the payment to Medicaid before billing the other insurance.

Providers should bill their usual and customary fee except for vaccines obtained from TVFC. Providers may not charge Medicaid or clients for the vaccine received from TVFC. Providers may charge a usual and customary fee not to exceed $14.85 for vaccine administration when providing immunizations to a client eligible for TVFC. Providers are reimbursed the lesser of the billed amount or the maximum allowable fee.

THSteps medical checkups may be billed electronically or on a CMS-1500 paper claim form. Providers may purchase CMS-1500 paper claim forms from the vendor of their choice. TMHP does not supply the forms. Providers may request information about electronic billing or the paper claim form by contacting the TMHP THSteps Contact Center at 1-800-757-5691.

All procedures, including the informational-only procedures, must have a billed amount associated with each procedure listed on the claim. Informational-only procedure codes must be billed in the amount of at least $.01.

Providers must record the following on the CMS-1500 claim form to receive reimbursement for a medical checkup, exception to periodicity checkup, or follow-up visit:

- The provider identifier and benefit code EP1 (exception: FQHC providers do not use benefit code EP1)

- The appropriate Texas Health Steps medical checkup procedure code (all ages) with diagnosis code Z0000, Z0001, Z00110, Z00111, Z00121, or Z00129. Diagnosis code Z23 may also be included.

- The condition indicator codes, which must be placed in 24C (ST, S2, or NU only to identify a checkup resulting in a referral)
- The provider type modifiers
- The exception-to-periodicity modifier, when applicable

Refer to: Subsection 5.3.6, “THSteps Medical Checkups” in this handbook for a listing of modifiers.
- The immunization administration and vaccine procedure codes if any were administered (all ages)
- The place of service must be 72 for RHCs
- The EP modifier must be used for FQHCs

Immunizations performed outside of a THSteps medical checkup must be billed without the benefit code EP1.

5.5.2 Reimbursement

As with all Medicaid services, providers acknowledge compliance with all Texas Medicaid requirements when they submit a claim for reimbursement. THSteps-enrolled providers are reimbursed for THSteps medical checkups and administration of immunizations in accordance with 1 TAC §355.8441.

Note: NP, CNS, and PA providers who are enrolled in Texas Medicaid as THSteps providers may receive 92 percent of the rate paid to a physician for THSteps services.

FQHCs are reimbursed using visit rates calculated in accordance with 1 TAC §355.8261.

RHCs are reimbursed using visit rates calculated in accordance with 1 TAC §355.8101.

Providers may refer to the OFL or the applicable fee schedule on the TMHP website at www.tmhp.com.

6 Claims Resources

Refer to the following sections or forms when filing claims:

<table>
<thead>
<tr>
<th>Resource</th>
<th>Location</th>
</tr>
</thead>
<tbody>
<tr>
<td>Acronym Dictionary</td>
<td>“Appendix C: Acronym Dictionary” (Vol. 1, General Information)</td>
</tr>
<tr>
<td>Automated Inquiry System (AIS)</td>
<td>Subsection A.10, “TMHP Telephone and Fax Communication” in “Appendix A: State, Federal, and TMHP Contact Information” (Vol. 1, General Information)</td>
</tr>
<tr>
<td>CMS-1500 Paper Claim Filing Instructions</td>
<td>Subsection 6.5, “CMS-1500 Paper Claim Filing Instructions” in “Section 6: Claims Filing” (Vol. 1, General Information)</td>
</tr>
<tr>
<td>State, federal, and TMHP contact information</td>
<td>“Appendix A: State, Federal, and TMHP Contact Information” (Vol. 1, General Information)</td>
</tr>
<tr>
<td>TMHP electronic claims submission information</td>
<td>Subsection 6.2, “TMHP Electronic Claims Submission” in “Section 6: Claims Filing” (Vol. 1, General Information)</td>
</tr>
<tr>
<td>TMHP Electronic Data Interchange (EDI) information</td>
<td>“Section 3: TMHP Electronic Data Interchange (EDI)” (Vol. 1, General Information)</td>
</tr>
</tbody>
</table>

7 Contact TMHP

For a complete list of TMHP communications, refer to subsection A.10, “TMHP Telephone and Fax Communication” in “Appendix A: State, Federal, and TMHP Contact Information” (Vol. 1, General Information).
7.1 Automated Inquiry System (AIS)
AIS (1-800-925-9126, Option 1) is available 7 days a week, 23 hours a day, with scheduled downtime between 3 a.m. and 4 a.m., and is the main point of contact for client eligibility information. AIS requires the use of a touch-tone telephone in order to access the system.

7.2 TMHP Website
Additional information about Medicaid enrollment, general customer service, and provider education/training is available on the TMHP website at www.tmhp.com.

7.3 Dental Information and Assistance
For assistance with claims, dental providers may contact a TMHP Contact Center representative on the Dental Inquiry Line (1-800-568-2460).

7.3.1 Dental Inquiry Line
The Dental Inquiry Line (1-800-568-2460) is available Monday through Friday, 7 a.m. to 7 p.m., Central Time, and is the main point of contact for information about dental services and appeals.

Any dental service claim denial may be appealed by telephone if it was not denied as an incomplete claim and does not require one of the following items or conditions:

- Narratives
- Radiographs
- Models
- Other tangible documentation
- Review by the TMHP Dental Director

7.4 THSteps Information and Assistance
Providers with questions, concerns, or problems about claims should contact the TMHP Contact Center (1-800-925-9126). For contact information for their regional TMHP Provider Representative, providers can refer to the TMHP website at www.tmhp.com.

7.4.1 THSteps Inquiry Line
The THSteps Medical Inquiry Line at 1-800-757-5691 is available Monday through Friday, 7 a.m. to 7 p.m., Central Time, and is the main point of contact for information about THSteps medical services.

7.5 Assistance with Program
Providers with questions, concerns, or problems with program rules, policies, or procedures should contact DSHS regional program staff. THSteps staff contact numbers can be found in “Appendix A: State, Federal, and TMHP Contact Information” (Vol. 1, General Information), on the THSteps website at hhs.texas.gov/doing-business-hhs/provider-portals/health-services-providers/texas-health-steps or by calling THSteps at 1-512-776-7745.

THSteps regional staff make routine contact with providers to educate and assist them with THSteps policies and procedures.

Clients who are eligible for Medicaid and have questions about THSteps, need to locate medical or dental providers, or need assistance with arranging transportation to appointments should call the THSteps toll-free helpline (1-877-847-8377). Clients with questions about their Medicaid eligibility for THSteps should be directed to their caseworker at the local HHSC office or site.
## 8 Forms

The following linked forms can also be found on the [Forms](#) page of the Provider section of the TMHP website at [www.tmhp.com](http://www.tmhp.com):

<table>
<thead>
<tr>
<th>Forms</th>
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<tr>
<td>CCP Prior Authorization Request Form</td>
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<tr>
<td>Instructions</td>
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<tr>
<td>CCP Prior Authorization Request for Non-Face-to-Face Clinician-Directed Care Coordination Services</td>
</tr>
<tr>
<td>CCP Prior Authorization Request Form</td>
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<tr>
<td>CCP Prior Authorization Private Duty Nursing 6-Month Authorization</td>
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<td>Criteria for Dental Therapy Under General Anesthesia</td>
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<tr>
<td>DME Certification and Receipt Form</td>
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<tr>
<td>Home Health Plan of Care (POC)</td>
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<tr>
<td>Home Health Plan of Care (POC) Instructions</td>
</tr>
<tr>
<td>Nursing Addendum to Plan of Care for Private Duty Nursing and/or Prescribed Pediatric Extended Care Center</td>
</tr>
<tr>
<td>Texas Medicaid Handicapping Labio-Lingual Deviation (HLD) Index Score Sheet</td>
</tr>
<tr>
<td>THSteps Dental Mandatory Prior Authorization Request Form</td>
</tr>
<tr>
<td>THSteps Referral Form Instructions</td>
</tr>
<tr>
<td>THSteps Referral Form</td>
</tr>
<tr>
<td>Specialist or Subspecialist Telephone Consultation Form for Non-Face-to-Face Clinician-Directed Care Coordination Services–Comprehensive Care Program (CCP)</td>
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</table>

## 9 Claim Form Examples

The following linked claim form examples can also be found on the [Claim Form Examples](#) page of the Provider section of the TMHP website at [www.tmhp.com](http://www.tmhp.com):

<table>
<thead>
<tr>
<th>Claim Form Examples</th>
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</thead>
<tbody>
<tr>
<td>Acute Care Visit on the Same Day as a THSteps Preventive Visit Checkup</td>
</tr>
<tr>
<td>Diagnosis and Treatment (Referral from THSteps Checkup)</td>
</tr>
<tr>
<td>Durable Medical Equipment (CCP Only)</td>
</tr>
<tr>
<td>Inpatient Rehabilitation Facility (Freestanding) (CCP Only)</td>
</tr>
<tr>
<td>Medical Checkup Follow-up Visit with Immunization Administration</td>
</tr>
<tr>
<td>Medical Checkup Follow-up Visit with TB Skin Test (TST)</td>
</tr>
<tr>
<td>Medical Nutrition Counseling (CCP Only)</td>
</tr>
<tr>
<td>Orthotic and Prosthetic Services (CCP Only)</td>
</tr>
<tr>
<td>Postpartum Depression Screening During an Infant THSteps Checkup</td>
</tr>
<tr>
<td>Private Duty Nurses (CCP Only)</td>
</tr>
<tr>
<td>THSteps New Patient, Immunization Without Counseling no Referral and by an NP</td>
</tr>
<tr>
<td>THSteps Established Patient Exception to Periodicity and Referral, Immunizations with Counseling, and by a Physician</td>
</tr>
<tr>
<td>THSteps Established Patient and Referral, Tuberculin Skin Test (TST), and Physical Examination by a Physician</td>
</tr>
</tbody>
</table>
Claim Form Examples

THSteps Preventive Visit Checkup with Immunization and Vaccine Administration
APPENDIX A. THSteps FORMS

A.1 Claim Forms

Providers must order CMS-1500 and American Dental Association (ADA) Dental Claims Forms from the vendor of their choice. Copies cannot be used. Claims filing instructions and examples of the claim forms are located in “Section 6: Claims Filing” (Vol. 1, General Information).

Refer to:


A.2 THSteps Medical Checkup Forms

The use of the child health clinical records is optional. These forms were developed to help providers document all components of the medical checkup. Unless required to be submitted to another program, one of the following forms of documentation must be included in the client’s medical record: The completed screening tools with results, the completed questions to the tools within a provider-created medical record, and the results of the completed screening tools. Providers may be asked to provide the screening tool used to complete the screening. Texas Health Steps (THSteps) requires the following forms: Tuberculosis (TB) Questionnaire and the Texas Department of State Health Services (DSHS) State Laboratory forms. These forms can be downloaded from the THSteps website at https://hhs.texas.gov/doing-business-hhs/provider-portals/health-services-providers/texas-health-steps/forms. The Parent Hearing Checklist and Lead Risk Questionnaire are optional forms. Lead poisoning screening questionnaires can be downloaded from the Texas Childhood Lead Poisoning Prevention Program (TX CLPPP) website at www.dshs.texas.gov/lead/providers.shtm.

Links to growth charts may be found on the THSteps website at https://hhs.texas.gov/doing-business-hhs/provider-portals/health-services-providers/texas-health-steps/forms.

<table>
<thead>
<tr>
<th>Form Number</th>
<th>Form Name</th>
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<tr>
<td>ECH-1</td>
<td>Child Health History Form</td>
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<td>ECHR-5D</td>
<td>Discharge to 5 day Visit Child Health Record</td>
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<tr>
<td>ECHR-2W</td>
<td>2 Week Visit Child Health Record</td>
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<tr>
<td>ECHR-2M</td>
<td>2 Month Visit Child Health Record</td>
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<tr>
<td>ECHR-4M</td>
<td>4 Month Visit Child Health Record</td>
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<tr>
<td>ECHR-6M</td>
<td>6 Month Visit Child Health Record</td>
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<tr>
<td>ECHR-9M</td>
<td>9 Month Visit Child Health Record</td>
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<td>ECHR-12M</td>
<td>12 Month Visit Child Health Record</td>
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<td>ECHR-15M</td>
<td>15 Month Visit Child Health Record</td>
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<tr>
<td>ECHR-18M</td>
<td>18 Month Visit Child Health Record</td>
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<tr>
<td>ECHR-24M</td>
<td>24 Month Visit Child Health Record</td>
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<tr>
<td>ECHR-30M</td>
<td>30 Month Visit Child Health Record</td>
</tr>
<tr>
<td>ECHR-3Y</td>
<td>3 Year Visit Child Health Record</td>
</tr>
<tr>
<td>ECHR-4Y</td>
<td>4 Year Visit Child Health Record</td>
</tr>
<tr>
<td>ECHR-5Y</td>
<td>5 Year Visit Child Health Record</td>
</tr>
</tbody>
</table>
Providers should refer to sources such as *Bright Futures: Guidelines for Health Supervision of Infants, Children, and Adolescents* (3rd edition), located at www.brightfutures.org or the Guidelines for Adolescent Preventive Services (GAP) Implementation Materials located at https://brightfutures.aap.org/materials-and-tools/PerfPrevServ/Pages/default.aspx. For nutritional screening for all ages, refer to Bright Futures.

### A.3 Laboratory Forms

For information on procedures for submission of laboratory forms, refer to the DSHS Laboratory Services Section’s web page at www.dshs.texas.gov/lab/MRS_forms.shtm.

### A.4 Guidelines for Tuberculosis Skin Testing

For information on procedures for tuberculosis skin testing, refer to the DSHS tuberculosis web page at www.dshs.texas.gov/idcu/disease/tb/.

### A.5 Tuberculosis Screening and Guidelines

The screening tool for tuberculosis (TB) exposure risk is to be used annually to determine the need for tuberculin skin testing.

The questions in the screening tool are intended as a minimum screen. Follow-up questions may be necessary to clarify hesitant or ambiguous responses. Questions specific to TB exposure risks in the client’s community may need to be added.

The following applies for tuberculin screening and skin testing:

- If all the answers are unqualified negatives, the client is considered at low risk for exposure to TB and will not need tuberculin skin testing.
- If the answer to any question is “Yes” or “I don’t know,” the client should be tuberculin skin tested.

<table>
<thead>
<tr>
<th>Form Number</th>
<th>Form Name</th>
</tr>
</thead>
<tbody>
<tr>
<td>ECHR-6 Year</td>
<td>6 Year Visit Child Health Record</td>
</tr>
<tr>
<td>ECHR-7 Year</td>
<td>7 Year Visit Child Health Record</td>
</tr>
<tr>
<td>ECHR-8 Year</td>
<td>8 Year Visit Child Health Record</td>
</tr>
<tr>
<td>ECHR-9 Year</td>
<td>9 Year Visit Child Health Record</td>
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<tr>
<td>ECHR-10 Year</td>
<td>10 Year Visit Child Health Record</td>
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<td>ECHR-11 Year</td>
<td>11 Year Visit Child Health Record</td>
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<tr>
<td>ECHR-12 Year</td>
<td>12 Year Visit Child Health Record</td>
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<tr>
<td>ECHR-13 Year</td>
<td>13 Year Visit Child Health Record</td>
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<td>ECHR-14 Year</td>
<td>14 Year Visit Child Health Record</td>
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<td>ECHR-15 Year</td>
<td>15 Year Visit Child Health Record</td>
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<td>ECHR-16 Year</td>
<td>16 Year Visit Child Health Record</td>
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<td>ECHR-18 Year</td>
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<tr>
<td>ECHR-19 Year</td>
<td>19 Year Visit Child Health Record</td>
</tr>
<tr>
<td>ECHR-20 Year</td>
<td>20 Year Visit Child Health Record</td>
</tr>
<tr>
<td>ECHR-19-20 Year</td>
<td>Form Pb-110, Lead Risk Questionnaire</td>
</tr>
<tr>
<td></td>
<td>TB Questionnaire</td>
</tr>
</tbody>
</table>
• In the case of the client for whom an answer in the past of “Yes” or “I don’t know” prompted a skin test, which was negative, the skin test may not have to be repeated annually.

• The decision to administer a skin test must be made by the medical provider based upon an assessment of the possibility of exposure. A negative tuberculin skin test never excludes tuberculosis infection or active disease.

• Bacillus of Calmette and Guérin (BCG) vaccinated clients should also have the screening tool administered annually. Previous BCG vaccination is not a contraindication to tuberculin skin testing. Positive tuberculin skin tests in BCG vaccinated children are interpreted using the same guidelines used for non-BCG vaccinated children.

• Clients who have had a positive TB skin test in the past (whether treated or not), should be re-evaluated at least annually by a physician for signs and symptoms of TB.

Care of clients who are newly discovered to be tuberculin skin test positive includes:

• An evaluation for signs and symptoms of TB.

• A chest X-ray to rule out active disease.

• Oral medications to prevent progression to active disease or multi-drug therapy if active disease is present.

• Referral for consultation by a pediatric TB specialist is recommended if active disease is present.

• A report to the local health authority for investigation to find the source of the infection.

Refer to: The TB screening tool on the THSteps website at https://hhs.texas.gov/doing-business-hhs/provider-portals/health-services-providers/texas-health-steps/forms.

“Frequently Asked Questions About TB” on the DSHS website at www.dshs.texas.gov/idcu/disease/tb/faqs for information about assessing a child’s risk of TB exposure, providers can refer to

The TVFC Provider Manual, Forms, & Resources web page on the DSHS website at www.dshs.texas.gov/immunize/tvfc/ProviderResources.shtm#forms for the Patient Eligibility Screening Record (Bilingual), Questions & Answers about TVFC, and additional information.
APPENDIX B. IMMUNIZATIONS

B.1 Immunizations Overview

Clients who are 17 years of age and younger must be immunized according to the Recommended Childhood Immunization Schedule for the United States. If the immunizations are due as part of a Texas Health Steps (THSteps) medical checkup, the medical checkup provider is responsible for the administration of immunizations for clients who are birth through 20 years of age and may not refer clients to local health departments. The Department of State Health Services (DSHS) requires that immunizations be administered during the THSteps medical checkup, unless they are medically contraindicated or excluded from immunization for reasons of conscience, including a religious belief.

Providers, in both public and private sectors, are required by federal mandate to provide a Vaccine Information Statement (VIS) to the responsible adult accompanying a client for an immunization. These statements are specific to each vaccine and inform the responsible adult about the risks and benefits. It is important that providers use the most current VIS.

Providers interested in obtaining copies of current VISs and other immunization forms or literature may call the DSHS Immunization Branch at 1-512-458-7284. VISs may also be downloaded from the DSHS Immunization Branch website at www.immunizetexas.com.

B.1.1 Vaccine Adverse Event Reporting System (VAERS)

The National Childhood Vaccine Injury Act of 1986 (NCVIA) requires health-care providers to report:

- Any reaction listed by the vaccine manufacturer as a contraindication to subsequent doses of the vaccine.
- Any reaction listed in the Reportable Events Table that occurs within the specified time period after vaccination.

Clinically significant adverse events should be reported even if it is unclear whether a vaccine caused the event.

**Note:** Documentation of the injection site is recommended but not required.

For additional information about documentation, providers can refer to www.vaers.hhs.gov.

A copy of the Reportable Events Table can be obtained by calling VAERS at 1-800-822-7967 or by downloading it from http://vaers.hhs.gov/resources/vaersmaterialspublications.

B.1.2 TVFC Versus Non-TVFC Vaccines/Toxoids

When single antigen vaccines/toxoids or comparable antigen vaccines/toxoids are available for distribution through the Texas Vaccines for Children (TVFC) Program, but the provider chooses to use a different Advisory Committee on Immunization Practices (ACIP)-recommended product, the vaccine/toxoid will not be reimbursed; however, the administration fee will be considered.

**Note:** All administered vaccines/toxoids must be reported to DSHS. DSHS submits all vaccines/toxoids reported with consent to a centralized immunization registry, known as ImmTrac2.

**Refer to:** Subsection B.3.5, “How to Report Immunization Records to ImmTrac2, the Texas Immunization Registry” in this handbook.

B.1.3 Exemption from Immunization for School and Child-Care Facilities

Parents may obtain an exemption from immunization requirements for school and childcare entry for reasons of conscience or religious beliefs. An exemption is also available for clients who are medically contraindicated from receiving a vaccine. For more information on exemptions call 1-512-458-7284, or visit www.immunizetexas.com.

**Refer to:** Section 5, “THSteps Medical” in this handbook.
B.2 Recommended Childhood Immunization Schedule

The Recommended Childhood Immunization Schedule indicates the recommended age for routine administration of currently licensed childhood vaccines. This schedule was developed and approved by ACIP, the American Academy of Pediatrics (AAP), and the American Academy of Family Physicians (AAFP).

Some combination vaccines are available and may be used whenever any component of the combination is indicated and its other components are not contraindicated. Providers should consult the manufacturer’s package insert for detailed recommendations.

Vaccines should be administered at the recommended ages. Any dose not given at the recommended age should be given as a catch-up immunization on any subsequent visit when indicated and feasible.

A current copy of the Recommended Childhood Immunization Schedule can be accessed at www.cdc.gov/vaccines/schedules.

Refer to: Recommended Childhood and Adolescent Immunization Schedule on the THSteps page of the TMHP website at www.tmhp.com.

B.3 General Recommendations

For information about vaccine administration, dosing, and contraindications, immunization providers should consult vaccine package inserts and the January 28, 2011, issue of the Center for Disease Control and Prevention Morbidity and Mortality Weekly Report (MMWR). For copies of the MMWR, contact the Immunization Branch at 1-512-458-7284.

B.3.1 How to Obtain Vaccines at No Cost to the Provider

TVFC provides routinely recommended ACIP vaccines for immunization of THSteps and other Medicaid- and TVFC-eligible clients free of charge to providers who are enrolled in TVFC. The local health department/district or DSHS regional office provides information on how to order, account for, and inventory vaccines. Local and public health departments that are not otherwise enrolled as a provider that is authorized to receive reimbursement for vaccine administration fees should enroll as a Comprehensive Care Program (CCP) provider. Monthly reports are required in order to receive state-purchased vaccines. Physicians who request and accept state-supplied vaccines must complete and sign the provider enrollment and profile forms annually. The provider may not charge Medicaid or the client for vaccines obtained from TVFC.

Additional information is available at www.immunizetexas.com.

B.3.2 Administrations and Immunizations

B.3.2.1 Administrations

The following administration procedure codes must be submitted in combination with an appropriate vaccine/toxoid procedure code:

<table>
<thead>
<tr>
<th>Administration Procedure Code</th>
<th>90460</th>
<th>90461</th>
<th>90471</th>
<th>90472</th>
<th>90473</th>
<th>90474</th>
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Procedure codes 90460 and 90461 are benefits for services rendered to clients who are birth through 18 years of age when counseling is provided for the immunization administered. Documentation of counseling by the physician or other qualified health-care professional must be noted in the client’s medical record.
Procedure codes 90471 and 90472 are benefits for services rendered to clients of any age when counseling is not provided for the immunization administered. Procedure codes 90473 and 90474 are benefits for services rendered to clients who are birth through 20 years of age when counseling is not provided for the immunization administered.

Referto: Subsection 5.3.11.3 *, “Immunizations” in this handbook for appropriate diagnosis codes for immunization administration.

B.3.3 * Immunizations (Vaccine/Toxoids)

The following vaccines and toxoids are a benefit of Texas Medicaid:

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</table>

* TVFC-distributed vaccine/toxoid
** The number of components applies if counseling is provided and procedure code 90460 and 90461 are submitted.

Procedure codes 90655, 90657, 90685, and 90687 are limited to clients who are 6 through 35 months of age.

Procedure codes 90656 and 90658 are limited to clients who are 3 years of age and older.

Procedure codes 90686 and 90688 are limited to clients who are 6 months of age and older.

Procedure code 90682 is limited to clients who are 18 years of age and older. Procedure code 90756 is limited to clients who are 4 years of age and older.

Providers may use the state-defined modifier U1 in addition to the associated administered vaccine procedure code for clients who are birth through 18 years of age and the vaccine was unavailable through TVFC.
<table>
<thead>
<tr>
<th>Modifier</th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td>U1</td>
<td>State-defined modifier: Vaccines/toxoids privately purchased by provider when TVFC vaccine/toxoid is unavailable</td>
</tr>
</tbody>
</table>

**Note:** “Unavailable” is defined as a new vaccine approved by ACIP that has not been negotiated or added to a TVFC contract, funding for new vaccine that has not been established by TVFC, or national supply or distribution issues. Providers will be informed if a vaccine meets the definition of 'not available' from TVFC and when the provider’s privately purchased vaccine may be billed with modifier U1.

Modifier U1 may not be used for failure to enroll in TVFC, maintain sufficient TVFC vaccine/toxoid inventory, or for clients who are 19 through 20 years of age.

**B.3.4 Requirements for TVFC Providers**

By enrolling, public and private providers agree to:

- Screen patients for TVFC eligibility at all immunization encounters, and administer TVFC-purchased vaccines only to clients who are 18 years of age and younger who meet one or more of the following criteria:
  - Is an American Indian or Alaska Native.
  - Is enrolled in Medicaid.
  - Has no health insurance.
  - Is underinsured: clients who have other health insurance but the coverage does not include vaccines, clients whose insurance covers only selected vaccines (TVFC-eligible for noncovered vaccines only), clients whose insurance capitates vaccine coverage at a certain amount (once that coverage amount is reached, these clients are categorized as underinsured).
  - Is a client who receives benefits from the Children’s Health Insurance Program (CHIP) and the provider bills CHIP for the administration fee.

- Maintain all records related to the TVFC program, including parent, guardian, or authorized representative’s responses to screening for patient’s eligibility for at least five years. If requested, the provider will make such records available to DSHS, the local health department authority, or the U.S. Department of Health and Human Services (HHS).

- Comply with the appropriate vaccination schedule, dosage, and contraindications, as established by ACIP, unless (a) in making a medical judgment in accordance with accepted medical practice, the provider deems such compliance to be medically inappropriate, or (b) the particular requirement is not in compliance with Texas law, including laws relating to religious and medical exemptions.

- Provide VISs to the responsible adult, parent, or guardian, and maintain records in accordance with the NCVIA which include reporting clinically significant adverse events to VAERS. Signatures are required for informed consent. (The Texas Addendum portion of the VIS may be used to document informed consent.)

- Not charge for vaccines supplied by DSHS and administered to a client who is eligible for TVFC.

- Charge a vaccine administration fee to Texas Medicaid but not impose a charge for the administration of the vaccine in any amount higher than the maximum administration fee established by DSHS (providers may charge a vaccine administration fee to Medicaid, but not a fee for the vaccine). Medicaid clients cannot be charged any out-of-pocket expense for the vaccine or the administration of the vaccine.

- Not deny administration of a TVFC vaccine to a client because of the inability of the client’s parent or guardian/individual of record to pay an administration fee.
• Comply with the state’s requirements for ordering vaccines and other requirements as described by DSHS, and operate within the TVFC program in a manner intended to avoid fraud and abuse.

• Allow DSHS (or its contractors) to conduct scheduled and unannounced storage and handling visits.

The provider or the state may terminate the agreement at any time for failure to comply with the requirements listed above. If the agreement is terminated for any reason, the provider agrees to properly return any unused vaccine.

B.3.5 How to Report Immunization Records to ImmTrac2, the Texas Immunization Registry

Texas law requires all medical providers and payors to report all immunizations administered to clients who are 17 years of age and younger, to ImmTrac2, the Texas immunization registry operated by DSHS (Texas Health and Safety Code §§161.007-161.009). Providers must report all immunization information within 30 days of administration of the vaccine, and payors must report within 30 days of receipt of data elements from a provider. Prior to reporting immunizations to ImmTrac2, providers must first register for registry participation and access.

ImmTrac2 is a centralized repository of immunization histories for clients of all ages and is a free service and benefit available to all Texans. Registry information is confidential, and by law, may be released only to:

• The client or client’s parent, legal guardian, or managing conservator.

• The client’s physician, school, or licensed child-care facility in which the client is enrolled.

• Public health districts or local health departments.

• The insurance company, health maintenance organization, or other organization that pays for the provision of the client’s health-care benefits.

• A health-care provider authorized to administer a vaccine.

• A state agency that has legal custody of the client.

ImmTrac2 offers two methods for reporting immunizations to DSHS: direct internet entry into ImmTrac2’s internet application and electronic data transfer (import).

B.3.5.1 Direct Internet Entry

This method allows providers to access and review clients’ immunization histories prior to administering vaccines. Providers then update their client’s immunization record directly into the ImmTrac2 web application after administering vaccines to the patient.

B.3.5.2 Electronic Data Transfer (Import)

This method allows providers to report immunizations from an electronic medical record (EMR) software application by extract file for import into ImmTrac2. Providers may still have access to the ImmTrac2 web application to access and review their clients’ immunization histories after administering any vaccines.

Regardless of reporting option selected, all providers must first register for ImmTrac2 access and receive login credentials from ImmTrac2 Customer Support. To register for ImmTrac2 access, providers may obtain and complete an ImmTrac2 Registration Packet (for providers and schools) from www.dshs.state.tx.us/immunize/immtrac/default.shtm or request it from ImmTrac2 Customer Support at 1-800-348-9158.
B.3.5.3 Obtaining Parental Consent for Registry Participation

Before including a client’s immunization information in ImmTrac2, DSHS must verify that written consent for registry participation has been granted by the client’s parent, legal guardian, or managing conservator. Most parents grant consent for ImmTrac2 participation during the birth certificate registration process. Written parental consent for ImmTrac2 participation applies to all past, present, and future immunizations. Texas law also permits a parent, managing conservator, or guardian to withdraw consent for ImmTrac2 participation at any time.

Providers may offer parents the opportunity to grant consent for their child’s participation in ImmTrac2 using the pre-filled, ImmTrac2-generated Immunization Registry (ImmTrac2) Consent Form or the manual version (#C-7) of this form, also available from the ImmTrac2 application. Providers should retain the consent form and affirm parental consent through ImmTrac2 to establish the client’s ImmTrac2 record and report all immunizations administered and add any historical immunization information to the client’s record. Entering administered immunizations and historical immunization information to the client’s record constitutes “reporting” to ImmTrac2 as required by current Texas law.

B.4 Texas Vaccines for Children Program Packet

Refer to: The DSHS website for TVFC information at www.dshs.texas.gov/immunize/tvfc/default.shtm.
APPENDIX C.  LEAD SCREENING

C.1 Blood Lead Screening Procedures and Follow-up Testing
For all children enrolled in Texas Health Steps (THSteps) blood lead testing is mandatory when they are 12 months of age and 24 months of age, or whenever they receive their first checkup after these ages if blood testing was not completed (up to and including the 6-year checkup). Lead-risk assessment should be done at all other checkups through age 6, and may be performed using Form PB 110, Lead Risk Questionnaire. A “yes” or “don’t know” answer to any question on the questionnaire indicates that a blood lead test should be administered. All blood lead levels in clients who are birth through 14 years of age must be reported to the Department of State Health Services (DSHS). Reports should include all information as required on the Texas Child Blood Lead Level Report Form F09-11709, which is available at www.dshs.texas.gov/lead/providers.shtm or by calling 1-800-588-1248. Information related to blood lead screening and reporting for clients who are 15 years of age or older is available on the DSHS Blood Lead Surveillance Group’s website at www.dshs.texas.gov/lead/default.shtm.

C.2 Symptoms of Lead Poisoning
Children who have EBLLs in the range of 5–45µg/dL may be asymptomatic, although impairment of neurodevelopment may become evident as they get older. Very high lead levels may cause colic, constipation, anorexia, or vomiting. Children with venous blood lead levels (BLLs) over 44µg/dL are eligible for medical intervention. However, it is important not to equate the absence of symptoms with the absence of toxicity.

C.3 Measuring Blood Lead Levels
A blood lead test is the only definitive method to detect exposure. BLLs are measured as micrograms of lead per deciliter of whole blood (µg/dL). In Texas, a BLL requires medical case management and follow-up testing if the level is greater than or equal to 5 µg/dL.

Blood lead tests, in order of occurrence:

- Screening test—A blood lead test that indicates whether a client may have an EBLL. This test must be sent to the DSHS lab, or may be done using point-of-care technology in the provider’s office.

- Diagnostic test—A venous blood lead test that is performed within recommended guidelines to determine the status of a client who has previously had an EBLL on a screening test (See subsection C.5, “Lead Poisoning Prevention Educational Materials and Forms” in this handbook for a link to form Pb-109 and recommended guidelines).

- Follow-up test—A venous blood lead test to monitor the status of a client with a previously elevated diagnostic test for lead.

Note: A follow-up test is not related to the THSteps follow-up visit. A visit to monitor a child with EBLL would be submitted as an acute care evaluation and management (E/M) visit.

Providers are responsible for conducting a diagnostic test when a screening test finds a lead level of 5 µg/dL or greater. Blood for a screening test may be drawn from a venous or capillary site. A venous blood draw is strongly recommended and preferred. To order venous sample supplies from the DSHS Laboratory, call 1-888-963-7111, Ext. 7661.

Note: The capillary lead screen analysis is subject to a false positive result from skin lead contamination during collection. A soap and water wash of the patient’s hands or feet and the collector’s hands (or the wearing of gloves) must be performed to minimize the chance of contamination. Alcohol cleansing alone is not sufficient.
If the screening test is 5 µg/dL or above, recalling a client for a diagnostic sample may be billed as a THSteps follow-up visit. If the screening test was rejected due to clotting, insufficient quantities, or perceived contamination, the provider must repeat the sample as a diagnostic test. Again, the provider may bill the visit and analysis as an E/M visit. Providers can submit the specimen to the DSHS Clinical Chemistry Laboratory using the appropriate DSHS Laboratory Specimen Submission form (the same way as for all other THSteps laboratory blood specimens). If the initial blood lead test is collected as part of a THSteps medical checkup, it must either be sent to the DSHS lab or performed in the provider’s office using point-of-care. The diagnostic and follow-up test for the same client may be sent to a private laboratory.

Refer to: **Pb-109: Reference for Blood Lead Retesting and Medical Case Management** on the DSHS website for interpretation of laboratory test results and guidelines for follow-up for clients with elevated blood lead levels.

Subsection 5.3.11.6.6, “Required Laboratory Tests Related to Medical Checkups” in this handbook.

Subsection 5.3.9, “Newborn Examination” in this handbook.

Providers can find more information about the medical and environmental management of lead-poisoned children on the DSHS Texas Childhood Lead Poisoning Prevention Program (TX CLPPP) website at www.dshs.texas.gov/lead or by calling 1-800-588-1248.

### C.4 Environmental Lead Investigation Services

#### C.4.1 Enrollment

State and local health departments that employ or contract certified lead risk assessors must be enrolled with Texas Medicaid as a THSteps provider to perform environmental lead investigation (ELI) services.

- State and local health departments that are currently enrolled in Texas Medicaid must complete the THSteps Provider Enrollment Application.

- State and local health departments that are not currently enrolled in Texas Medicaid must complete the Texas Medicaid Provider Enrollment Application and the THSteps Provider Enrollment Application.

#### C.4.1.1 Services, Benefits, Limitations, and Prior Authorization

ELI services must be billed with procedure code T1029, which is restricted to the following diagnosis codes:

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<thead>
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<th>Diagnosis Codes</th>
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<tr>
<td>T560X1A</td>
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<td>T560X3S</td>
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<td>T560X4S</td>
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Texas Medicaid may only reimburse a state or local health department for the certified lead risk assessor’s time and activities during an onsite investigation of a client’s home or primary residence. Laboratory analysis of environmental substances (e.g., water, paint, or soil) is not a benefit of Texas Medicaid.

Children who have confirmed and persistent EBLLs may require an ELI to determine the source of the lead exposure. An ELI is completed in a client’s home or primary residence by a certified lead risk assessor to determine whether a lead hazard exists and, if so, whether the lead source could be the cause of the EBLL.
C.4.1.2 Requesting an Environmental Lead Investigation

For the purpose of requesting an ELI, a health-care provider is a physician, nurse practitioner (NP), clinical nurse specialist (CNS), or physician assistant (PA) who conducts blood lead tests for a THSteps client. Health-care providers may submit a request for an ELI after a blood lead test has been conducted and there is evidence of persistent and confirmed EBLLs for the client. An EBLL is defined as a BLL of 10µg/dL or higher.

An ELI may be considered medically necessary if the results of the most recent blood lead test indicate any of the following:

- A venous BLL result of 10µg/dL to 19µg/dL from two separate specimens conducted at least 12 weeks apart
- A venous BLL result of 20µg/dL or greater from one specimen

*Note: The ELI must be requested as soon as possible and no later than 30 days after obtaining the most recent BLL that indicates medical necessity. The health-care provider must maintain in the client’s medical record the ELI request and the documentation of the BLL that indicates medical necessity.*

The health-care provider can request an ELI by completing Form Pb-101 “Environmental Lead Investigation Request” and submitting it to the TX CLPPP. TX CLPPP will review the request and determine whether the criteria for an ELI have been met. If an ELI request meets the TX CLPPP criteria, TX CLPPP sends a referral for an ELI to a state or local health department that is enrolled as a THSteps provider so that it can be assigned to a certified lead risk assessor. A certified lead risk assessor conducts an ELI using a completed Pb-103 Texas Elevated Blood Lead Level Investigation Questionnaire (all pages).

An ELI can be performed under one of the following circumstances:

- No previous investigation of the current home or primary residence has been performed.
- There is a change in the client’s current home or primary residence.

If a previous investigation of the current home or primary residence has been performed and there has been a change in the client’s residential environment, TX CLPPP will determine whether the criteria have been met for an additional ELI.

C.4.1.3 Prior Authorization

Prior Authorization is not required for ELI services.

C.4.2 Documentation Requirements

The state or local health department that is responsible for conducting the investigation must maintain the following documentation in the client’s medical record:

- The TX CLPPP fax transmittal cover sheet that refers the ELI request to the local health department. The cover sheet must include:
  - The site to be assessed.
  - A statement that identifies the site as the client’s primary place of residence.
- A completed Form Pb-101: Environmental Lead Investigation Request (two pages) that includes the:
  - Name of the referring health-care provider.
  - BLLs that indicate medical necessity.
  - Client’s diagnosis.
- A completed Form Pb-103: Elevated Blood Lead Level Investigation Questionnaire (all pages) that includes the:
• Date and location of the investigation.
• Name of the client who received the investigation.
• Identifying information and signature of the certified lead risk assessor who conducted the investigation. The person listed as the assessor must be the same person who signs the report.

Note: Forms Pb-101 and Pb-103 are located on the TX CLPPP website at www.dshs.texas.gov/lead/providers.shtm.

C.4.3 Claims Filing and Reimbursement

C.4.3.1 Claims Filing
ELI services must be submitted to Texas Medicaid & Healthcare Partnership (TMHP) in an approved electronic format or on the CMS-1500 paper claim form. Providers can purchase CMS-1500 paper claim forms from the vendor of their choice. TMHP does not supply the forms.

The following documentation must be submitted with the claim:
• The TX CLPPP fax transmittal cover sheet that refers the ELI request to the state or local health department. The cover sheet must include:
  • The site to be assessed.
  • A statement that identifies the site as the client’s primary place of residence.
• A completed Form Pb-101: Environmental Lead Investigation Request.
• The first and last page of Form Pb-103: Elevated Blood Lead Level Investigation Questionnaire, which has been completed by the lead risk assessor.

An ELI is subject to retrospective review and may be recouped if the documentation maintained by the health-care and ELI providers does not support medical necessity.

Refer to:
“Section 3: TMHP Electronic Data Interchange (EDI)” (Vol. 1, General Information) for information about electronic claims submissions.
“Section 6: Claims Filing” (Vol. 1, General Information) for general information about claims filing.
Subsection 6.5, “CMS-1500 Paper Claim Filing Instructions” in “Section 6: Claims Filing” (Vol. 1, General Information) for instructions on completing paper claims.

C.4.3.2 Managed Care Clients
ELI services are carved-out of the Medicaid Managed Care Program and must be billed to TMHP for payment consideration. Carved-out services are those that are rendered to Medicaid Managed Care clients but are administered by TMHP and not the client’s managed care organization (MCO).

C.4.3.3 Reimbursement
Providers can refer to the online fee lookup (OFL) or the applicable fee schedule on the TMHP website at www.tmhp.com.

C.5 Lead Poisoning Prevention Educational Materials and Forms
Providers may download lead poisoning prevention education materials and forms from the Texas CLPPP website at www.dshs.texas.gov/lead.
The following table lists materials available to providers for download:

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<td>Pb-110</td>
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APPENDIX D.  TEXAS HEALTH STEPS STATUTORY STATE REQUIREMENTS

D.1 Legislative Requirements
Several specific legislative requirements affect Texas Health Steps (THSteps) and the provider’s participation in Texas Medicaid. The legislation includes, but is not limited to, those included in this Appendix.

D.2 Texas Health Steps (THSteps) Program
The Early and Periodic Screening, Diagnosis, and Treatment (EPSDT) Program is mandated by Title XIX of the Social Security Act. EPSDT is a program of prevention, diagnosis, and treatment for Medicaid-eligible clients who are birth through 20 years of age.

In Texas, EPSDT is known as THSteps. The Texas Department of State Health Services (DSHS), by authorization of Texas Department of Health and Human Services (HHSC), operates and administers the outreach and informing, medical and dental checkup, dental treatment utilization components of this program. State authority is found in Title 25 Texas Administrative Code (TAC), Part 1, Chapter 33, Subchapter A, Rule §33.1.

D.3 Communicable Disease Reporting
Diagnosis of sexually transmitted diseases (STDs), including human immunodeficiency virus (HIV), are reportable conditions under 25 TAC, Chapter 97, Subchapter F. Providers must report confirmed diagnosis of STDs as required by 25 TAC §§97.132-134.

D.4 Early Childhood Intervention (ECI) Referrals
All health-care professionals are required by federal and state regulations to refer children who are birth through 35 months of age to the Texas HHS ECI program as soon as possible, but no longer than 7 days after identifying a disability or suspected delay in development.

Referrals can be based on professional judgment or a family’s concern. A medical diagnosis or a confirmed developmental delay is not required for referrals.

To refer families for services, providers should use the ECI referral form available on the Texas Pediatric Society website at https://txpeds.org/sites/txpeds.org/files/documents/eci-referral-form-icd-10.pdf. Providers may also refer families for services by calling their local ECI program or the Health and Human Services Office of the Ombudsman at 1-877-787-8999, select a language, and then select Option 3. The Ombudsman staff will ask for your ZIP code, county, or city and provide the name and number of the local ECI program. Callers that are deaf or hard of hearing may use the relay option of their choice or dial 7-1-1 to connect with Relay Texas.

To facilitate referrals for ECI services an optional form is available on the Texas Pediatric Society website at https://txpeds.org/sites/txpeds.org/files/documents/ECI-Referral-Form.pdf.

For additional ECI information, providers can visit the HHS ECI website at https://hhs.texas.gov/services/disability/early-childhood-intervention-services. Persons who are deaf or hard of hearing may use the relay option of their choice or dial 7-1-1 to connect with Relay Texas.

D.5 Parental Accompaniment
Texas Human Resource Code (HRC) §§32.024(s)-(2) requires that, as a condition for provider reimbursement, a client who is 14 years of age or younger be accompanied by the client’s parent, legal guardian, or other authorized adult during medical and dental checkups and dental treatment. The authorized adult can be the client’s relative. DSHS implemented this requirement through rules found in 25 TAC §33.2 (Definitions) and 25 TAC §33.6 (THSteps Provider Responsibilities).
The DSHS rules require that the parent, legal guardian, or authorized adult accompany the client to the checkup, and that the parent, legal guardian, or authorized adult must wait for the client while the checkup, treatment, or service takes place.

Providers will not be required to submit documentation to TMHP to verify compliance with this policy in order for TMHP to process claims. By submitting the claim for reimbursement, the provider acknowledges compliance with all Medicaid requirements. Additional assurances are not necessary.

**Exception:** School health clinics, Head Start programs, and childcare facilities are exempt from this policy if the clinic, program, or facility encourages parental involvement in the health care of the client and obtains written consent for the services. The consent from the client's parent or guardian must have been received within the one-year period before the date on which the services are provided and must not have been revoked.

**Refer to:** HRC §§32.024(s)-(s-1) and 25 TAC §33.2 and §33.6.

### D.6 Newborn Blood Screening

The Health and Safety Code (HSC), Chapter 33, Section §33.011, implemented by the rules found at 25 TAC, Part 1, Chapter 37, Subchapter D, requires testing of all newborns. A current list of disorders can be found at [www.dshs.texas.gov/newborn/screened_disorders.shtm](http://www.dshs.texas.gov/newborn/screened_disorders.shtm).

This testing is the responsibility of the physician who is attending a newborn client (defined as up to 30 days of age by rule in 25 TAC, Chapter 37, Subchapter D, §37.52) or the person who is attending the delivery of a newborn client who is not attended by a physician to screen for the disorders within 24 to 48 hours of birth.

All infants must be tested a second time at 1 to 2 weeks of age. If there is any doubt that a client who is 12 months of age or younger was properly tested, the provider should submit a blood sample with the appropriate DSHS Form NBS3 to the DSHS Newborn Screening Laboratory.

### D.7 Abuse and Neglect

#### D.7.1 Requirements for Reporting Abuse or Neglect

Providers are required to report abuse or neglect as outlined in subsection 1.7, “Provider Responsibilities” in “Section 1: Provider Enrollment and Responsibilities” (Vol. 1, General Information).

Additionally, the General Appropriations Act, Article II, Rider 23 under DSHS, and Rider 13 under HHSC, of S.B. 1, 79th Legislative Regular Session, 2007, require that DSHS and HHSC distribute or provide appropriated funds only to recipients who show good faith efforts to comply with all child abuse and reporting requirements set forth in the Texas Family Code (TFC), Chapter 261, relating to investigations of reports of child abuse and neglect.

#### D.7.2 Procedures for Reporting Abuse or Neglect

Professionals, as defined in TFC §261.101 (b), are required to report abuse or neglect no later than the 48th hour after the hour in which the professional first has cause to believe the client has been or may be abused or is the victim of the offense of indecency with a child.

Nonprofessionals shall immediately make a report when the nonprofessional has cause to believe that the client’s physical or mental health or welfare has been adversely affected by abuse.

A report must be made regardless of whether the provider staff suspects that a report may have previously been made. Reports of abuse or indecency with a child should be made to one of the following:

- Texas Department of Family and Protective Services (DFPS), if the alleged or suspected abuse involves a person responsible for the care, custody, or welfare of the child (DFPS Texas Abuse Hotline, 1-800-252-5400, 24 hours a day, 7 days a week).
• Call the DFPS Texas Abuse Hotline if:
  • You believe your situation requires action in less than 24 hours.
  • You prefer to remain anonymous.
  • You have insufficient data to complete the required information on the report.
  • You do not want an email to confirm your report.

  **Note:** Providers can also report nonemergency abuse online at [www.txabusehotline.org](http://www.txabusehotline.org).

• Any local or state law enforcement agency or the state agency that operates, licenses, certifies, or registers the facility in which the alleged abuse or neglect occurred.

• The agency designated by the court to be responsible for the protection of children.

The law requires that the report include the following:

• Name and address of the minor, if known.

• Name and address of the minor’s parent or the person responsible for the care, custody, or welfare of the child if not the parent, if known.

• Any other pertinent information concerning the alleged or suspected abuse, if known.

A provider may not reveal whether the client has been tested or diagnosed with HIV or acquired immunodeficiency syndrome (AIDS). If the minor’s identity is unknown (e.g., the minor is at the provider’s office to receive testing for HIV or an STD anonymously), no report is required.

**D.7.2.1 Staff Training on Reporting Abuse and Neglect**

All providers shall develop training for all staff on the policies and procedures in regard to reporting child abuse, including sexual abuse and neglect. New staff shall receive this training as part of their initial training or orientation.

Training shall be documented. As part of the training, staff shall be informed that the staff person who conducts the screening and has cause to suspect abuse has occurred is legally responsible for reporting. A joint report may be made with the supervisor.

Several specific legislative requirements affect THSteps and the provider’s participation in Texas Medicaid. The legislation includes, but is not limited to those included in this appendix.
APPENDIX E. HEARING SCREENING INFORMATION

E.1 Texas Early Hearing Detection and Intervention (TEHDI) Process

The following processes for early hearing detection and intervention are addressed in this section:

- Birth screen
- Outpatient rescreen
- Evaluation using Texas Pediatric Protocol for Audiology
- Referral to an Early Childhood Intervention (ECI) program
- Periodic monitoring by the physician or medical home

Refer to: The 1-3-6 Month Practitioner’s Guide on the DSHS website.

E.1.1 Birth Screen

The hearing screen at birth will be either screening auditory brainstem response (ABR) or transient or distortion product otoacoustic emissions (OAE). The following items apply:

- A newborn’s hearing is screened at the birth facility. If a newborn does not pass the screen, hearing is rescreened before discharge.
- The birth facility reports results to the Department of State Health Services (DSHS) using the web-based eScreener Plus (eSP™) system.
- The newborn’s family and physician/medical home receive a written report of the hearing screen outcome.
- If a newborn passes the screen, the physician monitors hearing as part of well child checkups.
- If a newborn does not pass the second screen, a referral is made to a local resource who is experienced with the pediatric population for outpatient rescreen.

E.1.2 Outpatient Rescreen

If an outpatient rescreen is necessary, either ABR or OAE will be used. The following items apply:

- The physician/medical home receives the written report of results from the birth facility.
- The screener/physician reports results to the DSHS contractor, OZ Systems, using the web-based eSP™ system, by calling 1-866-427-5768 or faxing 1-817-385-3939.
- If the newborn passes the outpatient rescreen, the physician monitors hearing as part of well child checkups.
- If a newborn does not pass the outpatient rescreen, a referral is made to an audiologist for evaluation using the Texas Pediatric Protocol for Evaluation. Visit www.dshs.texas.gov/tehdi for more information.
- Hearing services for clients who are birth through 20 years of age are administered through the Texas Medicaid hearing services benefit. Clients may use the Online Provider Lookup (OPL) to locate a Texas Medicaid provider who provides hearing services for children (clients who are birth through 20 years of age).
E.1.3 Evaluation using Texas Pediatric Protocol for Audiology
These evaluations will include a diagnostic ABR and, if not previously done, a diagnostic OAE will be performed to determine cochlear involvement. The following items apply:

- Audiologists use equipment norms for newborns, preferably ones that they have collected on their equipment.
- Protocols include air and bone conduction testing using tone burst ABR, as well as click ABR, so the amplification may be appropriately fit.
- The physician/medical home receives results and makes the referral to ECI using the web-based eSP™ system or by using the ECI program search web page at https://citysearch.hhsc.state.tx.us.
- The physician/medical home monitors the child. See the American Academy of Pediatrics Position Statement at http://pediatrics.aappublications.org/cgi/content/full/113/Supplement_4/1545.
- The audiologist reports results to the DSHS contractor as noted above and makes the referral to ECI.
- Fitting of hearing aids by an audiologist when appropriate.
- Continued audiological assessment and monitoring as needed (usually monitor each three months for the first year of hearing aid use).

E.1.4 Referral to an ECI Program
The client will be referred to an ECI program by an audiologist or physician as soon as possible, but no longer than 7 days of identification of hearing loss as required by law. The following items apply:

- Service coordination provided by ECI.
- ECI will refer to the Local Education Agency (LEA) for auditory impairment (AI) services as outlined in the Memorandum of Understanding between TEA and HHS ECI.
- An evaluation and Individual Family Service Plan (IFSP) will occur within 45 days of referral to ECI.
- ECI services are available to clients birth through 35 months of age when determined by an IFSP.
- ECI and LEA will coordinate transition services upon the child’s third birthday.

E.1.5 Periodic Monitoring by the Physician or Medical Home
The physician/medical home will continue to monitor the client periodically and may consult or use the following:

- Providers may refer to the Joint Committee on Infant Hearing (JCIH) 2007 Position Statement for suggested monitoring protocols at http://pediatrics.aappublications.org/cgi/content/full/120/4/898.
- Deaf education and other special education services available from birth through 20 years of age when determined by an individualized education program.

E.2 JCIH 2007 Position Statement
The JCIH 2007 Position Statement is available on the JCIH website at www.jcih.org/posstatems.htm. The 2007 Position Statement lists the indicators that are associated with permanent congenital, delayed-onset or progressive hearing loss in childhood.
APPENDIX F. TEXAS HEALTH STEPS QUICK REFERENCE GUIDE
The Texas Health Steps Quick Reference Guide (THSteps-QRG) is available on the TMHP website.

APPENDIX G. AMERICAN ACADEMY OF PEDIATRIC DENTISTRY PERIODICITY GUIDELINES
The American Academy of Pediatric Dentistry (AAPD) periodicity guidelines are available on the AAPD website at www.aapd.org.
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1 General Information

This information is intended for Federally Qualified Health Centers (FQHCs) renal dialysis facilities, Rural Health Clinics (RHCs) and tuberculosis (TB) clinics. This handbook provides information about Texas Medicaid’s benefits, policies, and procedures applicable to these providers. This handbook contains information about Texas Medicaid fee-for-service benefits. For information about managed care benefits, refer to the Texas Medicaid Managed Care Handbook. Managed care carve-out services are administered as fee-for-service benefits. A list of all carve-out services is available in Subsection 17, “Carve-Out Services” in the Texas Medicaid Managed Care Handbook.

Important: All providers are required to read and comply with “Section 1: Provider Enrollment and Responsibilities” (Vol. 1, General Information). In addition to required compliance with all requirements specific to Texas Medicaid, it is a violation of Texas Medicaid rules when a provider fails to provide health-care services or items to Medicaid clients in accordance with accepted medical community standards and standards that govern occupations, as explained in Title 1 Texas Administrative Code (TAC) §371.1659. Accordingly, in addition to being subject to sanctions for failure to comply with the requirements that are specific to Texas Medicaid, providers can also be subject to Texas Medicaid sanctions for failure, at all times, to deliver health-care items and services to Medicaid clients in full accordance with all applicable licensure and certification requirements including, without limitation, those related to documentation and record maintenance.

1.1 National Drug Codes (NDC)

Refer to: Subsection 6.3.4, “National Drug Code (NDC)” in “Section 6: Claims Filing” (Vol. 1, General Information).

1.2 Revenue Codes for UB-04 Submissions

Claims that are submitted on the CMS-1450 UB-04 paper claim form or electronic equivalent by non-hospital facility or other non-hospital providers must be submitted with a revenue code for correct processing.

If the non-hospital provider is required to submit a procedure code for reimbursement, the provider must include the procedure code and an appropriate corresponding revenue code on the same detail, even if the chosen revenue code does not require a procedure code for claims processing.

Refer to: Subsection 4.5.5, “Outpatient Hospital Revenue Codes” in the Inpatient and Outpatient Hospital Services Handbook (Vol. 2, Provider Handbooks) for a list of revenue codes that do and do not require procedure codes.

1.3 Payment Window Reimbursement Guidelines for Services Preceding an Inpatient Admission

According to the three-day and one-day payment window reimbursement guidelines, most professional and outpatient diagnostic and nondiagnostic services that are rendered within the designated timeframe of an inpatient hospital stay and are related to the inpatient hospital admission will not be reimbursed separately from the inpatient hospital stay if the services are rendered by the hospital or an entity that is wholly owned or operated by the hospital.

These reimbursement guidelines do not apply in the following circumstances:

• The services were FQHC, RHC, THSteps, or some renal dialysis services.

• The hospital and the physician office or other entity are both owned by a third party, such as a health system.
• The hospital is not the sole or 100-percent owner of the entity.

Refer to: Subsection 3.7.4.14, “Payment Window Reimbursement Guidelines” in the Inpatient and Outpatient Hospital Services Handbook (Vol. 2, Provider Handbooks) for additional information about the payment window reimbursement guidelines.

2 Birthing Center

2.1 Provider Enrollment

A birthing center is a place, facility, or institution where a woman is scheduled to give birth following a normal, uncomplicated (low-risk) pregnancy. This term does not include a hospital, an ambulatory surgical center, or the residence of the woman giving birth.

A birthing center must be licensed as a birthing center by the Department of State Health Services (DSHS) and meet the minimum standards as required by the Texas Health and Safety Code, Chapter 244.010. To enroll in Texas Medicaid, a birthing center must be licensed to provide a level of service commensurate with the professional services of a doctor of medicine (MD), doctor of osteopathy (DO), certified nurse-midwife (CNM), or licensed midwife (LM) who acts as birth attendant. Texas Medicaid may reimburse birthing center providers only for those services that the attending physician or CNM determines to be reasonable and necessary for the care of the mother or newborn child.

Providers cannot be enrolled if their license is due to expire within 30 days. A current license must be submitted.

Birthing centers are encouraged to refer clients for Texas Health Steps (THSteps) services.

Refer to: “Section 1: Provider Enrollment and Responsibilities” (Vol. 1, General Information) for more information about enrollment procedures.

Section 2, “Medicaid Title XIX Family Planning Services” in the Gynecological, Obstetrics, and Family Planning Title XIX Services Handbook (Vol. 2, Provider Handbooks) for information on setting up referral procedures for family planning services.

The HHSC website at www.healthytexaswomen.org for information about family planning and the locations of family planning clinics receiving HHSC Family Planning Program funding from HHSC.


2.2 Services, Benefits, Limitations, and Prior Authorization

Birthing centers may only be reimbursed by Texas Medicaid for their facility labor and delivery services using the following procedure codes:

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<tr>
<th>Service</th>
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<td>Delivery</td>
<td>59409</td>
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<tr>
<td>Labor only</td>
<td>S4005</td>
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Note: Deliveries at a facility licensed as a birthing center by DSHS must be billed with procedure code 59409.

If the client is discharged prior to delivery, procedure code S4005 may be billed by the facility for labor services only.
Refer to: Subsection 9.2.34, “Immunization Guidelines and Administration” in the Medical and Nursing Specialists, Physicians, and Physician Assistants Handbook (Vol. 2, Provider Handbooks) for additional information about immunization administration.

2.2.1 Newborn Hearing Screening
The Texas Health and Safety Code, Chapter 47, requires birthing centers to offer all newborns a hearing screening as a part of the obstetrical care at delivery.

Refer to: Subsection 5.3.9, “Newborn Examination” in the Children’s Services Handbook (Vol. 2, Provider Handbooks) for more information about the newborn hearing screening.

Subsection 2.2.2.3, “Abnormal Hearing Screening Results” in the Vision and Hearing Services Handbook (Vol. 2, Provider Handbooks) for more information about abnormal hearing screens.

2.2.2 Newborn Eligibility Process
If the mother of the newborn is eligible for Medicaid, the newborn may be assigned his or her own Medicaid number. The birthing center must complete form GN.4, “Birthing Center Report (Newborn Child or Children) (Form 7484)” to provide information about each child born to a mother who is eligible for Medicaid.

Refer to: Hospital Report (Newborn Child or Children) (Form 7484) on the TMHP website at www.tmhp.com.

If the newborn’s name is known, the name must be on the form. The use of “Baby Boy” or “Baby Girl” delays the assignment of a number.

The form must be completed by the birthing center no later than five days after the child’s birth. Birthing centers that submit the birth certificate information using the HHSC, Vital Statistics Unit (VSU) Texas Electronic Registrar for Birth software and the HHSC Form 7484 receive a rapid and efficient assignment of a newborn Medicaid identification number. This process expedites reimbursement to hospitals and other providers that are involved in the care of the newborn.

Additional information about obtaining a newborn Medicaid identification number can be found on the agency website at https://hhs.texas.gov/services/health/medicaid-chip/provider-information/chip-perinatal-coverage/chip-perinatal-faqs. Providers may also call 1-888-963-7111, Ext. 7368 or 1-512-458-7368 for additional information or comments about this process.

Upon receipt of a completed 7484 form, DSHS verifies the mother’s eligibility and, within ten days of the receipt, sends notification letters to the hospital or birthing center, attending physician (if identified), mother, and caseworker. The notice includes the child’s Medicaid identification number and the effective date of coverage. After the child has been added to the eligibility file, HHSC issues a Medicaid Identification card (Your Texas Benefits Medicaid card) to the client.

The attending physician’s notification letter is sent to the address on file (by license number) at the Texas Medical Board. This address must be kept current to ensure timely notification. Physicians must submit address changes to the following address:

Texas Medical Board
Customer Information, MC-240
PO Box 2018
Austin, TX 78767-2018

2.2.3 Prior Authorization
Prior authorization is not required for services rendered in birthing centers.
2.2.4 Services Rendered in the Birthing Center Setting

Maternity clinic, physician, CNM, LM, nurse practitioner (NP), clinical nurse specialist (CNS), and physician assistant (PA) providers who render prenatal or family planning services in the birthing center setting must submit separate claims.

Refer to: Section 4, “Obstetric Services” in the Gynecological, Obstetrics, and Family Planning Title XIX Services Handbook (Vol. 2, Provider Handbooks) for information about birthing center providers.

2.3 Documentation Requirements

All services require documentation to support the medical necessity of the service rendered.

Birthing center services are subject to retrospective review and recoupment if documentation does not support the service billed.

2.4 Claims Filing and Reimbursement

2.4.1 Claims Information

Claims for birthing center services must be submitted to Texas Medicaid & Healthcare Partnership (TMHP) in an approved electronic format or on the CMS-1500 paper claim form. Providers may purchase CMS-1500 paper claim forms from the vendor of their choice. TMHP does not supply the forms.

When completing a CMS-1500 paper claim form, providers must include all required information on the claim, as TMHP does not key any information from attachments. Superbills or itemized statements are not accepted as claim supplements.

Refer to: “Section 3: TMHP Electronic Data Interchange (EDI)” (Vol. 1, General Information) for information on electronic claims submissions.

“Section 6: Claims Filing” (Vol. 1, General Information) for general information about claims filing.

Subsection 6.5, “CMS-1500 Paper Claim Filing Instructions” in “Section 6: Claims Filing” (Vol. 1, General Information) for instructions on completing paper claims. Blocks that are not referenced are not required for processing by TMHP and may be left blank.

2.4.2 Reimbursement

Birthing centers are reimbursed in accordance with 1 TAC §355.8181. See the applicable fee schedule on the TMHP website at www.tmhp.com. Texas Medicaid implemented mandated rate reductions for certain services.

Additional information about rate changes is available on the TMHP website at www.tmhp.com/resources/rate-and-code-updates/rate-changes.

2.4.2.1 National Correct Coding Initiative (NCCI) and Medically Unlikely Edit (MUE) Guidelines

The Healthcare Common Procedure Coding System (HCPCS) and Current Procedural Terminology (CPT) codes included in the Texas Medicaid Provider Procedures Manual are subject to National Correct Coding Initiative (NCCI) relationships, which supersede any exceptions to NCCI code relationships that may be noted in the manuals. The Centers for Medicare & Medicaid Services (CMS) NCCI and medically unlikely edits (MUE) guidelines can be found in the NCCI Policy and Medicaid Claims Processing manuals, which are available on the CMS NCCI web page. Providers should refer to the CMS NCCI web page for correct coding guidelines and specific applicable code combinations.
In instances when Texas Medicaid limitations are more restrictive than NCCI MUE guidance, Texas Medicaid limitations prevail.

3 Comprehensive Health Center (CHC)

CHCs or physician-operated clinics are funded by federal grants. To apply for participation in Texas Medicaid, they must be certified and participate as health centers under Medicare (Title XVIII).

CHC claims are paid according to each center’s encounter rates as established by CMS. Medicaid payments to CHCs are limited to Medicare deductible or coinsurance according to current guidelines. CHC providers that supply laboratory services in an office setting must comply with the rules and regulations for the Clinical Laboratory Improvement Amendments (CLIA). Providers that do not comply with CLIA are not reimbursed for laboratory services.

Refer to:
- Subsection 2.7, “Medicare Crossover Claim Reimbursement” in “Section 2: Texas Medicaid Fee-for-Service Reimbursement” (Vol. 1, General Information).
- “Section 1: Provider Enrollment and Responsibilities” (Vol. 1, General Information).
- Section 4, “Federally Qualified Health Center (FQHC)” in this handbook.
- Section 7, “Rural Health Clinic” in this handbook.

4 Federally Qualified Health Center (FQHC)

4.1 Enrollment

To enroll in Texas Medicaid, an FQHC must be receiving a grant under Section 329, 330, or 340 of the Public Health Service Act or designated by the U.S. Department of Health and Human Services (HHS) to have met the requirements to receive this grant. FQHCs and their satellites are required to enroll in Medicare to be eligible for Medicaid enrollment. The CMS has granted a waiver for the Medicare prerequisite at the time of initial enrollment of FQHC parents and satellites. FQHC look-alikes are not required to enroll in Medicare but may elect to do so to receive reimbursement for crossovers.

Refer to: Subsection 4.3.2.1, “Medicare Crossover Claims Pricing” in this handbook.

A copy of the Public Health Service’s Notice of Grant Award reflecting the project period and the current budget period must be submitted with the enrollment application. A current notice of grant award must be submitted to TMHP Provider Enrollment annually.

FQHCs are required to notify TMHP of all satellite centers that are affiliated with the parent FQHC and their actual physical addresses. All FQHC satellite centers billing Texas Medicaid for FQHC services must also be approved by the United States Department of Health and Human Services Health Resources and Services Administration (HRSA). For accounting purposes, centers may elect to enroll the HRSA-approved satellites using a Federally Qualified Satellite (FQS) provider identifier that ties back to the parent FQHC provider identifier and tax ID number (TIN). This procedure allows for the parent FQHC to have one provider agreement and one cost report that combines all costs from all approved satellites and the parent FQHC. If an approved satellite chooses to submit claims to Texas Medicaid directly, the center must have a provider identifier separate from the parent FQHC and will be required to file a separate cost report.

All providers are required to read and comply with “Section 1: Provider Enrollment and Responsibilities” (Vol. 1, General Information). In addition to required compliance with all requirements specific to Texas Medicaid, it is a violation of Texas Medicaid rules when a provider fails to provide health-care
services or items to Medicaid clients in accordance with accepted medical community standards and standards that govern occupations, as explained in 1 TAC §371.1659. Accordingly, in addition to being subject to sanctions for failure to comply with the requirements that are specific to Texas Medicaid, providers can also be subject to Texas Medicaid sanctions for failure, at all times, to deliver health-care items and services to Medicaid clients in full accordance with all applicable licensure and certification requirements including, without limitation, those related to documentation and record maintenance.

FQHC providers do not need to apply for a separate physician or agency number to provide family planning services.

Refer to: Subsection 1.1, “Provider Enrollment” in “Section 1: Provider Enrollment and Responsibilities” (Vol. 1, General Information) for more information about enrollment procedures.

FQHCs must identify and attest to all contractual agreements for those medical services in which the FQHC is receiving Prospective Payment System (PPS) reimbursement. This is a mandate from the 2012 to 2013 General Appropriations Act, H.B. 1, 82nd Legislature, Regular Session, 2011 (Article II, Health and Human Services Commission, Rider 78).

The attestation shall be made using the Federally Qualified Health Center Affiliation Affidavit, which is available on the TMHP website at www.tmhp.com.

4.1.1 Initial Cost Reporting

New FQHCs must file a projected cost report within 90 days of their designation as an FQHC to establish an initial payment rate. The cost report will contain the FQHC’s reasonable costs anticipated to be incurred during the FQHC’s initial fiscal year. The FQHC must file a cost report within five months of the end of the FQHC’s initial fiscal year. The cost settlement must be completed within 11 months of the receipt of a cost report. The cost per visit rate established by the cost settlement process will be the base rate. Any subsequent increases will be calculated as provided herein.

FQHC providers are required to submit a copy of their Medicare-audited cost report for the provider’s fiscal year within 30 days of receipt from Medicare to:

Texas Medicaid & Healthcare Partnership
Medicaid Audit
PO Box 200345
Austin, TX 78720-0345

A new FQHC location established by an existing FQHC participating in Texas Medicaid will receive the same effective rate as the FQHC establishing the new location. An FQHC establishing a new location may request an adjustment to its effective rate as provided herein if its costs have increased as a result of establishing a new location.

Refer to: Subsection 1.1, “Provider Enrollment” in “Section 1: Provider Enrollment and Responsibilities” (Vol. 1, General Information) for more information about enrollment procedures.

4.1.2 Services, Benefits, Limitations, and Prior Authorization

The services listed in the following tables may be reimbursed using the FQHC’s National Provider Identifier (NPI). Any additional physician services must be submitted for reimbursement using the physician’s Medicaid provider identifier. Hospital services are not considered for reimbursement to FQHC providers, and cannot be billed using the facility provider number assigned to the FQHC.

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<td>T1015</td>
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### FQHC Encounter Services

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Adult preventative care must be submitted with diagnosis codes Z0000, Z0001, Z01411, and Z01419.

### Case Management

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Comprehensive visit must be submitted using modifiers U2 and U5.
Follow-up face-to-face visit must be submitted using modifiers TS and U5.
Follow-up telephone visit must be submitted using modifier TS.

### Family Planning Services

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Annual family planning examination must be submitted with modifier FP.

### Mental Health Services

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* Procedures cannot be performed by Psychologist. Mental health services must be submitted using one of the appropriate modifiers AH, AJ, AM, U1, or U2.

### THSteps Dental Services

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<td>D7962</td>
</tr>
<tr>
<td>D7970</td>
<td>D7971</td>
<td>D7997</td>
<td>D7999</td>
<td>D8050</td>
<td>D8060</td>
<td>D8080</td>
<td>D8210</td>
<td>D8220</td>
<td>D8660</td>
</tr>
<tr>
<td>D8670</td>
<td>D8680</td>
<td>D8690</td>
<td>D9110</td>
<td>D9211</td>
<td>D9212</td>
<td>D9223</td>
<td>D9230</td>
<td>D9243</td>
<td>D9248</td>
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<tr>
<td>D9930</td>
<td>D9944</td>
<td>D9974</td>
<td>D9999</td>
<td></td>
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<td></td>
</tr>
</tbody>
</table>

Procedure codes D8210, D8220, and D8080 must be submitted with Diagnostic Procedure Code (DPC) remarks codes for correct claims processing.
**THSteps Medical Services**

<table>
<thead>
<tr>
<th>Procedure Code</th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td>99211</td>
<td>CNM services</td>
</tr>
<tr>
<td>99381</td>
<td>Clinical psychologist services</td>
</tr>
<tr>
<td>99382</td>
<td>Clinical social worker services; other mental health services</td>
</tr>
<tr>
<td>99383</td>
<td>Dental services</td>
</tr>
<tr>
<td>99384</td>
<td>CNM services</td>
</tr>
<tr>
<td>99385</td>
<td>Clinical psychologist services</td>
</tr>
<tr>
<td>99386</td>
<td>Clinical social worker services; other mental health services</td>
</tr>
<tr>
<td>99387</td>
<td>Dental services</td>
</tr>
<tr>
<td>99389</td>
<td>CNM services</td>
</tr>
<tr>
<td>99390</td>
<td>Clinical psychologist services</td>
</tr>
<tr>
<td>99391</td>
<td>Clinical social worker services; other mental health services</td>
</tr>
<tr>
<td>99392</td>
<td>Dental services</td>
</tr>
<tr>
<td>99393</td>
<td>CNM services</td>
</tr>
<tr>
<td>99394</td>
<td>Clinical psychologist services</td>
</tr>
<tr>
<td>99395</td>
<td>Clinical social worker services; other mental health services</td>
</tr>
<tr>
<td>99396</td>
<td>Dental services</td>
</tr>
<tr>
<td>96160</td>
<td>THSteps Medical services must be submitted using modifier EP in addition to one of the appropriate modifiers AM, SA, or U7.</td>
</tr>
<tr>
<td>96161</td>
<td>THSteps Medical services must be submitted using modifier EP in addition to one of the appropriate modifiers AM, SA, or U7.</td>
</tr>
</tbody>
</table>

**Note:** Procedure codes 96160 and 96161 are a benefit for Texas Medicaid clients who are 12 through 18 years of age and is limited to once per calendar year, any provider. Only one procedure code (96160 or 96161) may be reimbursed for the mental health screening per client per calendar year based on the description of the procedure code and the service rendered.

**Vision Care Services**

<table>
<thead>
<tr>
<th>Procedure Code</th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td>92002</td>
<td>Vision Care Services</td>
</tr>
<tr>
<td>92004</td>
<td>Vision Care Services</td>
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<tr>
<td>92012</td>
<td>Vision Care Services</td>
</tr>
<tr>
<td>92014</td>
<td>Vision Care Services</td>
</tr>
<tr>
<td>92015</td>
<td>Vision Care Services</td>
</tr>
<tr>
<td>92020</td>
<td>Vision Care Services</td>
</tr>
<tr>
<td>92025</td>
<td>Vision Care Services</td>
</tr>
<tr>
<td>92060</td>
<td>Vision Care Services</td>
</tr>
<tr>
<td>92065</td>
<td>Vision Care Services</td>
</tr>
<tr>
<td>92081</td>
<td>Vision Care Services</td>
</tr>
<tr>
<td>92082</td>
<td>Vision Care Services</td>
</tr>
<tr>
<td>92083</td>
<td>Vision Care Services</td>
</tr>
<tr>
<td>92100</td>
<td>Vision Care Services</td>
</tr>
<tr>
<td>92201</td>
<td>Vision Care Services</td>
</tr>
<tr>
<td>92202</td>
<td>Vision Care Services</td>
</tr>
<tr>
<td>92230</td>
<td>Vision Care Services</td>
</tr>
<tr>
<td>92235</td>
<td>Vision Care Services</td>
</tr>
<tr>
<td>92240</td>
<td>Vision Care Services</td>
</tr>
<tr>
<td>92242</td>
<td>Vision Care Services</td>
</tr>
<tr>
<td>92250</td>
<td>Vision Care Services</td>
</tr>
<tr>
<td>92260</td>
<td>Vision Care Services</td>
</tr>
<tr>
<td>92265</td>
<td>Vision Care Services</td>
</tr>
<tr>
<td>92270</td>
<td>Vision Care Services</td>
</tr>
<tr>
<td>92273</td>
<td>Vision Care Services</td>
</tr>
<tr>
<td>92274</td>
<td>Vision Care Services</td>
</tr>
<tr>
<td>92285</td>
<td>Vision Care Services</td>
</tr>
<tr>
<td>92286</td>
<td>Vision Care Services</td>
</tr>
<tr>
<td>92287</td>
<td>Vision Care Services</td>
</tr>
<tr>
<td>95060</td>
<td>Vision Care Services</td>
</tr>
<tr>
<td>95930</td>
<td>Vision Care Services</td>
</tr>
<tr>
<td>95933</td>
<td>Vision Care Services</td>
</tr>
<tr>
<td>95935</td>
<td>Vision Care Services</td>
</tr>
</tbody>
</table>

**Copayments**

<table>
<thead>
<tr>
<th>Copayment Code</th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td>CP001</td>
<td>Copayments</td>
</tr>
<tr>
<td>CP002</td>
<td>Copayments</td>
</tr>
<tr>
<td>CP003</td>
<td>Copayments</td>
</tr>
<tr>
<td>CP004</td>
<td>Copayments</td>
</tr>
<tr>
<td>CP005</td>
<td>Copayments</td>
</tr>
<tr>
<td>CP006</td>
<td>Copayments</td>
</tr>
<tr>
<td>CP007</td>
<td>Copayments</td>
</tr>
<tr>
<td>CP008</td>
<td>Copayments</td>
</tr>
</tbody>
</table>

**Referto:** Subsection 6.3.5, “Modifiers” in “Section 6: Claims Filing” (Vol. 1, General Information) for a definition of modifiers.

Section 4 *, “Texas Health Steps (THSteps) Dental” in the Children’s Services Handbook (Vol. 2, Provider Handbooks).

Section 5, “THSteps Medical” in the Children’s Services Handbook (Vol. 2, Provider Handbooks).


Subsection 9.2.56.3.2, “Preventive Care Visits” in the Medical and Nursing Specialists, Physicians, and Physician Assistants Handbook (Vol. 2, Provider Handbooks).

Subsection 8.8.2.2, “HMO Copayments” in “Section 8: Third Party Liability (TPL)” (Vol. 1, General Information) for information about HMO copayments.


Medicaid coverage is limited to FQHC services that are covered by Texas Medicaid and are reasonable and medically necessary. When furnished to a client of the FQHC, medically necessary services include the following:

- CNM services
- Clinical psychologist services
- Clinical social worker services; other mental health services
- Dental services
• NP services
• Other ambulatory services included in Medicaid such as family planning, THSteps, and maternity service clinic (MSC)
• PA services
• Physician services
• Services and supplies necessary for services that would be covered otherwise, if furnished by a physician or a physician service
• Vision care services
• Visiting nurse services to a homebound individual, in the case of those FQHCs located in areas with a shortage of home health agencies

Types of FQHC visits are defined in 1 TAC §355.8261. A visit is a face-to-face encounter between an FQHC client and a physician, PA, NP, CNM, visiting nurse, qualified clinical psychologist, clinical social worker, other health-care professional for mental health services, dentist, dental hygienist, or optometrist. Encounters that take place on the same day at a single location with more than one health-care professional or multiple encounters with the same health-care professional constitute a single visit, except where one of the following conditions exists:

• After the first encounter, the client suffers illness or injury requiring additional diagnosis or treatment.
• The FQHC client has a medical visit and an other health visit such as a qualified clinical psychologist, clinical social worker, other health professional for mental health services, a dentist, a dental hygienist, an optometrist, or a THSteps medical checkup.

All services provided that are incidental to the encounter, including developmental screening, must be included in the total charge for the encounter. They are not billable as a separate encounter.

Registered nurses may not be the sole provider of a medical checkup in an FQHC. If immunizations are given outside of a THSteps medical checkup, procedure codes given in the THSteps section of this manual should be used. These procedure codes are informational only, and are not payable.

To be reimbursed for Case Management for Children and Pregnant Women, an FQHC must be approved as a case management services provider by the DSHS Case Management Branch.

An annual family planning examination is allowed once per state fiscal year (September 1 through August 31), per client, per provider. An FQHC may be reimbursed for up to three family planning encounters per client, per year, regardless of the reason for the encounter. The three encounters may include any combination of general family planning encounters, an annual family planning examination, or intrauterine devices.

Family planning services must be submitted with the most appropriate evaluation and management (E/M) procedure code and one of the following family planning diagnosis codes:

<table>
<thead>
<tr>
<th>Family Planning Diagnosis Codes</th>
</tr>
</thead>
<tbody>
<tr>
<td>Z30011</td>
</tr>
<tr>
<td>Z3041</td>
</tr>
<tr>
<td>Z309</td>
</tr>
</tbody>
</table>

Procedure code 58300 must be submitted on the same claim as J7296, J7297, J7298, J7300, and J7301. Procedure code 58300 will process as informational only. Only the annual family planning examination requires modifier FP. All other family planning visits do not require the FP modifier. Claims filed incorrectly may be denied.
Laboratory and radiology services or the services of a licensed vocational nurse (LVN), registered nurse (RN), nutritionist, or dietitian are not considered an encounter, because they are incidental to an encounter with one of the previously-mentioned payable health-care professionals. Providers should continue to include the cost associated with these services on their cost report (they are allowable but do not constitute an encounter).

Per federal regulations, the provider cannot submit claims to Medicaid or bill the client for vaccines obtained from the Texas Vaccine for Children (TVFC) Program.

Refer to: Section 5, “THSteps Medical” in the Children’s Services Handbook (Vol. 2, Provider Handbooks).

4.1.3 After-Hours Care

After-hours care for FQHCs is defined as care provided on weekends, on federal holidays, or before 8 a.m. and after 5 p.m., Monday through Friday. After-hours care provided by FQHCs does not require a referral.

4.1.4 Prior Authorization

Prior authorization or authorization may be required for FQHC services. Refer to the individual sections referenced in subsection 4.1.2, “Services, Benefits, Limitations, and Prior Authorization” in this handbook.

4.1.5 Referral Requirements

Texas Medicaid fee-for-service limited clients, are allowed to choose any enrolled family planning provider.

4.2 Documentation Requirements

All services require documentation to support the medical necessity of the service rendered. All services provided are subject to retrospective review and recoupment if documentation does not support the service that was submitted for reimbursement.

4.3 Claims Filing and Reimbursement

4.3.1 Claims Information

All services (except for family planning, THSteps medical, THSteps dental, copayments, vision, mental health services, and case management for high-risk pregnant women and infants) provided during an encounter must be submitted for reimbursement using procedure code T1015. All services provided that are incidental to the encounter must be included in the total charge for the encounter and are not billable as a separate encounter. For example, if an office visit was provided at a charge of $30 and a lab test for $15, the center would submit a claim to TMHP for procedure code T1015 for $45 and would be reimbursed at the center’s encounter rate.

All providers of laboratory services must comply with the rules and regulations of CLIA. Providers who do not comply with CLIA are not reimbursed for laboratory services.

Refer to: Subsection 2.1.1, “Clinical Laboratory Improvement Amendments (CLIA)” in the Radiology and Laboratory Services Handbook (Vol. 2, Provider Handbooks).

To obtain the encounter rate when submitting claims for family planning services that are provided under Title XIX or HTW, FQHCs must use the most appropriate E/M procedure code, or procedure code J7296, J7297, J7298, J7300, and J7301, or J7307 with a family planning diagnosis code. Providers must use procedure code J7296, J7297, J7298, J7300, J7301, or J7307 if the visit is for the insertion of an intrauterine device (IUD) or implantable contraceptive capsule. These procedure codes must be submitted in conjunction with the most appropriate informational procedure codes for services that were rendered. Procedure codes J7296, J7297, J7298, J7300, J7301, and J7307 may be reimbursed in
addition to the FQHC encounter payment. When seeking reimbursement for an IUD or implantable contraceptive capsule, providers must submit on the same claim the procedure code for the family planning service provided and the procedure code for the contraceptive device. The contraceptive device is not subject to FQHC limitations. Providers must use modifier U8 when submitting claims for a contraceptive device purchased through the 340B Drug Pricing Program. Providers must use modifier FP only to submit claims for the annual family planning examination.

If an employed physician of an FQHC provides a service in the hospital (e.g., a delivery), the service may be billed using the physician provider number if the terms of the FQHC and physician agreement indicate this occurrence. Physicians must be enrolled in Medicaid separately from the FQHC facility. Physicians are not allowed to bill through their FQHC group number for hospital services. The services will be reimbursed at the physician fee-for-service (FFS) fee schedule rate. The costs that are associated with these physician services must be excluded from the FQHC’s cost report and will not be considered during the FQHC cost settlement or encounter rate setting process.

Services rendered in the (inpatient or outpatient) hospital setting are not considered a reimbursable FQHC encounter and are not payable to the FQHC. FQHC services for clients who have only Medicaid must be submitted to TMHP in approved electronic format or on a UB-04 CMS-1450, CMS-1500, or 2017 paper claim form. Providers may purchase UB-04 CMS-1450 or CMS-1500 paper claim forms from the vendor of their choice. TMHP does not supply the forms. When completing a UB-04 CMS-1450 or CMS-1500 paper claim form, all required information must be included on the claim, as TMHP does not key any information from claim attachments. Superbills, or itemized statements, are not accepted as claim supplements.

The ADA Dental Claim Form can be downloaded at www.ada.org/7119.aspx.

The 2017 Claim Form can be found in the Forms section of this manual.

**Refer to:** 2017 Claim Form on the TMHP website at www.tmhp.com.

“Section 3: TMHP Electronic Data Interchange (EDI)” (Vol. 1, General Information) for information on electronic claims submissions.

“Section 6: Claims Filing” (Vol. 1, General Information) for general information about claims filing.

Section 11, “Forms” in this handbook.

Claims must be filed as follows:

<table>
<thead>
<tr>
<th>Services</th>
<th>Claim Form</th>
</tr>
</thead>
<tbody>
<tr>
<td>THSteps medical services</td>
<td>UB-04 CMS-1450 or CMS-1500 paper claim form or approved electronic format</td>
</tr>
<tr>
<td>Family planning claims filed by FQHC providers who have contracted with HHSC</td>
<td>2017 Claim Form or approved electronic format</td>
</tr>
<tr>
<td>Family planning claims filed by FQHC providers not contracted with HHSC</td>
<td>UB-04 CMS-1450 or 2017 paper claim form or approved electronic format</td>
</tr>
<tr>
<td>THSteps dental services</td>
<td>American Dental Association (ADA) Dental Claim Form or approved electronic format</td>
</tr>
<tr>
<td>Case Management for Children and Pregnant Women services</td>
<td>UB-04 CMS-1450 or CMS-1500 paper claim form or approved electronic format</td>
</tr>
</tbody>
</table>

When filing for a client who has Medicare and Medicaid coverage, providers must file on the same claim form that was filed with Medicare.
Services provided by a health-care professional require one of the following modifiers with procedure code T1015, to designate the health-care professional providing the services: AH, AJ, AM, SA, TD, TE, U1, U2, or U7.

- If more than one health-care professional is seen during the encounter, the modifier must indicate the primary contact. The primary contact is defined as the health-care professional who spends the greatest amount of time with the client during that encounter.
- If the encounter is for antepartum care or postpartum care, the modifier TH must be indicated on the claim in addition to any other appropriate modifier.
- If the antepartum or postpartum care is provided by a CNM, the modifier SA must be indicated on the claim in addition to any other appropriate modifiers.

Use modifier TD or TE for home health services provided in areas with a shortage of home health agencies.

Refer to:
“Section 3: TMHP Electronic Data Interchange (EDI)” (Vol. 1, General Information) for information on electronic claims submissions.
“Section 6: Claims Filing” (Vol. 1, General Information) for general information about claims filing.
The Claim Form Examples page of the TMHP website at www.tmhp.com.

4.3.2 Reimbursement
FQHCs are reimbursed provider-specific prospective payment system encounter rates in accordance with 1 TAC §355.8261.
FQHCs are exempt from the mandated rate reductions except for HHSC Family Planning services.
Texas Medicaid implemented mandated rate reductions for certain services. Additional information about rate changes is available on the TMHP website at www.tmhp.com/resources/rate-and-code-updates/rate-changes.

Refer to: Subsection 2.2, “Fee-for-Service Reimbursement Methodology” in “Section 2: Texas Medicaid Fee-for-Service Reimbursement” (Vol. 1, General Information) for more information about reimbursement.

Subsection 2.3, “Reimbursement Reductions” in “Section 2: Texas Medicaid Fee-for-Service Reimbursement” (Vol. 1, General Information) for more information about reimbursement.

4.3.2.1 Medicare Crossover Claims Pricing
For Medicare Part B cost sharing obligations, all deductible obligations will be reimbursed at 100 percent of the deductible amount owed, even if the cost sharing comparison results in a lower payment. For all other cost sharing obligations (including Medicare Part A, B, and C), the cost sharing comparison is performed according to current guidelines.
For FQHC Medicare crossover claims, Texas Medicaid will reimburse the lesser of the following:
- The coinsurance and full deductible payment.
- The amount remaining after the Medicare payment amount is subtracted from the allowed Medicaid fee or encounter rate for the service. If this amount is less than the deductible, then the full deductible is reimbursed instead.
If the Medicare payment is equal to, or exceeds the Medicaid allowed amount or encounter payment for the service, Texas Medicaid will not make a payment for coinsurance.

The client has no liability for any balance or Medicare coinsurance and deductible related to Medicaid-covered services.

**Refereto:** Subsection 2.7, “Medicare Crossover Claim Reimbursement” in “Section 2: Texas Medicaid Fee-for-Service Reimbursement” (Vol. 1, General Information).

**4.3.2.2 NCCI and MUE Guidelines**

The HCPCS and CPT codes included in the Texas Medicaid Provider Procedures Manual are subject to NCCI relationships, which supersede any exceptions to NCCI code relationships that may be noted in the manuals. Providers should refer to the CMS NCCI web page for correct coding guidelines and specific applicable code combinations.

In instances when Texas Medicaid limitations are more restrictive than NCCI MUE guidance, Texas Medicaid limitations prevail.

**5 Maternity Service Clinic (MSC)**

MSCs are limited provider clinics that are unrelated to a hospital and that only provide maternity services. An MSC will be reimbursed for antepartum care and/or postpartum care visits only. Hemoglobin, hematocrit and urinalysis procedures are included in the charge for antepartum care and not separately reimbursed. Services other than antepartum and postpartum care visits will be denied.


**6 Renal Dialysis Facility**

**6.1 Enrollment**

To enroll in Texas Medicaid, a renal dialysis facility must be Medicare-certified in the state where it is located. Facilities must also adhere to the appropriate rules, licensing, and regulations of the state where they operate.

**Refereto:** Subsection 1.1, “Provider Enrollment” in “Section 1: Provider Enrollment and Responsibilities” (Vol. 1, General Information) for more information.

**6.2 Services, Benefits, Limitations, and Prior Authorization**

Renal dialysis is a benefit of Texas Medicaid for the following acute renal failure or end stage renal disease (ESRD) diagnosis codes:

<table>
<thead>
<tr>
<th>Diagnosis Codes</th>
</tr>
</thead>
<tbody>
<tr>
<td>N170</td>
</tr>
<tr>
<td>N1831</td>
</tr>
</tbody>
</table>

All of the services, except for ultrafiltration (revenue code B-881), are diagnosis-restricted to the diagnoses in the above table.

Subsection 3.2.5, “Organ and Tissue Transplant Services” in the Inpatient and Outpatient Hospital Services Handbook (Vol. 2, Provider Handbooks) for information on organ transplant and facility services.

Dialysis treatments are a benefit for clients in an inpatient or outpatient hospital or a renal dialysis facility according to the guidelines for outpatient maintenance dialysis approved through CMS. Dialysis treatments may also be a benefit in the client’s home. Outpatient dialysis includes:

- Staff-assisted dialysis performed by the staff of the center or facility.
- Self-dialysis performed by a client with little or no professional assistance (the client must have completed an appropriate course of training).
- Home dialysis performed by an appropriately trained client (and the client’s caregiver) at home.
- Dialysis furnished in a facility on an outpatient basis at an approved renal dialysis facility.

6.2.1 Physician Supervision

Physician reimbursement for supervision of ESRD clients on dialysis is based on a monthly capitation payment (MCP) that is calculated by Medicare. The MCP is a comprehensive payment that covers all of the physician services that are associated with the continuing medical management of a maintenance dialysis client for treatments received in the facility. An original onset date of dialysis treatment must be included on claims for all renal dialysis procedures in all places of service except inpatient hospital.

Physician supervision of outpatient ESRD dialysis includes services that are rendered by the attending physician in the course of office visits during which any of the following occur:

- The routine monitoring of dialysis
- The treatment or follow-up of complications of dialysis, including:
  - The evaluation of related diagnostic tests and procedures
  - Services that are involved in the prescription of therapy for illnesses that are unrelated to renal disease, if the treatment occurs without increasing the number of physician-client contacts

The following physician services are a benefit for physician supervision of outpatient ESRD dialysis services:

<table>
<thead>
<tr>
<th>Procedure Codes</th>
</tr>
</thead>
<tbody>
<tr>
<td>90951 90952 90953 90954 90955 90956 90957 90958 90959 90960</td>
</tr>
<tr>
<td>90961 90962 90963 90964 90965 90966 90967 90968 90969 90970</td>
</tr>
</tbody>
</table>

Procedure codes 90935, 90937, 90945, and 90947 are a benefit for:

- ESRD or non-ERSD services in the inpatient setting when the physician is present during dialysis treatment. The physician must be physically present and involved during the course of dialysis. These codes are not payable for a cursory visit by the physician. Hospital visit procedure codes must be used for a cursory visit.
- Non-ERSD services when provided by a physician, nurse practitioner, clinical nurse specialist, or physician assistant in an office or outpatient setting.

Only one of the following procedure codes 90935, 90937, 90945, or 90947 may be reimbursed per day by any provider.
If the physician sees the client only when the client is not dialyzing, the physician must submit the appropriate hospital visit procedure code. The inpatient dialysis procedure code must not be submitted for payment.

Providers must use one of the following procedure codes to submit claims for services when the client:

- Is not on home dialysis.
- Has had a complete assessment visit during the calendar month.
- Has received a full month of ESRD related services

<table>
<thead>
<tr>
<th>Procedure Codes</th>
</tr>
</thead>
<tbody>
<tr>
<td>90951</td>
</tr>
<tr>
<td>90961</td>
</tr>
</tbody>
</table>

When a full calendar month of ESRD-related services are submitted for clients on home dialysis, providers must use procedure code 90963, 90964, 90965, or 90966.

Providers must submit claims with procedure code 90967, 90968, 90969, or 90970 if ESRD-related services are provided for less than a full month, per day, under the following conditions:

- Partial month during which a client who is not on home dialysis received one or more face-to-face visits but did not receive a complete assessment.
- A client who is on home dialysis received less than a full month of services.
- Transient client.
- Client was hospitalized during a month of services before a complete assessment could be performed.
- Dialysis was stopped due to recovery or death of a client.
- Client received a kidney transplant.

Procedure codes 90967, 90968, 90969, and 90970 are limited to one per day by any provider. When submitting claims for these procedure codes, providers must indicate the dates of service on which supervision was provided.

Procedure codes 90967, 90968, 90969, and 90970 will be denied if they are submitted with dates of service in the same calendar month by any provider as the following procedure codes:

<table>
<thead>
<tr>
<th>Procedure Codes</th>
</tr>
</thead>
<tbody>
<tr>
<td>90951</td>
</tr>
<tr>
<td>90961</td>
</tr>
</tbody>
</table>

Only one of the procedure codes in the previous table will be reimbursed per calendar month by any provider.

The following services may be provided in conjunction with physician supervision of outpatient ESRD dialysis but are considered nonroutine and may be submitted for reimbursement separately.

- Declotting of shunts when performed by the physician.
- Physician services to inpatients.

If one of the following occurs:

- A client is hospitalized during a calendar month of ESRD-related services before a complete assessment is performed.
• The client receives one or more face-to-face assessments, but the timing of inpatient admission prevents the client from receiving a complete assessment.

Then the physician must submit both of the following:

• Procedure code 90967, 90968, 90969, or 90970 for each date of outpatient supervision.

• The appropriate hospital evaluation and management code for individual services provided on the days during which the client was hospitalized.

If a client has a complete assessment in the month during which the client is hospitalized, one of the following procedure codes must be submitted for the month of supervision:

<table>
<thead>
<tr>
<th>Procedure Codes</th>
</tr>
</thead>
<tbody>
<tr>
<td>90951</td>
</tr>
<tr>
<td>90961</td>
</tr>
</tbody>
</table>

The appropriate inpatient evaluation and management codes must be reported for procedures provided during the hospitalization.

• Dialysis at an outpatient facility other than the usual dialysis setting for a client of a physician who bills the MCP. The physician must submit procedure code 90967, 90968, 90969, or 90970 for each date on which supervision is provided. The physician may not submit claims for days that the client dialyzed elsewhere.

• Physician services beyond those that are related to the treatment of the client’s renal condition that cause the number of physician-client contacts to increase. Physicians may submit claims on a fee-for-service basis if they supply documentation on the claim that the illness is not related to the renal condition and that additional visits are required.

Inpatient services that are provided to a hospitalized client for whom the physician has agreed to submit monthly claims, may be reimbursed in one of the following ways:

• The physician may elect to continue monthly billing, in which case the physician may not bill for individual services that were provided to the hospitalized client.

• The physician may reduce the monthly amount submitted by 1/30th for each day of hospitalization and may charge fees for individual services that were provided on the hospitalized days.

• The physician may submit a claim for inpatient dialysis services using the inpatient dialysis procedure codes. The physician must be present and involved with the client during the course of the dialysis.

Clients may receive dialysis at an outpatient facility other than the client’s usual dialysis setting, even if their physician bills for monthly dialysis coordination. The physician must reduce the monthly amount submitted for reimbursement by 1/30th for each day the client is dialyzed elsewhere.

Physician services beyond those related to the treatment of the client’s renal condition may be reimbursed on a fee-for-service basis. The physician must provide medical documentation with the claim that identifies how the illness is not related to the renal condition and added visits are required.

Payment is made for physician training services in addition to the MCP for physician supervision rendered to maintenance facility clients.

6.2.1.1 Unscheduled or Emergency Dialysis in a Non-Certified ESRD Facility

For some medical situations in which ESRD clients cannot obtain their regularly scheduled dialysis treatment at a certified ESRD facility, Texas Medicaid will allow for non-routine dialysis treatments furnished in the outpatient department of a hospital that does not have a certified dialysis facility.
Unscheduled dialysis for clients may be a benefit for one of the following reasons:

- Dialysis was performed following, or in connection with, a vascular access procedure.
- Dialysis was performed following treatment for an unrelated medical emergency (e.g., a client goes to the emergency room and, as a result, misses a regularly scheduled dialysis treatment that cannot be rescheduled).
- Emergency dialysis was performed for clients who would otherwise have to be admitted as inpatient in order for the hospital to receive payment.

Providers must submit claims using procedure code G0257 with revenue code 880 in order to receive payment for unscheduled outpatient dialysis.

Procedure code G0257 is only reimbursed to clients with ESRD and must be billed with revenue code 880 on the same claim. If procedure code G0257 is not on the same claim as revenue code 880, it will be denied.

Procedure code G0257 is limited to diagnosis codes N185 and N186 and is limited to one service per day, any provider.

Erythropoietin (procedure code Q4081) may be billed separately and must be billed with revenue code 634 or 635 on the same claim.

Texas Medicaid will provide a single payment to reimburse unscheduled or emergency dialysis treatments furnished to ESRD clients in the outpatient department of a hospital that does not have a certified ESRD facility.

Reimbursement for procedure code G0257 is limited to the same services included in the Method 1 composite. Providers will not be reimbursed for individual services related to dialysis.

Repeated billing of this service by the same provider for the same clients may indicate routine dialysis treatments are being performed and providers will be subject to recoupment upon medical record review.

Reimbursement of other outpatient hospital services are only reimbursed if they are not related to the dialysis services and are determined to be medically necessary with supporting documentation.

### 6.2.2 Renal Dialysis Facilities-Method I Composite Rate

The composite rate includes all necessary equipment, supplies, and services for the client receiving dialysis whether in the home or in a facility. The facility’s charge must not include the charge for the physician’s routine supervision. Examples of services included in the composite rate include, but are not limited to:

- Cardiac monitoring—procedure code 93040 or 93041.
- Catheter changes—procedure code 36000 or 49421.
- Crash cart usage for cardiac arrest.
- Declotting of shunt (procedure code 36593) and any supplies used to declot shunts performed by facility staff in the dialysis unit.
- Dialysate—procedure code A4720, A4722, A4723, A4724, A4725, A4726, or A4765.
- Oxygen—procedure code E0424, E0431, E0434, E0439, E0441, E0442, E0443, E0444, or E0447.
- Routine laboratory services for dialysis.

**Note:** When one of these laboratory services is required more frequently, renal dialysis facility providers must submit the appropriate procedure code with modifier 91 for separate reimbursement.
• Staff time to administer blood, separately billable drugs, and blood collection for laboratory—procedure code 36430 or 36591.

• Suture removal or dressing changes.

• Certain drugs such as those to elevate or decrease blood pressure, antiarrhythmics, blood thinners or expanders, antihistamines or antibiotics to treat infections or peritonitis related to peritoneal dialysis are included in the composite rate. Examples include, but are not limited to:
  • Hydralazine—procedure code J0360
  • Diphenhenhydramine—procedure code J1200
  • Heparin—procedure code J1642 or J1644
  • Dopamine—procedure code J1265
  • Etelcalcetide—procedure code J0606
  • Ferric pyrophosphate citrate solution—procedure code J1443
  • Glucose
  • Propranolol—procedure code J1800
  • Insulin
  • Digoxin—procedure code J1160
  • Norepinephrine bitartrate
  • Mannitol—procedure code J2150
  • Procaine
  • Protamine—procedure code J2720
  • Saline—procedure code A4216 or A4217
  • Hydrocortisone sodium succinate—procedure code J1720
  • Verapamil

Medically necessary drugs that are not included in the composite rate may be separately reimbursed when provided by and administered in the dialysis facility by facility staff. Staff time and supplies used to administer the drugs are included in the composite rate. Examples include, but are not limited to, the following:

• Antibiotics, except when prescribed for clients to treat infections or peritonitis related to peritoneal dialysis
• Hematinics
• Anabolics
• Muscle relaxants
• Analgesics
• Sedatives
• Tranquilizers
• Erythropoietin
• Thrombolytics used to declot central venous catheters
• Intravenous levocarnitine (procedure code J1955), for ESRD clients who have been on dialysis for a minimum of three months with one of the following indications (All other indications for levocarnitine are not covered.):

• Carnitine deficiency, defined as a plasma free carnitine level less than 40 micromoles per liter.
• Signs and symptoms of erythropoietin-resistant anemia that has not responded to standard erythropoietin with iron replacement, and for which other causes have been investigated and adequately treated.
• Hypotension on hemodialysis that interferes with delivery of the intended dialysis despite application of usual measures deemed appropriate (e.g., fluid management) (such episodes of hypotension must have occurred during at least two dialysis treatments in a 30-day period).

Note: Continued use of levocarnitine is not covered if improvement has not been demonstrated within six months of the initiation of treatment.

The ordering physician must maintain documentation in the client’s medical record to support medical necessity.

6.2.3 Method II Dealing Direct-Support Services

With Method II, the client selects and works with a single supplier to obtain supplies and equipment to dialyze at home. The selected supplier cannot be a dialysis facility, although the supplier must maintain a written agreement with a support dialysis facility to provide backup and support services. Method II support services are reimbursed under revenue codes 845 and 855.

Support services reimbursed monthly under Method II are limited to clients who are 20 years of age and younger, and include, but are not limited to:

• Periodic monitoring of a client’s adaptation to home dialysis and performance of dialysis, including provisions for visits to the home or the facility.
• Visits by trained personnel for the client with a qualified social worker and a qualified dietitian, made in accordance with a plan prepared and periodically reviewed by a professional team, which includes the physician.
• Individual unscheduled visits to a facility made on an as-needed basis; (e.g., assistance with difficult access situations).
• ESRD-related laboratory tests covered under the composite rate.
• Providing, installing, repairing, testing, and maintaining home dialysis equipment, including appropriate water testing and treatment.
• Ordering of supplies on an ongoing basis.
• A record keeping system that assures continuity of care.
• Support services specifically applicable to chronic ambulatory peritoneal dialysis (CAPD) also include, but are not limited to:
  • Changing the connecting tube and administration set.
  • Monitoring the client’s performance of CAPD, assuring that it is done correctly, and reviewing proper techniques with the client or informing the client of modifications to apparatus or technique.
  • Documenting whether the client has or has had peritonitis that requires physician intervention or hospitalization (unless there is evidence of peritonitis, a culture for peritonitis is not necessary).
  • Inspecting the catheter site.
Routine laboratory services are included in the support services and are not reimbursed separately.

Equipment and supplies are:
- Reimbursed under Method II to only one provider per month who must agree to submit claims once per month for only one month’s quantity per claim.
- Limited to clients who are 20 years of age and younger.
- Reimbursed separately up to the total monthly allowable as determined by HHSC.

The following equipment, supply, and services procedure codes are benefits of Texas Medicaid under Method II:

<table>
<thead>
<tr>
<th>Procedure Codes</th>
</tr>
</thead>
<tbody>
<tr>
<td>36000 36430 36591 36593 49421 93040 93041 A4216 A4217 A4651</td>
</tr>
<tr>
<td>A4652 A4657 A4660 A4663 A4670 A4680 A4690 A4706 A4707 A4708</td>
</tr>
<tr>
<td>A4709 A4714 A4719 A4720 A4721 A4722 A4723 A4724 A4725 A4726</td>
</tr>
<tr>
<td>A4730 A4736 A4737 A4740 A4750 A4755 A4760 A4765 A4766 A4772</td>
</tr>
<tr>
<td>A4773 A4774 A4802 A4860 A4911 A4913 A4918 A4927 A4928 A4929</td>
</tr>
<tr>
<td>A4930 A4931 A4932 E0424 E0431 E0434 E0439 E0441 E0442 E0443</td>
</tr>
<tr>
<td>E0444 E1510 E1520 E1530 E1540 E1550 E1560 E1570 E1575 E1580</td>
</tr>
<tr>
<td>E1590 E1592 E1594 E1600 E1620 E1630 E1632 E1635 E1637 E1639</td>
</tr>
<tr>
<td>J2150 J2720</td>
</tr>
</tbody>
</table>

Installation and repair of home hemodialysis machines are not a benefit of Texas Medicaid. Home modifications for use of medical equipment are not a benefit of Texas Medicaid.

A Medicaid client may receive CAPD and continuous cycle peritoneal dialysis (CCPD) support services furnished by the facility on a monthly basis. Charges for support services in excess of this frequency must include documentation of medical necessity.

Clients may have a one month reserve of supplies available for use. Renal dialysis services beyond these limitations may be considered for clients who are 20 years of age and younger through the Comprehensive Care Program (CCP) with prior authorization.

### 6.2.4 Facility Revenue Codes

The following services are a benefit for renal dialysis centers billing under reimbursement methodology I composite rate or II dealing direct:

<table>
<thead>
<tr>
<th>Service</th>
<th>Revenue Code</th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td>Maintenance</td>
<td>821</td>
<td>Hemodialysis (outpatient/home)–composite or other rate</td>
</tr>
<tr>
<td></td>
<td>831</td>
<td>Peritoneal Dialysis (outpatient/home)–composite or other rate</td>
</tr>
<tr>
<td></td>
<td>841</td>
<td>CAPD (outpatient/home)–composite or other rate</td>
</tr>
<tr>
<td></td>
<td>851</td>
<td>CCPD (outpatient/home)–composite or other rate</td>
</tr>
<tr>
<td>Training</td>
<td>829</td>
<td>Hemodialysis (outpatient/home)–other</td>
</tr>
<tr>
<td></td>
<td>839</td>
<td>Peritoneal Dialysis (outpatient/home)–other</td>
</tr>
<tr>
<td></td>
<td>849</td>
<td>CAPD (outpatient/home)–other</td>
</tr>
<tr>
<td></td>
<td>859</td>
<td>CCPD (outpatient/home)–other</td>
</tr>
</tbody>
</table>
Renal dialysis facilities should not use a HCPCS/CPT code when submitting a claim with a revenue code. Method II is limited to clients who are 20 years of age or younger.

The facility charge must not include the charge for the physician’s routine supervision.

### 6.2.5 Training for Hemodialysis, Intermittent Peritoneal Dialysis (IPD), Continuous Cycle Peritoneal Dialysis (CCPD), and Chronic Ambulatory Peritoneal Dialysis (CAPD)

Most self-dialysis training for hemodialysis, IPD, CCPD, and CAPD is provided in an outpatient setting. Dialysis training provided in an inpatient setting will be reimbursed at the same rate as the facility’s outpatient training rate.

Reimbursement for hemodialysis, IPD, CCPD, and CAPD training services and supplies provided by the dialysis facility includes personnel services, parenteral items routinely used in dialysis, training manuals and materials, and routine dialysis laboratory tests.

No frequency limitation is applied to routine laboratory tests during the training period because these tests commonly are given during each day of training. Nonroutine laboratory tests performed during the training period may be reimbursed when documentation of medical necessity is submitted with the claim.

It may be necessary to supplement the patient’s dialysis during CAPD training with intermittent peritoneal dialysis or hemodialysis because the client has not mastered the CAPD technique.

Training is limited to once per day. The composite rate will be denied as part of dialysis training when submitted for the same date of service.

### 6.2.6 Maintenance Hemodialysis

The facility composite rate applies when a chronic renal dialysis client receives hemodialysis in an approved renal dialysis facility. Reimbursement is based on the facility’s per-treatment composite rate, as calculated by Medicare. Services included in the facility’s charge are routine laboratory tests, personnel services, equipment, supplies, and other services associated with the treatment.

For hospitals to be reimbursed for maintenance hemodialysis, they must be enrolled as an approved dialysis facility with the appropriate provider identifier. When a client is admitted for hospitalization for no reason other than to receive maintenance renal dialysis, the dialysis services are considered outpatient services and are covered if the hospital has been designated as a CMS certified renal dialysis center.

### 6.2.7 Maintenance IPD

Maintenance IPD is usually performed in sessions of 10 to 12 hours duration, three times per week. It may also be performed in fewer sessions that are longer in duration. If more than three sessions occur in one week, the provider must supply documentation of medical necessity with the claim.

### 6.2.8 Maintenance CAPD and CCPD

Support services for maintenance furnished to clients receiving CAPD or CCPD in the home may be reimbursed to dialysis facilities. Home dialysis support services must be furnished by the facility in either the home or the facility. CAPD and CCPD support services are limited to once per day.
6.2.9 Laboratory and Radiology Services

All providers of laboratory services must comply with the rules and regulations of CLIA. Providers who do not comply with CLIA will not be reimbursed for laboratory services.

Refer to: Subsection 2.1.1, “Clinical Laboratory Improvement Amendments (CLIA)” in the Radiology and Laboratory Services Handbook (Vol. 2, Provider Handbooks).

6.2.9.1 In-Facility Dialysis—Routine Laboratory

Laboratory testing may be obtained and processed in the renal dialysis facility or by an outside laboratory. Charges for routine laboratory tests performed according to the established frequencies in the following tables are included in the facility’s composite rate submitted to Texas Medicaid regardless of where tests were processed. If the routine laboratory testing is processed by an outside laboratory, the outside laboratory will bill the renal dialysis facility. The renal dialysis facility will then submit a claim to Texas Medicaid unless the test results are inclusive tests.

If additional in-facility laboratory testing is medically necessary beyond the following routine frequencies, providers must bill with modifier 91 to indicate the billed laboratory procedure is medically necessary. The billing provider must also submit documentation supporting the medical necessity with the claim and maintain the documentation in the client’s medical record.

Modifier 91 is used to indicate that a test was performed more than once on the same day for the same client only when it is necessary to obtain multiple results in the course of the treatment. This modifier may not be used to indicate any of the following:

- When tests are rerun to confirm initial results
- Testing problems with specimens or equipment
- When a normal one-time, reportable result is all that is required
- When there are standard Healthcare Common Procedure Coding System (HCPCS) codes available that describe the series of results (e.g., glucose tolerance tests, evocative/suppression testing, etc.).

Modifier 91 may only be used for laboratory tests paid under the clinical diagnostic laboratory fee schedule.

Per Dialysis

<table>
<thead>
<tr>
<th>Procedure Codes</th>
</tr>
</thead>
<tbody>
<tr>
<td>85014^ 85018^ 85345 85347</td>
</tr>
</tbody>
</table>

^ QW modifier is required.

Per Week

<table>
<thead>
<tr>
<th>Procedure Codes</th>
</tr>
</thead>
<tbody>
<tr>
<td>82565^ 84520^ 85610^</td>
</tr>
</tbody>
</table>

^ QW modifier is required.

Per Month

<table>
<thead>
<tr>
<th>Procedure Codes</th>
</tr>
</thead>
<tbody>
<tr>
<td>82040^ 82310^ 82374^ 82435^ 83615 84075^ 84100 84132^ 84155^ 84450^</td>
</tr>
</tbody>
</table>

^ QW modifier is required.
The routine tests listed in the previous tables are frequently performed as an automated battery of tests such as the sequential multi-channel analysis with computer (SMAC)-12 (chemistry panels). These tests are considered routine and are included in the charge for dialysis, unless there is an additional diagnosis to document medical necessity for performing the tests in excess of the recommended frequencies.

6.2.9.2 In-Facility Dialysis—Nonroutine Laboratory

The following procedure codes are considered necessary, nonroutine tests. They must be submitted separately from the dialysis charge when performed in the renal dialysis facility or by an outside laboratory that bills the facility for laboratory services. All nonroutine laboratory and radiology tests beyond the following recommended frequencies must be medically necessary.

If additional in-facility laboratory testing is medically necessary beyond the following nonroutine frequencies, providers must submit the claim with modifier 91 to indicate the billed laboratory procedure is medically necessary. The billing provider must also submit documentation supporting the medical necessity with the claim and maintain the documentation in the client’s medical record.

Once a Month

<table>
<thead>
<tr>
<th>Procedure Codes</th>
</tr>
</thead>
<tbody>
<tr>
<td>87340</td>
</tr>
</tbody>
</table>

Every 3 Months

<table>
<thead>
<tr>
<th>Procedure Codes</th>
</tr>
</thead>
<tbody>
<tr>
<td>93005</td>
</tr>
</tbody>
</table>

Every 6 Months

<table>
<thead>
<tr>
<th>Procedure Codes</th>
</tr>
</thead>
<tbody>
<tr>
<td>71045 71046 71047 71048 95907 95908 95909 95910 95911 95912</td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th>Procedure Codes</th>
</tr>
</thead>
<tbody>
<tr>
<td>95913</td>
</tr>
</tbody>
</table>

Annually

<table>
<thead>
<tr>
<th>Procedure Codes</th>
</tr>
</thead>
<tbody>
<tr>
<td>78300 78305 78306</td>
</tr>
</tbody>
</table>

A handling fee (procedure code 99001) for nonroutine laboratory services may be submitted to Texas Medicaid only if the specimen is obtained by venipuncture or catheterization and sent to an outside lab. The claim form must document that the handling fee is for nonroutine laboratory services.

6.2.9.3 CAPD Laboratory

The following laboratory tests are routine for home maintenance CAPD clients when performed according to the indicated frequency. These laboratory tests may be reimbursed separately when the client is dialyzing in the home and is not undergoing IPD or hemodialysis in the facility. The provider must indicate the client’s diagnosis and the type of dialysis on the claim form. Tests in excess of this frequency or tests not listed in the tables require documentation of medical necessity for payment to be made.

Every Month

<table>
<thead>
<tr>
<th>Procedure Codes</th>
</tr>
</thead>
<tbody>
<tr>
<td>82040^ 82310^ 82374^ 82565^ 83615 83735 84075^ 84100 84132^ 84155^</td>
</tr>
</tbody>
</table>

^ QW modifier is required.
6.2.9.4 Hematopoietic Injections

Medicaid reimbursement is allowed for hematopoietic injections that are administered to clients who have anemia that is associated with chronic renal failure.

Providers must submit the client’s most recent dated hemoglobin or hematocrit levels in the comments section of the claim form when billing with procedure code Q4081. Frequency and quantity limitations apply.

Refer to: The Outpatient Drug Services Handbook (Vol. 2, Provider Handbooks) for more information about benefit and limitation criteria.

6.2.9.5 Blood Transfusions

Whole blood transfusions may be reimbursed separately to dialysis facilities when medically indicated for a Medicaid eligible client.

6.2.10 Prior Authorization

Prior authorization is not required for renal dialysis services. Prior authorization must be obtained for transplant-related services provided to clients who are not eligible for Medicare and are eligible only for Medicaid.

6.3 Documentation Requirements

All services require documentation to support the medical necessity of the service rendered, including renal dialysis services. Renal dialysis services are subject to retrospective review and recoupment if documentation does not support the service submitted for reimbursement. All physicians’, renal dialysis centers’, and medical suppliers’ supporting documentation is subject to retrospective review.

6.4 Claims Filing and Reimbursement

6.4.1 Claims Information

Renal dialysis facility services must be submitted to TMHP in an approved electronic claims format or on a UB-04 CMS-1450 paper claim form. Providers may purchase UB-04 CMS-1450 paper claim forms from the vendor of their choice. TMHP does not supply them.
When completing a UB-04 CMS-1450 paper claim form, all required information must be included on the claim, as information is not keyed from attachments. Superbills, or itemized statements, are not accepted as claim supplements.

**Refer to:** “Section 3: TMHP Electronic Data Interchange (EDI)” (*Vol. 1, General Information*) for information on electronic claims submissions.

“Section 6: Claims Filing” (*Vol. 1, General Information*) for general information about claims filing.

Subsection 6.6, “UB-04 CMS-1450 Paper Claim Filing Instructions” in “Section 6: Claims Filing” (*Vol. 1, General Information*) for instructions on completing paper claims. Blocks that are not referenced are not required for processing by TMHP and may be left blank.

**Reminder:** *The original onset date must be included on the claim form to prevent claim denial. The original onset date must be the same date entered on Form CMS-2728 sent to the Social Security office.*

### 6.4.2 Reimbursement

Renal dialysis facilities are reimbursed according to composite rates, which are based on the CMS-specified calculations and the Texas Medicaid Reimbursement Methodology (TMRM). Texas Medicaid may reimburse for dialysis services through either Method I or Method II as defined by CMS.

The hemodialysis, IPD, CAPD and CCPD laboratory and radiology services and the physician supervision of dialysis clients limitations pertain to both Method I and Method II reimbursement.

Texas Medicaid implemented mandated rate reductions for certain services. Additional information about rate changes is available on the TMHP website at [www.tmhp.com/resources/rate-and-code-updates/rate-changes](http://www.tmhp.com/resources/rate-and-code-updates/rate-changes).

**Refer to:** Section 2.3, “Reimbursement Methodology” (*Vol. 1, General Information*) for more information about reimbursement.

### 6.4.2.1 NCCI and MUE Guidelines

The HCPCS and CPT codes included in the *Texas Medicaid Provider Procedures Manual* are subject to NCCI relationships, which supersede any exceptions to NCCI code relationships that may be noted in the manuals. Providers should refer to the CMS NCCI web page for correct coding guidelines and specific applicable code combinations.

In instances when Texas Medicaid limitations are more restrictive than NCCI MUE guidance, Texas Medicaid limitations prevail.

### 6.5 Medicare and Medicaid

Medicaid coverage of a renal dialysis client who may be eligible for Medicare coverage begins with the original onset date of the dialysis treatments and may continue for a period of three months. During this period, Medicare eligibility is reviewed through the Health and Human Services Commission (HHSC). If HHSC determines that the client is Medicare-eligible, Medicaid coverage begins with the original onset date and continues until Medicare coverage begins.

If HHSC determines that the client is not eligible for Medicare, Medicaid coverage of eligible clients begins with the original onset date and continues as long as the dialysis treatments are medically necessary and the client is eligible for Medicaid. The date of onset is the date of the first dialysis treatment and does not change even if the client sees another provider.

Medicare eligibility usually begins after a three-month waiting period has been served. Medicare eligibility begins before the waiting period has expired if the individual receives a transplant or participates in a self-dialysis training program during the waiting period.
6.5.1 Facility Providers
Texas Medicaid pays the Medicare coinsurance less 5 percent and full Medicare deductible for Medicare crossover claims that are submitted by nephrology (hemodialysis, renal dialysis) and renal dialysis facility providers.

6.5.2 Physician Providers
The five percent reduction does not apply to physician-billed services. Nephrologists that are enrolled in Texas Medicaid as physician providers may be reimbursed according to the current payment guidelines.

Refer to: Subsection 2.7, “Medicare Crossover Claim Reimbursement” in “Section 2: Texas Medicaid Fee-for-Service Reimbursement” (Vol. 1, General Information) for additional information about Medicare coinsurance and deductible reimbursement for professional and outpatient services.

7 Rural Health Clinic

7.1 Enrollment
To enroll in Texas Medicaid and qualify for participation as a Title XIX RHC, RHCs must be enrolled in Medicare. A nine-digit provider identifier is issued to the RHC after a certification letter from Medicare is received, stating that the clinic qualifies for Medicaid participation. An RHC can also apply for enrollment as a family planning agency.

All providers of laboratory services must comply with the rules and regulations of CLIA. Providers who do not comply with CLIA are not reimbursed for laboratory services.

7.1.1 Initial Cost Reporting
New RHCs must file a projected cost report within 90 days of their designation as an RHC to establish an initial payment rate. The cost report will contain the RHC’s reasonable costs anticipated to be incurred during the RHC’s first full fiscal year. The projected cost report must contain a minimum of six months of information. The RHC must file a cost report within five months of the end of the RHC’s initial fiscal year. The cost settlement must be completed within six months of the receipt of a cost report. The cost per visit rate established by the cost settlement process shall be the base rate. Any subsequent increases or decreases shall be calculated as provided herein. A new RHC location established by an existing RHC participating in Texas Medicaid will receive the same effective rate as the RHC establishing the new location. An RHC establishing a new location may request an adjustment to its effective rate as provided herein if its costs have increased as a result of establishing a new location.

Providers must submit initial cost reports to the following address:

Texas Medicaid & Healthcare Partnership
Medicaid Audit
PO Box 200345
Austin, TX 78720-0345

Providers can refer to 1 TAC §355.8101 for more information about reimbursement.

Refer to: Subsection 1.1, “Provider Enrollment” in “Section 1: Provider Enrollment and Responsibilities” (Vol. 1, General Information) for more information.

7.2 Services, Benefits, Limitations, and Prior Authorization

7.2.1 Services Rendered by the RHC Facility Provider

General services and copayments are billed using the RHC’s National Provider Identifier (NPI). All other services billed using the RHC’s NPI are processed as informational only.

**Important:** If an RHC facility provider submits a claim for THSteps and Family Planning services, the services will process as informational only and will not be reimbursed.

The following services are benefits of Texas Medicaid when provided in an RHC:

- Physician services
- Services and supplies furnished as incidental to physician services
- Services provided by an NP, a CNM, a clinical social worker, or a PA’s services
- Services and supplies furnished as incidental to the NP’s or PA’s services
- Visiting nurse services on a part-time or intermittent basis to homebound clients in areas determined to have a shortage of home health agencies (A homebound client is someone who is permanently or temporarily confined to his place of residence, not including a hospital or skilled nursing facility (SNF), because of a medical condition.)

When an RHC bills for visiting nurse services, the written plan of treatment to be used for the visiting nurse must be developed by the RHC supervising physician. It must be approved and ordered by the client’s treating physician if different from the supervising physician. The plan of treatment must be reviewed and approved by the supervising physician of the clinic at least every 60 days.

A visit is a face-to-face encounter between an RHC client and a physician, PA, NP, CNM, visiting nurse, or clinical NP. Encounters with more than one health professional and multiple encounters with the same health professional that take place on the same day and at a single location constitute a single visit, except where one or the other of the following conditions exists:

- After the first encounter, the client suffers illness or injury requiring additional diagnosis or treatment.
- The RHC client has a medical visit and an other health visit.

An other health visit includes, but is not limited to, a face-to-face encounter between an RHC client and a clinical social worker.

For freestanding RHCs, all laboratory services provided in the RHC’s laboratory are included in the encounter. This includes the basic laboratory tests as well as any other laboratory tests provided in the RHC laboratory. Consequently, there is no separate billing for laboratory services. However, if the RHC laboratory becomes a certified Medicare laboratory with its own supplier number, and enrolls in Medicaid as an independent laboratory, all laboratory tests (except the basic laboratory tests) performed for RHC and non-RHC clients can be billed to Medicaid. The claim must be filed under their independent laboratory Medicaid provider identifier and using the appropriate HCPCS codes.

**Refs:**
- The Medicare website at [www.cms.gov](http://www.cms.gov) for more information about Medicare RHC laboratory requirements.
7.2.1.1 Encounter Rates

An encounter rate may be reimbursed to the RHC facility only for the following services:

<table>
<thead>
<tr>
<th>General Medical Services (encounter may be reimbursed to the RHC facility only)</th>
</tr>
</thead>
<tbody>
<tr>
<td>T1015</td>
</tr>
</tbody>
</table>

General medical services must be submitted using one of the appropriate modifiers AJ, AM, SA, TD, TE, or U7. Adult preventative care must be submitted with diagnosis codes Z0000, Z0001, Z01411, and Z01419.

**Note:** If the encounter is for antepartum or postpartum care, use modifier TH in addition to the modifier required to clarify the service that was performed.

7.2.1.2 Medicaid Fee-for-Service Reimbursement Rates

The following copayments may be reimbursed to RHC providers billing under their own NPI, and are reimbursed at the Medicaid fee-for-service rate.

<table>
<thead>
<tr>
<th>Copayments</th>
</tr>
</thead>
<tbody>
<tr>
<td>CP001</td>
</tr>
</tbody>
</table>

7.2.1.3 Freestanding Rural Health Clinic Services

The following services cannot be reimbursed to freestanding RHCs using only the RHC provider identifier. Use of the RHC provider identifier for billing these services causes claims to be processed as informational only. Services in any of these categories must be billed using the professional (non-RHC) provider identifier and the appropriate benefit code:

- THSteps medical checkups, which includes immunizations
- Family planning services (including implantable contraceptive capsules provision, insertion, or removal)

These services must be billed with an AM, SA, or U7 modifier.

Physician supplies are not a benefit of Texas Medicaid. Costs of supplies are included in the reimbursement for office visits. Outpatient hospital services (including emergency room services) and inpatient hospital services provided outside the RHC setting are to be billed using the individual or group physician provider identifier.

**Exception:** If later in the same day the client suffers an additional illness or injury requiring diagnosis or treatment, the clinic may submit a claim for a second visit.

Freestanding RHCs submit an all-inclusive encounter for services provided. All services provided that are incidental to the encounter, including developmental screening, must be included in the total charge for the encounter. A claim for these services may not be submitted as a separate encounter.

If immunizations are given outside of a THSteps medical checkup, procedure codes given in the THSteps section of this manual should be identified on the claim. These procedure codes are informational only, and are not payable.

All services provided during a freestanding RHC encounter must be submitted using procedure code T1015. The total submitted amount should be the combined charges for all services provided during that encounter.

One of the following modifiers must be reported with procedure code T1015 to designate the health-care professional providing the services: AH, AJ, AM, SA, TD, TE, or U7. If the encounter is for antepartum or postpartum care, use modifier TH in addition to the modifier required to designate the health-care professional providing the service.

**Reminder:** The primary initial contact is defined as “the health-care professional who spends the greatest amount of time with the client during that encounter.”
If more than one health-care professional is seen during the encounter, the modifier (if appropriate) must indicate the primary contact. For example, if an NP or a PA performs an antepartum exam, modifiers SA or U7, and TH, must be entered. A maximum of two modifiers may be reported with each encounter.

Providers who render services in an RHC setting for THSteps Medical services or Family Planning services may be reimbursed an encounter rate.

**THSteps Medical Services**

RHC facility providers may be reimbursed for THSteps medical services using their RHC NPI with the appropriate benefit code.

If the appropriate benefit code is not included, the service will process as informational only and will not be reimbursed.

<table>
<thead>
<tr>
<th>THSteps Medical Services</th>
</tr>
</thead>
<tbody>
<tr>
<td>99381 99382 99383 99384 99385 99391 99392 99393 99394 99395</td>
</tr>
</tbody>
</table>

**7.2.1.4 Family Planning Services**

RHC facility providers may be reimbursed for family planning services using their RHC NPI with the appropriate benefit code.

If the appropriate benefit code is not included, the service will process as informational only and will not be reimbursed.

Family planning services must be submitted with the most appropriate evaluation and management (E/M) procedure code and the most appropriate family planning diagnosis code:

<table>
<thead>
<tr>
<th>Family Planning Services*</th>
</tr>
</thead>
<tbody>
<tr>
<td>99202 99203 99204 99205 99211 99212 99213 99214 99215 J7296</td>
</tr>
</tbody>
</table>

Family planning services performed in the RHC setting must be billed with the appropriate modifier: AM, SA, or U7.

RHC providers may receive an encounter rate when submitting claims with procedure code T1015, in addition to a flat “add on” fee for the Long-Acting Reversible Contraception (LARC) procedure codes listed above.

**Refereto:** Subsection 4.1.2, “Services, Benefits, Limitations, and Prior Authorization” in this handbook for the list of family planning diagnosis codes.

Subsection 6.3.5, “Modifiers” in “Section 6: Claims Filing” (Vol. 1, General Information) for a definition of modifiers.


Section 5, “THSteps Medical” in the Children’s Services Handbook (Vol. 2, Provider Handbooks).

Section 2, “Medicaid Title XIX Family Planning Services” in the Gynecological, Obstetrics, and Family Planning Title XIX Services Handbook (Vol. 2, Provider Handbooks).
Subsection 8.8.2.2, “HMO Copayments” in “Section 8: Third Party Liability (TPL)” (Vol. 1, General Information) for information about HMO copayments.

Subsection 9.2.56.3.2, “Preventive Care Visits” in the Medical and Nursing Specialists, Physicians, and Physician Assistants Handbook (Vol. 2, Provider Handbooks).

7.2.2 Services Rendered by Non-RHC Providers In An RHC Setting

Non-RHC providers (i.e., physicians and non-physician practitioners) submitting claims for THSteps and Family Planning services in the RHC setting must use their own NPI and the appropriate benefit code.

The following services, when rendered in an RHC setting by a non-RHC provider, will process as an encounter and will be reimbursed to the non-RHC provider as an encounter rate equivalent to the host facility:

- THSteps medical services
- Family planning services

Non-RHC providers rendering services in an RHC setting must use the appropriate national place of service (72) in order for claims to process as encounters.

7.2.3 Hospital-Based Rural Health Clinic Services

Hospital-based RHCs must use the encounter code T1015. A hospital-based RHC is paid based on an all-inclusive encounter rate. One of the following modifiers must be submitted for general medical services: AH, AJ, AM, SA, TD, TE, or U7.

The following services must be submitted using the physician’s Texas Provider Identifier (TPI) and the appropriate benefit code:

- THSteps medical checkups
- Family planning services (including implantable contraceptive capsules provision, insertion, or removal)
- Immunizations provided in hospital-based RHCs

Note: Refer to the tables in the above sections for procedure codes.

These services must be submitted with an AM, SA, or U7 modifier if performed in an RHC setting. Claims are paid under the PPS reimbursement methodology.

Outpatient hospital services (including emergency room services) and inpatient hospital services provided outside the RHC setting are to be submitted using the individual or group physician provider identifier. Hospital-based RHCs must submit claims for pneumococcal and influenza vaccines as non-RHC services, under their hospital provider identifier.

7.2.3.1 After-Hours Care

After-hours care for RHCs is defined as care provided on weekends, federal holidays, or before 8 a.m. and after 5 p.m., Monday through Friday.

7.3 Prior Authorization

Prior authorization or authorization is not required for RHC services.

7.4 Documentation Requirements

All services require documentation to support the medical necessity of the service rendered, including RHC services. RHC services are subject to retrospective review and recoupment if documentation does not support the service billed.
7.4.1 Record Retention
Freestanding RHCs must retain their records for a minimum of six years. Hospital-based RHCs must retain their records for a minimum of ten years.

7.5 Claims Filing and Reimbursement
7.5.1 Claims Information
General services and copayments are billed using the RHC’s NPI. For all other services, providers must submit claims using their NPI and the appropriate benefit code.


Place of service 72 must be used on all claims when billing for services other than general medical. Benefit code EP1 must be used on claims for THSteps medical services.

Freestanding and hospital-based RHC services must be submitted to TMHP in an approved electronic format or on a UB-04 CMS-1450 paper claim form. Providers may purchase UB-04 CMS-1450 paper claim forms from the vendor of their choice. TMHP does not supply the forms.

When completing a UB-04 CMS-1450 paper claim form, all required information must be included on the claim, as TMHP does not key any information from claim attachments. Superbills, or itemized statements, are not accepted as claim supplements.

7.5.2 Reimbursement
Freestanding and hospital-based RHCs are reimbursed provider-specific per visit rates calculated in accordance with 1 TAC §355.8101. Texas Medicaid implemented mandated rate reductions for certain services. Additional information about rate changes is available on the TMHP website at www.tmhp.com/resources/rate-and-code-updates/rate-changes.

7.5.2.1 Medicare Crossover Claims Pricing
For Medicare Part B cost sharing obligations, all deductible obligations will be reimbursed at 100 percent of the deductible amount owed, even if the cost sharing comparison results in a lower payment. For all other cost sharing obligations (including Medicare Part A, B, and C), the cost sharing comparison is performed according to current guidelines.

For RHC Medicare crossover claims, Texas Medicaid will reimburse the lesser of the following:
- The coinsurance and full deductible payment.
- The amount remaining after the Medicare payment amount is subtracted from the allowed Medicaid fee or encounter rate for the service. If this amount is less than the deductible, then the full deductible is reimbursed instead.

If the Medicare payment is equal to, or exceeds the Medicaid allowed amount or encounter payment for the service, Texas Medicaid will not make a payment for coinsurance.

The client has no liability for any balance or Medicare coinsurance and deductible related to Medicaid-covered services.

Refer to: Subsection 2.7, “Medicare Crossover Claim Reimbursement” in “Section 2: Texas Medicaid Fee-for-Service Reimbursement” (Vol. 1, General Information).

7.5.2.2 NCCI and MUE Guidelines
The HCPCS and CPT codes included in the Texas Medicaid Provider Procedures Manual are subject to NCCI relationships, which supersede any exceptions to NCCI code relationships that may be noted in the manuals. Providers should refer to the CMS NCCI web page for correct coding guidelines and specific applicable code combinations.
In instances when Texas Medicaid limitations are more restrictive than NCCI MUE guidance, Texas Medicaid limitations prevail.

8 Tuberculosis Services

TB clinics must be enrolled in Texas Medicaid and provide services in accordance with 1 TAC, §354.1371.

8.1 Enrollment

To enroll in Texas Medicaid, a TB clinic must be either:

• A public entity operating under an HHSC tax identification number (TB regional clinic)
• A public entity operating under a non-HHSC tax identification number (city/county/local clinic)
• A non-hospital-based entity for private providers

Providers of TB-related clinic services must complete a provider application from the TB Services Branch within DSHS. Per Texas DSHS policy, TB clinics must develop and operate under a set of written policies and procedures that specify the criteria for licensed and non-licensed staff to provide services. The policies and procedures must include the following:

• The personnel file requirements for staff who provide directly observed therapy (DOT).
• The training and supervision that are required for outreach workers to be considered qualified to perform the assigned services.
• The written delegation protocol for services that are not performed by a physician, advanced practice registered nurse (APRN), or PA.
• The documentation that is required for all client encounters.

Upon written notice of approval by TB Services Branch, Medicaid enrollment applications from TMHP Provider Enrollment are sent to HHSC-approved providers of TB-related clinic services.

TMHP is responsible for issuing a group or individual a nine-digit provider identifier. Providers that list additional (satellite) clinics in the TB Services Branch provider application will receive nine-digit performing provider identifiers for each off-site clinic. TB off-site clinics operating under the jurisdiction of the applying provider must use the assigned group provider identifier and their nine-digit performing provider identifier.

Enrollment as a Medicaid provider is not complete until the TMHP enrollment packet has been finalized and a nine-digit provider identifier number is issued to the provider.

The effective date for participation is the date an approved provider application with the TB Services Branch is established.

To receive a TB Services Branch provider application form or provider supplement, send a request to the following address:

Texas Department of State Health Services
TB/HIV/STD/Viral Hepatitis Unit
Tuberculosis Services Branch
Mail Code 1939
1100 West 49th Street
PO Box 149347
Austin, TX 78714-9347
Refer to: Subsection 1.1, “Provider Enrollment” in “Section 1: Provider Enrollment and Responsibilities” (Vol. 1, General Information) for more information about enrollment procedures related to the TMHP Medicaid enrollment applications.

8.1.1 Managed Care Program Enrollment

TB clinics do not need to enroll with the Medicaid managed care health plans. All services provided by TB clinics are submitted to TMHP for all Medicaid clients, including Medicaid managed care clients.

8.2 Services, Benefits, Limitations, and Prior Authorization

The level of service provided varies depending on whether the services are delivered by a nonphysician or physician and if medications are prescribed.

8.2.1 TB-Related Clinic Services

The following services may be performed by a physician, APRN, or PA in the TB clinic:

<table>
<thead>
<tr>
<th>Procedure Codes</th>
</tr>
</thead>
<tbody>
<tr>
<td>99202</td>
</tr>
</tbody>
</table>

A physician’s presence is not required to perform procedure code 99211; however, the physician must provide direct supervision by being present in the clinic and immediately available to furnish assistance and direction at the time service is provided.

Before TB treatment can be initiated, an initial screening (procedure code T1023) by an RN, LPN, or LVN, or a new patient physician E/M visit (procedure code 99202, 99203, 99204, or 99205) must be performed. If the treatment is initiated by a nursing screening, a new patient physician E/M visit must be completed within 90 days, or subsequent reimbursement for DOT (procedure code H0033) will be denied.

Following the initial new patient physician E/M visit, an established patient physician E/M visit (procedure code 99212, 99213, 99214, or 99215) must be billed every 90 days throughout the course of treatment, or subsequent reimbursement for DOT (procedure code H0033) will be denied.

Clients with latent TB infection, including those in a high-risk group (children who are 4 years of age and younger, those who are immunocompromised, and clients who are HIV-positive), and those with active TB disease, must be seen by a physician every 90 days throughout the course of treatment.

A physician must evaluate each client with active or latent TB disease prior to discharge from TB treatment.

Procedure codes H0033, T1002, T1003, and T1023 may be provided under established clinic protocols. The initial TB screening (procedure code T1023), performed by an RN, LPN, or LVN includes, but is not limited to the following:

- Brief mental and physical assessment
- Exposure history
- Referral for lab or X-ray per protocol
- Referral for social or other medical services
- Other assessment

Procedure code T1023 may be reimbursed prior to the client being seen by a physician, and no more often than once per 12 months. One RN or LVN/LPN (procedure codes T1023, T1002, and T1003) service may be reimbursed per day, per client, when physician services are not performed.
Subsequent nursing services (Procedure code T1002 and T1003) may be a benefit when not provided the same day as a physician E/M visit.

Reimbursement for DOT services (procedure code H0033) provided in the clinic or other places of service, excluding inpatient hospitals, SNFs, intermediate care facilities (ICFs), outpatient hospitals, independent laboratories, birthing centers, and extended care facilities will be limited to one per day, and a maximum of five per week, per client, throughout the course of treatment.

Procedure codes T1002 and T1003 are limited to a maximum of eight 15-minute units per day, per client.

- Minutes of nursing services cannot be accumulated over multiple days. Minutes of nursing services can only be billed per calendar day.
- If the total number of minutes of nursing services per procedure code is less than 8 minutes for a calendar day, then no unit of service can be billed for that day. The minutes cannot be added to minutes of nursing services from any previous or subsequent days for billing purposes.
- If more than 1 unit of service is billed, every unit except the last must be for the complete 15 minutes, with the last unit being no less than 8 minutes of nursing service.
- Time spent in contact investigations in not reimbursable.

Reimbursement for new client examinations (procedure code 99202, 99203, 99204, and 99205) are limited to new clients who have not received services in the same clinic for a period of three years. One physician E/M service may be reimbursed per day, per client.

### 8.2.2 Ancillary Services

The following ancillary TB services are a benefit of Texas Medicaid:

<table>
<thead>
<tr>
<th>Procedure Codes</th>
</tr>
</thead>
<tbody>
<tr>
<td>71045</td>
</tr>
<tr>
<td>96366</td>
</tr>
<tr>
<td>J2280**</td>
</tr>
</tbody>
</table>

* CLIA waived test  
^ Must be billed with QW modifier  
** Must be billed with KX modifier when oral formulation is not appropriate for the client

Procedure code 99000 is limited to specimens transported from the office setting.

Procedure code 99001 is limited to specimens transported from any setting except the office.

Certain injectable TB medications (procedure codes J2020, J2280, and J3000), which also have an oral formulation, must be billed with modifier KX to indicate that the oral formulation is not appropriate for the client.

All drugs for which Medicaid is billed must have been purchased by the TB clinic. In the event that the clinic received the drug at no cost through DSHS or another source, it cannot be billed to Texas Medicaid. All medication claims are subject to retrospective review.

Handling or conveyance of a specimen from the patient in the clinic to a laboratory (procedure code 99000) will be reimbursed only when submitted with one of the following professional or nursing services performed on the same date of service. Prior authorization is not required for procedure code 99000 or 99001.
Texas Medicaid follows the Medicare categorization of tests for CLIA certificate holders.


For waived tests, providers must use modifier QW as indicated on the CMS website.

### 8.2.3 Prior Authorization

Prior authorization is not required for TB-related services.

### 8.3 Documentation Requirements

All services require documentation to support the medical necessity of the service rendered, including TB services. TB services are subject to retrospective review and recoupment if documentation does not support the service billed.

### 8.4 Provider Responsibilities

If approved to submit claims as a TB clinic under Texas Medicaid, the provider must adhere to the following requirements:

- Be a facility that is not an administrative, organizational, or financial part of a hospital, but is organized and operated to provide medical care to outpatients.
- Comply with all applicable federal, state, and local laws and regulations.
- Employ or have a contract or formal arrangement with a licensed physician (M.D. or D.O.) who is responsible for providing medical direction and supervision over all services provided to the clinic’s clients. To meet this requirement, a physician must see the client at least once every 90 days to prescribe the type of care provided and, if the services are not limited by the prescription, periodically review the need for continued care.
- Adhere to the guidelines issued by HHSC, under the authority of the Texas Health and Safety Code, and ensure that services are consistent with the recommendations of the American Thoracic Society and the Centers for Disease Control and Prevention (CDC). For more information, visit the website at [www.cdc.gov/tb/default.htm](http://www.cdc.gov/tb/default.htm).
- Maintain complete and accurate medical records of each recipient’s care and treatment and accurately document all services provided and the medical necessity for the services.
- Ensure that services provided to each client are commensurate with the client’s medical needs based on the client’s assessment or evaluation, diagnostic studies, plan of care, and physician direction. These services must be documented in the client’s medical records.
- Be enrolled and approved for participation in Texas Medicaid.
- Sign a written provider agreement with HHSC or its designee. By signing the agreement, the provider of TB-related clinic services agrees to comply with the terms of the agreement and all requirements of Texas Medicaid including regulations, rules, handbooks, standards, and guidelines published by HHSC or its designee.
- Submit claims for services covered by Texas Medicaid in the manner and format prescribed by HHSC or its designee.

<table>
<thead>
<tr>
<th>Procedure Codes</th>
</tr>
</thead>
<tbody>
<tr>
<td>99202</td>
</tr>
</tbody>
</table>

* Reimbursed for individual or group TB clinics for services rendered in the home or other setting.
• Be organized and operated to provide TB-related services, which include, but are not limited to, the covered services as indicated in subsection 8.2, “Services, Benefits, Limitations, and Prior Authorization” in this handbook.

• Not provide services within an SNF, ICF, or intermediate care facility for persons with intellectual disability (ICF-ID)

Refer to: Subsection 1.1, “Provider Enrollment” in “Section 1: Provider Enrollment and Responsibilities” (Vol. 1, General Information) for more information.

8.5 Claims Filing and Reimbursement

8.5.1 Claims Information

TB-related clinic services must use benefit code TB1 on all claims and authorization requests. All TB-related clinic services must be submitted to TMHP in an approved electronic claims format or on a CMS-1500 paper claim form from the vendor of their choice. TMHP does not supply them. When completing a CMS-1500 paper claim form, all required information must be included on the claim, as information is not keyed from attachments. Superbills, or itemized statements, are not accepted as claim supplements.

Refer to: “Section 3: TMHP Electronic Data Interchange (EDI)” (Vol. 1, General Information) for information on electronic claims submissions.

“Section 6: Claims Filing” (Vol. 1, General Information).

Subsection 6.5, “CMS-1500 Paper Claim Filing Instructions” in “Section 6: Claims Filing” (Vol. 1, General Information) for instructions on completing paper claims. Blocks that are not referenced are not required for processing by TMHP and may be left blank.

8.5.1.1 Managed Care Clients

TB-related services are carved out of the Medicaid Managed Care Program and must be billed to TMHP for payment consideration. Carved-out services are those that are rendered to Medicaid Managed Care clients, but are administered by TMHP and not the client’s MCO.

8.5.2 Reimbursement

The Medicaid reimbursement rates for TB clinics are calculated in accordance with 1 TAC §355.8085. X-ray services are reimbursed in accordance with 1 TAC §355.8085 and are listed in the current physician fee schedule on the TMHP website at www.tmhp.com. Texas Medicaid implemented mandated rate reductions for certain services. The Online Fee Lookup (OFL) and static fee schedules include a column titled “Adjusted Fee” to display the individual fees with all mandated percentage reductions applied. Additional information about rate changes is available on the TMHP website at www.tmhp.com/resources/rate-and-code-updates/rate-changes.

8.5.2.1 NCCI and MUE Guidelines

The HCPCS and CPT codes included in the Texas Medicaid Provider Procedures Manual are subject to NCCI relationships, which supersede any exceptions to NCCI code relationships that may be noted in the Texas Medicaid Provider Procedures Manual. Providers should refer to the CMS NCCI web page for correct coding guidelines and specific applicable code combinations.

In instances when Texas Medicaid limitations are more restrictive than NCCI MUE guidance, Texas Medicaid limitations prevail.
9 Claims Resources

Refer to the following sections or forms when filing claims:

<table>
<thead>
<tr>
<th>Resource</th>
<th>Location</th>
</tr>
</thead>
<tbody>
<tr>
<td>Acronym Dictionary</td>
<td>“Appendix C: Acronym Dictionary” (Vol. 1, General Information)</td>
</tr>
<tr>
<td>American Dental Association (ADA) Dental Claim Filing Instructions</td>
<td>Subsection 6.7, “American Dental Association (ADA) Dental Claim Filing Instructions” in “Section 6: Claims Filing” (Vol. 1, General Information)</td>
</tr>
<tr>
<td>Automated Inquiry System (AIS)</td>
<td>Subsection A.10, “TMHP Telephone and Fax Communication” in “Appendix A: State, Federal, and TMHP Contact Information” (Vol. 1, General Information)</td>
</tr>
<tr>
<td>CMS-1500 Paper Claim Filing Instructions</td>
<td>Subsection 6.5, “CMS-1500 Paper Claim Filing Instructions” in “Section 6: Claims Filing” (Vol. 1, General Information)</td>
</tr>
<tr>
<td>State, federal, and TMHP contact information</td>
<td>“Appendix A: State, Federal, and TMHP Contact Information” (Vol. 1, General Information)</td>
</tr>
<tr>
<td>TMHP electronic claims submission information</td>
<td>Subsection 6.2, “TMHP Electronic Claims Submission” in “Section 6: Claims Filing” (Vol. 1, General Information)</td>
</tr>
<tr>
<td>TMHP Electronic Data Interchange (EDI) information</td>
<td>“Section 3: TMHP Electronic Data Interchange (EDI)” (Vol. 1, General Information)</td>
</tr>
<tr>
<td>Tuberculosis Screening and Guidelines</td>
<td>Subsection A.5, “Tuberculosis Screening and Guidelines” in the Children’s Services Handbook (Vol. 2, Provider Handbooks)</td>
</tr>
</tbody>
</table>

10 Contact TMHP

The TMHP Contact Center at 1-800-925-9126 is available Monday through Friday from 7 a.m. to 7 p.m., Central Time.

11 Forms

The following linked forms can also be found on the Forms page of the Provider section of the TMHP website at www.tmhp.com:

<table>
<thead>
<tr>
<th>Forms</th>
<th></th>
</tr>
</thead>
<tbody>
<tr>
<td>Federally Qualified Health Center Affiliation Affidavit</td>
<td></td>
</tr>
<tr>
<td>Hospital Report (Newborn Child or Children) (Form 7484)</td>
<td></td>
</tr>
</tbody>
</table>
## 12 Claim Form Examples

<table>
<thead>
<tr>
<th>Claim Form Examples</th>
</tr>
</thead>
<tbody>
<tr>
<td>Birthing Center</td>
</tr>
<tr>
<td>2017 Claim Form</td>
</tr>
<tr>
<td>Family Planning Services for Hospitals, FQHCs</td>
</tr>
<tr>
<td>FQHC Encounter (T1015)</td>
</tr>
<tr>
<td>FQHC Follow-Up</td>
</tr>
<tr>
<td>Renal Dialysis Facility CAPD Training</td>
</tr>
<tr>
<td>Renal Dialysis Facility CAPD/CCPD</td>
</tr>
<tr>
<td>Renal Dialysis CMS-1500 Example</td>
</tr>
<tr>
<td>Rural Health Clinic Freestanding</td>
</tr>
<tr>
<td>Rural Health Clinic Freestanding (Immunization)</td>
</tr>
<tr>
<td>Rural Health Clinic Hospital-Based</td>
</tr>
<tr>
<td>Tuberculosis</td>
</tr>
</tbody>
</table>
CERTIFIED RESPIRATORY CARE PRACTITIONER (CRCP) SERVICES

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1 General Information

This handbook contains information about Texas Medicaid fee-for-service benefits. For information about managed care benefits, refer to the Medicaid Managed Care Handbook.

Managed care carve-out services are administered as fee-for-service benefits. A list of all carve-out services is available in Section 8, “Carve-Out Services” in the Medicaid Managed Care Handbook (Vol. 2, Provider Handbooks).

The information in this handbook is intended for CRCP services and provides information about Texas Medicaid’s benefits, policies, and procedures applicable to these therapies.

**Important:** All providers are required to read and comply with Section 1: Provider Enrollment and Responsibilities. In addition to required compliance with all requirements specific to Texas Medicaid, it is a violation of Texas Medicaid rules when a provider fails to provide healthcare services or items to Medicaid clients in accordance with accepted medical community standards and standards that govern occupations, as explained in Title 1 Texas Administrative Code (TAC) §371.1659. Accordingly, in addition to being subject to sanctions for failure to comply with the requirements that are specific to Texas Medicaid, providers can also be subject to Texas Medicaid sanctions for failure, at all times, to deliver health-care items and services to Medicaid clients in full accordance with all applicable licensure and certification requirements including, without limitation, those related to documentation and record maintenance.

**Refer to:** “Section 1: Provider Enrollment and Responsibilities” (Vol. 1, General Information) for more information about enrollment procedures.

2 Enrollment

**Refer to:** Subsection 1.7.16, “Certified Respiratory Care Practitioner (CRCP) Services” in “Section 1: Provider Enrollment and Responsibilities” (Vol. 1, General Information) for additional information about enrollment.

3 CRCP Services

3.1 Services, Benefits, Limitations, and Prior Authorization

Certified respiratory care practitioner services are a benefit of Texas Medicaid when provided in the home setting for ventilator-dependent clients by providers enrolled in Texas Medicaid as respiratory therapists.

Subject to the following specifications, conditions, and limitations, in-home certified respiratory care practitioner services are available to eligible clients who:

- Are ventilator-dependent for life support at least six hours per day.
- Have been ventilator-dependent for at least 30 consecutive days as an inpatient in one or more hospitals, skilled nursing facilities (SNF), or intermediate care facilities (ICF).
- But for the availability of these respiratory care services at home, would require respiratory care as an inpatient in a hospital, SNF, or ICF.
- Would be eligible to have reimbursement made for such inpatient care under the state Medicaid plan.
- Have adequate social support services to be cared for at home.
• Wish to be cared for at home.

• Require professional respiratory therapy services in addition to those respiratory therapy services that are provided through the Home Health durable medical equipment (DME) lease of a ventilator.

  **Note:** For clients who are birth through 20 years of age who do not meet the criteria above, services to be performed by a certified respiratory care practitioner may be considered through the Comprehensive Care Program (CCP).

Benefits include:

• Respiratory therapy services and treatments prescribed by a physician who is familiar with the client’s medical history and care, and who has medically determined that in-home care is safe and feasible for the client.

• Education of the client, the appropriate family members, or support persons regarding the in-home respiratory care. Education must include the use and maintenance of required supplies, equipment, and techniques appropriate to the situation.

Providers must use procedure code 99504 for in-home respiratory services.

Providers of respiratory therapy services must meet the following requirements:

• Be certified by the Texas Medical Board to practice under Chapter 604 of the Texas Occupations Code.

• Be enrolled and approved for participation in the Texas Medical Assistance Program.

• Bill for benefit services in the manner and format prescribed by HHSC or its designee.

• Comply with all applicable federal, state, and local laws and regulations.

The professional service may be billed by the certified respiratory care practitioner for services provided in the client’s home (procedure code 99504). The professional service will be allowed once per day up to a limit of 24 visits per year. The recommended schedule includes 7 visits during the first week, a total of 6 visits during the second through fourth week, and 11 monthly visits for the second through the 12th month.

Providers will not be reimbursed for procedure codes 99503 and 99504 on the same date of service, any provider.

Disposable respiratory supplies are a benefit through Texas Medicaid Title XIX Home Health Services and are not reimbursed to the certified respiratory therapist.

### 3.1.1 Authorization Requirements

Prior authorization requests may be submitted to the TMHP Prior Authorization Department via mail, fax, or the electronic portal. Prescribing or ordering providers, dispensing providers, clients’ responsible adults, and clients may sign prior authorization forms and supporting documentation using electronic or wet signatures.

  **Refer to:** Subsection 5.5.1.2, “Document Requirements and Retention” in “Section 5: Fee-for-Service Prior Authorizations” (*Vol. 1, General Information*) for additional information about electronic signatures.

Prior authorization is required for in-home certified respiratory care practitioner services (procedure code 99504).

To avoid unnecessary denials, the provider must submit correct and complete information including documentation of medical necessity for the service requested. The prescribing physician and provider must maintain documentation of medical necessity in the client’s medical record. The requesting provider may be asked for additional information to clarify or complete a request for the service.
Prior authorization requests for traditional Medicaid clients must be submitted by the physician or the certified respiratory care practitioner to the Special Medical Prior Authorization (SMPA) Department by approved electronic method using the “Special Medical Prior Authorization (SMPA) Request Form.”

Refer to: Special Medical Prior Authorization (SMPA) Request Form on the TMHP website at www.tmhp.com.

When required, the requests must include the physician’s signature and the date signed. Without this information, requests will be considered incomplete.

The SMPA Request Form must be submitted with the following documentation supporting medical necessity for the requested procedure:

- The client is on a ventilator at least six hours per day.
- The client has been ventilator dependent for 30 consecutive days or more as an inpatient in one or more hospitals, SNF, or ICF.
- The respiratory therapy services are in lieu of respiratory services requiring the client to remain in an inpatient care setting.
- Identification of the adequate support services in place that allow the client to be cared for at home
- The respiratory services and goals for the services that will be provided by the certified respiratory care practitioner.
- The frequency and number of home visits requested by the certified respiratory care practitioner
- The client’s wish to be cared for at home.
- Documentation supporting why the respiratory therapy visits included in the Home Health DME rental of a ventilator would not meet the client’s medical needs.

The request may be authorized for up to a 12-month period. Requests for more than 24 visits in a 12-month period will be referred for the medical director to review and a determination will be based on the individual client's medical needs.

Retrospective review may be performed to ensure documentation supports the medical necessity of the service when billing the claim for the procedure codes listed within the policy.

4 CRCP-CCP Services

4.1 Services, Benefits, Limitations, and Prior Authorization

In-home respiratory services by a licensed certified respiratory care practitioner are a benefit of the Texas Medicaid Comprehensive Care Program (CCP) for non-ventilator dependent clients who are birth through 20 years of age when rendered by providers who are enrolled with Texas Medicaid as follows:

- Respiratory Therapist
- Physicians
- Home Health Agency

CRCP services are a benefit when provided in the home setting for a client with a chronic underlying respiratory illness, or a newly diagnosed long-term respiratory condition that is currently resulting in a suboptimal respiratory status.

The services provided through this policy are designed to maximize the client or caregiver’s ability to self-manage the client’s disease when the physician deems the client or caregiver will benefit from the expertise of a respiratory care practitioner for the provision of respiratory care or education. Respiratory
therapy care services that do not require the specialty of a certified respiratory care practitioner are not a benefit. The certified respiratory care practitioner’s services allow for the performance of pulmonary care, when required, and the education of the client or caregivers in:

- Disease management.
- Prevention of infections and/or complications.
- Proper use of medications and respiratory equipment which the client is using.

Refer to: Section 3, “CRCP Services” in this handbook for respiratory care practitioner services for clients who are ventilator-dependent.

Providers must use the following procedure codes for in-home respiratory services:

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<td>98960</td>
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<tr>
<td>99503</td>
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</table>

A certified respiratory care practitioner must hold a certificate or temporary permit in compliance with the Texas Occupations Code §604.105a.

A certified respiratory care practitioner must meet the following requirements:

- Be enrolled and approved for participation with Texas Medicaid as an independent practitioner or be employed by a physician, physicians group, or home health agency.
- Submit claims for covered services in the manner and format prescribed by HHSC or its designee
- Comply with all applicable federal, state, and local laws and regulations

Procedure codes S9441, 98960, and 99503 are each limited to once per day, by any provider, and twice per lifetime. Additional visits may be reimbursed when additional prior authorization criteria have been met.

Providers will not be reimbursed for procedure codes S9441, 98960, 99503 or 99504, in any combination, if submitted on the same date of service, by any provider.

Disposable respiratory supplies are a benefit through Texas Medicaid Title XIX Home Health Services and are not reimbursed to the certified respiratory therapist. Retrospective review may be performed to ensure documentation supports the medical necessity of the service when claims are submitted for the procedure codes listed within this handbook.

4.1.1 Authorization Requirements

Prior authorization requests may be submitted to the TMHP Prior Authorization Department via mail, fax, or the electronic portal. Prescribing or ordering providers, dispensing providers, clients’ responsible adults, and clients may sign prior authorization forms and supporting documentation using electronic or wet signatures.

Refer to: Subsection 5.5.1.2, “Document Requirements and Retention” in “Section 5: Fee-for-Service Prior Authorizations” (Vol. 1, General Information) for additional information about electronic signatures.

Prior authorization is required for in-home certified respiratory care practitioner’s services.

To avoid unnecessary denials, the provider must submit correct and complete information, including documentation for medical necessity of the service requested. The provider ordering the service and the provider performing the service must maintain documentation of medical necessity in the client’s medical record. The requesting provider may be asked for additional information to clarify or complete a request for the service.
A completed CRCP Prior Authorization Request Form requesting these services must be signed and dated by the treating physician familiar with the client before requesting prior authorization. A copy of the completed, signed, and dated CRCP Prior Authorization Request Form must be maintained by the provider in the client’s medical record. The completed CRCP Prior Authorization Request Form with the original dated signature must be maintained by the physician in the client’s medical record.

Refer to: CRCP Prior Authorization Request Form on the TMHP website at www.tmhp.com.

To complete the prior authorization process electronically, the provider must complete the prior authorization requirements through any approved electronic methods and retain a copy of the signed and dated CRCP Prior Authorization Request Form in the client’s medical record at the provider’s place of business.

To complete the prior authorization process by paper, the provider must fax or mail the completed CRCP Prior Authorization Request Form to the CCP Prior Authorization Unit and retain a copy of the signed and dated CRCP Prior Authorization Request Form in the client’s medical record at the provider’s place of business.

The following documentation must be submitted to the CCP Prior Authorization Unit for prior authorization:

- A physician’s order
- Client’s primary diagnosis with details of current suboptimal respiratory status and history of more than one emergency room or acute care clinic visits within the last three months
- The services the certified respiratory care practitioner will provide
- Reason this service or education needs to be provided in the home setting and cannot be provided in the office or facility setting. These may include, but are not limited to:
  - Testing of home equipment.
  - Evaluation of the patient or caregiver’s technique with home respiratory care equipment.
  - Evaluation of caregiver’s ability to assess the client’s respiratory status and intervene appropriately if necessary.
  - Home environment assessment.
- The goals of the services to be provided in the home and the estimated length of time to attain these goals

Prior authorization is required for procedure code S9441, and services must be performed by a certified respiratory care practitioner who has been certified by the National Asthma Educator Certification Board (NAECB) as a certified asthma educator. Certification documentation must be provided with the CRCP Prior Authorization Request Form in order to be considered for prior authorization.

Asthma conditions may include, but are not limited to:

- Extrinsic asthma
- Intrinsic asthma
- Chronic obstructive asthma
- Exercise-induced asthma

Prior authorization is required for procedure codes 98960 and 99503. Respiratory conditions may include, but are not limited to:

- Cystic fibrosis
- Obstructive sleep apnea (use of CPAP or BiPAP)
• Chronic respiratory insufficiency

Prior authorization will not be considered for certified respiratory care practitioners to perform routine respiratory treatment or services in the home.

Prior authorization requests for conditions or quantities beyond those limits established in this policy (two per lifetime) will be considered on a case-by-case basis upon review by the Texas Medicaid & Healthcare Partnership (TMHP) Medical Director. The following additional information must be provided:

• Documentation of how the objectives of prior visits have not been achieved to support the need for additional visits beyond those limits established in this policy.
• Reason these additional services need to be provided in the home setting.
• The goals of these services and the estimated length of time to attain these goals.
• The frequency and number of home visits requested by the certified respiratory care practitioner.

5 Documentation Requirements

All supporting documentation must be included with the request for prior authorization. Providers can submit requests to TMHP as follows:

• TMHP Prior Authorization on the Portal: www.tmhp.com/topics/prior-authorization

  Note: The above link is to the TMHP Prior Authorization web page. From the TMHP Prior Authorization web page, click "PA On the Portal" to access the TMHP PA on the Portal application.

• CCP requests, Fax to: 1-512-514-4212
• Non-CCP requests, Fax to: 1-512-514-4213

6 Prescribed Pediatric Extended Care Center (PPECC) Services

CRCP services may be reimbursed when provided in a PPECC setting. When services are rendered in a PPECC setting, the PPECC’s name and NPI number must appear on the professional claim, in addition to the certified respiratory care provider’s NPI. The certified respiratory care practitioner must also indicate outpatient hospital as the place of service when rendering services in a PPECC, or the claim will be denied.

When certified respiratory care services are rendered in a PPECC, the certified respiratory care provider must document coordination with the PPECC.

The PPECC and the certified respiratory care practitioner must have a written agreement for each client related to the provision of respiratory care practitioner services provided at the PPECC. The written agreement must address responsibilities of both parties, and how the parties will coordinate related to the client’s plan of care. The written agreement must be maintained in the client’s medical record.

7 Claims Filing and Reimbursement

7.1 Claims Information

CRCP services must be submitted to the Texas Medicaid & Healthcare Partnership (TMHP) in an approved electronic format or on the CMS-1500 paper claim form. Providers may purchase CMS-1500 paper claim forms from the vendor of their choice. TMHP does not supply the forms.
When completing a CMS-1500 paper claim form, all required information must be included on the claim, as TMHP does not key any information from claim attachments. Superbills, or itemized statements, are not accepted as claim supplements.

**Refer to:** “Section 3: TMHP Electronic Data Interchange (EDI)” (Vol. 1, General Information) for information on electronic claims submissions.

“Section 6: Claims Filing” (Vol. 1, General Information) for general information about claims filing.

Subsection 6.5, “CMS-1500 Paper Claim Filing Instructions” in “Section 6: Claims Filing” (Vol. 1, General Information). Blocks that are not referenced are not required for processing by TMHP and may be left blank.

Electronic billers must submit the prior authorization number (PAN) on the electronic claim form.

Providers should consult the software vendor for the location of this field in the software.

### 7.2 Reimbursement

Respiratory therapy services provided by a participating CRCP are reimbursed the lesser of the provider’s billed charges or the rate calculated in accordance with 1 TAC §355.8089.

Providers can refer to the Online Fee Lookup (OFL) or the applicable fee schedule on the TMHP website at [www.tmhp.com](http://www.tmhp.com).

Texas Medicaid implemented mandated rate reductions for certain services. The OFL and static fee schedules include a column titled “Adjusted Fee” to display the individual fees with all mandated percentage reductions applied. Additional information about rate changes is available on the TMHP website at [www.tmhp.com/resources/rate-and-code-updates/rate-changes](http://www.tmhp.com/resources/rate-and-code-updates/rate-changes).

**Refer to:** Subsection 2.2, “Fee-for-Service Reimbursement Methodology” in “Section 2: Texas Medicaid Fee-for-Service Reimbursement” (Vol. 1, General Information) for more information about reimbursement.

Providers will not be reimbursed for procedure codes 99503 and 99504 on the same date of service, any provider.

The professional service may be billed by the CRCP for services provided in the client’s home (procedure code 99504). The professional service will be allowed once per day up to a limit of 24 visits per year. The recommended frequency for CRCP services is as follows: 7 visits during the first week, a total of 6 visits during the second through fourth weeks, and 11 monthly visits for the second through the 12th month.

Disposable respiratory supplies and respiratory equipment rental or purchase are a home health services benefit and are not reimbursed to the certified respiratory therapist.

**Refer to:** Subsection 2.2, “Services, Benefits, Limitations and Prior Authorization” in the [Durable Medical Equipment, Medical Supplies, and Nutritional Products Handbook](http://www.tmhp.com/resources/policy-handbooks) (Vol. 2, Provider Handbooks) for more information about DME or medical supplies prior authorization information.

Retrospective review may be performed to ensure documentation supports the medical necessity of the service when claims are submitted for the procedure codes listed within this handbook.

The Healthcare Common Procedure Coding System (HCPCS)/Current Procedural Terminology (CPT) codes included in this policy are subject to National Correct Coding Initiative (NCCI) relationships. Any exceptions to NCCI code relationships are specifically noted in the policy. Providers should refer to NCCI for correct coding guidelines and specific applicable code combinations.
DURABLE MEDICAL EQUIPMENT, MEDICAL SUPPLIES, AND NUTRITIONAL PRODUCTS HANDBOOK

TEXAS MEDICAID PROVIDER PROCEDURES MANUAL: VOL. 2

MAY 2021
DURABLE MEDICAL EQUIPMENT, MEDICAL SUPPLIES, AND NUTRITIONAL PRODUCTS

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1 General Information

The information in this handbook is intended for Texas Medicaid durable medical equipment (DME) supplier and medical supply company providers. This handbook provides information about the Texas Medicaid benefits, policies, and procedures that are applicable to these providers.

This handbook contains information about Texas Medicaid fee-for-service benefits. For information about managed care benefits, refer to the Texas Medicaid Managed Care Handbook.

Managed care carve-out services are administered as fee-for-service benefits. A list of all carve-out services is available in section 8, “Carve-Out Services” in the Medicaid Managed Care Handbook (Vol. 2, Provider Handbooks).

All providers are required to report suspected child abuse or neglect as outlined in subsection 1.7.1.2, “Reporting Child Abuse or Neglect” in “Section 1: Provider Enrollment and Responsibilities” (Vol. 1, General Information).

Important: All providers are required to read and comply with “Section 1: Provider Enrollment and Responsibilities” (Vol. 1, General Information). In addition to required compliance with all requirements specific to Texas Medicaid, it is a violation of Texas Medicaid rules when a provider fails to provide health-care services or items to Medicaid clients in accordance with accepted medical community standards and standards that govern occupations, as explained in 1 Texas Administrative Code (TAC) §371.1659. Accordingly, in addition to being subject to sanctions for failure to comply with the requirements that are specific to Texas Medicaid, providers can also be subject to Texas Medicaid sanctions for failure, at all times, to deliver health-care items and services to Medicaid clients in full accordance with all applicable licensure and certification requirements including, without limitation, those related to documentation and record maintenance.

Refer to: “Section 1: Provider Enrollment and Responsibilities” (Vol. 1, General Information) for more information about enrollment procedures.

1.1 Payment Window Reimbursement Guidelines for Services Preceding an Inpatient Admission

According to the three-day and one-day payment window reimbursement guidelines, most professional and outpatient diagnostic and nondiagnostic services that are rendered within the designated timeframe of an inpatient hospital stay, and are related to the inpatient hospital admission, will not be reimbursed separately from the inpatient hospital stay if the services are rendered by the hospital or an entity that is wholly owned or operated by the hospital.

These reimbursement guidelines do not apply in the following circumstances:

- The professional services are rendered in the inpatient hospital setting.
- The hospital and the physician office or other entity are both owned by a third party, such as a health system.
- The hospital is not the sole or 100-percent owner of the entity.

Refer to: Subsection 3.7.4.14, “Payment Window Reimbursement Guidelines” in the Inpatient and Outpatient Hospital Services Handbook (Vol. 2, Provider Handbooks) for additional information about the payment window reimbursement guidelines.
2 Texas Medicaid (Title XIX) Home Health Services

2.1 Enrollment
All DME providers must be Medicare-certified before applying for enrollment in Texas Medicaid.

Providers that render custom DME wheeled mobility systems to Texas Medicaid clients must enroll in Texas Medicaid as a specialized/custom wheeled mobility group provider and must have at least one qualified rehabilitation professional (QRP) performing provider.

Certified QRP providers must enroll in Texas Medicaid as performing providers under DME provider groups.

To enroll in Texas Medicaid as a QRP performing provider, individual professionals must be certified by the National Registry of Rehabilitation Technology Suppliers (NRRTS) or Rehabilitation Engineering and Assistive Technology Society of North America (RESNA) and must enroll as a performing provider under a Specialized /Custom Wheeled Mobility group.

Providers may download the Texas Medicaid Provider Enrollment Application at www.tmhp.com or request a paper application form by contacting Texas Medicaid & Healthcare Partnership (TMHP) directly at 1-800-925-9126.

Providers may also obtain the paper enrollment application by writing to the following address:

Texas Medicaid & Healthcare Partnership
Provider Enrollment
PO Box 200795
Austin, TX 78720-0795
1-800-925-9126
Fax: 1-512-514-4214

Providers may request prior authorization for home health services by contacting:

Texas Medicaid & Healthcare Partnership
Home Health Services
PO Box 202977
Austin, TX 78720-2977
1-800-925-8957
Fax: 1-512-514-4209

2.1.1 Pending Agency Certification
DME providers that submit claims before the enrollment process is complete or without prior authorization for services issued by the TMHP Home Health Services Prior Authorization Department will not be reimbursed. The effective date of enrollment is the date on which all Medicaid provider enrollment forms have been received and approved by TMHP.

Upon the receipt of notice of Medicaid enrollment, the supplier must contact the TMHP Home Health Services Prior Authorization Department before rendering to a Medicaid client, services that require a prior authorization number. Prior authorization cannot be issued before Medicaid enrollment has been completed. Regular prior authorization procedures are followed at that time.

Providers must not submit home health services claims for payment until they have received their Medicaid certification and a prior authorization number has been assigned.

Refer to: Subsection 2.1.1, “Clinical Laboratory Improvement Amendments (CLIA)” in the Radiology and Laboratory Services Handbook (Vol. 2, Provider Handbooks).
2.1.2 Surety Bond Requirements

All newly enrolling and re-enrolling durable medical equipment (DME) providers must, as a condition of enrollment and continued participation into Texas Medicaid, obtain a surety bond that complies with Title 1, Texas Administrative Code (TAC) §352.15.

Important: Surety bonds obtained for the purpose of accreditation in the Medicare program, which lists the Centers for Medicare & Medicaid Services (CMS) as obligee, do not fulfill the surety bond requirement for Texas Medicaid.

The surety bond submitted to Texas Medicaid must meet the following requirements:

- A bond in an amount of no less than $50,000 must be provided for each enrolled location.
  
  Note: Only one surety bond is required if the provider has multiple Medicaid DME provider numbers related to the same location. For example, if the provider has a TPI with a suffix for DME and a second suffix for Specialized Custom Wheeled Mobility all for the same practice location, only one surety bond is required.
  
- The bond must be submitted on the State of Texas Medicaid Provider Surety Bond Form. No other form will be accepted. The use of this form designates the Health and Human Services Commission (HHSC) as the sole obligee of the bond. Instructions are included with the form.
  
- The bond must be issued for a term of 12 months. Bonds for longer or shorter terms are not acceptable.
  
- The bond must be in effect on the date that the provider enrollment application is submitted to TMHP for consideration. The effective date stated on the bond must be:
    - No later than the date that the provider enrollment application is submitted.
    - No earlier than 12 months before the date that the provider enrollment application is submitted.
  
- The bond must be a continuous bond. A continuous bond remains in full force and effect from term to term unless the bond is canceled.
  
Important: An annual bond that specifies effective and expiration dates for the bond, is not acceptable.

At the time of enrollment or re-enrollment, providers must submit the surety bond form with original signatures and a copy of the Power of Attorney document from the surety company that issued the bond.

Note: Surety companies may refer to Texas Department of Insurance (TDI) file #9212562912 or TDI link #132456 when filing the bond.

2.1.2.1 Proof of Continuation

DME providers must maintain a current surety bond to continue participation in Texas Medicaid. Each year, providers must submit documentation that shows proof of continuation of the bond for a new 12-month term. The document may be submitted on the surety bond company’s form and must include the following components:

- Bond number
- Principal’s name, address, and Tax ID or Medicaid provider number (Texas Provider Identifier)
- Surety’s company name and address
- Date of the original bond
- New “good through” date
To avoid losing Medicaid enrollment status, providers must submit the proof of continuation to the TMHP Provider Enrollment before the expiration date of the bond that is currently on file. The completed proof of continuation document must include the original signatures of the authorized corporate representative of the DME provider (principal), and the attorney-in-fact of the surety company. Providers may submit a copy of the proof of continuation (i.e., scan, FAX, photocopy) pending the submission of the original document.

Submission Information

The surety bond can be uploaded to PIMS or submitted to the TMHP Provider Enrollment Department at the following address:

Texas Medicaid & Health Partnership
ATTN: Provider Enrollment
PO Box 200795
Austin, TX 78720-0795
Fax: 1-512-514-4214

2.2 Services, Benefits, Limitations and Prior Authorization

Home health services include home health skilled nursing (SN), home health aide (HHA), physical therapy (PT) and occupational therapy (OT) services; DME; and expendable medical supplies that are provided to eligible Medicaid clients at their place of residence.

Refer to:
- The Physical Therapy, Occupational Therapy, and Speech Therapy Services Handbook (Vol. 2, Provider Handbooks) for more information about therapy services.
- The Home Health Nursing and Private Duty Nursing Services Handbook (Vol. 2, Provider Handbooks) for more information about nursing services.

2.2.1 Home Health Services

The benefit period for home health professional services is up to 60 days with a current plan of care (POC). For all DME and medical supplies with or without prior authorization requirements, providers must complete a Home Health Services (Title XIX) DME/Medical Supplies Physician Order Form except as outlined in subsection 2.2.12, “Diabetic Equipment and Supplies” in this handbook. In chronic and stable situations, the Home Health Services (Title XIX) DME/Medical Supplies Physician Order Form is valid for up to, but no more than, 6 months from the date of the physician’s signature on the form, unless otherwise noted in this handbook. If necessary, DME and supplies that are ordered on a Home Health Services (Title XIX) DME/Medical Supplies Physician Order Form may be prior authorized for up to 6 months with medical necessity determination. Because Medicaid clients have a one-month eligibility period, providers must bill for a one month supply at a time, even though prior authorization may be granted for up to 6 months. This extended prior authorization period begins on the date that clients receive their first prior-authorized home health service. Texas Medicaid allows additional DME or supplies that have been determined to be medically necessary and have been prior authorized by TMHP Home Health Services Prior Authorization Department. Durable medical equipment providers must retain all orders; copies of completed, signed, and dated Title XIX forms; delivery slips; and corresponding invoices for all supplies provided to a client. Durable medical equipment providers must disclose these records to HHSC or its designee on request. These records and claims must be retained for a minimum of five years from the date of service (DOS) or until audit questions, appeals, hearings, investigations, or court cases are resolved. Use of these services is subject to retrospective review.

2.2.1.1 Client Eligibility

Home health clients do not have to be homebound to qualify for services.
To qualify for home health services, the Medicaid client must be eligible on the DOS and must:

- Have a medical need for home health professional services, DME, or supplies that is documented in the client’s POC and considered a benefit under home health services.
- Receive services that meet the client’s existing medical needs and can be safely provided in the client’s home.
- Receive prior authorization from TMHP for most home health professional services, DME, and medical supplies.

Unless otherwise noted in this handbook, certain DME/supplies may be obtained without prior authorization. Durable medical equipment providers must retain a copy of the completed Home Health Services (Title XIX) DME/Medical Supplies Physician Order Form that has been reviewed, signed, and dated by the treating physician for these clients.


2.2.1.2 Prior Authorization Requests for Clients with Retroactive Eligibility

Retroactive eligibility occurs when the effective date of a client’s Medicaid coverage is before the date on which the client’s Medicaid eligibility is added to TMHP’s eligibility file, which is called the “add date.” For clients with retroactive eligibility, prior authorization requests must be submitted after the client’s add date and before a claim is submitted to TMHP.

For services provided to fee-for-service Medicaid clients during the client’s retroactive eligibility period (i.e., the period from the effective date to the add date), prior authorization must be obtained within 95 days of the client’s add date and before a claim for those services is submitted to TMHP. For services provided on or after the client’s add date, the provider must obtain prior authorization within three business days of the date of service.

The provider is responsible and strongly encouraged to verify client eligibility frequently. Eligibility can be verified electronically by checking:

- TexMedConnect.
- The Medicaid Client Portal for Providers.

Additionally, providers can verify eligibility by calling the TMHP Contact Center at 1-800-925-9126.

If services are discontinued before the client’s add date, the provider must still obtain prior authorization within 95 days of the add date to be able to submit claims.

Refer to: “Section 4: Client Eligibility” (Vol. 1, General Information).

2.2.1.3 Prior Authorization

Prior authorization must be obtained for some supplies and most DME from TMHP within three business days of the DOS. Although durable medical equipment providers may supply some DME and medical supplies to a client without prior authorization, they must still retain a copy of the Home Health Services (Title XIX) DME/Medical Supplies Physician Order Form that has Section B completed, signed, and dated by the client’s attending physician, unless otherwise noted in this handbook.

The following prior authorization requests can be submitted on the TMHP website at www.tmhp.com:

- External Insulin Pump
- Home Health Services (Title XIX) DME/Medical Supplies Physician Order Form
- Home Health Services POC
- Statement for Initial Wound Therapy System In-Home Use
- Statement for Recertification of Wound Therapy System In-Home Use
- Wheelchair/Scooter/Stroller Seating Assessment Form (CCP/Home Health Services) (Attachments will be sent separately due to size and detailed information)

Refer to:
- Subsection 5.5.1, “Prior Authorization Requests Through the TMHP Website” in “Section 5: Fee-for-Service Prior Authorizations” (Vol. 1, General Information) for more information, including mandatory documentation requirements.

If a client’s primary coverage is private insurance and Medicaid is secondary, prior authorization is required for Medicaid reimbursement. If the primary coverage is Medicare, Medicare approves the service, and Medicaid is secondary, prior authorization is not required. TMHP will pay only the coinsurance or deductible according to current payment guidelines. If Medicare denied the service, TMHP must receive a prior authorization request within 30 days of the date of Medicare’s final disposition. The Medicare Remittance Advice Notice (MRAN) containing Medicare’s final disposition must accompany the prior authorization request. If the service is a Medicaid-only service, prior authorization is required within three business days of the DOS.

The provider must contact the TMHP Home Health Services Prior Authorization Department within three business days of the DOS to obtain prior authorization for DME and medical supplies.

If inadequate or incomplete information is provided or medical necessity is lacking, the provider will be asked to furnish any required or additional documentation so that a decision about the request can be made. Because the documentation must often be obtained from the client’s physician, providers have two weeks to submit the requested documentation. If the additional documentation is received within the two-week period, prior authorization can be considered for the original date of contact. If the additional documentation is received more than two weeks after the request for the documentation, prior authorization is not considered before the date on which the additional documentation is received. It is the DME supplier’s responsibility to contact the physician to obtain the requested additional documentation. The physician must maintain documentation of medical necessity in the client’s record.

TMHP Home Health Services toll-free number is 1-800-925-8957.

Refer to:
- Subsection 5.5.1, “Prior Authorization Requests Through the TMHP Website” in “Section 5: Fee-for-Service Prior Authorizations” (Vol. 1, General Information) for more information, including mandatory documentation requirements.

If a client’s primary coverage is private insurance and Medicaid is secondary, prior authorization is required for Medicaid reimbursement. If the primary coverage is Medicare, Medicare approves the service, and Medicaid is secondary, prior authorization is not required. TMHP will pay only the coinsurance or deductible according to current payment guidelines. If Medicare denied the service, then Medicaid prior authorization is required. TMHP must receive a prior authorization request within 30 days of the date of Medicare’s final disposition. The Medicare Remittance Advice Notice (MRAN) containing Medicare’s final disposition must accompany the prior authorization request. If the service is a Medicaid-only service, prior authorization is required within three business days of the DOS.

The provider must contact the TMHP Home Health Services Prior Authorization Department within three business days of the DOS to obtain prior authorization for DME and medical supplies.

If inadequate or incomplete information is provided or medical necessity is lacking, the provider will be asked to furnish any required or additional documentation so that a decision about the request can be made. Because the documentation must often be obtained from the client’s physician, providers have two weeks to submit the requested documentation. If the additional documentation is received within the two-week period, prior authorization can be considered for the original date of contact. If the additional documentation is received more than two weeks after the request for the documentation, prior authorization is not considered before the date on which the additional documentation is received. It is the DME supplier’s responsibility to contact the physician to obtain the requested additional documentation. The physician must maintain documentation of medical necessity in the client’s record.

TMHP Home Health Services toll-free number is 1-800-925-8957.

Refer to:
- Subsection 2.2.2.2, “Prior Authorization” in this handbook for DME prior authorization information.

Subsection 2.3.1, “Medicaid Relationship to Medicare” in this handbook.

Client eligibility for Medicaid is for one month at a time. Providers should verify their client’s eligibility every month. Prior authorization does not guarantee payment.

### 2.2.2 Durable Medical Equipment (DME) and Supplies

Texas Medicaid defines DME as:

*Medical equipment or appliances that are manufactured to withstand repeated use, ordered by a physician for use in the home, and required to correct or ameliorate a client’s disability, condition, or illness.*

Since there is no single authority, such as a federal agency, that confers the official status of “DME” on any device or product, HHSC retains the right to make such determinations with regard to Texas Medicaid DME benefits.

Requested DME may be a benefit when it meets the Medicaid definition of DME. The majority of DME and expendable supplies are covered home health services. If a service cannot be provided for a client who is 20 years of age or younger through home health services, these services may be covered through CCP if they are determined to be medically necessary.
To be reimbursed as a home health benefit:

- The client must be eligible for home health benefits.
- The criteria listed for the requested equipment or supply must be met.
- The requested equipment or supply must be medically necessary, and Federal Financial Participation (FFP) must be available.
- The client’s health status would be compromised without the requested equipment or supply.
- The requested equipment or supplies must be safe for use in the home.
- The client must be seen by a physician no more than 6 months prior to the start of service.

For any purchased DME, the DME provider and the client must sign the DME Certification and Receipt Form that is available on the TMHP website at www.tmhp.com before the claim is submitted for payment. The client’s signature on the certification form verifies that the DME is the property of the client. The certification form must include:

- The date that the client received the DME
- The name of the item
- The printed name of the client or primary caregiver
- The printed name of the provider
- The signature of the client or primary caregiver
- The signature of the provider

The provider must maintain a completed copy of the certification form in the client’s record.

The completed, signed, and dated DME Certification and Receipt Form must be submitted to TMHP for claims and appeals for DME that meet or exceed a billed amount of $2,500.00. The form must also be submitted when multiple items that meet or exceed a total billed amount of $2,500.00 are billed for the same DOS. The form is required in addition to obtaining prior authorization, when applicable.

If the DME Certification and Receipt Form is not submitted to TMHP, the claim payment or appeal will be reviewed and will be eligible for recoupment. Incomplete forms will be returned to the provider for correction and resubmission.

TMHP will contact clients that received DME that meets or exceeds a billed amount of $2,500.00 to verify that services were rendered. If the delivery of the equipment cannot be verified by the client, the claim payment will be eligible for recoupment.

DME providers must maintain all Home Health Services (Title XIX) DME/Medical Supplies Physician Order Forms and Addendum to Home Health Services (Title XIX) DME/Medical Supplies Physician Order Forms in client files. To document the item and date of delivery for all DME that is provided to a client, DME providers must also retain the following documentation:

- All delivery slips
- Corresponding invoices
- The completed, signed, and dated DME Certification and Receipt Form

DME providers must disclose this documentation to HHSC or its designee upon request.

The DME must be used for medical or therapeutic purposes, and supplied through an enrolled DME provider in compliance with the client’s POC.
These records and claims must be retained for a minimum of five years from the DOS or until audit questions, appeals, hearings, investigations, or court cases are resolved. Use of these services is subject to retrospective review.

**Note:** All purchased equipment must be new upon delivery to client. Used equipment may be utilized for lease, but when purchased, must be replaced with new equipment.

HHSC/TMHP reserves the right to request the Home Health Services (Title XIX) DME/Medical Supplies Physician Order Form or Addendum to Home Health Services (Title XIX) DME/Medical Supplies Physician Order Form at any time.

DME must meet the following requirements to qualify for reimbursement under Home Health Services:

- The client received the equipment as prescribed by the physician.
- The equipment has been properly fitted to the client or meets the client’s needs.
- The client, the parent or guardian of the client, or the primary caregiver of the client, has received training and instruction regarding the equipment’s proper use and maintenance.

DME must:

- Be medically necessary due to illness or injury or to improve the functioning of a body part, as documented by the physician in the client’s POC or the Home Health Services (Title XIX) DME/Medical Supplies Physician Order Form.
- Be prior authorized by the TMHP Home Health Services Prior Authorization Department for rental or purchase of most equipment. Some equipment does not require prior authorization. Prior authorization for equipment rental can be issued for up to six months based on diagnosis and medical necessity. If an extension is needed, requests can be made up to 60 days before the start of the new prior authorization period with a new Home Health Services (Title XIX) DME/Medical Supplies Physician Order Form.
- Meet the client’s existing medical and treatment needs.
- Be considered safe for use in the home.
- Be provided through an enrolled DME provider or supplier.

**Note:** Texas Health Steps (THSteps)-eligible clients who qualify for medically necessary services beyond the limits of this home health benefit will receive those services through CCP.

DME that has been delivered to the client’s home and then found to be inappropriate for the client’s condition will not be eligible for an upgrade within the first six months following purchase unless there has been a significant change in the client’s condition, as documented by the physician familiar with the client. All adjustments and modifications within the first six months after delivery are considered part of the purchase price.

All DME purchased for a client becomes the Medicaid client’s property upon receipt of the item. This property includes equipment delivered which will not be prior authorized or reimbursed in the following instances:

- Equipment delivered to the client before the physician signature date on the Home Health Services (Title XIX) DME/Medical Supplies Physician Order Form or Addendum to Home Health Services (Title XIX) DME/Medical Supplies Physician Order Form.
- Equipment delivered more than three business days before obtaining prior authorization from the TMHP Home Health Services Prior Authorization Department and meets the criteria for purchase.
Additional criteria:

- A determination as to whether the equipment will be rented, purchased, replaced, repaired, or modified will be made by HHSC or its designee based on the client’s needs, duration of use, and age of the equipment.
- Periodic rental payments are made only for the lesser of either the period of time the equipment is medically necessary, or when the total monthly rental payments equal the reasonable purchase cost for the equipment.
- Purchase is justified when the estimated duration of need multiplied by the rental payments would exceed the reasonable purchase cost of the equipment or it is otherwise more practical to purchase the equipment.
- If a DME/medical supply provider is unable to deliver a prior authorized piece of equipment or supply, the provider should allow the client the option of obtaining the equipment or supplies from another provider.

Items or services are reimbursed at the lesser of:

- The provider’s billed charges
- The published fee determined by HHSC
- Manual pricing as determined by HHSC based on one of the following:
  - The manufacturer’s suggested retail price (MSRP) less 18 percent
  - The provider’s documented invoice cost

If an item is manually priced, providers must submit documentation of one of the following for consideration of purchase or rental with the appropriate procedure codes:

- The MSRP or average wholesale price (AWP), whichever is applicable
- The provider’s documented invoice cost

*Note: Handwritten alterations (crossing out of information or changing values) of the invoice render the invoice invalid.*

### 2.2.2.1 Modifications, Adjustments, and Repairs

Modifications are the replacement of components because of changes in the client’s condition, not replacement because the component is no longer functioning as designed. All modifications and adjustments within the first six months after delivery are considered part of the purchase price.

Modifications to custom equipment may be prior authorized should a change occur in the client’s needs, capabilities, or physical and mental status which cannot be anticipated.

Documentation must include the following:

- All projected changes in the client’s mobility needs
- The date of purchase, and serial number of the current equipment
- The cost of purchasing new equipment versus modifying the current equipment

All modifications within the first six months after delivery are considered part of the purchase price.

Adjustments do not require supplies. Adjustments made within the first six months after delivery will not be prior authorized. Adjustments made within the first six months after delivery are considered part of the purchase price. A maximum of one hour of labor for adjustments may be prior authorized as needed after the first six months following delivery.
Repairs to client-owned equipment may be prior authorized as needed with documentation of medical necessity. Technician fees are considered part of the cost of the repair. Repairs require the replacement of components that are no longer functional. Providers are responsible for maintaining documentation in the client’s medical record specifying the repairs and supporting medical necessity.

A DME repair will be considered based on the age of the item and cost to repair it.

A request for repair of DME must include an itemized estimated cost list from the vendor or DME provider of the repairs. Rental equipment may be provided to replace purchased medical equipment for the period of time it will take to make necessary repairs to purchased medical equipment.

Repairs will not be prior authorized in situations where the equipment has been abused or neglected by the client, client’s family, or caregiver. Routine maintenance of rental equipment is the provider’s responsibility. For clients requiring wheelchair repairs only, the date last seen by physician does not need to be filled in on the Home Health Services (Title XIX) DME/Medical Supplies Physician Order Form.

2.2.2.1.1 Accessories
Equipment accessories including, but not limited to, pressure support cushions, may be prior authorized with documentation of medical necessity.

2.2.2.2 Prior Authorization
Prior authorization is required for most DME and supplies provided through Home Health Services. These services include accessories, modifications, adjustments, and repairs for the equipment.

Providers must submit a completed Home Health Services (Title XIX) DME/Medical Supplies Physician Order Form to the TMHP Home Health Services Prior Authorization Department.

Unless otherwise noted in this handbook, a completed Home Health Services (Title XIX) Durable Medical Equipment (DME) or Medical Supplies Physician Order Form prescribing the DME or supplies must be signed and dated by a physician and by the representative of the DME/Medical Supply provider familiar with the client before requesting prior authorization for all DME equipment and supplies. A current signature and date is valid for no more than 90 days prior to the date of the requested prior authorization or the initiation of service. The completed Home Health Services (Title XIX) DME/Medical Supplies Physician Order Form must include the procedure codes and numerical quantities for services requested.

A copy of the completed, signed, and dated Home Health Services (Title XIX) DME/Medical Supplies Physician Order Form must be maintained by the DME provider. The ordering physician must maintain the completed, originally signed, and dated Home Health Services (Title XIX) DME/Medical Supplies Physician Order Form in the client’s medical record.

To complete the prior authorization process by paper, the provider must fax or mail the completed Home Health Services (Title XIX) DME/Medical Supplies Physician Order Form to the Home Health Services Prior Authorization Department and retain a copy of the completed, signed, and dated form in the client’s medical record at the provider’s place of business.

To complete the prior authorization process electronically, the provider must submit the prior authorization requirements through any approved electronic methods and retain a copy of the completed, signed, and dated Home Health Services (Title XIX) DME/Medical Supplies Physician Order Form in the client’s medical record at the provider’s place of business.

Retrospective review may be performed to ensure that the documentation included in the client’s medical record supports the medical necessity of the requested services.

The date last seen by the physician must be within the past 6 months. The physician’s signature on the Home Health Services (Title XIX) DME/Medical Supplies Physician Order Form is only valid for 90 days before the initiation of services. The requesting provider may be asked for additional information to clarify or complete the request.
Providers must obtain prior authorization within three business days of providing the service by calling the TMHP Home Health Services Prior Authorization Department or faxing the Home Health Services (Title XIX) DME/Medical Supplies Physician Order Form.

To facilitate a determination of medical necessity and avoid unnecessary denials when requesting prior authorization, the physician must provide correct and complete information supporting the medical necessity of the equipment or supplies requested, including:

- Accurate diagnostic information pertaining to the underlying diagnosis or condition as well as any other medical diagnoses or conditions, to include the client’s overall health status.
- Diagnosis or condition causing the impairment resulting in a need for the equipment or supplies requested.

Purchased DME is anticipated to last a minimum of five years, unless otherwise noted, and may be considered for replacement when the time has passed or the equipment is no longer functional or repairable. A copy of the police or fire report, when appropriate, and the measures to be taken to prevent reoccurrence must be submitted.

Prior authorization for equipment replacement is considered within five years of equipment purchase when one of the following occurs:

- There has been a significant change in the client’s condition such that the current equipment no longer meets the client’s needs.
- The equipment is no longer functional and either cannot be repaired or it is not cost-effective to repair.

Replacement of equipment is also considered when loss or irreparable damage has occurred. The following must be submitted with the prior authorization request:

- A copy of the police or fire report, when appropriate
- A statement about the measures to be taken in order to prevent reoccurrence

Payment may be prior authorized for repair of purchased DME. Maintenance of rental equipment (including repairs) is the supplier’s responsibility. The toll-free number for the TMHP Home Health Services Prior Authorization Department is 1-800-925-8957. Requests for repairs must include the cost estimate, reasons for repairs, age of equipment, and serial number.

### 2.2.3 Home Health DME and Supplies Exceptional Circumstances Provision

The Home Health Durable Medical Equipment (DME) and Supplies Exceptional Circumstances provision is made available in accordance with Title 42 Code of Federal Regulations (CFR) §440.70 and Title 1 Texas Administrative Code (TAC) §354.1039. Under the Exceptional Circumstances provision, Texas Medicaid is obligated to consider coverage of medically necessary DME and supplies that are not currently listed as benefits of Texas Medicaid for clients who are 21 years of age or older.

#### 2.2.3.1 Prior Authorization Requests for Home Health DME and Supplies under the Exceptional Circumstances Provision

All Exceptional Circumstances DME and supplies must be prior authorized. The Home Health DME and Supplies Exceptional Circumstances provision is not an available process to pursue for clients who receive prior authorization denials for medical necessity or technical reasons (e.g., missing essential fields, incomplete documentation). Clients that have been denied prior authorization under these circumstances may appeal the decision through the regular fair hearing process.
To request prior authorization for home health DME and supplies under the Exceptional Circumstances provision, providers must submit a written notice to TMHP. The written notice must include:

- Completed copies of all of the necessary forms for the requested home health DME or supplies, such as the Home Health Services (Title XIX) DME/Medical Supplies Physician Order Form, Special Medical Prior Authorization (SMPA) Request Form, Prior Authorization Request for Oxygen Therapy Devices and Supplies, Wound Care Equipment and Supplies Order Form etc. The forms must be signed and dated by the prescribing physician along with a cover letter indicating the forms are being submitted under the Home Health DME and Supplies Exceptional Circumstances provision.

- The client’s specific diagnosis, medical needs and the reasons why they can only be met by the requested home health DME or supply.

- A clear, concise description of the requested DME or supply.

- The manufacturer’s suggested retail price (MSRP) for the requested DME or supply or an invoice documenting the provider’s cost.

- Letters of Medical Necessity (LOMN) from the client’s prescribing physician and other clinical professionals, as appropriate, documenting the alternative measures and alternative DME or supplies that have been tried and have failed to meet the client’s medical needs, or have been ruled out and an explanation of why they have failed or have been ruled out.

**Important:** TMHP may request additional supporting documentation after reviewing the initial request.

### 2.2.3.2 Reimbursement and Billing

TMHP will only consider reimbursement for home health DME and supplies under the Exceptional Circumstances provision if:

- Providers request DME and supplies using the most appropriate procedure code available.

- TMHP approves the prior authorization request.

### 2.2.4 Medical Supplies

Medical supplies are benefits of the Home Health Services Program if they meet the following criteria:

- Unless otherwise noted in this handbook, the representative of the DME/medical supply provider and a physician who is familiar with the client must sign and date a completed Home Health Services (Title XIX) DME/Medical Supplies Physician Order Form that prescribes the DME or supplies before requesting prior authorization for the DME or supplies. A current signature and date is valid for no more than 90 days prior to the date of the requested prior authorization or the initiation of service. The completed Home Health Services (Title XIX) DME/Medical Supplies Physician Order Form must include the procedure codes and numerical quantities for the services requested.

- The provider must contact TMHP within three business days of providing the supplies to the client and obtain prior authorization, if required.

- The durable medical equipment provider must keep all completed copies of the Home Health Services (Title XIX) DME/Medical Supplies Physician Order Forms and Addendum to Home Health Services (Title XIX) DME/Medical Supplies Physician Order Forms on file. The ordering physician must maintain copies of the completed, originally signed, and dated Home Health Services (Title XIX) DME/Medical Supplies Physician Order Form and Addendum to Home Health Services (Title XIX) DME/Medical Supplies Physician Order Forms in their records.

- Providers must retain individual delivery slips or invoices for each DOS that documents the date of delivery for all supplies provided to a client and must disclose them to HHSC or its designee upon request. Documentation of delivery must include one of the following:
• Delivery slip or corresponding invoice signed and dated by client or caregiver, or

• A dated carrier tracking document with shipping date and delivery date must be printed from the carrier’s website as confirmation that the supplies were shipped and delivered. The dated carrier tracking document must be attached to the delivery slip or corresponding invoice.

• The dated delivery slip or invoice must include the client’s full name, the address to which supplies were delivered, and an itemized list of goods that includes the descriptions and numerical quantities of the supplies delivered to the client and the corresponding tracking number from the carrier. This document could also include prices, shipping weights, shipping charges, or other descriptions.

• All claims submitted for medical supplies must include the same quantities or units that are documented on the delivery slip or corresponding invoice and on the Home Health Services (Title XIX) DME/Medical Supplies Physician Order Form. They must reflect the number of units by which each product is measured. For example, diapers are measured as individual units. If one package of 300 diapers is delivered, the delivery slip or corresponding invoice and the claim must reflect that 300 diapers were delivered and not that one package was delivered. Diaper wipes are measured as boxes or packages. If one box of 200 wipes is delivered, the delivery slip or invoice and the claim must reflect that one box was delivered and not that 200 individual wipes were delivered. There must be one dated delivery slip or invoice for each claim submitted for each client. All claims submitted for medical supplies must reflect either one business day before or one business day after the date of service as documented on the delivery slip or corresponding invoice and the same time frame covered by the Home Health Services (Title XIX) DME/Medical Supplies Physician Order Form. The DME Certification and Receipt Form is still required for all equipment delivered.

  Note: These records and claims must be retained for a minimum of five years from the DOS or until audit questions, appeals, hearings, investigations, or court cases are resolved. Use of these services is subject to retrospective review.

• The ordering physician must document medical supplies as medically necessary in the client’s POC or on a completed Home Health Services (Title XIX) DME/Medical Supplies Physician Order Form and Addendum to Home Health Services (Title XIX) DME/Medical Supplies Physician Order Form.

  Note: Client eligibility can change monthly. Providers are responsible for verifying eligibility before providing supplies.

The DOS is the date on which supplies are delivered to the client or shipped by a carrier to the client as evidenced by the dated tracking document attached to the invoice for that date. The provider must maintain the signed and dated records supporting documentation that an item was not billed before delivery. These records are subject to retrospective review.

Refer to: Home Health Services (Title XIX) DME/Medical Supplies Physician Order Form on the TMHP website at www.tmhp.com.

Home Health Services (Title XIX) DME/Medical Supplies Physician Order Form Instructions on the TMHP website at www.tmhp.com.

Subsection 2.7, “Durable Medical Equipment (DME) Supplier (CCP)” in the Children’s Services Handbook (Vol. 2, Provider Handbooks) for specific information about certain DME and medical supplies.

Subsection 2.2.1.1, “Client Eligibility” in this handbook.
2.2.4.1 Supply Procedure Codes

When submitting supplies on the CMS-1500 claim form, itemize the supplies, including quantities, and also provide the Healthcare Common Procedure Coding System (HCPCS) national procedure codes.

Refer to: Subsection 6.3.3, “Procedure Coding” in “Section 6: Claims Filing” (Vol. 1, General Information) for more information about HCPCS procedure codes.

2.2.4.2 Prior Authorization

TMHP must prior authorize most medical supplies. They must be used for medical or therapeutic purposes, and supplied through an enrolled DME provider in compliance with the client’s POC.

Some medical supplies may be obtained without prior authorization; however, the provider must retain a copy of the completed POC or Home Health Services (Title XIX) DME/Medical Supplies Physician Order Form in the client’s file. Unless otherwise noted in this handbook, a completed Home Health Services (Title XIX) DME/Medical Supplies Physician Order Form for medical supplies not requiring prior authorization may be valid for a maximum of six months, unless the physician indicates the duration of need is less. If the physician indicates the duration of need is less than six months, then a new Home Health Services (Title XIX) DME/Medical Supplies Physician Order Form is required at the end of the determined duration of need.

For a list of DME/medical supplies that do not require prior authorization, providers can refer to subsection 2.2.30, “Procedure Codes That Do Not Require Prior Authorization” in this handbook.

Clients with ongoing needs may receive up to six months of prior authorizations for some expendable medical supplies under Home Health Services when requested on a Home Health Services (Title XIX) DME/Medical Supplies Physician Order Form. Providers may deliver medical supplies as ordered on a Home Health Services (Title XIX) DME/Medical Supplies Physician Order Form for up to six months from the date of the physician’s signature. In these instances, a review of the supplies requested by the physician familiar with the client’s condition, and a new Home Health Services (Title XIX) DME/Medical Supplies Physician Order Form is required for each new prior authorization request. Requests for prior authorization can be made up to 60 days before the start of the new prior authorization period. Professional Home Health Services prior authorization requests require a review by the physician familiar with the client’s condition and a physician signature every 60 days when requested on a POC.

Note: These records and claims must be retained for a minimum of five years from the DOS or until audit questions, appeals, hearings, investigations, or court cases are resolved. Use of these services is subject to retrospective review.

2.2.4.3 Cancelling a Prior Authorization

The client has the right to choose his DME/medical supply provider and change providers. If the client changes providers, TMHP must receive a change of provider letter with a new Home Health Services (Title XIX) DME/Medical Supplies Physician Order Form. The client must sign and date the letter, which must include the name of the previous provider and the effective date for the change. The client is responsible for notifying the original provider of the change and the effective date. Prior authorization for the new provider can only be issued up to three business days before the date TMHP receives the change of provider letter and the new Home Health Services (Title XIX) DME/Medical Supplies Physician Order Form.

2.2.5 Augmentative Communication Device (ACD) System

An ACD system, also known as an augmentative and alternative communication (AAC) device system, allows a client with an expressive speech language disorder to electronically represent vocabulary and express thoughts or ideas in order to meet the client’s functional speech needs.

Digitized speech devices and synthesized speech devices are benefits of Texas Medicaid Title XIX Home Health Services.
A digitized speech device, sometimes referred to as a “whole message” speech output device, uses words or phrases that have been recorded by someone other than the ACD system user for playback upon command by the ACD system user.

Providers must use procedure codes E2500, E2502, E2504, and E2506 when billing for a digitized speech device.

A synthesized speech device uses technology that translates a user’s input into device-generated speech using algorithms representing linguistic rules. Users of synthesized speech ACD systems are not limited to prerecorded messages, but can independently create messages as their communication needs dictate. Some synthesized speech devices require the user to make physical contact with a keyboard, touch screen, or other display containing letters.

Providers must use procedure code E2508 when billing for a synthesized speech device.

Other synthesized devices allow for multiple methods of message formulation and multiple methods of device access. Multiple methods of message formulation must include message selection by two or more of the following methods:

- Letters
- Words
- Pictures
- Symbols

Multiple methods of access must include the capability to access the device by direct physical contact with a keyboard or touch screen and one or more of the following indirect selection techniques:

- Joystick/switches
- Head mouse
- Optical head pointer
- Light pointer
- Infrared pointer
- Scanning device
- Morse code

**Note:** ACD systems that do not meet the criteria through Title XIX Home Health Services may be considered for clients who are birth through 20 years of age under CCP.

Providers must use procedure code E2510 when billing for other synthesized speech devices.

Items included in the reimbursement for an ACD system and not reimbursed separately include, but are not limited to, the following:

- ACD
- Basic, essential software (except for software purchased specifically to enable a client-owned computer or personal digital assistant [PDA] to function as an ACD system)
- Batteries
- Battery charger
- Power supplies
- Interface cables
- Interconnects
• Sensors
• Moisture guard
• Alternating current (A/C) or other adapters
• Adequate memory to allow for system expansion within a three-year timeframe
• All basic operational training necessary to instruct the client and family/caregivers in the use of the ACD system
• Manufacturer’s warranty

2.2.5.1 ACD System Accessories
Accessories are a benefit of Texas Medicaid if the criteria for ACD system prior authorization are met and the medical necessity for each accessory is clearly documented in the speech language pathologist (SLP) evaluation.

All accessories necessary for proper use of an ACD system, including those necessary for the potential growth and expansion of the ACD system (such as a memory card), must be included in the initial prescription/Title XIX form. The following accessories for an ACD system may be covered:

• Access devices for an ACD system include, but are not limited to, devices that enable selection of letters, words, or symbols by direct or indirect selection techniques such as optical head pointers, joysticks, and ACD scanning devices.
• Gross motor access devices, such as switches and buttons, may be considered for clients with poor fine motor and head control.
• Fine motor, head control access devices, such as laser or infrared pointers, may be considered for clients with poor hand control and good head control.

Mounting systems are devices necessary to place the ACD system, switches and other access devices within the reach of the client. Mounting devices may be considered for reimbursement when used to attach an ACD system or access device to a wheelchair or table.

A request for prior authorization of a wheelchair mounting device must include the manufacturer name, model, and purchase date of the wheelchair. One additional mounting device, separate from the one included in the system, may be considered for prior authorization for the same client.

Providers must use procedure codes E2512 and E2599 when billing for ACD system accessories.

2.2.5.1.1 Carrying Case
Carrying cases may be considered for separate reimbursement with supporting documentation of medical necessity.

Providers must use procedure code E2599 and modifier U1 when billing for the carrying case. Carrying cases are limited to one every three years.

Carrying cases may be considered for prior authorization. The prior authorization request must include the make, model, and purchase date of the ACD system.

2.2.5.1.2 Nonwarranty Repairs
Nonwarranty repairs of an ACD system may be considered for prior authorization using procedure code V5336 with documentation from the manufacturer explaining why the repair is not covered by the warranty.
2.2.5.1.3 Trial Period
In order to ensure the client’s needs are met in the most cost effective manner and to ascertain the most appropriate system and access device for the client, the ACD system is prior authorized for purchase only after the client has completed a three-month trial period that includes experience with the requested system.

The ACD system for the trial period may be obtained through the rental, the school setting, or another setting determined by the licensed SLP.

In the situation where an ACD system is not available for rental and the client has recent documented experience with the requested ACD system, purchase can be considered.

A trial period is not required when replacing an existing ACD system, unless the client’s needs have changed and another ACD system or access device is being considered.

2.2.5.1.4 Rental
Prior authorization may be provided for rental during this trial period. All components necessary for use of the device, such as access devices, mounting devices, and lap trays, must be evaluated during this trial period.

2.2.5.1.5 Purchase
Purchase of an ACD system may be considered for prior authorization when all of the following ACD system criteria are met:

- The evaluation/re-evaluation includes documentation that the client has had sufficient experience with the requested ACD system through trial, rental, school, or another setting. When the SLP has confirmed the appropriateness of a specific device for the client, the trial/rental period may be cancelled.

- A Home Health Services (Title XIX) DME/Medical Supplies Physician Order Form listing the prescribed ACD system, access device, and accessories (such as a mounting device) must be completed, signed by the physician, and dated.

ACD systems, equipment, and accessories that have been purchased are anticipated to last a minimum of three years.

2.2.5.1.6 Replacement
Prior authorization for replacement may be considered within three years of purchase when one of the following occurs:

- There has been a significant change in the client’s condition such that the current device no longer meets his or her communication needs.

- The ACD system is no longer functional and either cannot be repaired or it is not cost effective to repair.

- Three years have passed and the equipment is no longer repairable.

  Note: Replacements for clients who are birth through 20 years of age that do not meet the criteria above may be considered through CCP.

2.2.5.1.7 Software
Computer software that enables a client’s computer or PDA to function as an ACD system may be covered as an ACD system. Providers must use procedure code E2511 when billing for a speech generating software. Requests for ACD software may be considered for prior authorization if the software is more cost effective than an ACD system.

If an ACD system is more cost effective than adapting the client’s computer or PDA, an ACD system may be prior authorized instead of the ACD software.
Laptop or desktop computers, PDAs, or other devices that are not dedicated ACD systems are not a benefit of Texas Medicaid, because they do not meet the definition of DME.

### 2.2.5.2 Non-Covered ACD System Items

Noncovered items that are not necessary to operate the system and are unrelated to the ACD system or software components are not benefits of Texas Medicaid. These items include, but are not limited to:

- Printer
- Wireless Internet access devices

**Note:** For clients who are 21 years of age or older, requests for ACD systems that do not meet the criteria through Title XIX Home Health Services may be considered under the Texas Medicaid Home Health—Durable Medical Equipment (DME) Exceptional Circumstances process.

### 2.2.5.3 Prior Authorization

Prior authorization is required for ACD systems provided through Home Health Services. The prior authorization also includes all related accessories and supplies. The physician must provide information supporting the medical necessity of the equipment or supplies requested, including:

- Accurate diagnostic information pertaining to the underlying diagnosis or condition and any other medical diagnoses or conditions, including the client's overall physical and cognitive limitations.
- Diagnosis or condition causing the impairment of speech.

Prior authorization for an ACD system and accessories (rental or purchase) must be requested using the following information:

- Medical diagnosis and how it relates to the client’s communication needs.
- Any significant medical information pertinent to ACD system use.
- Limitations of the client’s current communication abilities, systems, and devices.
- Statement as to why the prescribed ACD system is the most effective, including a comparison of benefits using other alternatives.
- Complete description of the ACD system with all accessories, components, mounting devices, or modifications necessary for client use (must include manufacturer’s name, model number, and retail price).
- Documentation that the client is mentally, emotionally, and physically capable of operating the device.
- An evaluation and assessment must be conducted by a licensed SLP in conjunction with other disciplines, such as physical or occupational therapies. The prescribing physician must base the prescription on the professional evaluation and assessment.

The prior authorization request must include the specifications for the ACD system, all component accessories necessary for the proper use of the ACD, and all necessary therapies or training. It is recommended that the preliminary evaluation for an ACD system include the involvement of an occupational therapist or physical therapist to address the client’s seating/postural needs and the motor skills required to utilize the ACD system.

The prescribing physician familiar with the client must review the SLP evaluation of the client’s cognitive and language abilities and base the prescription and treatment plan on the SLP’s recommendations.
An evaluation and assessment by a licensed SLP must be signed and dated before the date on the physician’s prescription or the Title XIX form and include the following information:

- Documentation of medical necessity for an ACD system, including a formal written evaluation performed by a licensed SLP.
- Medical status or condition and medical diagnoses underlying the client’s expressive speech-language disorder that justifies the need for an ACD system.
- Current expressive speech-language disorder, including the type, severity, anticipated course, and present language skills.
- Description of the practical limitations of the client’s current aided and unaided modes of communication.
- Other forms of therapy or intervention that have been considered and ruled out.
- Rationale for the recommended ACD system and each accessory, including a statement as to why the recommended device is the most appropriate and least costly alternative for the client and how the recommended system will benefit the client.
- Documentation that the client possesses the cognitive and physical abilities to use the recommended system.
- Comprehensive description of how the ACD system will be integrated into the client’s everyday life, including home, school, or work.
- Treatment plan that includes training in the basic operation of the recommended ACD system necessary to ensure optimal use by the client (if appropriate, the client’s caregiver) and a therapy schedule for the client to gain proficiency in using the ACD system.
- Description of the client’s speech-language goals and how the recommended ACD system will assist the client in achieving these goals.
- Description of the anticipated changes, modifications, or upgrades with projected time frames of the ACD system necessary to meet the client’s short- and long-term speech-language needs.
- Identification of the assistance or support needed by, and available to, the client to use and maintain the ACD system.
- Statement that the licensed SLP is financially independent of the ACD system manufacturer/vendor.
- Speech- and language- skills assessment that includes the prognosis for speech or written communication.
- Interactional/behavioral and social abilities.
- Capabilities, including intellectual, postural, sensory (visual and auditory), and physical status.
- Motivation to communicate.
- Residential, vocational, and educational setting.
- Alternative ACD system considered with comparison of capabilities.
- Ability to meet projected communication needs, growth potential, and length of time it will meet the client’s needs.
2.2.6  **Bath and Bathroom Equipment**

Bath and bathroom equipment is DME that is included in a treatment protocol, serves as a therapeutic agent for life and health maintenance, and is required to treat an identified medical condition. Bath and bathroom equipment may be considered for reimbursement for those clients who have physical limitations that do not allow for bathing, showering, or bathroom use without assistive equipment.

Bath seats are not considered for clients who are younger than one year of age or weighing less than 30 pounds.

*Note:* For clients who are 21 years of age or older, requests for bath and bathroom equipment that do not meet the criteria through Title XIX Home Health Services may be considered under the Texas Medicaid Home Health—Durable Medical Equipment (DME) Exceptional Circumstances process.

Rental of equipment includes all necessary supplies, adjustments, repairs, and replacement parts.

2.2.6.1  **Hand-Held Shower Wand**

A hand-held shower wand with attachments may be considered for prior authorization only if the client currently owns or meets the criteria for a bath or shower chair, tub stool or bench, or tub transfer bench. Prior authorization of a hand-held shower wand includes all attachments and accessories. Providers must use procedure code E1399 when billing for a hand-held shower wand. Hand-held shower wands with attachments are limited to one every five years.

2.2.6.2  **Bath Equipment**

2.2.6.2.1  **Bath or Shower Chairs, Tub Stool or Bench, Tub Transfer Bench**

A bath or shower chair is a stationary or mobile seat with or without upper body or head support used to support a client who is unable to stand or sit independently in the shower or tub.

Bath/shower chairs are grouped into three levels of design to assist the client based on their physical condition and mobility status:

- Level 1 - standard bath or shower chair is defined as stationary equipment.
- Level 2 - intermediate bath or shower chair is defined as mobile equipment with or without a commode cut out.
- Level 3 - complex bath or shower chair is defined as custom equipment (either stationary or mobile) with or without a commode cut out.

A tub stool or bench is a stationary seat or bench used to support a client who is unable to stand or sit independently in the shower or tub.

A tub transfer bench is a stationary bench that sits in the tub and extends outside the tub. It is used to support a client who is unable to stand or sit independently in the shower or tub and allows the client to scoot or slide over the side of the tub.

Bath or shower chairs, tub stools or benches, and tub transfers are limited to one every five years.

A custom bath or shower chair may be considered for prior authorization only if the client does not also have any type of commode chair.

**Level 1 Group**

A Level 1 device may be considered if the client:

- Is either unable to stand independently or is unstable while standing, or
- Is unable to independently enter or exit the shower or tub due to limited functional use of the upper or lower extremities, and
- Maintains the ability to ambulate short distances (with or without assistive device), or


- Has a condition that is defined as a short-term disability without a concomitant long-term disability (including, but not limited to postoperative status).

Providers must use procedure code E0240 without a modifier when billing for Level 1 group bath or shower chairs.

**Level 2 Group**

A Level 2 device may be considered if the client:

- Has good upper body stability, and
- Has impaired functional ambulation, including, but not limited to, lower body paralysis, osteoarthritis, or
- Is nonambulatory.

The client must have a shower that is adapted for rolling equipment; ramps will not be prior authorized for access to showers.

Providers must use procedure code E0240 and modifier TF (Intermediate Level) when billing for Level 2 group bath or shower chairs.

**Level 3 Group**

A Level 3 device may be considered if the client requires:

- Trunk or head or neck support, or
- Positioning to accommodate conditions, including, but not limited to, spasticity, or frequent and uncontrolled seizures.

Providers must use procedure code E0240 and modifier TG (Complex/high Level) when billing for Level 3 group bath/shower chairs.

A bath or shower chair may be prior authorized for clients who meet the Level 1, 2, or 3 criteria. A Level 3 custom bath or shower chair may be prior authorized only if the client does not also have any type of commode chair. A Level 3 mobile bath or shower chair may be considered for clients who have a shower that is adapted for rolling equipment; ramps will not be prior authorized for access to showers.

A tub stool or bench may be prior authorized for clients who meet the Level 1 criteria. Providers must use procedure code E0245 when billing for a tub stool or bench.

A tub transfer bench may be considered for clients who meet the Level 1 or 2 criteria. Providers must use procedure code E0247 when billing for a tub transfer bench.

A heavy duty tub transfer bench may be considered for clients who meet the Level 1 or 2 criteria and who weigh more than 200 pounds. Providers must use procedure code E0248 when billing for a heavy duty tub transfer bench.

### 2.2.6.3 Bathroom Equipment

#### 2.2.6.3.1 Non-fixed Toilet Rail, Bathtub Rail Attachment, and Raised Toilet Seat

Nonfixed toilet rails are limited to two every five years. A bathtub rail is limited to one every five years.

Raised toilet seats are limited to one every five years. Nonfixed toilet rails and bathtub rail attachments may be considered for prior authorization for a client who has decreased functional mobility and is unable to safely self-toilet or self-bathe without assistive equipment. Raised toilet seats do not require prior authorization. Providers must use procedure code E0243 when billing for non-fixed toilet rails, procedure code E0244 when billing for raised toilet seats, and procedure code E0246 when billing for bathtub rails.
2.2.6.3.2 Toilet Seat Lifts

A toilet seat lift mechanism is designed for the top of the toilet to assist lifting the body from a sitting position to a standing position.

A toilet seat lift mechanism must be prior authorized. To qualify for prior authorization, clients must meet all the following criteria:

- The client must have severe arthritis of the hip or knee or have a severe neuromuscular disease.
- The toilet seat lift mechanism must be a part of the physician’s course of treatment and be prescribed to correct or ameliorate the client’s condition.
- Once standing, the client must have the ability to ambulate.
- The client must be completely incapable of standing up from a regular armchair or any chair in the client’s home.

The client’s difficulty or incapability of getting up from a chair is not sufficient justification for a toilet seat lift mechanism. Almost all clients who are capable of ambulating can get out of an ordinary chair if the seat height is appropriate and the chair has arms.

Prior authorization will be given for either mechanical or powered toilet assist devices, not for both. If a client already owns one or more mechanical toilet-assist devices, a powered toilet seat lift mechanism will not be prior authorized unless there has been a documented change in the client’s condition such that the client can no longer use the mechanical equipment.

Toilet seat lift mechanisms are limited to those types that operate smoothly, can be controlled by the client, and effectively assist a client in standing up and sitting down without other assistance. A toilet seat lift operated by a spring release mechanism with a sudden, catapult-like motion that jolts the client from a seated to a standing position is not a benefit of Texas Medicaid.

Providers must use procedure code E0172 when billing for a toilet seat lift mechanism. A toilet seat lift mechanism is limited to one purchase every five years.

2.2.6.3.3 Commode Chairs and Foot Rests

Commode chairs, foot rests, and replacement commode pails or pans may be considered as benefits, depending on the client’s level of need. The client must meet the criteria for the level of commode chair or foot rest requested.

A commode chair with or without a foot rest may be considered a benefit for the client who also has a stationary bath chair without a commode cutout.

Documentation must support medical necessity for a customized commode chair or the addition of attachments to a standard commode chair.

**Level 1: Stationary Commode Chair**

A Level 1 commode chair is defined as a stationary commode chair with fixed or removable attachments to support the arms.

A stationary commode chair with fixed or removable arms may be considered for prior authorization when the client has a medical condition that results in an inability to ambulate to the bathroom safely (with or without mobility aids).

Providers must use procedure code E0163 or E0165 when billing for a stationary and mobile commode chair.

**Level 2: Mobile Commode Chair**

A Level 2 commode chair is defined as a mobile commode chair with fixed or removable attachments to support the arms.
A mobile commode chair with fixed or removable arms may be considered for prior authorization when the following criteria are met:

- In addition to meeting the criteria for a Level 1 commode chair, the client must be on a bowel program and require a combination commode or bath chair for performing the bowel program and bathing after.
- A mobile commode chair will be considered for reimbursement with prior authorization only if the client does not also have any type of bath chair. If the client meets the criteria for a stationary bath chair, prior authorization of a stationary chair may be considered.

**Level 3: Custom Commode Chair**

A Level 3 commode chair is defined as a custom commode chair with all of the following characteristics:

- Is stationary or mobile
- Has fixed or removable attachments to support the arms, head, neck, or trunk.

A custom stationary or mobile commode chair with fixed or removable arms and head, neck, and/or trunk support attachments may be considered for prior authorization when the following criteria are met:

- In addition to meeting the criteria for a Level 1 or 2 commode chair, the client must have a medical condition that results in an inability to support their head, neck, or trunk without assistance.
- A mobile custom commode chair may be considered for reimbursement only if the client does not also have any type of bath chair.

Providers must use procedure code E0163 or E0165 with modifier TG when billing for a custom stationary or mobile commode chair.

**Extra-wide and Heavy-Duty Commode Chair**

An extra-wide, heavy-duty commode chair is defined as one with a width greater than or equal to 23 inches, and capable of supporting a client who weighs 300 pounds or more.

An extra-wide or heavy-duty commode chair may be considered for prior authorization when the client meets the criteria for a Level 1, 2, or 3 commode chair and weigh 300 pounds or more.

Providers must use procedure code E0168 and the appropriate modifiers when billing for an extra-wide or heavy-duty commode chair. Use modifier TF when billing for a mobile extra-wide, heavy-duty commode chair. Use modifier TG when billing for a custom extra-wide, heavy-duty commode chair.

**Commode Chair With Integrated Seat Lift**

A commode chair with integrated seat lift is designed to assist lifting the body from a sitting position to a standing position.

A commode chair with integrated seat lift mechanism for top of the commode must be prior authorized for clients who meet all the following criteria:

- The client must have severe arthritis of the hip or knee or have a severe neuromuscular disease.
- The client must be completely incapable of standing up from a regular toilet, commode, or any chair in their home.
- The commode chair with integrated seat lift must be a part of the physician’s course of treatment and be prescribed to correct or ameliorate the client’s condition.
• Once standing, the client must have the ability to ambulate independently for a short distance of no more than ten feet.

**Note:** The fact that a client has difficulty or is even incapable of getting up from a chair, particularly a low chair, is not sufficient justification for a seat lift mechanism. Almost all clients who are capable of ambulating can get out of an ordinary chair if the seat height is appropriate and the chair has arms.

Providers must use procedure code E0170 or E0171 when billing for a commode chair with integrated seat lift. The purchase of a commode chair with integrated seat lift is limited to one every five years.

**Replacement Commode Pail or Pan**

Replacement commode pails or pans are a benefit through Title XIX Home Health Services and are limited to one per year. Additional quantities may be considered for prior authorization with documentation of medical necessity.

Providers must use procedure code E0167 when billing for a commode pail or pan.

**Foot Rest**

A foot rest is used to support feet during use of the commode chair.

A foot rest may be considered for prior authorization if the client meets the criteria for a Level 1, 2, or 3 commode chair and the foot rest is necessary to support contractures of the lower extremities of clients who are paraplegic or quadriplegic.

Providers must use procedure code E0175 when billing for a foot rest.

**2.2.6.3.4 Portable Sitz Bath**

Portable sitz baths that fit over commode seats are limited to two per year for clients requiring any of the following:

• Cleaning, irrigation, or pain relief of a perianal wound.
• Relief of pain associated with the pelvic area (hemorrhoids, bladder, vaginal infections, prostate infections, herpes, testicle disorders).
• Muscle toning for bowel and bladder incontinence.

Providers must use procedure codes E0160 or E0161 when billing for portable sitz baths.

**2.2.6.3.5 Bath Lifts**

The purchase of a bath lift is limited to one every five years. The rental of a bath lift is limited to one per month.

The two types of bath lifts that are considered for reimbursement include:

• An outside the tub bath lift which is a portable transfer system used to move a nonambulatory client a short distance from bed or chair to bath and is designed to accommodate the smaller space. This type of lift is either hydraulic or electric and consists of a base with wheels or casters and a sling which can transfer the client in and out of the bath.
• An inside the tub bath lift is a portable transfer system used to lower and raise a nonambulatory client into and out of the bath tub. This type of lift is either hydraulic or electric and consists of a base which adheres to the tub surface using suction cups and a seat that will lower and raise the client into and out of the tub.

Providers must use procedure code E0625 with the appropriate modifier (U1, U2, or U3) if necessary when billing for a bath lift.

The bath lift must be free standing, it cannot be attached to the floor, walls, or ceiling. Home adaptation for use of medical equipment is not a benefit of Home Health Services.
A hydraulic bath lift is for a client who is unable to assist in their own transfers and is operated by the weight or pressure of a liquid.

An electric bath lift is operated by electricity and may be considered when a hydraulic lift will not meet the client’s needs.

A bath lift is not a benefit for the convenience of a caregiver.

There are four levels of bath lifts:

- **Level 1** - an outside the tub bath lift (hydraulic or electric) and must accommodate a client weighing up to 300 pounds. Providers must use procedure code E0625 when billing for the purchase of a Level 1 bath lift.

- **Level 2** - an in-tub bath lift (hydraulic or electric) and must accommodate a client weighing up to 300 pounds. Providers must use procedure code E0625 and the U1 modifier when billing for the purchase of a Level 2 bath lift.

- **Level 3** - a bariatric lift (hydraulic or electric, out of tub type) designed to lift a client weighing greater than 300 pounds. Providers must use procedure code E0625 and the U2 modifier when billing for the purchase of a Level 3 bath lift.

- **Level 4** - a bariatric lift (hydraulic or electric, in tub type) designed to lift a client weighing greater than 300 pounds. Providers must use procedure code E0625 and the U3 modifier when billing for the purchase of a Level 4 bath lift.

A bath lift may be considered for prior authorization if the client:

- Has an inability to transfer to the bathtub or shower independently using assistive devices (including, but not limited to, a cane, walker, bathtub rails).

- Requires maximum assistance by the caregiver to transfer to the bathtub or shower.

- Has bathroom and tub or shower that meets the manufacturer’s recommended depth, width, and height for safe bath lift installation and operation.

Providers must use procedure code E0621 when billing for a lift sling. The purchase of a lift sling is limited to one every five years.

The following are payable procedure codes for bath and bathroom equipment:

<table>
<thead>
<tr>
<th>Procedure Code</th>
<th>Maximum Limitation</th>
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<tbody>
<tr>
<td>E0160</td>
<td>2 per year</td>
</tr>
<tr>
<td>E0161</td>
<td>2 per year</td>
</tr>
<tr>
<td>E0163</td>
<td>1 every 5 years</td>
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<tr>
<td>E0165</td>
<td>1 every 5 years</td>
</tr>
<tr>
<td>E0167</td>
<td>1 per year</td>
</tr>
<tr>
<td>E0168</td>
<td>1 every 5 years</td>
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<tr>
<td>E0170</td>
<td>1 every 5 years</td>
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<tr>
<td>E0244</td>
<td>1 every 5 years</td>
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<tr>
<td>E0245</td>
<td>1 every 5 years</td>
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</tbody>
</table>
2.2.6.4 Prior Authorization

Except as otherwise indicated in this section, prior authorization is required for all bath and bathroom equipment and related supplies, including any accessories, modifications, adjustments, replacements and repairs to the equipment.

Bathroom and toilet lift rentals may be prior authorized during the period of repair up to a maximum of four months per lifetime per client.

Prior authorization will not be considered for modifications, adjustments, or repairs to bath or bathroom equipment delivered to a client’s home and then found to be inappropriate for the client’s condition within the first six months after delivery. This applies unless there is a significant change in the client’s condition that is documented by a physician familiar with the client.

2.2.6.5 Documentation Requirements

2.2.6.5.1 Bath and Bathroom Equipment

To request prior authorization for all bath or bathroom equipment, the following documentation must be provided:

- Accurate diagnostic information pertaining to the underlying diagnosis or condition, including the client’s overall health status, any other medical needs, developmental level, and functional mobility skills and why regular bath or bathroom equipment will not meet the client’s needs.
- The age, height, and weight of the client.
- Assessment of the client’s home to ensure the requested equipment can be safely accommodated.
- Anticipated changes in the client’s needs, including anticipated modifications or accessory needs and the growth potential of any custom shower and bath equipment.

2.2.6.5.2 Toilet Seat Lifts

In addition to the above documentation, the submitted documentation for a toilet seat lift must include an assessment completed by a physician, physical therapist, or occupational therapist that includes all of the following:

- A description of the client’s current level of function without the device
- An explanation why a nonmechanical toilet elevation device, such as toilet rails or elevated toilet seat, will not meet the client’s needs
- Documentation that identifies how the toilet seat lift mechanism will improve the client’s function
- A list of the mobility related activities of daily living (MRADLs) the client will be able to perform with the toilet seat lift mechanism that the client is unable to perform without the toilet seat lift mechanism and how the device will increase the client’s independence
- The client’s goals for use of the toilet seat lift mechanism

<table>
<thead>
<tr>
<th>Procedure Code</th>
<th>Maximum Limitation</th>
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<tbody>
<tr>
<td>E0246</td>
<td>1 every 5 years</td>
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<tr>
<td>E0247</td>
<td>1 every 5 years</td>
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<tr>
<td>E0248</td>
<td>1 every 5 years</td>
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<tr>
<td>E0621</td>
<td>1 per 5 years</td>
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<tr>
<td>E0625</td>
<td>1 purchase every 5 years; 1-month rental</td>
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<tr>
<td>E0630</td>
<td>1 purchase every 5 years; 1-month rental</td>
</tr>
<tr>
<td>E1399</td>
<td>1 every 5 years</td>
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</tbody>
</table>
Supporting documentation must be kept in the client’s record that all appropriate therapeutic modalities (e.g., medication or physical therapy) have been tried and that they failed to enable the client to transfer from a chair to a standing position.

2.2.7 Blood Pressure Devices (Manual and Automated)

Blood pressure devices are benefits in the home setting for self-monitoring when:

- The devices are medically necessary and appropriate.
- The devices are prescribed by a physician.

A manual blood pressure device requires manual cuff inflation with real-time visualization of the results displayed on the manometer and does not require prior authorization for purchase when provided for one of the diagnosis codes listed in the table below. Providers must use procedure code A4660 when billing for a manual blood pressure device.

An automated blood pressure device inflates the cuff manually or automatically, displays the blood pressure results on a small screen, and does not require prior authorization for purchase when provided for one of the diagnosis codes listed in the table below. Providers must use procedure code A4670 when billing for an automated blood pressure device.

Repair of equipment may be considered with documentation of why the equipment needs repair. Providers must use procedure code A4660 when billing for the replacement of other components or repair of equipment.

Finger cuff automated blood pressure devices and ambulatory blood pressure devices for diagnostic purposes are not a benefit of Texas Medicaid.

2.2.7.1 Prior Authorization

Procedure codes A4660 and A4670 do not require prior authorization if they are billed with one of the following diagnosis codes:

<table>
<thead>
<tr>
<th>Diagnosis Codes</th>
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<tbody>
<tr>
<td>I10</td>
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<tr>
<td>I132</td>
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<tr>
<td>I169</td>
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<td>I2692</td>
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<td>I2722</td>
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<tr>
<td>I5082</td>
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<tr>
<td>I953</td>
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<td>N013</td>
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</table>
Manual and automated blood pressure devices should last at least one year and may be considered for replacement after one year has passed. If it is medically necessary to replace nonfunctional and irreparable equipment before one year has passed, providers can submit prior authorization requests with documentation of medical necessity that explains the need for the replacement.

Prior authorization is required in the following situations:

- Another blood pressure device is medically necessary within the same year. Replacement of equipment within the same year as the purchase requires prior authorization. If equipment must be replaced before the end of the anticipated lifespan, the provider must submit a copy of the police or fire report, when appropriate, and the measures that will be taken to prevent reoccurrence.
- The diagnosis code is not in the table above. If the diagnosis code is not one of those listed in the table above, providers must submit a request for the prior authorization of the initial or replacement device and must include all of the documentation necessary to support the medical necessity of the blood pressure device.

2.2.7.2 Hospital-Grade Blood Pressure Devices

Hospital-grade blood pressure devices and their components are benefits of CCP in the home setting for self-monitoring when the equipment is prescribed by a physician. A hospital-grade blood pressure device includes memory for continuous recording, has an alarm system to notify the caregiver of abnormal readings, and is capable of frequent or continuous automatic blood pressure and heart rate monitoring with correction of motion artifact.

Documentation that supports medical necessity of the requested equipment, including the diagnosis, must be maintained in the client’s medical record and is subject to retrospective review.


Providers must use procedure code A9279 with modifier U1 when billing for hospital-grade blood pressure devices.

Hospital-grade blood pressure devices that have been purchased are anticipated to last a minimum of three years and may be considered for replacement when three years have passed or when the equipment is not functional and not repairable.

For clients who are birth through 11 months of age, the rental or purchase of a hospital-grade blood pressure device is a benefit when documentation supports medical necessity and includes an explanation of why the client cannot use a standard automated blood pressure device.

For clients who are 12 months of age and older, the rental or purchase of a hospital-grade blood pressure device is a benefit on a case-by-case basis. Supporting documentation of medical necessity must be provided.
The following indications are recognized by Texas Medicaid for hospital-grade blood pressure devices:

- Hypotension
- Essential hypertension
- Hypertensive heart disease
- Hypertensive renal disease
- Acute pulmonary heart disease
- Chronic pulmonary heart disease
- Cardiomyopathy
- Conduction disorders
- Cardiac dysrhythmias
- Heart failure
- Acute kidney failure
- Chronic kidney disease
- Hydronephrosis
- Vesicoureteral reflux with neuropathy
- Bulbus cordis anomalies and anomalies of cardiac septal closure

All rental costs of the hospital-grade blood pressure device apply toward the purchase price.

2.2.7.3 Components, Replacements, and Repairs

The following may be considered for reimbursement of blood pressure device replacement and repairs with prior authorization:

- Replacement of blood pressure cuffs (procedure code A4663)
- Replacement of other components (procedure code A4660)
- Repairs of the equipment (procedure code A4660)

2.2.7.4 Prior Authorization

Prior authorization is required for the rental or purchase of a hospital-grade blood pressure device. A determination will be made by HHSC or its designee as to whether the equipment will be rented, purchased, repaired, or modified based on the client’s needs, duration of use, and age of the equipment. Repairs and modifications can only be performed on purchased equipment.

Documentation of medical necessity for the hospital-grade blood pressure device must support the client’s need for self-monitoring and address why an automated blood pressure device will not meet the client’s needs. The documentation must include:

- All pertinent diagnoses.
- Initial evaluation.
- Symptoms.
- Duration of symptoms.
- Any recent hospitalizations (within past 12 months).
- Comorbid conditions.
- How frequent or continuous self-monitoring will affect treatment.
• All pertinent laboratory and radiology results.
• Client’s weight.
• A family or caregiver(s) who has an understanding of cause and effect and object permanence and who has agreed to accept the responsibility to be trained to use the hospital-grade monitor.

Prior authorization may be granted for a six-month rental period when the request is submitted with documentation of medical necessity supporting the client’s need for self-monitoring and addressing why an automated blood pressure device will not meet the client’s needs.

Recertification for an additional six-month period may be considered when the physician provides current documentation that supports the ongoing medical necessity for self-monitoring and confirms the client or family is compliant with its use.

A hospital-grade blood pressure device will not be considered for prior authorization of purchase until the client has completed a six-month trial period.

Purchase of a hospital-grade blood pressure device may be prior authorized when all of the following criteria are met:

• The client is 12 months of age or older.
• Documentation of medical necessity supports the client’s need for ongoing self-monitoring and addresses why an automated blood pressure device will not meet the client’s needs.

2.2.7.4.1 Components, Replacements, and Repairs
Replacement of blood pressure cuffs and other components may be considered for purchase with prior authorization and documentation of medical necessity that explains the need for the replacement.

Repair of equipment must be prior authorized when irreparable damage has occurred and documentation exists that supports the need for repair. Repair of equipment will be considered after the factory warranty has expired.

2.2.8 Bone Growth Stimulators
Internal and external bone growth (osteogenic) stimulators are a benefit of Texas Medicaid. Bone growth stimulators are a benefit for skeletally-mature individuals only.

Note: Bone growth stimulators that do not meet criteria for coverage through Title XIX Home Health Services may be considered through Texas Health Steps—Comprehensive Care Program (THSteps-CCP) for clients who are birth through 20 years of age.

Note: For clients who are 21 years of age or older, requests for bone growth stimulators that do not meet the criteria through Title XIX Home Health Services may be considered under the Texas Medicaid Home Health—Durable Medical Equipment (DME) Exceptional Circumstances process.

Electromagnetic bone growth stimulators promote healthy bone growth and repair by low intensity electrical stimulation. Electrical stimulation is provided by implanting low-voltage electrodes within the tissue surrounding the bone (internal) or by external placement of a device that transmits low-voltage currents through the soft tissue to the bone (external).

Ultrasonic bone growth stimulators promote healthy bone growth and repair through low-intensity, pulsed ultrasound waves.

A noninvasive electrical bone growth stimulator (procedure codes E0747 and E0748) and noninvasive ultrasound bone growth stimulator (procedure code E0760) are benefits of Texas Medicaid for DME providers when provided in the home setting. An invasive electrical bone growth stimulator (procedure code E0749) is a benefit of Texas Medicaid for freestanding and hospital-based ambulatory surgical centers when provided in the outpatient setting.
Electrical and ultrasonic bone growth stimulator devices for the treatment of orthopedic and neurosurgical conditions are a benefit for Texas Medicaid clients when the client experiences nonunion of a fracture, requires an adjunct to spinal fusion surgery, or experiences congenital pseudarthrosis.

Nonunion is defined as a fractured bone that fails to heal completely. Diagnosis of nonunion is established when a minimum of six months has passed since the injury and the fracture site shows no progressive signs of healing for a minimum of three months and is not complicated by a synovial pseudoarthrosis. Serial radiographs must confirm that fracture healing has ceased for three months or longer before the client begins treatment with the bone growth stimulator.

### 2.2.8.1 Professional Services

Procedure codes 20974, 20975, and 20979 are a benefit of Texas Medicaid and limited to one per six months. During the six-month limitation period, a subsequent fracture that meets the above criteria for a bone growth stimulator may be reimbursed after the submission of an appeal with documentation of medical necessity that demonstrates the criteria have been met.

### 2.2.8.2 Prior Authorization Criteria and Documentation Requirements for Bone Growth Stimulators

Procedure codes E0747, E0748, E0749, and E0760 require prior authorization. Additional bone growth stimulators may be considered for prior authorization with documentation that supports treatment of a different fracture.

For DME that requires prior authorization, an ordering physician who is familiar with the client must submit a completed, signed, and dated Home Health Services (Title XIX) DME/Medical Supplies Physician Ordering Form prior to requesting authorization. The ordering physician must maintain the complete original Home Health Services (Title XIX) DME/Medical Supplies Physician Ordering Form in the client’s file. The DME provider must maintain a copy of the completed, original Home Health Services (Title XIX) DME/Medical Supplies Physician Ordering Form in the client’s file.

To avoid unnecessary authorization denials, the physician must provide correct and complete information, including documentation for medical necessity of the equipment or supplies prescribed. Either provider may be asked for additional information to clarify or complete a request for the bone growth stimulator.

The ordering physician must maintain all original, completed documentation that supports medical necessity for a bone growth stimulator in the client’s file. The DME provider also must maintain copies of documentation that supports medical necessity for a bone growth stimulator in the client’s file. All documentation is subject to retrospective review.

#### 2.2.8.2.1 Documentation for Noninvasive Electrical Bone Growth Stimulator

Documentation of one of the following is required for prior authorization of the external, electromagnetic bone stimulator (procedure code E0747):

- Nonunions, failed fusions, and congenital pseudarthrosis where there is no evidence of progression of healing for three or more months despite appropriate fracture care.
- Delayed unions of fractures of failed arthrodesis at high risk sites (e.g., open or segmental tibial fractures, carpal navicular fractures).

Documentation must also indicate all of the following:

- Serial radiographs have confirmed that no progressive signs of healing have occurred.
- The fractured gap is 1 cm or less.
- The individual can be adequately immobilized and is likely to comply with non-weight-bearing restrictions.
Documentation of one of the following is required for prior authorization of the external, electromagnetic bone stimulator for spinal application (procedure code E0748):

- One or more failed fusions.
- Grade II or worse spondylolisthesis.
- A multiple-level fusion with extensive bone grafting is required.
- Other risk factors for fusion failure are present, including gross obesity, degenerative osteoarthritis, severe spondylolisthesis, current smoking, previous fusion surgery, previous disc surgery, or gross instability.

2.2.8.2.2 Documentation for Invasive Electrical Bone Growth Stimulators

Documentation of one of the following is required for prior authorization of the surgically implanted bone growth stimulator (procedure code E0749):

- Nonunion of long bone fractures (i.e., clavicle, humerus, radius, ulna, femur,ibia, fibula, and metacarpal, metatarsal, carpal, and tarsal bones). Nonunion of long bone fractures is considered to exist only when serial radiographs have confirmed that fracture healing has ceased for three or more months prior to starting treatment with the bone growth stimulator. Serial radiographs must include a minimum of two sets of radiographs separated by a minimum of 90 days. Each set of radiographs must include multiple views of the fracture site.
- Failed fusion of a joint other than the spine when a minimum of three months has elapsed since the joint fusion was performed.
- Congenital pseudoarthrosis.
- An adjunct to spinal fusion surgery for patients at high risk for pseudoarthrosis due to previously failed spinal fusion at the same site.
- An adjunct to multiple-level fusion, which involves three or more vertebrae (e.g., L3-L5, L4-S1, etc).

2.2.8.2.3 Documentation for Ultrasound Bone Growth Stimulator

Documentation of the following is required for prior authorization of the external, low-intensity ultrasound bone growth stimulator device (procedure code E0760):

- Nonunion of a fracture, other than the skull or vertebrae, in a skeletally mature person, which is documented by a minimum of two sets of radiographs that were:
  - Obtained prior to starting treatment with the bone growth stimulator.
  - Separated by a minimum of 90 days.
  - Taken with multiple views of the fracture site.
  - Accompanied by a written interpretation by a physician who states that there has been no clinically significant evidence of fracture healing between the two set of radiographs.
- Evidence of all of the following:
  - The fracture is not tumor-related.
  - The fracture is not fresh (less than seven days), closed or grade I open, tibial diaphyseal fractures, or closed fractures of the distal radius (Colles fracture).
2.2.8.3 Claims Reimbursement for Professional Services

Professional claims that are submitted for bone growth stimulation (procedure codes 20974, 20975, and 20979) may be reimbursed if the claim includes documentation of one of the following:

- Documentation of medical necessity as outlined in subsection 2.2.8.2, “Prior Authorization Criteria and Documentation Requirements for Bone Growth Stimulators” in this handbook.
- The corresponding bone growth stimulator device was submitted within 95 days of the date the bone growth stimulation procedure was performed.

The appropriate evaluation and management (E/M) procedure code must be billed for monitoring the effectiveness of bone growth stimulation treatment.

2.2.9 Breast Feeding Support Services

Refer to: Section 3, “Breastfeeding Support Services” in the Gynecological, Obstetrics, and Family Planning Title XIX Services Handbook (Vol. 2, Provider Handbooks) for information about breastfeeding support services.

2.2.10 Cochlear Implants

Cochlear implant services (procedure codes L8499, L8615, L8616, L8617, L8618, L8619, L8623, and L8624) may be reimbursed in the home setting to DME providers.

Refer to: Subsection 9.2.22, “Cochlear Implants” in the Medical and Nursing Specialists, Physicians, and Physician Assistants Handbook (Vol. 2, Provider Handbooks) for more information about cochlear implant services.

2.2.11 Continuous Passive Motion (CPM) Device

A CPM device is reimbursed on a daily basis and is limited to once per day. Reimbursement includes delivery, set-up and all supplies. Providers must use procedure code E0935 when billing for a CPM machine.

2.2.11.1 Prior Authorization

A CPM device may be considered for prior authorization through Home Health Services. Reimbursement for a CPM device is considered after joint surgery, such as knee replacement, when prescribed by a physician and submitted with clinical documentation of medical necessity and appropriateness.

2.2.12 Diabetic Equipment and Supplies

Diabetic equipment and supplies are a benefit through Title XIX Home Health Services and do not require prior authorization unless otherwise specified.

Note: For clients who are 21 years of age or older, requests for diabetic equipment and supplies that do not meet the criteria through Title XIX Home Health Services may be considered under the Texas Medicaid Home Health—Durable Medical Equipment (DME) Exceptional Circumstances process.

Diabetic equipment and supplies may be obtained through one of the following methods:

- A Home Health Services (Title XIX) DME/Medical Supplies Physician Order Form prescribing the DME or medical supplies. The Title XIX Form must be signed and dated by the prescribing physician who is familiar with the client prior to supplying any medical equipment or supplies.
- A verbal or a detailed written order provided by a physician, physician assistant (PA), nurse practitioner (NP), clinical nurse specialist (CNS), or a certified nurse midwife (CNM).
2.2.12.1 Obtaining Equipment and Supplies Through a Title XIX Form

The completed Title XIX Form must be maintained by the dispensing provider and the prescribing physician in the client’s medical record. The physician must maintain the original signed and dated copy of the Title XIX Form. The completed Title XIX Form is valid for a period up to six months from the physician’s signature date.

2.2.12.2 Obtaining Equipment and Supplies Through a Verbal or Detailed Written Order

If the dispensing provider does not have a detailed written order then a verbal order is required to be on file until the written order is received from the prescribing provider and before providing diabetic equipment and supplies. The prescribing provider’s order may be a written, fax, electronic, or verbal order and must include:

- A description of the item(s).
- The client’s name.
- The name of the physician or authorized prescribing provider.
- The date of the order.

A detailed written order must be received by the DME supplier within 90 days from the date of the prescribing provider’s signature. The detailed written order for diabetic equipment and supplies is valid for six months from the date of the order or the date of the prescribing provider’s signature, whichever is earlier, for initial orders, and from the start date of renewal orders. In the absence of a start date, then the authorized prescribing signature date will be the beginning date of service.

A completed, detailed written order must be signed and dated by the authorized prescribing provider. The prescribing provider is required to retain a copy of the signed and dated detailed written order in the client’s medical record. The DME provider must retain the original, faxed, photocopied, or electronic, signed and dated detailed written order in the client’s medical record.

A completed detailed written order must contain all the following components:

- The client’s name
- The date of the verbal order if different from the date the authorized prescribing provider signed the written order
- Description of item(s) to be provided
- Quantity to dispense (quantity required per day or month)
- Diagnosis code or description supporting the medical necessity

Before submitting a claim to Texas Medicaid, DME providers must have on file a detailed written order with the required information. No other documentation is required.

Prior Authorization

Prior authorization, when necessary, may be considered with documentation of medical necessity, which must include one of the following:

- A completed Title XIX Form that has been signed and dated by the physician who is familiar with the client
- Or all the following:
  - A completed and signed detailed written or verbal order.
  - A Title XIX Form with section A completed.
### 2.2.12.3 Glucose Testing Equipment and Other Supplies

The prescribing provider must indicate on a completed, signed and dated Title XIX Form, or a signed and dated detailed written order how many times a day the client is required to test blood glucose or ketone levels when applicable (not all supplies are related to testing glucose or urine, e.g., batteries).

Glucose tablets or gel (procedure code A9150) may be considered with prior authorization when provided to a client with a diagnosis from the diagnosis code table below. Procedure code A9150 is limited to one per six months.

The procedure codes for the diabetic supplies listed in the following table do not require prior authorization, up to the quantities listed in the table, when provided to a client with a diagnosis from the diagnosis code table below. These limitations are not dependent on the client’s use of insulin:

<table>
<thead>
<tr>
<th>Procedure Code</th>
<th>Limitation</th>
</tr>
</thead>
<tbody>
<tr>
<td>A4233</td>
<td>1 per 6 months</td>
</tr>
<tr>
<td>A4234</td>
<td>1 per 6 months</td>
</tr>
<tr>
<td>A4235</td>
<td>1 per 6 months</td>
</tr>
<tr>
<td>A4236</td>
<td>1 per 6 months</td>
</tr>
<tr>
<td>A4252</td>
<td>10 strips per month</td>
</tr>
<tr>
<td>A4256</td>
<td>2 per year</td>
</tr>
<tr>
<td>A4258</td>
<td>2 per year</td>
</tr>
</tbody>
</table>

### Insulin-Dependent Clients

The following procedure codes for diabetic supplies do not require prior authorization up to the quantities listed when the supplies are provided to an insulin-dependent client with a valid diagnosis. If the client is insulin-dependent, providers must submit claims for these procedure codes with modifier U9:

<table>
<thead>
<tr>
<th>Procedure Code</th>
<th>Limitation</th>
</tr>
</thead>
<tbody>
<tr>
<td>A4253*</td>
<td>2 boxes per month</td>
</tr>
<tr>
<td>A4259</td>
<td>1 box per month</td>
</tr>
<tr>
<td>A9275*</td>
<td>2 per month</td>
</tr>
</tbody>
</table>

*A client may receive a combined total of two per calendar month of procedure codes A4253 and A9275, either two or one procedure code or one of each procedure code

### Non-Insulin-Dependent Clients

The following procedure codes for diabetic supplies do not require prior authorization up to the quantities listed when they are provided to a non-insulin-dependent client with a valid diagnosis. If the client is not insulin-dependent, providers must submit claims for these procedure codes with no modifier:

<table>
<thead>
<tr>
<th>Procedure Code</th>
<th>Limitation</th>
</tr>
</thead>
<tbody>
<tr>
<td>A4253*</td>
<td>1 box per month</td>
</tr>
<tr>
<td>A4259</td>
<td>1 box every 2 months</td>
</tr>
<tr>
<td>A9275*</td>
<td>1 per month</td>
</tr>
</tbody>
</table>

*A client may receive only one per calendar month of either procedure code A4253 or A9275.*
The following diagnosis codes apply to the tables listed above:

<table>
<thead>
<tr>
<th>Diabetic Diagnosis Codes</th>
</tr>
</thead>
<tbody>
<tr>
<td>E0800</td>
</tr>
<tr>
<td>E08319</td>
</tr>
<tr>
<td>E083299</td>
</tr>
<tr>
<td>E083399</td>
</tr>
<tr>
<td>E083499</td>
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<tr>
<td>E083529</td>
</tr>
<tr>
<td>E083549</td>
</tr>
<tr>
<td>E083599</td>
</tr>
<tr>
<td>E0841</td>
</tr>
<tr>
<td>E08610</td>
</tr>
<tr>
<td>E08641</td>
</tr>
<tr>
<td>E0910</td>
</tr>
<tr>
<td>E093212</td>
</tr>
<tr>
<td>E093312</td>
</tr>
<tr>
<td>E093412</td>
</tr>
<tr>
<td>E093512</td>
</tr>
<tr>
<td>E093532</td>
</tr>
<tr>
<td>E093552</td>
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<td>E0937X1</td>
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<td>E0943</td>
</tr>
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<td>E09621</td>
</tr>
<tr>
<td>E0969</td>
</tr>
<tr>
<td>E10311</td>
</tr>
<tr>
<td>E103293</td>
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<tr>
<td>E103493</td>
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<tr>
<td>E103523</td>
</tr>
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<td>E103543</td>
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<td>E103593</td>
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<td>E1059</td>
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<tr>
<td>E10638</td>
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<td>E1101</td>
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<td>E113212</td>
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<tr>
<td>E113312</td>
</tr>
<tr>
<td>E113412</td>
</tr>
<tr>
<td>E113512</td>
</tr>
<tr>
<td>E113532</td>
</tr>
<tr>
<td>E113552</td>
</tr>
</tbody>
</table>
Non-diabetic Diagnosis Codes

<table>
<thead>
<tr>
<th>Non-diabetic Diagnosis Codes</th>
</tr>
</thead>
<tbody>
<tr>
<td>E161</td>
</tr>
<tr>
<td>E7132</td>
</tr>
</tbody>
</table>

Note: THSteps-eligible clients who qualify for medically necessary services beyond the limits of this home health benefit will receive those services through CCP.

Alcohol wipes (procedure code A4245) and urine test or reagent strips or tablets (procedure code A4250) are a benefit of Texas Medicaid when they are necessary for the treatment of some diabetic conditions or other conditions and therefore are not limited to the diagnoses listed in the diagnosis code table above.

Procedure code A4245 is limited to four boxes per month and procedure code A4250 is limited to one box per six months. Prior authorization is not required for these procedure codes up to the quantities listed.

The quantity of glucose testing supplies billed for a one-month supply should relate to the number of tests ordered per day by the prescribing provider.

Glucose testing supplies may be reimbursed for the quantities prescribed or the quantity prior authorized.

Blood glucose test or reagent strips (procedure code A4253) and home glucose disposable monitors with test strips (procedure code A9275) are limited to a combined total of two per month.

2.2.12.3.1 Prior Authorization

Glucose tablets or gel (procedure code A9150) requires prior authorization with documentation supporting medical necessity.
Glucose testing supplies for quantities beyond the limits listed in the procedure code table above or for diagnoses other than those listed in the diagnosis code table above in subsection 2.2.12.3, “Glucose Testing Equipment and Other Supplies” in this handbook may be considered for prior authorization with documentation of medical necessity. Quantities will be prior authorized based on the documentation of medical necessity related to the number of tests ordered per day by the physician.

2.2.12.4 Blood Glucose Monitors

Blood glucose monitors with integrated voice synthesizers (procedure code E2100) and blood glucose monitors with integrated lancing blood sample (procedure code E2101) may be considered for prior authorization with documentation of medical necessity. Glucose monitors that have been purchased are anticipated to last a minimum of three years and may be considered for replacement when three years have passed or the equipment is no longer repairable.

Standard home glucose monitors (procedure code E0607) are not a benefit of Texas Medicaid.


2.2.12.5 Therapeutic Continuous Glucose Monitors

Therapeutic continuous glucose monitor (CGM) (procedure code K0554) and its related supplies (procedure codes K0553) are used to measure glucose levels in real-time throughout the day and night. A tiny electrode called a glucose sensor is inserted under the skin to measure glucose levels in tissue fluid. It is connected to a transmitter that sends the information via wireless radio frequency to a monitoring and display device.

A therapeutic CGM device is used as a replacement for self-blood glucose monitoring (SBGM), for clients with diabetes to use at home.

Therapeutic CGM may be considered as a Medicaid DME benefit with prior authorization of medical necessity. When a therapeutic CGM device (procedure code K0554) is covered, the related supplies (procedure code K0553) may also be covered.

Procedure codes K0553 and K0554 will be a benefit in the home setting when services are provided by home health DME and medical supplier (DME) providers.

The short-term CGM is used for diagnostic purposes to assist the clinician in establishing or modifying the client’s treatment plan; it is a distinct and separate Medicaid benefit in Texas.

Other home glucose monitors (procedure codes E2100 and E2101) will be denied when submitted within three calendar years of procedure code K0554.

The following procedure codes for an SBGM and its related supplies will be denied when submitted during the same calendar month by any provider, as procedure codes K0553 and K0554:

<table>
<thead>
<tr>
<th>Procedure Codes</th>
</tr>
</thead>
<tbody>
<tr>
<td>A4233</td>
</tr>
<tr>
<td>E2101</td>
</tr>
</tbody>
</table>

The supply allowance (procedure code K0553) for supplies used with the therapeutic CGM system encompasses all items necessary for the use of the device. The DME provider is responsible for delivering the appropriate items and quantities for the client to initiate and continue use of the therapeutic CGM.
2.2.12.5.1 Prior Authorization
The initial order from a health-care provider who manages the client’s diabetes is valid for the first six months. If the client continues to be compliant with the use of the CGM and treatment plan, the physician may write an order for an additional six months. After the first year, an order for replacement sensors, transmitter, and receiver (following frequency rules) may be written for a 12-month period.

Therapeutic CGM Device
Therapeutic CGM must be prescribed by a health care provider who manages the client’s diabetes, and may be considered for prior authorization for clients who have type 1 or 2 diabetes. All of the following medical necessity criteria are met:

- The client has been using a SBGM and performing frequent (at least four times per day) testing.
- The client utilizes insulin injections three or more times per day or is on an insulin pump.
- The client’s insulin treatment regimen requires frequent adjustments, based on SBGM or CGM testing results.
- The client is able or has a caregiver who is able to learn to use the device, and to or view CGM alerts and respond appropriately.

A client with hypoglycemia unawareness or several episodes of hypoglycemia a day may also qualify for therapeutic CGM services if the client does not meet the above listed criteria.

The ordering provider should also verify that the client’s condition meets the manufacturers’ recommendations for appropriate age range, testing and calibration requirements, prior to prescribing the CGM device.

CGM devices that have been purchased are expected to last a minimum of three years and may be considered for replacement when three years have passed or the equipment is no longer repairable. The replacement of the equipment may also be considered when it has been lost or irreparably damaged. A copy of the police or fire report, when appropriate, and the measures to be taken to prevent a reoccurrence must be submitted. Additional services may be reimbursed with prior authorization based on documentation of medical necessity.

Associated Supplies
When a therapeutic CGM device (procedure code K0554) is covered, the related supplies (procedure code K0553) may also be covered.

A one-time prior authorization for the initiation of a therapeutic CGM monthly supplies (procedure code K0553) is only required when the client already owns the device. The physician must submit a statement with prior authorization request, verifying the following:

- The client owns a therapeutic CGM device (as defined by CMS).
- The client’s current condition meets the criteria for the therapeutic CGM coverage criteria.
- The client is compliant with using the CGM device to manage his or her diabetes.

CGM devices covered in this policy are defined as therapeutic CGMs, which is a device used as a replacement of SBGM for diabetes treatment decisions. Therapeutic CGM services replace any home blood glucose monitor used for SBGM and related supplies.

2.2.12.5.2 Claims Filing Requirements
Procedure codes K0554 and K0553 must be submitted with modifier KF when submitting a claim for a class III device, as designated by the U.S. Food and Drug Administration (FDA), and its related supplies. No modifier is required when submitting a claim for a class II device, as designated by the FDA, and its related supplies.
2.2.12.5.3 Non Covered Services

The following services are not benefits of Texas Medicaid:

- Non-therapeutic CGM devices used as an adjunct to SBGM
- Rental of therapeutic CGM devices
- Non-medical items, even if the items may be used to serve a medical purpose:
  - Smart devices (smart phones, tablets, personal computers, etc.) used as CGM monitors.
  - Medical supplies used with non-covered equipment. An exception would be for the transmission and receiving of data, using a smart device application, from a client’s personally-owned smart device, who meet the medical criteria for telemonitoring services.

2.2.12.5.4 Prior Authorization

Blood glucose monitors with special features (procedure code E2100 or E2101) may be considered for prior authorization with documentation supporting medical necessity for the special feature requested.

Purchase of a blood glucose monitor with integrated voice synthesizer (procedure code E2100) may be considered for prior authorization with documentation that includes a diagnosis of diabetes and significant visual impairment.

Purchase of a blood glucose monitor with integrated lancing and blood sample (procedure code E2101) may be considered for prior authorization with documentation that includes a diagnosis of diabetes and significant manual dexterity impairment related but not limited to neuropathy, seizure activity, cerebral palsy, or Parkinson’s disease.

2.2.12.6 * External Insulin Pump and Supplies

An external insulin infusion pump is a programmable, battery-powered mechanical syringe or reservoir device controlled by a microcomputer to provide a basal continuous subcutaneous insulin infusion (CSII) and release a "bolus" dose at meals and at programmed intervals. The pump is connected to an infusion set with an attached small needle or cannula that is inserted into the subcutaneous tissue. The purpose of the insulin pump is to provide an accurate, continuous, controlled delivery of insulin which can be regulated by the user to achieve intensive glucose control and prevent the metabolic complications of hypoglycemia, hyperglycemia and diabetic ketoacidosis. The typical external insulin pump capacity is two to three days of insulin.

Note: External insulin pumps that do not require tubing may be considered for clients who are birth through 20 years of age.

An external insulin pump must be ordered by, and the client’s follow-up care must be managed by, a prescribing provider with experience managing clients with insulin infusion pumps and who is knowledgeable in the use of insulin infusion pumps.

The external insulin pump (procedure code E0784) may be considered for prior authorization with documentation of medical necessity. Procedure code E0784 is limited to one purchase every three years, and one rental per month. External insulin pumps that have been purchased are anticipated to last a minimum of three years and may be considered for replacement when three years have passed or the equipment is no longer repairable.

The following procedure codes for external insulin pump supplies are a benefit through Title XIX Home Health Services and do not require prior authorization up the maximum quantities allowed. Additional quantities may be considered with documentation of medical necessity and prior authorization.

<table>
<thead>
<tr>
<th>Procedure Code</th>
<th>Limitation</th>
</tr>
</thead>
<tbody>
<tr>
<td>A4224</td>
<td>4 per month</td>
</tr>
<tr>
<td>A4225</td>
<td>15 per month</td>
</tr>
</tbody>
</table>

Note: External insulin pumps that do not require tubing may be considered for clients who are birth through 20 years of age.
[Revised] Providers must bill replacement batteries (procedure codes K0601, K0602, K0603, K0604, and K0605 with modifier U1).

When there is not an appropriate procedure code for supplies providers may request prior authorization using procedure code A9900.

The external insulin pump supplies (including batteries) are not included in the external insulin pump rental. Routine maintenance of rental equipment is the provider’s responsibility.

Infusion sets for the external insulin pump (procedure codes A4230 or A4231) are limited to clients with a previously billed external insulin pump device or supply. Infusion sets for clients who did not receive the external insulin pump through Texas Medicaid are considered for reimbursement on appeal with a physician’s statement documenting medical necessity.

An internal insulin pump will not be prior authorized as it is considered part of the surgery to place the pump.

2.2.12.6.1 Prior Authorization

Prior authorization is required for an external insulin pump (procedure code E0784) with carrying cases.

Rental of External Insulin Pump

An external insulin pump may be considered for prior authorization of rental with submission of clinical documentation indicating one of the following:

- A client who has a diagnosis of type 1 or 2 diabetes must meet at least two of the following criteria while on multiple daily injections of insulin:
  - Elevated glycosylated hemoglobin level (HbA1c) > 7.0 percent
  - History of dawn phenomenon with fasting blood sugars frequently exceeding 200 mg/dl
  - History of severe glycemic excursions with wide fluctuations in blood glucose
  - History of recurring hypoglycemia (less than 60 mg/dL) with or without hypoglycemic unawareness
  - Anticipation of pregnancy within three months
- A client with a diagnosis of gestational diabetes must meet at least one of the following criteria:

<table>
<thead>
<tr>
<th>Procedure Code</th>
<th>Limitation</th>
</tr>
</thead>
<tbody>
<tr>
<td>A4230</td>
<td>15 per month</td>
</tr>
<tr>
<td>A4231</td>
<td>15 per month</td>
</tr>
<tr>
<td>A4232</td>
<td>10 per month</td>
</tr>
<tr>
<td>A4601</td>
<td>1 per 6 months</td>
</tr>
<tr>
<td>A4602</td>
<td>1 per 6 months</td>
</tr>
<tr>
<td>A6257</td>
<td>30 per month</td>
</tr>
<tr>
<td>A6258</td>
<td>30 per month</td>
</tr>
<tr>
<td>A6259</td>
<td>15 per month</td>
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<tr>
<td>K0601</td>
<td>4 per 2 months</td>
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<tr>
<td>K0602</td>
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<td>K0603</td>
<td>4 per 2 months</td>
</tr>
<tr>
<td>K0604</td>
<td>1 per 6 months</td>
</tr>
<tr>
<td>K0605</td>
<td>1 per 6 months</td>
</tr>
</tbody>
</table>
• Erratic blood sugars in spite of maximal compliance and split dosing
• Other evidence that adequate control is not being achieved by current methods

In addition to the clinical documentation the provider must submit the External Insulin Pump form indicating:

• The client or caregiver possess the following competencies:
  • The cognitive and physical abilities to use the recommended insulin pump treatment regimen
  • An understanding of cause and effect
  • The willingness to support the use of the external insulin pump

• The prescribing provider must attest that:
  • A training/education plan will be completed prior to initiation of pump therapy.
  • The client or caregiver will be given face-to-face education and instruction and will be able to demonstrate proficiency in integrating insulin pump therapy with their current treatment regimen for ambient glucose control.

**Purchase of External Insulin Pump**

An external insulin pump may be considered for prior authorization of purchase after it has been rented for a three-month trial and all of the following documentation is provided:

• The training/education plan has been completed
• The pump is the appropriate equipment for the specific client
• The client is compliant with the use of the pump

### 2.2.12.7 Tubeless External Insulin Infusion Pumps

The tubeless external insulin infusion pump and supplies are a benefit of Texas Medicaid for clients who are birth through 20 years of age.

The tubeless external insulin pump must be ordered by, and the client’s follow-up care must be managed by, a prescribing provider who has experience managing clients with insulin infusion pumps and who is knowledgeable in the use of insulin infusion pumps.

Providers must use procedure code E0784 and modifier U1 for the rental or purchase of the tubeless external insulin pump and procedure code A9274 for the tubeless external insulin pump supplies.

Procedure code A9274 is limited to 15 per month.

A tubeless external insulin pump that has been purchased is expected to last a minimum of three years and may be considered for replacement when three years have passed or the equipment is no longer repairable. The replacement of the equipment may also be considered when it has been lost or irreparably damaged. A copy of the police or fire report, when appropriate, and the measures to be taken to prevent a reoccurrence must be submitted. Additional services may be considered based on documentation of medical necessity.

Routine maintenance of rental equipment is the provider’s responsibility.

### 2.2.12.7.1 Prior Authorization and Documentation Requirements

Prior authorization is required for the tubeless external insulin pump with carrying cases and related supplies and repairs. The tubeless external insulin pump supplies may be considered separately when a tubeless external insulin pump is rented.
The tubeless external insulin pump and supplies may be obtained through one of the following methods:

- **CCP Prior Authorization Request Form**—the completed CCP Prior Authorization Request Form must be maintained by the dispensing provider and the prescribing physician in the client’s medical record. The physician must maintain the original signed and dated copy of the CCP Prior Authorization Request Form. The completed CCP Prior Authorization Request Form is valid for a period up to six months from the physician’s signature date.

- **Verbal or detailed written order**—the verbal or detailed written order must be provided by a physician, PA, NP, CNS, or a CNM.

If the dispensing provider does not have a detailed written order, a verbal order is required to be on file until the written order is received from the prescribing provider and before providing diabetic equipment and supplies. The prescribing provider’s order may be a written, fax, electronic, or verbal order and must include:

- A description of the item(s).
- The client’s name.
- The name of the physician or authorized prescribing provider.
- The date of the order.

A detailed written order must be received by the DME supplier within 90 days from the date of the prescribing provider's signature. For initial orders, the detailed written order for diabetic equipment and supplies is valid for six months from the date of the order or the date of the prescribing provider’s signature, whichever is earlier. For renewal orders the detailed written order is valid for six months from the start date, or in absence of a start date, the date of the authorized prescribing signature.

### 2.2.12.7.2 Tubeless External Insulin Pump Rentals

Tubeless external insulin pump rentals may be considered for prior authorization with the submission of clinical documentation that indicates one of the following:

- The client has a diagnosis of type 1 or type 2 diabetes and meets at least two of the following criteria while on multiple daily injections of insulin:
  - Elevated glycosylated hemoglobin level (HbA1c) > 7.0 percent.
  - A history of dawn phenomenon with fasting blood sugars frequently exceeding 200 mg/dl.
  - A history of severe glycemic excursions with wide fluctuations in blood glucose.
  - A history of recurring hypoglycemia (less than 60 mg/dL) with or without hypoglycemic unawareness.
  - Expectation of becoming pregnant within three months.
- The client has a diagnosis of gestational diabetes and meets at least one of the following criteria:
  - Erratic blood sugars in spite of maximal compliance and split dosing.
  - Other evidence that adequate control is not being achieved by current methods.

In addition to the clinical documentation, the provider must submit an External Insulin Pump form that indicates:

- The client or caregiver possesses:
  - The cognitive and physical abilities to use the recommended insulin pump treatment regimen.
  - An understanding of cause and effect.
  - The willingness to support the use of the external insulin pump
• The prescribing provider has attested that:
  • A training and education plan will be completed prior to initiation of pump therapy.
  • The client or caregiver will be given face-to-face education and instruction and will be able to demonstrate the necessary proficiency to integrate insulin pump therapy with their current treatment regimen for ambient glucose control.

2.2.12.7.3 Purchase of Tubeless External Insulin Pump
The purchase of a tubeless external insulin pump may be considered for prior authorization after it has been rented for a three-month trial and all of the following documentation has been provided:
• The training or education plan has been completed.
• The pump is the appropriate equipment for the specific client.
• The client is compliant with the use of the pump.

2.2.12.8 Insulin and Insulin Syringes
Insulin and insulin syringes (0.5 and 1.0 cc sizes only) that are prescribed to fee-for-service clients are reimbursed through the Medicaid Vendor Drug Program and are not covered under Title XIX Home Health Services. The Medicaid Vendor Drug Program (VDP) only enrolls pharmacies.

Refer to: The Outpatient Drug Services Handbook (Vol. 2, Provider Handbooks) for more information about VDP.

2.2.13 Donor Human Milk
Donor human milk is a benefit for clients who are birth through 11 months of age when documentation submitted clearly shows that it is medically necessary and will correct or ameliorate the client’s disability or physical or mental illness or condition. Documentation must include all of the following:
• The ordering physician has documented medical necessity and appropriateness.
• The parent or guardian has signed and dated an informed consent form indicating that the risks and benefits of using banked donor human milk have been discussed with them.
• The donor human milk bank adheres to quality guidelines consistent with the Human Milk Bank Association of North America or such other standards as may be adopted by HHSC.

Additional donor human milk benefits beyond the limitations listed in this handbook may be available to clients who are birth through 20 years of age with documentation of medical necessity.

Procedure code B9998 must be used when requesting or billing for donor human milk.

Donor human milk is reimbursed at a maximum fee determined by HHSC or manual pricing.

Donor human milk is only reimbursed to a Texas Medicaid-enrolled donor milk bank and only for children who are in the home setting.

The physician must address the benefits and risks of using donor human milk, such as HIV, freshness, effects of pasteurization, nutrients, and growth factors to the parent. The physician also must address donor screening, pasteurization, milk storage, and transport of the donor milk. The physician may obtain this information from the donor milk bank.

2.2.13.1 Prior Authorization and Documentation Requirements
Donor human milk may be considered for a maximum of six months per authorization. The authorization may be extended with documentation of medical necessity.

Prior authorization is required for donor human milk provided through Texas Medicaid CCP Services.
To obtain prior authorization, ordering providers must complete the CCP Prior Authorization Request Form and a Donor Human Milk Request Form every 180 days. The ordering physician must maintain an originally signed and completed form in the client’s medical records, and the providing milk bank must also maintain a copy of the completed form in the client’s medical records.

The physician ordering the donor human milk must complete all of the fields in Part A of the original form, including the documentation of medical necessity. This information must be substantiated by written documentation in the clinical report. The physician must specify the quantity and the time frame in the Quantity Requested field (e.g., cubic centimeters per day or ounces per month). All of the fields in Part B of the form must be completed by the donor milk bank providing the donor human milk.

The completed documentation of medical necessity and appropriateness and the signed and dated written informed consent form must be maintained in the client’s clinical records. The documentation of medical necessity must be completed by the physician ordering the donor human milk. The clinical records are subject to retrospective review. The documentation must address all of the following:

- Medical necessity, including why the particular client cannot survive and gain weight on any appropriate formula (e.g., elemental, special, or routine formula or food), or any enteral nutritional product other than donor human milk.
- Clinical feeding trial of an appropriate nutritional product has been considered with each authorization.
- The informed consent provided to the parent or guardian details the risks and benefits of using banked donor human milk.
- A copy of the CCP Prior Authorization Request Form and the Donor Human Milk Request Form.

Refer to: Donor Human Milk Request Form on the TMHP website at www.tmhp.com.

2.2.13.2 Donor Human Milk Services for Inpatient Clients
Donor human milk may be reimbursed to hospital providers for services rendered to inpatient clients. Hospital providers may receive reimbursement for the donor human milk service separate from the inpatient diagnosis-related group (DRG) payment.

The hospital may be reimbursed using the following revenue and procedure code combination as an outpatient hospital service using the CMS-1450 (UB-04) claim form with the most appropriate outpatient type of bill (TOB):

<table>
<thead>
<tr>
<th>Revenue Code</th>
<th>Procedure Code</th>
</tr>
</thead>
<tbody>
<tr>
<td>220 (special charges)</td>
<td>T2101</td>
</tr>
</tbody>
</table>

For proper reimbursement when billing procedure code T2101, the Units field on the claim form must indicate the number of ounces rendered to the client.

Procedure code T2101 may be reimbursed for donor human milk as medically necessary for clients who are 6 months of age and younger. Prior authorization is not required.

Hospitals must follow clinical recommendations for administering donor human milk to inpatient clients, and must maintain all applicable and appropriate medical necessity documentation in the client’s medical record.

2.2.14 Hospital Beds and Equipment
A hospital bed and related equipment are considered for reimbursement for the diagnosis or treatment of illness or injury or to improve the functioning of a malformed body member. A hospital bed is not one that is typically sold as home furniture.
The following items are a benefit of Home Health Services with prior authorization:

- Hospital bed
- Air-fluidized bed
- Pressure pads or a nonpowered pressure-reducing mattress overlay
- Nonpowered pressure-reducing mattress
- Powered pressure-reducing mattress overlay system
- Powered pressure-reducing mattress
- Advanced nonpowered pressure-reducing mattress overlay
- Powered pressure-reducing mattress overlay
- Advanced nonpowered pressure-reducing mattress
- Sheepskin and lamb’s wool pads
- Decubitus care accessories

**Note:** For clients who are 20 years of age and younger and do not meet criteria through Title XIX Home Health Services, hospital beds and equipment may be considered through the Texas Health Steps—Comprehensive Care Program (THSteps-CCP).

**Note:** For clients who are 21 years of age or older, requests for hospital beds and equipment that do not meet the criteria through Title XIX Home Health Services may be considered under the Texas Medicaid Home Health—Durable Medical Equipment (DME) Exceptional Circumstances process.

Side rails or mattresses may be considered for replacement only and may be considered if it is a client-owned hospital bed and the client’s condition requires a replacement of an innerspring mattress or side rails.

The following items may be considered for clients who are birth through 20 years of age when documentation submitted clearly shows that the equipment is medically necessary and will correct or ameliorate the client’s disability or physical or mental illness or condition:

- Pediatric hospital cribs and beds
- Enclosure frame, canopy, or bubble tops
- Positioning pillows or cushions
- Reflux wedges
- Reflux slings

Hospital beds, cribs, and equipment are a benefit when all the following criteria are met:

- FFP must be available.
- The requested equipment or supplies must be safe for use in the home.

### 2.2.14.1 Hospital Beds

A hospital bed is defined as a medical device with all of the following features:

- An articulating frame that allows adjustment of the head and foot of the bed
- A headboard
- A foot board
- A mattress
• Side rails of any type (A side rail is defined as a hinged or removable rail, board, or panel of any height.)

  **Note:** Without all the components listed above, Texas Medicaid will not consider a request for any hospital bed.

### 2.2.14.2 Pediatric Hospital Bed

A pediatric hospital bed or pediatric crib is defined as a fully enclosed bed with all of the following features:

• A bed that allows adjustment of the head and foot of the bed.

• A manual pediatric hospital bed (procedure code E0328) or pediatric crib (procedure code E0300) allows manual adjustment to the head and leg elevation.

• A semi-electric or fully electric hospital bed (procedure code E0329) allows manual or electric adjustments to height and electric adjustments to head and leg elevation.

• A headboard

• A footboard

• A mattress

• Side rails of any type (A side rail is defined as a hinged or removable rail, board, or panel.)

Pediatric hospital beds and pediatric cribs that do not have all of these features will not be considered for prior authorization.

A bed that has side rails that extend 24 inches or less above the mattress is considered a pediatric hospital bed (procedure code E0328 or E0329). A pediatric hospital bed may be fixed or variable height. Variable height beds may be adjusted manually or electrically as required for the client’s medical condition.

Procedure codes E0328 and E0329 are restricted to clients who are 20 years of age and younger.

A bed that has side rails that extend more than 24 inches above the mattress is considered a pediatric crib (procedure code E0300).

A pediatric hospital bed or pediatric crib of any width that has all of the features defined above may be considered for prior authorization using only procedure code E0300, E0328, or E0329.

Hospital beds that are not fully enclosed can be considered through Texas Medicaid home health services.

  **Note:** Texas Medicaid defines fully enclosed as having 360-degree side enclosures.

The following procedure codes are used when billing for the rental or purchase of pediatric hospital beds, cribs, and equipment:

<table>
<thead>
<tr>
<th>Procedure Codes</th>
</tr>
</thead>
<tbody>
<tr>
<td>E0190*</td>
</tr>
<tr>
<td>* Purchase only</td>
</tr>
</tbody>
</table>

  **Note:** Reflux slings must be billed using procedure code E1399.

The purchase of a safety enclosure frame, canopy, or bubble top (procedure code E0316) may be a benefit when the protective crib top or bubble top is for safety use. It is not considered a benefit when it is used as a restraint or for the convenience of family or caregivers.

Procedure code E0316 may be used in conjunction with procedure codes E0300, E0328, or E0329 to request a pediatric fully-enclosed bed with a canopy.
Reflux slings or wedges may be considered for clients who are birth through 11 months of age. Reflux slings or wedges may be used as positioning devices for infants who require elevation after feedings when prescribed by a physician as medically necessary and appropriate.

Procedure code E0190 with modifier UD must be used to bill the purchase of reflex wedges and positional devices (positioning pillows and cushions). This code and modifier will require manual pricing. Procedure code E0190 is limited to once per three years, per client, any provider.

Procedure code K0739 may be reimbursed for the repair of equipment.

2.2.14.3 Prior Authorization

Hospital beds may be considered for prior authorization for clients who cannot safely utilize a regular bed.

2.2.14.3.1 Fixed-Height Hospital Bed

A fixed-height bed (procedure code E0250), which allows for manual adjustment to the head and leg elevation but not height, may be considered for prior authorization if at least one of the following criteria exists:

- The client’s medical condition requires positioning of the body in ways that are not feasible in an ordinary bed.
- The client’s medical condition requires special positioning to alleviate pain.
- It is necessary to elevate the head of the bed 30 or more degrees most of the time due to, but not limited to, congestive heart failure, chronic pulmonary disease, or problems with aspiration, and alternative measures such as wedges or pillows, have been attempted but have failed to manage the client’s medical condition.

*Note:* Texas Medicaid defines a failed measure as having no clinically significant improvement after being introduced.

- The client requires traction equipment that can only be attached to a hospital bed.

2.2.14.3.2 Variable-Height Hospital Bed

A variable-height hospital bed (procedure E0255), which allows manual adjustments to height as well as to head and leg elevations, may be considered for prior authorization if the client meets the criteria for a fixed-height hospital bed and requires a bed height that is different from a fixed-height hospital bed to permit transfers in and out of the bed to a chair, wheelchair, or to a standing position. Medical conditions that require a variable-height hospital bed include, but are not limited to, the following:

- Severe arthritis and other injuries to lower extremities that require the variable height feature to assist in ambulation by enabling the client to place his or her feet on the floor while sitting on the edge of the bed.
- Severe cardiac conditions, where the client is able to leave the bed, but must avoid the strain of “jumping” up and down.
- Spinal cord injuries (including quadriplegia and paraplegia), multiple limb amputations, and stroke, where the client is able to transfer from a bed to a wheelchair with or without help.
- Other severely debilitating diseases and conditions if the client requires a bed height different than a fixed-height hospital bed to permit transfers to a chair, wheelchair, or to a standing position.
2.2.14.3.3 Semi-Electric Hospital Bed

A semi-electric hospital bed (procedure code E0260), which allows manual adjustments to height and electric adjustments to head and leg elevation, may be considered for prior authorization if the client meets the criteria for a fixed-height hospital bed and has a condition that requires frequent changes in body position or might require an immediate change in body position to avert a life-threatening situation.

2.2.14.3.4 Fully-Electric Hospital Bed

A fully-electric bed (procedure code E0265), which allows electric adjustments to height and head and leg elevation, may be considered for prior authorization when all of the following criteria are met:

- The client has paraplegia or hemiplegia.
- The fully-electric hospital bed will allow the client to have functional independence with self-care.

Documentation must include an attestation statement from the client’s physician or physical or occupational therapist that verifies a determination has been made that the fully-electric hospital bed will allow the client to independently meet their daily self-care needs.

The following hospital beds may be considered for prior authorization if the client meets the criteria for a hospital bed and the weight requirements for a bariatric bed as listed below:

- Heavy-duty, extra-wide hospital bed (procedure code E0303) capable of supporting a client who weighs more than 350 pounds, but no more than 600 pounds
- Extra heavy-duty, extra-wide hospital bed (procedure code E0304) capable of supporting a client who weighs more than 600 pounds

2.2.14.3.5 Pediatric Hospital Beds and Safety Enclosure

Pediatric hospital beds and pediatric cribs (procedure codes E0300, E0316, E0328, and E0329) may be considered for prior authorization when the documentation submitted clearly shows that the requested bed or crib will correct or ameliorate the client’s condition. The documentation must meet at least one of the following criteria:

- The client’s medical condition requires positioning of the body in ways that are not feasible in an ordinary bed, including, but not limited to, the need for positioning to alleviate pain.
- The head of the bed must be elevated 30 or more degrees most of the time due to, but not limited to, congestive heart failure, chronic pulmonary disease, or problems with aspiration, and alternative measures, such as wedges or pillows, have been attempted but have failed to manage the client’s medical condition.

**Note:** Texas Medicaid defines a failed measure as having no clinically significant improvement after being introduced.

- The client requires traction equipment that can only be attached to a hospital bed.

A semi-electric or fully electric hospital bed (procedure code E0329) may be considered for prior authorization when the submitted documentation shows that the client has a medical condition that requires frequent changes in body position or might require an immediate change in body position to avert a life-threatening situation.

The safety enclosure frame, canopy, or bubble top may be considered for prior authorization with documentation that the protective canopy top or bubble will provide for the client’s safety. Prior authorization will not be considered when it will be used as a restraint or for the convenience of family or caregivers.
Reflux slings or wedges may be considered for prior authorization for clients who are 11 months of age and younger. These may be used as positioning devices for infants who require the head of the bed or crib to be elevated greater than 30 degrees after feedings when prescribed by a physician as medically necessary and appropriate.

Positioning pillows and cushions may be considered for prior authorization with documentation of medical necessity that indicates the item will provide for or assist in the positioning needs of the client to maintain proper body alignment and skin integrity. Documentation must include what other devices have been used previously and why they proved to be ineffective.

Items used for PT or rehabilitation in the home are provided by the therapist. Requests for authorization for these purposes will not be considered.

2.2.14.4 Documentation Requirements

To request prior authorization for a hospital bed, the following documentation must be submitted:

- Accurate diagnostic information pertaining to the underlying medical diagnoses or conditions (e.g., gastrostomy feeding, suctioning, ventilator dependent, other respiratory equipment or ventilation assistance devices) to include the client’s overall health status
- Client height and weight
- Client functional mobility status
- Client use of any pressure-reducing support surfaces, if applicable

The following documentation must be submitted for clients who are birth through 20 years of age:

- The diagnosis, medical needs, treatments, developmental level, and functional skills of the child. A diagnosis alone is insufficient information to consider prior authorization of the requested equipment.
- The age, length, and weight of the child.
- Description of any other devices that have been used, the length of time used, and why they were ineffective.
- How the requested equipment will correct or ameliorate the client’s condition beyond that of a standard child’s crib, regular bed, or standard hospital bed.
- The name of the manufacturer and the manufacturer’s suggested retail price (MSRP).

A determination will be made by HHSC or its designee whether the equipment will be rented, purchased, repaired, or modified based on the client’s needs, duration of use, and age of equipment. All modifications, adjustments, and repairs within the first six months after delivery are considered to be part of the purchase price.

2.2.14.5 Mattresses and Support Surfaces

A pressure-reducing support surface includes three separate groups of mattress or mattress-like equipment designed to assist in the healing of wounds. These devices are used in conjunction with conventional wound care therapy to prevent the occurrence of said wounds in susceptible clients. Pressure-reducing support surfaces are designed to prevent skin breakdown or to promote the healing of pressure ulcers by reducing or eliminating tissue interface pressure. Most of these devices reduce interface pressure by conforming to the contours of the body so that pressure is distributed over a larger surface area rather than concentrated on a more circumscribed location.

For all types of pressure-reducing support surfaces, the support surface provided for the client should be one in which the client does not “bottom out.” The Centers for Medicare & Medicaid Services (CMS) define “bottoming out” as: when an outstretched hand, palm up, between the undersurface of the overlay or mattress and in an area under the bony prominence can readily palpate the bony prominence (coccyx
or lateral trochanter). This “bottoming out” criterion should be tested with the client in the supine position with head flat, in the supine position with head slightly elevated (no more than 30 degrees), and in the side-lying position.

Pressure-reducing support surfaces containing multiple components are categorized according to the clinically predominant component (usually the top-most layer of a multi-layer product) and the presence and stage of pressure ulcers.

The staging of pressure ulcers is as follows:

**Stage I:** Observable pressure related alteration of intact skin whose indicators are as follows:
- Compared to the adjacent or opposite area on the body may include changes in one or more of the following: skin temperature (warmth or coolness), tissue consistency (firm or boggy feel), or sensation (pain, itching).
- The ulcer appears as a defined area of persistent redness in lightly pigmented skin, whereas in darker skin tones, the ulcer may appear with persistent red, blue, or purple hues.

**Stage II:** Partial thickness skin loss involving epidermis, dermis, or both. The ulcer is superficial and presents clinically as an abrasion, blister, or shallow crater.

**Stage III:** Full thickness skin loss involving damage to, or necrosis of, subcutaneous tissue that may extend down to, but not through, underlying fascia. The ulcer presents clinically as a deep crater with or without undermining of adjacent tissue.

**Stage IV:** Full thickness skin loss with extensive destruction, tissue necrosis, or damage to muscle, bone, or supporting structures (e.g., tendon, joint capsule). Undermining and sinus tracts also may be associated with Stage IV pressure ulcers.

### 2.2.14.5.1 Documentation Requirements

A support surface that does not meet the characteristics specified in the criteria for grouping levels may be denied and considered to be not medically necessary.

To request prior authorization for a pressure-reducing support surface, the following documentation must be provided:

- Client’s overall health status and all other medical diagnoses or conditions (e.g., history of decubitus)
- Documentation of the client’s limited mobility or confinement to a bed
- History of previous use and results of pressure-reducing support surfaces, (e.g., wound improvement, stasis, or degradation)
- Current wound therapy, if any

### 2.2.14.5.2 Group 1 Support Surfaces

A group 1 Support Surface may be considered for prior authorization with documentation of medical necessity if the client is completely immobile without assistance, or the client has limited mobility or existing pressure ulcer on the pelvis or trunk and at least one of the following conditions:

- Impaired nutritional status
- Fecal or urinary incontinence
- Altered sensory perception
- Compromised circulatory status

All of the support surfaces described below are considered a benefit of the Home Health Services Program when medical necessity criteria for Group 1 support surfaces are met.
Pressure pads or a nonpowered pressure-reducing mattress overlay for mattresses with the following features may be considered for reimbursement with documentation of medical necessity:

- A gel or gel-like layer with a height of two inches or greater
- An air mattress overlay with interconnected air cells that are inflated with an air pump and a cell height of three inches or greater
- A water mattress overlay with a filled height of three inches or greater
- A foam mattress overlay with all the following features:
  - Base thickness of two inches or greater and peak height of three inches or greater if it is a convoluted overlay (e.g., eggcrate) or an overall height of at least three inches if it is a nonconvoluted overlay
  - Foam with a density and other qualities that provide adequate pressure reduction
  - Durable, waterproof cover

Nonpowered pressure-reducing mattresses, with the following features, may be considered for reimbursement with documentation supporting medical necessity:

- A foam mattress with all the following features may be considered with documentation supporting medical necessity. Documentation must include all of the following features:
  - A foam height of five inches or greater
  - Foam with a density and other qualities that provide adequate pressure reduction
  - Durable, waterproof cover
  - Can be placed directly on a hospital bed frame
- An air, water, or gel mattress with all the following features may be considered for reimbursement:
  - A height of five inches or greater
  - Durable, waterproof cover

A powered pressure reducing mattress overlay system, with all the following features, may be considered for reimbursement when documentation supports medical necessity:

- The system includes an air pump or blower which provides either sequential inflation and deflation of air cells, or a low interface pressure throughout the overlay.
- Inflated cell height of the air cells through which air is being circulated is 2.5 inches or greater.
- Height of the air chambers, proximity of the air chambers to one another, frequency of air cycling (for alternating pressure overlays), and air pressure provide adequate client lift, reduces pressure, and prevents bottoming out.

### 2.2.14.5.3 Group 2 Support Surfaces

A Group 2 support surface may be considered for prior authorization with documentation of medical necessity if the client has multiple stage II ulcers on the trunk or pelvis and has been on a comprehensive ulcer treatment program for at least the past month which has included the use of a Group 1 support surface.

The client must also have at least one of the following:

- The ulcers have remained the same or worsened over the past month.
- There are large or multiple stage III or IV pressure ulcers on the trunk or pelvis.
• Received a myocutaneous flap or skin graft for a pressure ulcer on the trunk or pelvis within the last 60 days, and have been prescribed or placed on a Group 2 or 3 support surface immediately before discharge (within the last 30 days) from the hospital or a nursing facility

All of the support surfaces described below are considered a benefit of the Home Health Services Program when medical necessity criteria for Group 2 support surfaces are met.

The powered pressure reducing mattress (alternating pressure low air loss, or powered flotation without air loss) device with all the following features may be considered for reimbursement when documentation supports medical necessity:

• The system includes an air pump or blower which provides either sequential inflation and deflation of the air cells or a low interface pressure throughout the mattress.

• Inflated cell height of the air cells through which air is being circulated is five inches or greater.

• Height of the air chambers, proximity of the air chambers to one another, frequency of air cycling (for alternating pressure mattress), and air pressure to provide adequate client lift, reduce pressure, and prevent bottoming out.

• A surface designed to reduce friction and shear.

A semi-electric hospital bed with fully integrated powered pressure-reducing mattress that has all of the features described above may be considered for reimbursement when documentation supports medical necessity.

The advanced nonpowered pressure-reducing mattress overlay device with all the following features may be considered for reimbursement when documentation supports medical necessity:

• Height and design of individual cells which provide significantly more pressure reduction than Group 1 overlay and prevent bottoming out

• Total height of 3 inches or greater

• A surface designed to reduce friction and shear

• Manufacturer product information that substantiates the product is effective for the treatment of conditions described by the coverage criteria for Group 2 support surfaces

The powered pressure-reducing mattress overlay device with all the following features may be considered for reimbursement when documentation supports medical necessity:

• The system includes an air pump or blower that provides either sequential inflation and deflation of the air cells or a low interface pressure throughout the overlay.

• Inflated cell height of the air cells through which air is being circulated is three and a half inches or greater.

• Height of the air chambers, proximity of the air chambers to one another, frequency of air cycling (for alternating pressure overlays), and air pressure to provide adequate client lift, reduce pressure and prevent bottoming out.

The advanced nonpowered pressure-reducing mattress device with all the following features may be considered for reimbursement when documentation supports medical necessity:

• Height and design of individual cells designed to provide significantly more pressure than a Group 1 mattress and prevent bottoming out

• Total height of 5 inches or greater

• A surface designed to reduce friction and shear

• Documented evidence substantiates that the product is effective for the treatment of conditions described by the coverage criteria for Group 2 support surfaces
Sheepskin and lambs wool pads are considered a benefit of the Home Health Services Program under the same conditions as alternating pressure pads and mattresses (Group 2 pressure-reducing support surfaces) when prior authorized.

### 2.2.14.5.4 Group 3 Support Surfaces

A Group 3 support surface may be considered for prior authorization with documentation of medical necessity when all the following criteria are met:

- There is a presence of a stage III or IV ulcer.
- Severely limited mobility rendering the client bed or chair bound.
- Without an air-fluidized bed, the client would be institutionalized.
- The client has been placed on a Group 2 support surface for at least a month before ordering the air-fluidized bed with the ulcers not improving or worsening.
- There has been at least weekly assessment of the wound by the physician, a nurse or other licensed health-care professional and the treating physician has done a comprehensive evaluation of the client’s condition within the week before ordering the air-fluidized bed.
- A trained adult caregiver is available to assist the client with activities of daily living, maintaining fluid balance, supplying dietary needs, aiding in repositioning and skin care, administering prescribed treatments, recognizing and managing altered mental status, and managing the air-fluidized bed system and its potential problems, such as leakage.
- The physician continues to re-evaluate and direct the home treatment regimen monthly.
- All other alternative equipment has been considered and ruled out.

The existence of any one of the following conditions may result in noncoverage of the air-fluidized bed:

- Coexisting pulmonary disease (the lack of firm back support can render coughing ineffective and dry air inhalation thickens pulmonary secretions).
- Wounds requiring moist wound dressings that are not protected with an impervious covering such as plastic wrap or other occlusive material (if wet-to-dry dressings are being utilized, dressing changes must be frequent enough to maintain their effectiveness).
- For clients who are 21 years of age and older, the caregiver is unwilling or unable to provide the type of care required by the client who uses an air-fluidized bed.
- The home’s structural support or electrical system cannot safely accommodate the air-fluidized bed.

Initial prior authorization for a Group 3 pressure-reducing support surface will be for no more than 30 days. Prior authorized extensions may be considered for reimbursement in increments of 30-day periods, up to a maximum of four months, when documentation supports continued significant improvement in wound healing. Coverage beyond four months will be on a case-by-case basis after review by the medical director or designee.

Air-fluidized beds may be considered for reimbursement when the medical necessity criteria for Group 3 support surfaces are met.

### 2.2.14.6 Equipment and Other Accessories

The following equipment or accessories may be considered with documentation of medical necessity:

- Positioning devices
- Bed cradle (keeps bed covers from touching affected skin)
- Trapeze bars
2.2.14.6.1 Accessories
A mattress of any size with innerspring may be considered for prior authorization with procedure code E0271.

Replacement rails and hospital bed frame padding or covers may be considered for prior authorization as a hospital bed accessory (procedure code E0315) with documentation that the padding, covers or rails are required to prevent injury (for example, related to seizure activity) or to prevent entrapment.

2.2.14.6.2 Prior Authorization
Heel or elbow protector (procedure code E0191) does not require prior authorization. Prior authorization is required for all other hospital beds, equipment, and services provided through Texas Medicaid Title XIX Home Health Services. Prior authorization also includes any accessories, modifications, adjustments, and repairs of the equipment. Positioning cushions or pillows (procedure code E0190) may be considered with documentation of medical necessity that the item will provide pressure relief and positioning in the treatment of decubiti, burns, or musculoskeletal injuries. Documentation must include a listing of other devices that have been used and why the devices proved ineffective.

A trapeze bar attached to a bed (procedure code E0910 or E0911) may be considered if the client requires this device to sit up, to change body position, to get in or out of bed, or for other medical reasons with documentation of medical necessity.

“Free-standing” trapeze equipment (procedure code E0940 or E0912) may be considered if the client does not have an eligible hospital bed, but the client needs this device to sit up, to change body position, to get in or out of bed, or for other medical reasons with documentation of medical necessity.

An over-bed table (procedure code E0315) may be considered if the client is bed-bound and needs the over-bed table for treatments.

2.2.14.7 Decubitus Care Accessories
For prior authorization of decubitus care accessories, the following documentation must be provided:

- Wound measurements including location, length, width, and depth
- Any undermining or tunneling
- Odor, if applicable

2.2.14.8 Replacement
Beds rails and frames that have been purchased are anticipated to last a minimum of five years.

2.2.14.8.1 Prior Authorization
Prior authorization for replacement may be considered within five years of purchase when one of the following occurs:

- There has been a significant change in the client’s condition, such that the current equipment no longer meets the client’s needs.
- The equipment is no longer functional and cannot be repaired or it is not cost effective to repair.

Replacement of equipment may be considered when loss or irreparable damage has occurred. A copy of the police or fire report, when appropriate, and the measures to be taken to prevent reoccurrence must be submitted.

In situations where the equipment has been abused or neglected by the client, the client’s family, or the caregiver, a referral to the Department of State Health Services (DSHS) Health Screening and Case Management unit will be made by the Home Health Services prior authorization unit for clients who are 20 years of age and younger. Providers will be notified that the state will be monitoring this client’s services to evaluate the safety of the environment for both the client and equipment.
Repairs require replacement of components that are no longer functional. Technician fees are considered to be part of the cost of the repair.

Repairs to client-owned equipment may be considered with documentation of medical necessity.

Providers are responsible for maintaining documentation in the client’s medical record specifying the repairs and supporting medical necessity.

Rental equipment may be considered during the period of repair. Routine maintenance of rented equipment is the provider’s responsibility.

Pediatric hospital cribs and beds, enclosed beds, and safety enclosure frames, canopies, or bubble tops that have been purchased are anticipated to last a minimum of five years.

2.2.14.9 Non-covered Items

A safety enclosure (procedure code E0316) used to prevent a client from leaving the bed is not a benefit of Home Health Services. A safety enclosure or enclosed bed may be considered through CCP for safety use.

Traction equipment (procedure codes E0890, E0947, and E0948) is not a benefit of Home Health Services.

The following types of beds will not be considered for prior authorization, because they are not considered medically necessary or are inappropriate for use in the home setting:

- Institutional type beds (procedure code E0270)
- An ordinary or standard bed typically sold as furniture (may consist of a frame, box spring, and mattress, and is of fixed height with no head or leg elevation adjustments). These types of beds are not primarily medical in nature, not primarily used in the treatment of disease of injury, and are normally of use in the absence of illness or injury. They are not considered durable medical equipment (DME) by Texas Medicaid.
- All non-hospital adjustable beds available to the general public as furniture. These types of beds are not primarily medical in nature, not primarily used in the treatment of disease or injury, and are normally of use in the absence of illness or injury. They are a comfort and convenience item and are not considered DME by Texas Medicaid.
- Hospital beds without rails. Texas Medicaid considers side rails an integral part of medically necessary bed.
- Beds with rails of any height that do not allow head and foot elevation (e.g., platform beds with rails), and are primarily used to prevent clients from leaving the bed. This types of beds are not primarily medical in nature.

2.2.14.10 Hospital Beds and Equipment Procedure Code Table

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<td>E0196</td>
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### 2.2.15 Incontinence Supplies

Incontinence supplies, such as diapers, briefs, pull-ons, liners, wipes, and underpads, may be considered for reimbursement through CCP for those clients who are birth through 3 years of age with a medical condition resulting in an increased urine or stool output beyond the typical output for this age group, such as celiac disease, short bowel syndrome, Crohn’s disease, thymic hypoplasia, Acquired Immunodeficiency Syndrome (AIDS), congenital adrenal hyperplasia, diabetes insipidus, Hirschsprung’s disease, or radiation enteritis.

For clients who are 4 years of age or older, incontinence supplies may be considered through Title XIX Home Health Services when their medical conditions result in an impairment of urination and/or stool. For clients who do not meet criteria through Title XIX Home Health Services, incontinence supplies may be considered through CCP with documentation of medical necessity.

**Note:** For clients who are 21 years of age or older, requests for incontinence supplies that do not meet the criteria through Title XIX Home Health Services may be considered under the Texas Medicaid Home Health—Durable Medical Equipment (DME) Exceptional Circumstances process.

Lack of bladder or bowel control is considered normal development for clients who are 4 years of age or younger.

Reusable diapers, briefs, pull-ons, liners, wipes, and underpads are not a benefit of CCP. Gloves used to change diapers, briefs, and pull-ons are not considered medically necessary unless the client has skin breakdown or a documented disease that may be transmitted through the urine.

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<td>E0940</td>
<td>1 purchase every 5 years; 1-month rental</td>
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Incontinence supplies billed for a one-month period must be based on the frequency or quantity ordered by the physician on the Home Health Services (Title XIX) DME/Medical Supplies Physician Order Form.

2.2.15.1 Skin Sealants, Protectants, Moisturizers, and Ointments for Incontinence-Associated Dermatitis

Incontinence-associated dermatitis is classified by category:

- **Category 1**—Small area of skin breakdown (<20 cm²) with mild redness (blotchy and non-uniform) and mild erosion involving the epidermis only.
- **Category 2**—Moderate area of skin breakdown (20-50 cm²) with moderate redness (severe in spots, but not uniform in appearance) and moderate erosion involving epidermis and dermis with no or little exudate.
- **Category 3**—Large area of skin breakdown (>50 cm²) with severe redness (uniformly severe in appearance) and severe erosion of epidermis with moderate involvement of the dermis and no or small volume of exudate.
- **Category 4**—Large area of skin breakdown (>50 cm²) with severe redness (uniformly severe in appearance) and extreme erosion of epidermis and dermis with moderate volume of persistent exudate.

Skin sealants, protectants, moisturizers, and ointments (procedure code A6250) may be considered for clients who are 4 years of age or older and have documented incontinence-associated dermatitis.

For clients who have Category 1 or Category 2 incontinence-associated dermatitis, prior authorization is not required for a maximum quantity of 2 containers (no less than 4 ounces per container) per month and 12 containers per year of skin sealants, protectants, moisturizers, and ointments. Providers must use procedure code A6250 with modifier UA to bill for these products.

For clients who have Category 3 or Category 4 incontinence-associated dermatitis, prior authorization and documentation of medical necessity is required for skin sealants, protectants, moisturizers, and ointments that are not used for Category 1 or Category 2 incontinence-associated dermatitis. Providers must use procedure code A6250 without a modifier to bill for these products.

Providers must use procedure code A6250 instead of procedure code A5120 when billing for skin sealants, protectants, moisturizers, and ointments.

*Note:* Skin sealants, protectants, moisturizers, ointments for diagnoses other than incontinence related dermatitis (i.e., wounds, decubitus ulcers, periwound skin complications, peristomal skin complications) may be considered for reimbursement with prior authorization.

2.2.15.2 Diapers, Briefs, Pull-ons, and Liners

Diapers and briefs are defined as incontinence items attached with tabs. Pull-ons are defined as incontinence items that do not attach with tabs and are slip-on items, such as “pull-ups.” Liners are intended to be worn inside diapers, briefs, and pull-ons to increase absorbency. Reusable diapers or briefs are not a benefit of Home Health Services.

For clients who are 4 years of age and older and have a medical condition that results in chronic incontinence, up to a maximum total combination of 240 per month of diapers, briefs, or liners may be considered without prior authorization. Quantities in excess of 240 per month may be considered with documentation of medical necessity and prior authorization.
The following procedure codes must be used when billing for diapers, briefs, and liners and are limited to a combined total of 240 per month:

<table>
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<th>Procedure Codes</th>
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<tr>
<td>T4521</td>
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<tr>
<td>T4531</td>
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</table>

**Note:** Gloves used to change diapers and briefs are not considered medically necessary unless the client has skin breakdown or a documented disease that may be transmitted through the urine or stool.

### 2.2.15.3 Diaper Wipes

For clients who are 4 years of age and older and are receiving diapers/briefs/pull-ons, up to 2 boxes of diaper wipes do not require prior authorization. Exceptions will not be considered through Title XIX Home Health Services. Quantities in excess of 2 boxes per month may be considered through CCP for clients who are 20 years of age and younger with documentation of medical necessity and prior authorization.

Providers must use procedure code A4335 with modifier U9 instead of procedure code A5120 when billing for diaper wipes.

If there is not an appropriate procedure code for supplies, providers may request prior authorization using procedure code A4335.

Diaper wipes may be considered for clients who are receiving diapers, briefs, or pull-ons through CCP.

### 2.2.15.4 Underpads

For clients who are 4 years of age and older and are receiving diapers/briefs/pull-ons/liners/urine collection devices/bowel management supplies, up to a maximum of 120 underpads per month may be considered without prior authorization. Quantities in excess of 120 per month may be considered with documentation of medical necessity and prior authorization.

Reusable underpads are not a benefit of Home Health Services.

Underpads may be considered for clients who are receiving diapers, briefs, or pull-ons through CCP.

Providers must use procedure code A4554 when billing for underpads. Procedure code A4554 is limited to 120 per month.

**Note:** The Home Health Services (Title XIX) DME/Medical Supplies Physician Order Form for the supplies listed above must reflect no more than a one-month’s supply of the incontinence product. The Home Health Services (Title XIX) DME/Medical Supplies Physician Order Form must not reflect more than the maximum allowed quantity per month without requesting prior authorization.

### 2.2.15.5 Ostomy Supplies

The physician must specify the type of ostomy device or system to be used and how often it is to be changed on the Home Health Services (Title XIX) DME/Medical Supplies Physician Order Form. The quantity of ostomy supplies billed for a one-month period must relate to the number of changes per month based on the frequency ordered by the physician.

Ostomy supplies may be considered for reimbursement without prior authorization.
2.2.15.6 Indwelling or Intermittent Urine Collection Devices

The home setting is considered a clean environment, not a sterile one. Sterile incontinence supplies, (including the supplies in procedure codes A4311, A4312, A4313, A4314, A4315, A4316, and A4353) are a benefit in the home setting when requested for the following:

- Indwelling urinary catheters
- Intermittent catheters for clients who:
  - Are immunosuppressed
  - Have radiologically documented vesico-ureteral reflux
  - Are pregnant and have a neurogenic bladder due to spinal cord injury
  - Have a history of distinct, recurrent urinary tract infections, defined as a minimum of two within the prior 12-month period, while on a program of clean intermittent catheterization

Nonsterile or sterile gloves for use by a health-care provider in the home setting, such as a registered nurse (RN), licensed vocational nurse (LVN), or attendant, are not a benefit of Home Health Services.

2.2.15.6.1 Indwelling Catheters and Related Insertion Supplies

Indwelling catheters and related supplies may be considered without prior authorization up to a maximum of 2 per month for clients who have a medical condition that results in an impairment of urination. Quantities in excess of 2 per month may be considered with documentation of medical necessity and prior authorization.

2.2.15.6.2 Intermittent Catheters and Related Insertion Supplies

Intermittent catheters and related supplies, up to a maximum of 150 per month, may be considered without prior authorization for clients who have a medical condition that results in an impairment of urination. Quantities in excess of 150 per month may be considered with documentation of medical necessity and prior authorization.

Procedure code A4351 denotes catheters used for intermittent catheterizations. Procedure code A4351 must be accompanied with modifier SC when a hydrophilic catheter is used.

A completed Home Health Services (Title XIX) DME/Medical Supplies Physician Order Form may be valid for up to 12 months for intermittent catheters and related insertion supplies for quantities within the stated benefit limits for clients who have one of the following chronic conditions:

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<th>Diagnosis Codes</th>
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<tbody>
<tr>
<td>G35</td>
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<tr>
<td>N311</td>
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<td>Q0703</td>
</tr>
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</table>

Note: The diagnosis codes R32 and R339 are not specific enough to allow for the extension of the prior authorization to 12 months.

For clients who have diagnoses other than those listed in the above table, the completed Title XIX Form may be valid for up to six months for intermittent catheters and related insertion supplies for quantities within the stated benefit limits.

For quantities greater than the stated benefit limits, prior authorization will be required and may be granted for up to six months regardless of diagnosis.

Nonsterile gloves are a benefit with prior authorization when a family member or friend is performing the catheterization.
Providers must use procedure codes A4351 or A4352 when billing for intermittent catheters. Providers must use procedure code A4353 when billing for intermittent catheters with insertion supplies. For hydrophilic catheters, procedure code A4351 must be accompanied with modifier SC.

### 2.2.15.6.3 External Urinary Collection Devices

For clients who are 4 years of age and older and have a medical condition that results in a permanent impairment of urination, external urinary collection devices, including, but not limited to, male external catheters, female collection devices, and related supplies may be considered without prior authorization. Male external catheters are limited to 31 per month. Female collection devices are limited to 4 per month. Male external catheters in excess of 31 per month and female collection devices in excess of 4 per month may be considered with documentation of medical necessity and prior authorization.

External urinary collection devices, including, but not limited to, male external catheters, female collection devices, and related supplies may be considered with a documented medical condition resulting in an increased urine or stool output beyond the typical output.

The following procedure codes must be used when billing for external urinary collection devices:

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<th>Procedure Code</th>
<th>Maximum Limitation</th>
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<td>4 per month</td>
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<tr>
<td>A4349</td>
<td>31 per month</td>
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### 2.2.15.6.4 Urinals and Bed Pans

Urinals and bed pans may be considered without prior authorization for clients who have a medical condition that results in an inability to ambulate to the bathroom safely (with or without mobility aids) up to a limit of 2 per year. Quantities in excess of 2 per year may be considered with documentation of medical necessity and prior authorization.

Urinals and bed pans are purchase only.

### 2.2.15.7 Prior Authorization

Prior authorization is required for incontinence supplies if amounts greater than the maximum limits are medically necessary.

Prior authorization is required for incontinence supplies through CCP.

A determination is made by HHSC or its designee as to the number of incontinence supplies prior authorized based on the client’s medical needs.

Additional quantities may be considered with documentation of medical necessity.

The quantity of incontinence supplies billed for a one-month period must be consistent with the number of times per day the physician has ordered the supply to be used on the CCP Prior Authorization Request Form.

To request prior authorization for incontinence supplies, the following documentation must be provided for the items requested:

- Accurate diagnostic information pertaining to the underlying diagnosis or condition as well as any other medical diagnoses or conditions, to include the client’s overall health status
- Diagnosis or condition causing increased urination or stooling
- Client’s height, weight, and waist size
- Number of times per day the physician has ordered the supply to be used
• Quantity of disposable supplies requested per month

Additional information may be requested to clarify or complete a request for the supplies and equipment.

2.2.15.8 Documentation Requirements

To request prior authorization for incontinence supplies and equipment, the following documentation must be provided:

• Diagnostic information pertaining to the underlying diagnosis or condition, the diagnosis causing incontinence, and any other medical diagnoses or conditions, including the client’s overall health status
• Weight and height or waist size, when applicable
• Number of times per day the physician has ordered the supply be used
• Quantity of disposable supplies requested per month by the physician

Additional information may be requested to clarify or complete a request for the supplies.

2.2.15.9 Incontinence Procedure Codes with Limitations

Any service or combination of services, except diaper wipes, requires prior authorization if the maximum limitation is exceeded. Requests for prior authorization of diaper wipes that exceed more than two boxes per month will not be considered through Home Health Services.

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Refer to: Subsection 2.2.15.2, "Diapers, Briefs, Pull-ons, and Liners" in this handbook for an explanation of the item limitations identified with an asterisk (*).

The following procedure codes always require prior authorization even if the maximum benefit limitation allowed has not been exceeded:

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2.2.16 Intravenous (IV) Therapy Equipment and Supplies

The following equipment and supplies are used in the delivery of IV therapy and are a benefit of Home Health Services. The following procedure codes require prior authorization unless otherwise specified:

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<td>A6250</td>
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Note: Additional supply procedure codes not listed may be considered with documentation of medical necessity.

The following IV supplies listed are available without prior authorization up to the stated quantity limitations. Prior authorization is required for any quantities exceeding the limitations with documentation supporting the medical necessity of the quantity requested.
Types of IV access devices include but are not limited to:

- Peripheral IV lines.
- Central IV lines, including but not limited to, peripherally-inserted central catheters, subclavian catheters, and vena cava catheters.
- Central venous lines, including but not limited to, tunneled and peripherally inserted central venous catheters.
- Implantable ports, including but not limited to, access devices with subcutaneous ports.

Stopcocks increase the risk of infection and should not be routinely used for infusion administration.

Routine use of in-line filters is not recommended for infection control.

Note: Nonsterile or sterile gloves for use by a health-care provider in the home setting, such as an RN, LVN, or attendant, are not a benefit of Home Health Services.

Stationary infusion pumps may be a benefit when the infusion rate must be more consistent and cannot be obtained with gravity drainage. Ambulatory infusion pumps may be a benefit when the length of infusion is greater than two hours, the client must be involved in activities away from home, and when the infusion rate must be more consistent and cannot be obtained with gravity drainage. Elastomeric infusion pumps may be a benefit for short-term use when the caregiver cannot administer the infusion by pump. Dial flow regulators are a benefit and are incorporated into IV extension sets or IV tubing. Elastomeric devices may be reimbursed using procedure codes A4305 and A 4306.

Rental of an infusion pump may be prior authorized on a monthly basis for a maximum of four months per lifetime. Purchase of an infusion pump (ambulatory or stationary) may be prior authorized with documentation of medical necessity that supports repeated IV administration for a chronic condition.

For clients who require cardiovascular medications, infusion pumps will be rented, but not purchased.

Repairs to client-owned equipment may be prior authorized as needed with documentation of medical necessity. Technician fees are considered part of the cost of the repair. Providers are responsible for maintaining documentation in the client’s medical record that specifies the repairs and supports medical necessity. All repairs and replacement parts within the first six months after delivery are considered part of the purchase price. Batteries for client-owned equipment require prior authorization. Additional documentation, such as the purchase date, serial number, and manufacturer’s information, may be required.

Note: For clients who are 21 years of age or older, requests for IV supplies and equipment that do not meet the criteria through Title XIX Home Health Services may be considered under the Texas Medicaid Home Health—Durable Medical Equipment (DME) Exceptional Circumstances process.

IV therapy, supplies, and equipment are not considered a benefit when the infusion or medication being administered:

- Is not considered medically necessary to the treatment of the client’s illness.
- Exceeds the frequency or duration ordered by the physician.
- Is a chemotherapeutic agent.
- Is not FDA-approved, unless the physician documents why the off-label use is medically appropriate and not likely to result in an adverse reaction. In order to consider coverage of an off-label (non-FDA approved) use of a drug, documentation must include why a drug usually indicated for the specific diagnosis or condition has not been effective for the client.

<table>
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<th>Procedure Code</th>
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<td>A6402</td>
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Routine maintenance of rental equipment is included in the rental price.

Repairs or replacement parts may be reimbursed with documentation of a client-owned device.

Replacement batteries (procedure codes K0601, K0602, K0603, K0604, and K0605) for client-owned pumps are limited to one battery per 180 days.

### 2.2.16.1 Prior Authorization

Additional replacement batteries for client-owned pumps (procedure codes K0601, K0602, K0603, K0604, and K0605) beyond the limit of 1 per 180 days may be considered for prior authorization with documentation of medical necessity.

All IV equipment and supplies, with the exception of implantable access catheter (A4300) require prior authorization. Prior authorization of IV equipment and supplies may be considered when administration of the drug in the home is medically necessary and is appropriate in the home setting. IV equipment may be prior authorized for rental or purchase depending on the clinician’s predicted length of treatment.

The following standards are used when considering prior authorization of IV supplies:

- The aseptic technique is acceptable for IV catheter insertion and site care; the sterile technique is not required:
  - Nonsterile gloves are acceptable for the insertion of a peripheral IV catheter and for changing any IV site dressing.
  - The sterile technique may be medically necessary. Examples of medical necessity include, but are not limited to, a client who is immuno-compromised.
- A peripheral IV site is rotated no more frequently than every 72 hours, but it is rotated at least weekly.
- The IV administration set (with or without dial flow regulator), extension set (with or without dial flow regulator), and any add-on devices are changed every 72 hours.
- One IV access catheter is used per insertion.
- Saline or heparin-locked catheters:
  - Use one syringe to flush the catheter before administration of an intermittent infusion to assess.
  - Use two syringes to flush the catheter after the intermittent infusion—one to clear the medication and one to infuse the anticoagulant or other medication used to maintain IV patency between doses, including, but not limited to, heparin.
- An injection port is cleaned before administering an intermittent infusion and capped after the infusion.
- IV catheter site care:
  - Disinfect the site with an appropriate antiseptic (including but not limited to 2 percent chlorhexidine-based preparation, tincture of iodine, or 70 percent alcohol).
  - Cover with sterile gauze, transparent dressing, or semi-permeable dressing.
  - Replace the dressing if it becomes damp, loosened, or visibly soiled.

Elastomeric devices and dial flow regulators are specialized infusion devices that may be considered for prior authorization when the device:

- Will be used for short-term medication administration (less than two weeks duration).
- Is expected to increase client compliance.
• Will better facilitate drug administration.
• Costs less than the cost of pump rental or tubing.
• The caregiver can not administer the infusion by pump.

The following criteria must be met for prior authorization of a stationary infusion pump:
• An infusion pump is required to safely administer the drug.
• The standard method of administration of the drug is through prolonged infusion or intermittent
  infusion, and the infusion rate must be more consistent than can be obtained with gravity drainage.
• The drug being administered requires IV infusion (i.e., the drug cannot be administered orally,
  intramuscularly, or by push technique).

The following criteria must be met for prior authorization of an ambulatory infusion pump:
• An infusion pump is required to safely administer the drug.
• The standard method of administration of the drug is through prolonged infusion or intermittent
  infusion and the infusion rate must be more consistent than can be obtained with gravity drainage.
• The drug being administered requires IV infusion (i.e., the drug cannot be administered orally,
  intramuscularly, or by push technique).
• The infusion administration is more than two hours and the client is involved in activities away
  from home, including but not limited to, physician visits.

2.2.16.2 Documentation Requirements
To request prior authorization for IV supplies and equipment, the following documentation must be
provided:
• Diagnostic information pertaining to the underlying diagnosis or condition
• A physician’s order and documentation supporting medical necessity
• The medication and dose being administered, the duration of drug therapy, and the frequency of
  administration

If additional supplies are needed beyond the standards listed, prior authorization may be considered
with documentation supporting medical necessity.

For additional IV access catheters, supporting documentation must have evidence that includes, but is
not limited to, the following:
• Dehydration
• Vein scarring
• Fragile veins, including but not limited to, clients who are infants or elderly

For more frequent IV site changes, supporting documentation must have evidence that includes, but is
not limited to, the following:
• Phlebitis
• Infiltration
• Extravasation

For more frequent IV tubing or add-on changes, supporting documentation must have evidence that
includes, but is not limited to, the following:
• Phlebitis
• IV catheter-related infection
• The administered infusion requires more frequent tubing changes

2.2.17 Mobility Aids

Mobility aids and related supplies, including, but not limited to canes, crutches, walkers, wheelchairs, and ramps are a benefit through Title XIX Home Health Services to assist clients to move about in their environment.

Mobility aids and related supplies, including, but not limited to, strollers, special-needs car seats, travel safety restraints, and thoracic-hip-knee-ankle orthoses (THKAO)/parapodiums are a benefit to assist clients to move about in their environment when medically necessary and Federal Financial Participation is available.

Mobility aids and related supplies may be considered for reimbursement through CCP for clients who are 20 years of age or younger who are CCP-eligible when documentation submitted clearly shows that the equipment is medically necessary and will correct or ameliorate the client’s disability or physical or mental illness or condition. Documentation must include the following:

• The client’s mobility status would be compromised without the requested equipment.

• The requested equipment or supplies are safe for use in the home.

Mobility aids may be considered through CCP if the requested equipment is not available through Title XIX Home Health Services or the client does not meet criteria through Title XIX Home Health Services.

Note: A mobility aid for a client who is birth through 20 years of age is medically necessary when it is required to correct or ameliorate a disability or physical illness or condition.

2.2.17.1 Canes, Crutches, and Walkers

Canes, crutches, and walkers are a benefit through Title XIX Home Health Services when medically necessary to assist clients to move about in their environment. Walkers require prior authorization. Prior authorization is not required for canes, crutches, or walker accessories. Documentation of medical necessity must be provided by a physician familiar with the client and must include information on the client’s impaired mobility.

2.2.17.2 Wheeled Mobility Systems

A wheeled mobility system is a manual or power wheelchair, or scooter that is a customized power or manual mobility device, or a feature or component of the mobility device, including, but not limited to, the following:

• Seated positioning components

• Manual seating options

• Adjustable frame

• Other complex or specialized components

A stroller (a multipositional client transfer system with integrated seat, operated by caregiver) for medical needs may be considered for clients who are CCP-eligible when documentation submitted clearly shows that the equipment is medically necessary and will correct or ameliorate the client’s disability or physical or mental illness or condition. Documentation must include the following:

• The client does not own another seating system, including, but not limited to, a wheelchair

• The client’s condition does not require another type of seating system, including, but not limited to, a wheelchair

If the client does not meet criteria for a stroller, a wheelchair may be considered through Texas Medicaid (Title XIX) Home Health Services.
Scooters may be considered for reimbursement through Texas Medicaid (Title XIX) Home Health Services.

A wheelchair is a non-customized chair mounted on four wheels that incorporates a non-adjustable frame, a sling or solid back and seat, and arm rests. Optional items included in this definition include, but are not limited, to the following:

- Handles at the back
- Foot rest
- Seat belt or safety restraint

A wheelchair includes all of the following:

- Standard (manual) wheelchairs
- Standard hemi (manual) wheelchairs
- Standard reclining (manual) wheelchairs
- Lightweight (manual) wheelchairs
- High strength lightweight (manual) wheelchairs

### 2.2.17.2.1 Prior Authorization

A wheelchair may be prior authorized for short-term rental or for purchase with documentation supporting medical necessity and an assessment of the accessibility of the client’s residence to ensure that the wheelchair is usable in the home (i.e., doors and halls wide enough, no obstructions).

### 2.2.17.2.2 Documentation Requirements

Documentation by a physician familiar with the client must include information on the client’s impaired mobility and physical requirements. In addition, the following information must be submitted with documentation of medical necessity:

- Why the client is unable to ambulate a minimum of 10 feet due to their condition (including, but not limited to, AIDS, sickle cell anemia, fractures, a chronic diagnosis, or chemotherapy)
- If the client is able to ambulate further than 10 feet, why a wheelchair is required to meet the client’s needs

### 2.2.17.3 Manual Wheelchairs-Standard, Standard Hemi, and Standard Reclining

A standard manual wheelchair is defined as a manual wheelchair that:

- Weighs more than 36 pounds.
- Does not have features to appropriately accept specialized seating or positioning.
- Has a weight capacity of 250 pounds or less.
- Has a seat depth of between 15 and 19 inches.
- Has a seat width of between 15 and 19 inches.
- Has a seat height of 19 inches or greater.
- Is fixed height only, fixed, swing away, or detachable armrest.
- Is fixed, swing away, or detachable footrest.

A standard hemi (low seat) wheelchair is defined as a manual wheelchair that:

- Has the same features as a standard manual wheelchair.
- Has a seat to floor height of less than 19 inches.
A standard reclining wheelchair is defined as a manual wheelchair that:

- Has the same features as a standard or standard hemi manual wheelchair.
- Has the ability to allow the back of the wheelchair to move independently of the seat to provide a change in orientation by opening the seat-to-back angle and, in combination with leg rests, open the knee angle.

2.2.17.3.1 Prior Authorization

A standard manual wheelchair may be considered for prior authorization for short-term rental or purchase when all the following criteria are met:

- The client has impaired mobility and is unable to consistently ambulate more than 10 feet.
- The client does not require specialty seating components.

A standard hemi wheelchair may be considered for prior authorization for short-term rental or purchase when the client meets criteria for a standard manual wheelchair and the following criteria is met:

- The client requires a low seat-to-floor height.
- The client must use their feet to propel the wheelchair.

A standard reclining wheelchair may be considered for prior authorization for short-term rental or purchase when the client meets criteria for a standard manual wheelchair and one or more of the following criteria are met:

- The client develops fatigue with longer periods of sitting upright.
- The client is at increased risk of pressure sores with prolonged upright position.
- The client requires assistance with respirations in a reclining position.
- The client needs to perform mobility related activities of daily living (MRADLs) in a reclining position.
- The client needs to improve venous return from lower extremity in a reclining position.
- The client has severe spasticity.
- The client has excess extensor tone of the trunk muscles.
- The client has quadriplegia.
- The client has a fixed hip angle.
- The client must rest in a reclining position two or more times per day.
- The client has the inability or has great difficulty transferring from wheelchair to bed.
- The client has trunk or lower extremity casts or braces that require the reclining feature for positioning.

2.2.17.4 Manual Wheelchairs-Lightweight and High-Strength Lightweight

A lightweight manual wheelchair is defined as a manual wheelchair that:

- Has the same features as a standard or hemi manual wheelchair.
- Weighs 34 to 36 pounds.
- Has available arm styles that are height adjustable.

A high-strength lightweight wheelchair is defined as a manual wheelchair that:

- Has the same features as a lightweight manual wheelchair.
- Weighs 30 to 34 pounds.
• Has a lifetime warranty on side frames and cross braces.

2.2.17.4.1 Prior Authorization

A lightweight manual wheelchair may be considered for prior authorization for rental or purchase when all the following criteria are met:

• The client is unable to propel a standard manual wheelchair at home.
• The client is capable of independently propelling a lightweight wheelchair to meet their MRADLs at home.

A high-strength lightweight wheelchair may be considered for prior authorization for rental or purchase when the client meets all of the criteria for a lightweight manual wheelchair and meets one or more of the following criteria:

• The high-strength lightweight wheelchair will allow the client to self-propel while engaging in frequently performed activities that cannot otherwise be completed in a standard or lightweight wheelchair.
• The client requires frame dimensions (seat width, depth, or height) that cannot be accommodated in a standard, lightweight, or hemi wheelchair and the wheelchair is used at least 2 hours a day.

2.2.17.5 Manual Wheelchairs-Heavy-Duty and Extra Heavy Duty

A heavy duty wheelchair is defined as a manual wheelchair that:

• Meets the standard manual wheelchair definition.
• Has a weight capacity greater than 250 pounds.

An extra heavy duty wheelchair is defined as a manual wheelchair that:

• Meets the standard manual wheelchair definition.
• Has a weight capacity greater than 300 pounds.

2.2.17.5.1 Prior Authorization

A heavy-duty wheelchair may be considered for prior authorization for short-term rental or purchase when the client has severe spasticity or all the following criteria are met:

• The client meets criteria for a standard manual wheelchair.
• The client weighs between 250 and 300 pounds.

An extra heavy duty wheelchair may be considered for prior authorization for short-term rental or purchase when all the following criteria are met:

• The client meets criteria for a standard manual wheelchair.
• The client weighs more than 300 pounds.

2.2.17.6 Wheeled Mobility Systems

A wheeled mobility system is a manual or power wheelchair, or scooter that is a customized power or manual mobility device, or a feature or component of the mobility device, including but not limited to, the following:

• Seated positioning components
• Powered or manual seating options
• Specialty driving controls for powered chairs
• Adjustable frame
• Other complex or specialized components
A wheeled mobility system includes all of the following:

- Tilt-in-space (manual) wheelchairs
- Pediatric size (manual) wheelchairs and strollers
- Custom ultra lightweight (manual) wheelchairs
- All power wheelchairs
- All scooters

2.2.17.6.1 Definitions and Responsibilities

The following definitions and responsibilities apply to the provision of wheeled mobility systems.

*Adjustments*—The adjustment of a component or feature of a wheeled mobility system.

Adjustments require labor only and do not include the addition, modification, or replacement of components or supplies needed to complete the adjustment.

Texas Medicaid will consider adjustments only to client-owned equipment that is considered a benefit of Texas Medicaid.

*Major Modification*—The addition of a custom or specialized feature or component of a wheeled mobility system that did not previously exist on the system due to changes in the client’s needs, including, but not limited to, the items listed in this paragraph. This definition also includes the modification of a custom or specialized feature or component due to a change in the client’s needs, including, but not limited to, the following:

- Seated positioning components, including, but not limited to, specialized seating or positioning components
- Powered or manual seating options, including, but not limited to, power tilt or recline seating systems and seat elevation systems
- Specialty driving controls, including, but not limited to, non-standard alternative power drive control systems
- Adjustable frame, including, but not limited to, non-standard seat frame dimensions
- Other complex or specialized components, including, but not limited to, power elevating leg rests and specialized electronic interfaces

The replacement of a previously existing custom or specialized feature or component with an identical or comparable component is considered a repair and not a major modification.

Texas Medicaid will consider major modifications only to client-owned equipment that is considered a benefit of Texas Medicaid.

*Minor Modification*—The addition or modification of non-custom or non-specialized features or components due to changes in the client’s needs, including but not limited to, the following:

- Armpads/armrests
- Legrests/Leg extensions
- Modification of seating and positioning components to accommodate for a change in the client’s size.

The replacement of a previously existing non-custom or non-specialized feature or component with an identical or comparable component is considered a repair and not a minor modification.

Texas Medicaid will consider minor modifications only to client-owned equipment that is considered a benefit of Texas Medicaid.
Mobility Related Activity of Daily Living (MRADL)—An activity of daily living, including, but not limited to, toileting, feeding, dressing, grooming, and bathing performed in customary locations in the residence.

Occupational Therapist—A person who is currently licensed by the Executive Council of Physical Therapy & Occupational Therapy Examiners to practice occupational therapy.

Physical Therapist—A person who is currently licensed by the Executive Council of Physical Therapy & Occupational Therapy Examiners to practice physical therapy.

Note: A physical or occupational therapist is responsible for completing the seating assessment of a client required for obtaining a wheeled mobility system.

Qualified Rehabilitation Professional (QRP)—A person who meets one or more of the following criteria:

- Holds a certification as an Assistive Technology Professional (ATP) or a Rehabilitation Engineering Technologist (RET) issued by, and in good standing with, the Rehabilitation Engineering and Assistive Technology Society of North America (RESNA).
- Holds a certification as a Seating and Mobility Specialist (SMS) issued by, and in good standing with, RESNA.
- Holds a certification as a Certified Rehabilitation Technology Supplier (CRTS) issued by, and in good standing with, the National Registry of Rehabilitation Technology Suppliers (NRRTS).

The QRP is responsible for:

- Being present at and involved in the seating assessment of the client for the rental or purchase of a wheeled mobility system.
- Being present at the time of delivery of the wheeled mobility system to direct the fitting of the system to ensure that the system functions correctly relative to the client.

Repairs—The replacement of a component or feature of a wheeled mobility system that is no longer functioning as designed, with an identical or comparable component that does not change the size or function of the system.

Texas Medicaid will consider repairs only to client-owned equipment that is considered a benefit of Texas Medicaid.

Additional Benefit Information

The initial purchase of all manual wheelchairs and wheeled mobility systems must include the wheelchair base or frame, and the following standard components, which will not be prior authorized separately:

- Complete set of standard propulsion and caster wheels, including all of the following:
  - Propulsion or caster tires of any size, made of solid rubber or plastic
  - Standard hand rims
  - Complete wheel lock assembly
  - Bearings
  - Standard footrest assembly (fixed, detachable, or swing away), including standard footplates, calf rests/pads, and ratchet assembly
  - Standard armrests (fixed non-adjustable or detachable non-adjustable), including standard foam or plastic arm pads
  - Standard seat and back upholstery
Medically necessary non-standard components may be considered for prior authorization with documentation of medical necessity for the requested component. Such components include, but are not limited to, the following:

- Flat-free inserts
- Foam filled propulsion or caster tires
- Pneumatic propulsion or caster tires
- Non-standard hand rims (including ergonomic and contoured)
- Non-standard length footrests
- Custom footrests
- Elevating footrests
- Angle adjustable footplates
- Adjustable height fixed armrests
- Adjustable height detachable armrests
- Custom size arm pads
- Gel arm pads
- Arm troughs
- Elevating leg rests

Claims for wheelchairs, components, and accessories must be submitted using the most appropriate procedure code that describes the item.

2.2.17.6.2 Prior Authorization

A wheeled mobility system may be prior authorized for short-term rental or for purchase with documentation supporting medical necessity and an assessment of the accessibility of the client’s residence to ensure that the wheelchair is usable in the home (i.e., doors and halls wide enough, no obstructions).

2.2.17.6.3 Documentation Requirements

Documentation by a physician familiar with the client must include information on the client’s impaired mobility and physical requirements. In addition, the following information must be submitted with documentation of medical necessity:

- Why the client is unable to ambulate a minimum of 10 feet due to their condition (including, but not limited to, AIDS, sickle cell anemia, fractures, a chronic diagnosis, or chemotherapy).
- If the client is able to ambulate further than 10 feet, why a wheelchair is required to meet the client’s needs.
- An itemized component list for custom manual or power wheeled mobility systems.
- A completed Wheelchair/Scooter/Stroller Seating Assessment Form with seating measurements that includes documentation supporting medical necessity, including:
  - For clients 12 years of age and younger, the wheelchair frame must accommodate a minimum of 3 inches of growth potential in width and depth.
  - For clients 13 years of age through 17 years of age, the wheelchair frame must accommodate a minimum of 2 inches of growth potential in width and depth.
• For clients 18 years of age and older, the wheelchair frame must accommodate a minimum of 1 inch in depth and 2 inches in width of growth potential.

Documentation of frame modifications or growth kits may be submitted to demonstrate growth allowances to the dimensions specified above.

When medically necessary, prior authorization may also be considered for the rental or purchase of an alternative wheelchair on a case-by-case basis, as follows:

• A manual wheelchair will be considered for a client who owns or is requesting a power wheeled mobility system with no custom features.

• A manual wheelchair or a manual wheeled mobility system will be considered for a client who owns or is requesting a power wheeled mobility system with custom features.

2.2.17.7 Manual Wheeled Mobility System - Tilt-in-Space

A tilt-in-space manual wheeled mobility system is defined as a manual wheelchair that meets the following requirements:

• Has the ability to tilt the frame of the wheelchair greater than or equal to 45 degrees from horizontal while maintaining a constant back to seat angle to provide a change of orientation and redistribute pressure from one area (such as the buttocks and the thighs) to another area (such as the trunk and the head)

• Adult size has a weight capacity of at least 250 pounds

• Pediatric size has a seat width or depth of less than 15 inches

2.2.17.7.1 Prior Authorization

A tilt-in-space wheeled mobility system may be considered for prior authorization for short-term rental or purchase when all the following criteria are met:

• The client meets criteria for a standard manual wheelchair.

• The client has a condition that meets criteria for a tilt-in-space feature, including but not limited to:
  • Severe spasticity
  • Hemodynamic problems
  • Quadriplegia
  • Excess extensor tone
  • Range of motion limitations prohibit a reclining system, such as hip flexors, hamstrings, or even heterotopic ossification
  • The need to rest in a recumbent position two or more times per day and the client has an inability to transfer between bed and wheelchair without assistance
  • Documented weak upper extremity strength or a disease that will lead to weak upper extremities
  • At risk for skin break down because of inability to reposition body in a chair to relieve pressure areas

2.2.17.8 Manual Wheeled Mobility System- Pediatric Size

A pediatric sized wheeled mobility system is defined as a manual standard/custom wheelchair (including those optimally configured for propulsion or custom seating) that has a seat width or depth of less than 15 inches.
2.2.17.9 Manual Wheeled Mobility System - Custom (Includes Custom Ultra-Lightweight)

Custom manual wheeled mobility systems may be considered for a client who meets criteria for a manual wheelchair, has a condition that requires specialized seating, and cannot safely utilize a standard manual wheelchair.

A custom ultra-lightweight wheeled mobility system is defined as an optimally configured wheelchair for independent propulsion which cannot be achieved in a standard, lightweight, or high-strength lightweight wheelchair that:

- Meets the high-strength lightweight definition and weighs less than 30 pounds.
- Has one or more of the following features to appropriately accept specialized seating or positioning:
  - Adjustable seat-to-back angle
  - Adjustable seat depth
  - Independently adjustable front and rear seat-to-floor dimensions
  - Adjustable caster stem hardware
  - Adjustable rear axle
  - Adjustable wheel camber
  - Adjustable center of gravity
- Has a lifetime warranty on side frames and cross braces

2.2.17.9.1 Prior Authorization

A custom ultra-lightweight wheeled mobility system may be considered for prior authorization for rental or purchase when the client meets all the criteria for a lightweight manual wheelchair and one or more of the following criteria:

- The client is able to self-propel, will have independent mobility with the use of an optimally configured chair, and meets all of the following criteria:
  - The client uses the wheelchair for a significant portion of their day to complete MRADLs.
  - The client uses the wheelchair in the community to complete MRADLs.
- The client is able to self-propel, will have independent mobility with the use of an optimally configured chair, has a medical condition that cannot be accommodated by the seating available on a standard, lightweight, or high-strength lightweight wheelchair and one or more of the following features needed by the client to ensure optimal independence with MRADLs:
  - Adjustable seat to back angle.
  - Adjustable seat depth.
  - Independently adjustable front and rear seat-to-floor dimensions.
  - Adjustable caster stem hardware.
  - Adjustable rear axle (adjustable center of gravity).
- The client meets all of the following criteria:
  - The client is unable to self-propel.
  - The client has a documented condition that requires custom seating, including, but not limited to:
    - Poor trunk control.
• Contractures of elbow or shoulders.
• Muscle spasticity.
• Tone imbalance through shoulders or back.
• Kyphosis or Lordosis.
• Lack of flexibility in pelvis or spine.
• The client requires custom seating that cannot be accommodated on a standard, lightweight, or hemi-wheelchair.

Prior authorization for labor to create a custom molded seating system is limited to a maximum of 15 hours.

A medical stroller does not have the capacity to accommodate the client’s growth. Strollers for medical use may be considered for prior authorization when all of the following criteria are met:

- The client weighs 30 pounds or more.
- The client does not already own another seating system, including, but not limited to, a standard or custom wheelchair.
- The stroller must have a firm back and seat, or insert.
- The client is expected to be ambulatory within one year of the request date or is not expected to need a wheelchair within two years of the request date.

To request prior authorization for the purchase of procedure code E1035, the criteria must be met for the level of stroller requested:

- Level One, Basic Stroller—The client meets the criteria for a stroller. Providers must use procedure code E1035.
- Level Two, Stroller with Tray for Oxygen or Ventilator—The client meets the criteria for a level-one stroller and is oxygen- or ventilator-dependent. Providers must use procedure code E1035 with modifier TF.
- Level Three, Stroller with Positioning Inserts—The client meets the criteria for a level-one or level-two stroller and requires additional positioning support. Providers must use procedure code E1035 with modifier TG.

The following supporting documentation must be submitted:

- A completed Wheelchair/Stroller Seating Assessment Form that includes documentation supporting medical necessity. This documentation must address why the client is unable to ambulate a minimum of 10 feet due to his or her condition (including, but not limited to, AIDS, sickle cell anemia, fractures, a chronic diagnosis, or chemotherapy), or if able to ambulate further, why a stroller is required to meet the client’s needs.
- If the client is three years of age or older, documentation must support that the client’s condition, stature, weight, and positioning needs allow adequate support from a stroller.

**Note:** A stroller may be considered on a case-by-case basis with documentation of medical necessity for a client who does not meet the criteria listed above.

A seating assessment must be completed by a physician or licensed occupational therapist or physical therapist, who is not employed by the equipment supplier, before requesting prior authorization.
2.2.17.10 Seating Assessment for Manual and Power Custom Wheelchairs

A seating assessment is required for:

- The rental or purchase of any device meeting the definition of a wheeled mobility system as defined in subsection 2.2.17.6, “Wheeled Mobility Systems” in this handbook.

- The purchase of any device meeting the definition of a wheelchair as defined under subsection 2.2.17.2, “Wheeled Mobility Systems” in this handbook for a client with a congenital or neurological condition, myopathy, or skeletal deformity, which requires the use of a wheelchair.

A seating assessment is required for the rental or purchase of any device meeting the definition of a wheeled mobility system or purchase of any device meeting the definition of a wheelchair for a client with a congenital or neurological condition, myopathy, or skeletal deformity that requires the use of a wheelchair as defined under subsection 2.6.9.1.2, “Wheeled Mobility Systems” in this handbook.

A seating assessment with measurements, including specifications for exact mobility/seating equipment and all necessary accessories, must be completed by a physician, licensed occupational therapist, or licensed physical therapist.

A QRP directly employed or contracted by the DME provider must be present at and participate in all seating assessments, including those provided by a physician.

Upon completion of the seating assessment, the QRP must attest to his or her participation in the assessment by signing the Wheelchair/Scooter/Stroller Seating Assessment Form. This form must be submitted with all requests for wheeled mobility systems.

When the practitioner completing the seating assessment is an occupational or physical therapist, the occupational or physical therapist may perform the seating assessment as the therapist, or as the QRP, but may not perform in both roles at the same time. If the occupational or physical therapist is attending the seating assessment as the QRP, the occupational or physical therapist must meet the credentialing requirements and be enrolled in Texas Medicaid as a QRP.

If the practitioner completing the seating assessment is a physician, the seating assessment is considered part of the evaluation and management service provided.

If the seating assessment is completed by a physician, reimbursement is considered part of the physician’s office visit and will not be reimbursed separately.

2.2.17.10.1 Prior Authorization

A seating assessment performed by an occupational therapist, physical therapist, or a physician, with the participation of a QRP, does not require prior authorization. A seating assessment performed by a physician is considered part of the physician evaluation and management service.

The QRP’s participation in the seating assessment requires authorization before the service can be reimbursed. Authorization must be requested at the same time and on the same prior authorization request form as the prior authorization request for the QRP fitting and the wheeled mobility system or major modification to the wheeled mobility system.

Prior authorization requests for the QRP’s participation in the seating assessment will be returned to the provider if the seating assessment is requested separately from the prior authorization for the QRP fitting and the wheeled mobility system or major modification to the wheeled mobility system.

The QRP participating in the seating assessment must be directly employed by or contracted with the DME provider requesting the wheeled mobility system or major modification to a wheeled mobility system.

An authorization for the QRP’s participation in the seating assessment for a wheeled mobility system or major modification to a wheeled mobility system may be issued to the QRP in 15-minute increments, for a time period of up to one hour (4 units).
If the seating assessment is completed by a physician, reimbursement is considered part of the physician office visit and will not be reimbursed separately.

If the seating assessment is completed by a physician, reimbursement is considered part of the physician office visit and will not be reimbursed separately.

The physical therapist completing the seating assessment must submit procedure code 97542 with the GP and UC modifiers to bill for the seating assessment.

The occupational therapist completing the seating assessment must submit procedure code 97542 with the GO and UC modifiers to bill for the seating assessment.

Services for the QRP’s participation in the seating assessment must be submitted for reimbursement by the DME provider billing for the wheeled mobility system using procedure code 97542 with modifier U1. The DME provider must include the QRP specialty as the performing provider on the claim for all components of the wheeled mobility system, including the QRP’s participation in the seating assessment.

Seating assessments are reimbursed in 15-minute increments (units) and are limited to four units (one hour).

2.2.17.10.2 Documentation Requirements

The seating assessment must clearly show that the equipment is medically necessary and will correct or ameliorate the client’s disability or physical or mental illness or condition.

Documentation must include the following:

- Explain how the client or family will be trained in the use of the equipment.
- Anticipate changes in the client’s needs and include anticipated modifications or accessory needs, as well as the growth potential of the wheelchair.
- Include significant medical information pertinent to the client’s mobility and how the requested equipment will accommodate these needs, including intellectual, postural, physical, sensory (visual and auditory), and physical status.
- Address trunk and head control, balance, arm and hand function, existence and severity of orthopedic deformities, as well as any recent changes in the client’s physical and/or functional status, and any expected or potential surgeries that will improve or further limit mobility.
- Include information on the client’s current mobility/seating equipment, how long the client has been in the current equipment and why it no longer meets the client’s needs.
- Include the client’s height, weight, and a description of where the equipment is to be used.
- Include seating measurements.
- Include the accessibility of client’s residence.
- Include manufacturer’s information, including the description of the specific base, any attached seating system components, and any attached accessories, as well as the manufacturer’s retail pricing information and itemized pricing for manually priced components.
- Include documentation supporting medical necessity for all accessories.
- Be documented on the Wheelchair/Scooter/Stroller Seating Assessment Form, which must be signed and dated by the qualified practitioner completing the assessment (occupational therapist, physical therapist, or physician), and the QRP who was present and participated in the assessment.
- Be submitted with the prior authorization request for the wheeled mobility system. The Form must be completed, signed and dated as outlined above.
2.2.17.11 Fitting of Custom Wheeled Mobility Systems

The fitting of a wheeled mobility system is defined as the time the QRP spends with the client fitting the various systems and components of the system to the client. It may also include time spent training the client or caregiver in the use of the wheeled mobility system. Time spent setting up the system, or travel time without the client present, is not included.

A fitting is required for any device meeting the definition of a wheeled mobility system as defined under subsection 2.2.17.6, “Wheeled Mobility Systems” in this handbook.

The fitting of a wheeled mobility system must be:

- Performed by the same QRP that was present for, and participated in, the seating assessment of the client.
- Completed prior to submitting a claim for reimbursement of a wheeled mobility system.

The QRP performing the fitting will:

- Verify the wheeled mobility system has been properly fitted to the client.
- Verify that the wheeled mobility system will meet the client’s functional needs for seating, positioning, and mobility.
- Verify that the client, parent, guardian of the client, and/or caregiver of the client has received training and instruction regarding the wheeled mobility system’s proper use and maintenance.

The QRP must complete and sign the DME Certification and Receipt form after the wheeled mobility system has been delivered and fitted to the client. Completion of this form by the QRP signifies that all components of the fitting as outlined above have been satisfied. The form must be completed prior to submission of a claim for a wheeled mobility system, and submitted to HHSC’s designee according to instructions on the form to allow for proper claims processing.

Services for fitting of a wheeled mobility system by the QRP must be submitted for reimbursement by the DME provider of the wheeled mobility system using procedure code 97542 with modifier U2. The DME provider must list the QRP who participated in the seating assessment as the performing provider on the claim for all components of the wheeled mobility system, including the fitting performed by the QRP.

All adjustments and modifications to the wheeled mobility system, as well as the associated services by the QRP for the seating assessment and fitting, within the first six months after delivery are considered part of the purchase price and will not be separately reimbursed.

Procedure code 97542 with modifier U2 must be billed on the same claim as the procedure code(s) for the wheeled mobility system in order for both services to be reimbursed.

2.2.17.11.1 Prior Authorization

Prior authorization is required for the QRP performing the fitting of a wheeled mobility system, and must be included with the request for the wheeled mobility system.

The QRP must be directly employed by or contracted with the DME company providing the system, and must be the same QRP who was present at and participated in the client’s seating assessment.
A prior authorization may be issued to the QRP in 15-minute increments, for a time period of up to two hours (8 units), for the fitting of any manual or power wheeled mobility system. Up to one additional hour (4 units) may be authorized to the QRP with documentation of medical necessity demonstrating that fitting of three or more major systems is required, or that additional client training is required for such systems. Major systems can include, but are not limited to, the following:

- Complete complex seating system (planar system with trunk supports and hip supports or abductor or custom contoured seating system such as a molded system) Off-the-shelf seat and back cushions do not constitute a complex seating system.
- Alternative drive controls (such as a head array, mini-proportional system, etc.).
- Additional specialty control features (such as infrared access).
- Power positioning features (such as power tilt, power recline).
- Specific purpose specialty features (such as power seat elevation systems, power elevating leg rests).

2.2.17.11.2 Documentation Requirements

When the QRP that participated in the assessment of the client is not available to conduct the fitting of the wheeled mobility system, the DME provider must update the prior authorization for the wheeled mobility system and fitting by submitting all of the following information:

- A letter written on the DME provider’s letterhead, signed and dated by a representative of the DME provider other than the new QRP.
- Documentation explaining why the original QRP could not conduct the fitting. Examples may include, but are not limited to, documentation that the QRP:
  - Is no longer associated with the DME provider requesting the wheeled mobility system.
  - Is on an extended leave from the DME provider requesting the wheeled mobility system.
  
  **Note:** For purposes of this policy, an extended leave is any leave of more than 30 consecutive calendar days.
- The name, TPI, and NPI of the original QRP who performed the initial assessment, and the date the assessment was completed.
- The name, TPI, and NPI of the QRP who will be performing the fitting.
- A copy of the original, physician-signed Home Health Services (Title XIX) DME/Medical Supplies Physician Order Form.

A copy of this documentation must be maintained by the provider in the client’s medical record and be available upon request by HHSC or its designee.

2.2.17.12 Power Wheeled Mobility Systems- Group 1 through Group 5

A power wheeled mobility system or powered mobility device (PMD) is a professionally manufactured device that provides motorized wheeled mobility and body support specifically for individuals with impaired mobility. PMDs are four- or six-wheeled motorized vehicles whose steering is operated by an electronic device or joystick to control direction, turning, and alternative electronic functions, such as seat controls.

Each PMD must include all of the following basic components that may not be billed separately:

- Lap belt or safety belt (This does not include multiple-attachment-point positioning belts or padded belts.)
- Battery charger, single mode
- Batteries (initial)
• Complete set of tires and casters, any type
• Leg rests
• Foot rests or foot platform
• Arm rests
• Any weight-specific components (braces, bars, upholstery, brackets, motors, gears, etc.) as required by client weight capacity
• Controller and input device

The following definitions apply to PMDs:

• No-Power Option - A category of PMDs that cannot accommodate a power tilt, recline, or seat elevation system. A PMD that can accept only power-elevating leg rests is considered to be a no-power option chair.

• Single-Power Option - A category of PMDs that can accept and operate a power tilt, power recline, or a power seat elevation system, but not a combination power tilt and recline seating system. A single-power option PMD might be able to accommodate power elevating leg rests, or seat elevator, in combination with a power tilt or power recline. A PMD does not have to be able to accommodate all features to meet this definition.

• Multiple-Power Option - A category of PMDs that can accept and operate a combination power tilt and recline seating system. A multiple-power option PMD might also be able to accommodate power elevating leg rests, or a power seat elevator. A PMD does not have to accommodate all features to qualify to meet this definition.

2.2.17.12.1 Prior Authorization

Prior authorization for a power wheeled mobility system/PMD requires the following documentation in addition to all documentation required for a custom manual wheelchair:

• The client’s physical and mental ability to receive and follow instructions related to responsibilities of using equipment. The client must be able to operate a PMD independently. The therapist must provide written documentation that the client is physically and cognitively capable of managing a PMD.

• How the PMD will be operated (i.e., joystick, head pointer, puff-and-go).

• The capability of the client to understand how the PMD operates.

• The capability of the caregiver or client to care for the PMD and accessories.

2.2.17.12.2 Group 1 PMDs

All Group 1 PMDs must have all the specified basic components and meet all the following requirements:

• Standard integrated or remote proportional joystick
• Nonexpandable controller
• Incapable of upgrade to expandable controller
• Incapable of upgrade to alternative control devices
• May have cross brace construction
• Accommodates nonpowered options and seating systems (e.g., recline-only backs, manually elevating leg rests [except captains chairs])
• Length - less than or equal to 40 inches
• Width - less than or equal to 24 inches
• Minimum top end speed - 3 mph
• Minimum range - 5 miles
• Minimum obstacle climb - 20 mm
• Dynamic stability incline - 6 degrees

Prior Authorization Requirements
A Group 1 PMD may be considered for prior authorization for rental or purchase when all the following criteria are met:
• The client will use the PMD for less than 2 hours per day.
• The client will use the PMD indoors on smooth, hard surfaces.
• The client will not encounter obstacles in excess of 0.75 inch.

2.2.17.12.3 Group 2 PMDs
All Group 2 PMDs must have all the specified basic components and meet all the following requirements:
• Standard integrated or remote proportional joystick
• May have cross brace construction
• Accommodates seating and positioning items (e.g., seat and back cushions, headrests, lateral trunk supports, lateral hip supports, medical thigh supports [except captains chairs])
• Length - less than or equal to 48 inches
• Width - less than or equal to 34 inches
• Minimum top end speed - 3 mph
• Minimum range - 7 miles
• Minimum obstacle climb - 40 mm
• Dynamic stability incline - 6 degrees

Prior Authorization Requirements
A Group 2 PMD may be considered for prior authorization for rental or purchase when the following criteria are met:
• The client will use the PMD for 2 or more hours per day.
• The client will not routinely use the PMD for MRADLs outside the home.
• The client will not encounter obstacles in excess of 1.5 inches.

2.2.17.12.4 Group 3 PMDs
All Group 3 PMDs must have all the specified basic components and meet all the following requirements:
• Standard integrated or remote proportional joystick
• Nonexpandable controller
• Capable of upgrade to expandable controller
• Capable of upgrade to alternative control devices
• May not have cross brace construction
• Accommodates seating and positioning items (e.g., seat and back cushions, headrests, lateral trunk supports, lateral hip supports, medial thigh supports [except captains chairs])
• Drive wheel suspension to reduce vibration
• Length - less than or equal to 48 inches
• Width - less than or equal to 34 inches
• Minimum top end speed - 4.5 mph
• Minimum range - 12 miles
• Minimum obstacle climb - 60 mm
• Dynamic stability incline - 7.5 degrees

**Prior Authorization Requirements**
A Group 3 PMD may be considered for prior authorization for rental or purchase when the following criteria are met:

• The client’s mobility limitation is due to a neurological condition, myopathy, or congenital skeletal deformity.
• The client may routinely use the PMD for MRADLs outside of the home.
• The client will use the PMD primarily on smooth or paved surfaces.
• The client will not encounter obstacles in excess of 2.5 inches.

### 2.2.17.12.5 Group 4 PMDs
All Group 4 PMDs must have all the specified basic components and meet all the following requirements:

• Standard integrated or remote proportional joystick
• Nonexpandable controller
• Capable of upgrade to expandable controller
• Capable of upgrade to alternative control devices
• May not have cross brace construction
• Accommodates seating and positioning items (e.g., seat and back cushions, headrests, lateral trunk supports, lateral hip supports, medial thigh supports [except captains chairs])
• Drive wheel suspension to reduce vibration
• Length - less than or equal to 48 inches
• Width - less than or equal to 34 inches
• Minimum top end speed - 6 mph
• Minimum range - 16 miles
• Minimum obstacle climb - 75 mm
• Dynamic stability incline - 9 degrees
Prior Authorization Requirements

A Group 4 PMD may be considered for prior authorization for rental or purchase when all the following criteria are met:

- In addition to using the PMD in the home, the client will routinely use the PMD for MRADLs outside the home.
- The client will routinely use the PMD on rough, unpaved or uneven surfaces.
- The client will encounter obstacles in excess of 2.25 inches.
- The client has a documented medical need for a feature that is not available on a lower level PMD.

Documentation Requirements

The submitted documentation for a Group 4 PMD must include a completed assessment that is signed and dated by a physician or a licensed occupational or physical therapist and includes the following:

- A description of the environment where the PMD will be used in the routine performance of MRADLs.
- A listing of the MRADLs that would be possible with the use of a Group 4 PMD that would not be possible without the Group 4 PMD.
- The distance the client is expected to routinely travel on a daily basis with the Group 4 PMD.

Note: The enhanced features found on a Group 4 PMD must be medically necessary to meet the client’s routine MRADL and will not be approved for leisure or recreational activities.

In addition to meeting criteria for Group 2 through Group 4 PMDs, the submitted documentation of medical necessity must demonstrate that the client requires the requested power option (e.g., the need for a power recline or tilt in space, or a combination power tilt and power recline), the no-power option, single-power option, or multiple-power option as defined in subsection 2.2.17.12, “Power Wheeled Mobility Systems- Group 1 through Group 5” in this handbook.

2.2.17.12.6 Additional Requirements - Group 2 through Group 4 No-Power Option

Group 2 through Group 4 no-power option PMDs must have all the specified basic components and meet all the following requirements:

- Nonexpandable controller
- Incapable of upgrade to expandable controller
- Incapable of upgrade to alternative control devices
- Meets the definition of no-power option
- Accommodates nonpowered options and seating systems (e.g., recline-only backs, manually elevating leg rests [except captains chairs])

2.2.17.12.7 Group 2 through Group 4 Single-Power Option

Group 2 through Group 4 single-power option PMDs must have all the specified basic components and meet all the following requirements:

- Nonexpandable controller
- Capable of upgrade to expandable controller
- Capable of upgrade to alternative control devices
- Meets the definition of single-power option
2.2.17.12.8 Group 2 through Group 4 Multiple-Power Option

Group 2 through Group 4 multiple-power option PMDs must have all the specified basic components and meet all the following requirements:

- Nonexpandable controller
- Capable of upgrade to expandable controller
- Meets the definition of multiple-power option
- Accommodates a ventilator

2.2.17.12.9 Group 5 PMDs

All Group 5 PMDs must have all the specified basic components and meet all the following requirements:

- Standard integrated or remote joystick
- Nonexpandable controller
- Capable of upgrade to expandable controller
- Seat width - minimum of 5 one-inch options
- Seat depth - minimum of 3 one-inch options
- Seat height - adjustment requirements = 3 inches
- Back height - adjustment requirements minimum of 3 options
- Seat-to-back angle range of adjustment - minimum of 12 degrees
- Accommodates nonpowered options and seating systems
- Accommodates seating and positioning items (e.g., seat and back cushions, headrests, lateral trunk supports, lateral hip supports, medial thigh supports)
- Adjustability for growth (minimum of 3 inches for width, depth, and back height adjustment)
- Special developmental capability (i.e., seat to floor, standing, etc.)
- Drive wheel suspension to reduce vibration
- Length - less than or equal to 48 inches
- Width - less than or equal to 34 inches
- Minimum top end speed - 4 mph
- Minimum range - 12 miles
- Minimum obstacle climb - 60 mm
- Dynamic stability incline - 9 degrees
- Passed crash test

Prior Authorization Requirements

A Group 5 pediatric PMD may be considered for prior authorization for rental or purchase when all the following criteria are met:

- The client weighs less than 125 pounds.
- The client is expected to grow in height.
- The client may require growth of up to 5 inches in width.
• The client may require a change in seat to floor height up to 3 inches.
• The client may require a seat to back angle range of adjustment in excess of 12 degrees.
• The client requires special developmental capability (i.e., seat to floor, standing, etc.).

2.2.17.10 Group 5 Single-PMDs
A group 5 single-power option PMD must have all the specified basic components and have the capability to accept and operate a power tilt or recline or seat elevation system, but not a combination power tilt and recline seating system, and may be able to accommodate power elevating leg rests, or seat elevator, in combination with a power tilt or power recline.

Prior Authorization Requirements
A Group 5 pediatric PMD with single power option may be considered for prior authorization for rental or purchase when all the following criteria are met:
• The client meets criteria for a Group 5 PMD.
• The client requires a drive control interface other than a hand or chin-operated standard proportional joystick (examples include but are not limited to head control, sip and puff, or switch control).

2.2.17.11 Group 5 Multiple-PMDs
Group 5 multiple-power option PMD must have all the specified basic components and meet all the following requirements:
• Has the capability to accept and operate a combination power tilt and recline seating system, and may also be able to accommodate power elevating leg rests, or a power seat elevator.
• Accommodates a ventilator.

Prior Authorization Requirements
A Group 5 pediatric PMD with multiple power option may be considered for prior authorization for rental or purchase when the following criteria are met:
• The client meets criteria for a Group 5 PMD.
• The client requires a drive control interface other than a hand or chin-operated standard proportional joystick (examples include but are not limited to head control, sip and puff, switch control).
• The client has a documented medical need for a power tilt and recline seating system and the system is being used on the wheelchair or the client uses a ventilator which is mounted on the wheelchair.

2.2.17.13 Wheelchair Ramp-Portable and Threshold
Portable and threshold ramps are a benefit of Texas Medicaid.
A portable ramp is defined as a unit that is able to be carried as needed to access a home, weighs no more than 90 pounds, or measures no more than 10 feet in length. A threshold ramp is defined as a unit that provides access over elevated thresholds.
Portable ramps exceeding the above criteria may be considered on a case-by-case basis with documentation of medical necessity and a statement that the requested equipment is safe for use.
One portable ramp and one threshold ramp for wheelchair or stroller access may be considered for prior authorization when documentation supports medical necessity. The following documentation supporting medical necessity is required:
• The date of purchase and serial number of the client’s wheelchair or documentation of a wheelchair request being reviewed for purchase
• Diagnosis with duration of expected need
• A diagram of the house showing the access points with the ground-to-floor elevation and any obstacles

Providers must use procedure code E1399 for the purchase of portable and threshold stroller ramps.

A request for prior authorization must include documentation from the provider to support the medical necessity of the service, equipment, or supply.

Note: Permanent ramps, vehicle ramps, and home modifications are not a benefit of Texas Medicaid.

Ramps may be considered for rental for short term disabilities and for purchase for long term disabilities. Mobility aid lifts for vehicles and vehicle modifications are not a benefit of Texas Medicaid.

### 2.2.17.14 Power Elevating Leg Lifts

A power elevation feature involves a dedicated motor and related electronics with or without variable speed programmability, which allows the leg rest to be raised and lowered independently of the recline and/or tilt of the seating system. It includes a switch control which may or may not be integrated with the power tilt and/or recline control(s).

#### 2.2.17.14.1 Prior Authorization

Power elevating leg lifts may be prior authorized for clients who have compromised upper extremity function that limits the client’s ability to use manual elevating leg rests. The client must meet criteria for a PMD with a reclining back and at least one of the following:

- The client has a musculoskeletal condition such as flexion contractures of the knees and legs, or the placement of a brace that prevents 90-degree flexion at the knee.
- The client has significant edema of the lower extremities that requires elevating the client’s legs.
- The client experiences hypotensive episodes that require frequent positioning changes.
- The client needs power tilt-and-recline and is required to maintain anatomically correct positioning and reduce exposure to skin shear.

#### 2.2.17.14.2 Documentation Requirements

The submitted documentation must include an assessment completed, signed, and dated by a physician or a licensed occupational or physical therapist that includes the following:

- A description of the client’s current level of function without the device
- Documentation that identifies how the power elevating leg lifts will improve the client’s function
- A list of MRADLs the client will be able to perform with the power elevating leg lifts that the client is unable to perform without the power elevating leg lifts and how the device will increase independence
- The duration of time the client is alone during the day without assistance
- The client’s goals for use of the power elevating leg lifts

### 2.2.17.15 Power Seat Elevation System

A power seat elevation system is used to raise and lower the client in their seated position without changing the seat angles to provide varying amounts of added vertical access.

The use of a power seat elevation system will:

- Facilitate independent transfers, particularly uphill transfers, to and from the wheelchair, and
- Augment the client’s reach to facilitate independent performance of MRADLs in the home.
2.2.17.15.1 Prior Authorization
A power seat elevation system may be prior authorized to promote independence in a client who meets all of the following criteria:

- The client does not have the ability to stand or pivot transfer independently.
- The client requires assistance only with transfers across unequal seat heights, and as a result of having the power seat elevation system, the client will be able to transfer across unequal seat heights unassisted.
- The client has limited reach and range of motion in the shoulder or hand that prohibits independent performance of MRADLs (such as, dressing, feeding, grooming, hygiene, meal preparation, and toileting).

2.2.17.15.2 Documentation Requirements
The submitted documentation must include an assessment completed, signed, and dated by a physician or a licensed occupational or physical therapist that includes the following:

- A description of the client’s current level of function without the device
- Documentation that identifies how the power seat elevation system will improve the client’s function
- A list of MRADLs the client will be able to perform with the power seat elevation system that the client is unable to perform without the power seat elevation system and how the device will increase independence
- The duration of time the client is alone during the day without assistance
- The client’s goals for use of the power seat elevation system

Note: A power seat elevation system option will not be authorized for the convenience of a caregiver, or if the device will not allow the client to become independent with MRADLs and transfers.

2.2.17.16 Seat Lift Mechanisms
A medically necessary seat lift mechanism is one that operates smoothly, can be controlled by the client, and effectively assists the client in standing up and sitting down without other assistance.

The payment for a recliner or chair with the incorporated seat lift mechanism is limited to the amount of the seat lift mechanism.

2.2.17.16.1 Prior Authorization
A seat lift mechanism may be prior authorized for clients who meet all the following criteria:

- The client must have severe arthritis of the hip or knee or have a severe neuromuscular disease.
- The seat lift mechanism must be a part of the physician’s course of treatment and be prescribed to correct or ameliorate the client’s condition.
- Once standing, the client must have the ability to ambulate.
- The client must be completely incapable of standing up from a regular armchair or any chair in their home.

Note: The fact that a client has difficulty or is even incapable of getting up from a chair, particularly a low chair, is not sufficient justification for a seat lift mechanism. Almost all clients who are capable of ambulating can get out of an ordinary chair if the seat height is appropriate and the chair has arms.
Seat lift mechanisms are limited to those types that operate smoothly, can be controlled by the client, and can effectively assist a client in standing up and sitting down without other assistance. A seat lift operated by a spring release mechanism with a sudden, catapult-like motion and jolts the client from a seated to a standing position is not a benefit of Texas Medicaid.

2.2.17.16.2 Documentation Requirements
The submitted documentation must include an assessment completed, signed, and dated by a physician or a licensed occupational or physical therapist that includes the following:

- A description of the client's current level of function without the device
- Documentation that identifies how the seat lift mechanism will improve the client’s function
- A list of MRADLs the client will be able to perform with the seat lift mechanism that the client is unable to perform without the seat lift mechanism and how the device will increase independence
- The duration of time the client is alone during the day without assistance
- The client’s goals for use of the seat lift mechanism

Supporting documentation must be kept in the client’s record that shows that all appropriate therapeutic modalities (such as medication, physical therapy) have been tried and that they failed to enable the client to transfer from a chair to a standing position.

2.2.17.17 Batteries and Battery Charger
A battery charger and initial batteries are included as part of the purchase of a PMD. Replacement batteries or a replacement battery charger may be considered for reimbursement if they are no longer under warranty.

A maximum of one hour of labor may be considered to install new batteries. Labor is not reimbursed with the purchase of a new PMD or with replacement battery chargers.

2.2.17.17.1 Prior Authorization
Batteries and battery chargers will not be prior authorized for replacement within six months of delivery. Batteries and battery chargers within the first six months after delivery are considered part of the purchase price.

A maximum of one hour of labor may be prior authorized to install new batteries. Labor will not be prior authorized for a new power wheelchair or for replacement battery chargers.

2.2.17.17.2 Documentation Requirements
To request prior authorization for replacement batteries or a replacement battery charger, the provider must document the date of purchase and serial number of the currently owned wheelchair as well as the reason for the replacement batteries or battery charger.

Documentation required supporting the need to replace the batteries or battery charger must include:

- Why the batteries are no longer meeting the client’s needs, or
- Why the battery charger is no longer meeting the client’s needs

2.2.17.18 Power Wheeled Mobility Systems- Scooter
A scooter is a professionally manufactured three- or four-wheeled motorized base operated by a tiller with a professionally manufactured basic seating system for clients who have little or no positioning needs.

A scooter must meet all the following requirements:

- Length- less than or equal to 48 inches
• Width- less than or equal to 28 inches
• Minimum top end speed- 3 mph
• Minimum range- 5 miles
• Minimum obstacle climb- 20 mm
• Radius pivot turn of less than or equal to 54 inches
• Dynamic stability incline- 6 degrees

Custom seating for scooters is not a benefit of Texas Medicaid Title XIX Home Health Services. Repairs to scooters will be considered only for a scooter purchased by the Texas Medicaid.

2.2.17.18.1 Prior Authorization
A scooter may be prior authorized for ambulatory-impaired clients with good head, trunk, and arm/hand control, without a diagnosis of progressive illness (including, but not limited to, progressive neuro-muscular diseases such as amyotrophic lateral sclerosis [ALS]).

To request prior authorization for a scooter, the client must not own a power wheelchair.

A scooter may be prior authorized for a short-term rental or an initial three-month trial rental period based on documentation supporting the medical necessity and appropriateness of the device.

Assessment of the accessibility of the client’s residence must be completed and included in the prior authorization documentation to ensure that the scooter is usable in the home (i.e., doors and halls wide enough, no obstructions).

2.2.17.18.2 Documentation Requirements
Prior authorization for a scooter requires all the documentation required for a standard power wheelchair and meets all the following criteria:

• The client’s physical and cognitive ability to receive and follow instructions related to the responsibilities of using the equipment.
• The ability of the client to physically and cognitively operate the scooter independently.
• The capability of the client to care for the scooter and understand how it operates.

2.2.17.19 Client Lifts
A lift is an item of DME that is a mechanical system used to lift or transfer a nonambulatory client between a bed, chair, wheelchair, bedside commode, bathroom, or other location. A lift may be medically necessary to ameliorate a client’s medical condition or disability that results in impaired functional mobility impacting mobility related activities of daily living (MRADLs).

Electric and hydraulic lifts are movable single-stand mechanisms, often on casters, with a lifting arm attached to a sling, and lifting power that is provided by a manual hydraulic pump or an electric motor. Electric and hydraulic lifts are benefits for clients of all ages through Title XIX Home Health Services with documented medical necessity.

2.2.17.20 Hydraulic Lifts
Hydraulic lifts require prior authorization.

2.2.17.20.1 Documentation Requirements
Prior authorization for a hydraulic lift may be considered with the following documentation:

• The inability of clients to assist in their own transfers
• The weight of the client and the weight capacity of the requested lift
• The availability of a caregiver to operate the lift
• Training by the provider to the client and the caregiver on the safe use of the lift

2.2.17.21 Electric Lifts
Prior authorization for an electric lift may be considered when the client meets criteria for a hydraulic lift and additional documentation explains why a hydraulic lift does not meet the client’s needs.

2.2.17.22 Overhead, Fixed, and Portable Lifts
Overhead and fixed client lifts will be considered for reimbursement through CCP for clients who are 20 years of age or younger and are CCP-eligible. In order to consider a client lift outside of those specified in the home health benefit, consideration must be given to the client’s medical needs (e.g., muscle tone, pain, fear, etc.), environmental factors, and caregiver abilities.

A client lift will not be prior authorized solely for the convenience of a caregiver. Prevention of caregiver injury or consideration of client safety during transfer due to caregiver factors, such as physical abilities, is not considered “caregiver convenience.”

Final set up and installation costs of client lifts, including labor costs associated with ceiling or other fixed lifts, are included in the initial purchase price of the client lift and will not be separately reimbursed. Components and accessories are also considered part of the initial purchase price of a client lift. Components and accessories include, but are not limited to, the following:

• Lift motor and gear box
• Any type of sling
• Hand controls and connectors
• Carry or spreader bar and sling attachments or straps
• Ceiling tracks or rails and components
• All mounting hardware and brackets
• Batteries
• Charger system
• Emergency stop and lowering systems
• Lifting tape
• Wheels or castors of any type
• Installation of the fixed lift systems

Repair or replacement of components, such as a sling, to client-owned equipment may be considered a benefit as needed with documentation of medical necessity. Rental of an electric or hydraulic lift may be considered during a repair of a client lift.

2.2.17.23 Overhead Lifts
The purchase of a free-standing overhead client lift (procedure code E0639) is a benefit in the home setting when services are provided by DME providers.

An overhead lift is anticipated to last a minimum of five years but may be replaced in less than five years with documentation of medical necessity.

Delivery and labor to assemble the overhead lift are not separately reimbursed.

2.2.17.23.1 Prior Authorization
Procedure code E0639 requires prior authorization.
A free-standing overhead lift may be considered for prior authorization when the client meets the criteria for a hydraulic or electric lift and additional documentation explains why a hydraulic or electric lift will not meet the client’s needs. Documentation that supports the medical necessity of the requested free-standing overhead lift must include all of the following:

- A written statement from a licensed physical therapist, licensed occupational therapist, or physician that clearly outlines the client’s medical need to be transferred with a free-standing overhead lift versus a hydraulic or electric lift.
- Diagrams of the home (rooms) indicating the location where the free-standing lift will be used. Diagrams must include dimensions of the rooms, including doorways, as well as the dimensions and placement of all furnishings and equipment (i.e., hospital bed, wheelchair, bedside commode, etc.) in the room.
- A list of all equipment the lift will interact with (i.e., wheelchair, hospital bed, or therapy equipment). Documentation should clearly indicate if the client owns other mobility aids, including any type of bath chair or bath lift, and explain why those pieces of equipment are not sufficient for mobility.

### 2.2.17.24 Fixed Lifts

The purchase of a fixed client lift (procedure code E0640) is a benefit in the home setting when the services are provided by home health medical supplier DME providers.

Home modifications that are necessary for the final set up and installation of a fixed lift are not benefits of Texas Medicaid. Suppliers must not submit claims for any structural changes or remodeling necessitated by the installation of a lift system.

**Note:** Home modifications are physical changes to the home to prepare the structure for the final set up and installation of the equipment.

A fixed lift is anticipated to last a minimum of five years but may be replaced in less than five years with documentation of medical necessity.

Delivery and labor to assemble and install the fixed lift are not separately reimbursed.

#### 2.2.17.24.1 Prior Authorization

Procedure code E0640 requires prior authorization.

A fixed lift may be considered for prior authorization when the client meets criteria for a hydraulic or electric lift and additional documentation explains why a hydraulic, electric, or free-standing overhead lift will not meet the client’s needs. Documentation that supports the medical necessity of the requested fixed lift must include all of the following:

- A signed and dated statement from the DME provider attesting that the home in which the lift will be installed meets the manufacturer’s requirements for installation, including documentation that the ceiling and wall structures of the residence are adequate to safely support the fixed lift.
- Documentation of whether the home is owned by the client, parent, guardian, or responsible party, such as a signed and dated document attesting to the ownership of the home. If the home is not owned by the client, parent, guardian, or responsible party, written consent of the installation from the home owner or property manager must be submitted.
- A written statement from a licensed physical therapist, licensed occupational therapist, or physician that clearly outlines the client’s medical need to be transferred with a fixed lift.
- Indication of what type of home the client lives in (i.e., traditional 1-story or 2-story home, mobile home, or apartment), including diagrams of the home where the fixed lift system will be installed. Diagrams must include dimensions of the rooms including doorways, as well as the dimensions and
placement of all furnishings and equipment (i.e. hospital bed, wheelchair, bedside commode, etc.) in the room. Diagrams must also include the proposed placement of all fixed components (i.e. railing or track) of the system.

- A list of all equipment the lift will interact with (i.e., wheelchair, hospital bed, or therapy equipment). Documentation should clearly indicate if the client owns other mobility aids, including any type of bath chair or bath lift, and explain why those pieces of equipment are not sufficient for mobility.
- Attestation from the DME provider stating that the provider has personnel who have been trained to install the lift system.
- Documentation confirming the home modifications necessary to allow for installation of the lift have been done or are scheduled to be completed prior to installation.

2.2.17.25 Portable Client Lifts for Outside Home Setting

Portable client lifts are benefits for clients who are 20 years of age and younger. Providers must bill procedure code E0635 with modifier TG for the purchase of the portable client lift. Procedure code E0635 is limited to once per lifetime, any provider.

2.2.17.25.1 Prior Authorization

Prior authorization is required and will be considered on a case-by-case basis for portable client electric lifts that fold-up for transport and are necessary for use outside the home setting.

The provider must submit a prior authorization request with the following documentation for consideration of medical necessity:

- An explanation of why a home-based portable lift will not meet the client’s needs
- A description of the circumstances, including duration of need
- The family member or caregivers who support the client with the use of the portable client lift when the client travels outside the home setting

2.2.17.26 Standers

A stander is a device used by a client with a neuromuscular condition who is unable to stand alone. Standers and standing programs can improve digestion, increase muscle strength, decrease contractures, increase bone density, and minimize decalcification (this list is not all inclusive).

2.2.17.26.1 Prior Authorization

Standers, including all accessories, require prior authorization. Standers and gait trainers will not be prior authorized for a client within one year of each other.

2.2.17.26.2 Documentation Requirements

Prior authorization may be considered for standers with the following documentation:

- Diagnoses relevant to the requested equipment, including functioning level and ambulatory status
- Anticipated benefits of the equipment
- Frequency and duration of the client’s standing program
- Anticipated length of time the client will require this equipment
- Client’s height, weight, and age
- Anticipated changes in the client’s needs, anticipated modifications, or accessory needs, as well as the growth potential of the stander
2.2.17.27  Gait Trainers

Gait trainers are devices with wheels that are used to train clients with ambulatory potential. They provide the same benefits as a stander, in addition to assisting with gait training.

2.2.17.27.1  Prior Authorization

Prior authorization for a gait trainer may be considered with the following documentation:

- Documentation of medical necessity
- An assessment of the accessibility of the client’s residence to ensure that a gait trainer is usable in the home (i.e., doors and halls are wide enough and have no obstructions)
- A physician familiar with the client documents that the client has ambulatory potential and will benefit from a gait training program
- The client meets the criteria for a stander

2.2.17.28  Accessories, Modifications, Adjustments and Repairs

Accessories, modifications, adjustments, and repairs are benefits of Texas Medicaid as outlined below.

- All major and minor modifications, adjustments, and repairs to standard mobility aid equipment within the first six months after delivery are considered part of the purchase price.
- All modifications and adjustments to a wheeled mobility system, as well as the associated services by the QRP for the seating assessment and fitting, within the first six months after delivery are considered part of the purchase price.

Mobility aids that have been purchased are anticipated to last a minimum of five years.

A major modification to a wheeled mobility system requires the completion of a new seating assessment by a qualified practitioner (physician, occupational therapist, or physical therapist), with the participation of a QRP.

Prior authorization for equipment replacement is considered within five years of equipment purchase when one of the following occurs:

- There has been a significant change in the client’s condition such that the current equipment no longer meets the client’s needs.
- The equipment is no longer functional and either cannot be repaired or it is not cost-effective to repair.

A wheeled mobility system that has been fitted and delivered to the client’s home by a QRP and then found to be inappropriate for the client’s condition will not be eligible for an upgrade, replacement, or major modification within the first six months following purchase unless there has been a significant change in the client’s condition. The significant change in the client’s condition must be documented by a physician familiar with the client.

2.2.17.28.1  Prior Authorization

Modifications

Modifications to custom equipment after the first six months from fitting and delivery may be considered for prior authorization if a change occurs in the client’s needs, capabilities or physical/mental capability, that cannot be anticipated.

Documentation supporting the medical necessity of the requested modification must include the following:

- Description of the change in the client’s condition that requires accommodation by different seating, drive controls, electronics, or other mobility base components.
• All projected changes in the client’s mobility needs.
• The date of purchase, the serial number of the current equipment, and the cost of purchasing new equipment versus modifying current equipment.

Major modifications to a wheeled mobility system also require that a new seating assessment be completed by a qualified practitioner (physician, physical therapist, or occupational therapist) and submitted with the prior authorization request. A request for authorization of the QRP’s participation in the seating assessment for the major modification must be included with the prior authorization request for the major modification.

Minor modifications to a wheeled mobility system do not require the completion of a new seating assessment.

Requests for equipment submitted as a minor modification or a repair to a wheeled mobility system must be submitted with modifier RB.

**Adjustments**

Adjustments within the first six months after delivery, including adjustments to a wheeled mobility system within the first six months after fitting and delivery by a QRP will not be prior authorized.

A seating or positioning component alteration that does not require replacement components to accommodate a change in the client’s size (height or weight) is considered an adjustment and not a major modification.

A maximum of one hour of labor for adjustments may be prior authorized as needed after the first six months from delivery.

Documentation must include the date of purchase, the serial number of the current equipment, and the reason for adjustments.

**Repairs**

Repairs to client-owned equipment may be considered for prior authorization as needed with documentation of medical necessity. Technician fees are considered part of the cost of the repair.

HHSC or its designee reserves the right to request additional documentation about the need for repairs when there is evidence of abuse or neglect to equipment by the client, client’s family, or caregiver. Requests for repairs when there is documented proof of abuse or neglect will not be authorized.

Requests for equipment submitted as a repair to a wheeled mobility system must be submitted with modifier RB.

Providers are responsible for maintaining documentation in the client’s medical record specifying the repairs and supporting medical necessity.

Documentation must include the date of purchase and serial number of the current equipment, the cause of the damage or need for repairs, the steps the client or caregiver will take to prevent further damage if repairs are due to an accident, and when requested, the cost of purchasing new equipment as opposed to repairing current equipment.

**2.2.17.29 Replacement**

Replacement of equipment is also considered when loss or irreparable damage has occurred. The following must be submitted with the prior authorization request:

• A copy of the police or fire report, when appropriate.
• A statement about the measures to be taken in order to prevent reoccurrence.
• Replacement equipment for clients who are birth through 20 years of age and do not meet the criteria in this handbook may be considered for prior authorization through CCP.
### 2.2.17.30 Procedure Codes and Limitations for Mobility Aids

For clients who are 20 years of age and younger, the limitations listed below may not apply with evidence of medical necessity. Procedures include, but are not limited to, the following:

<table>
<thead>
<tr>
<th>Procedure Code</th>
<th>Maximum Limit</th>
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<tbody>
<tr>
<td><strong>Canes</strong></td>
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<tr>
<td>E0100</td>
<td>1 per 5 years</td>
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<tr>
<td>E0105</td>
<td>1 per 5 years</td>
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<td><strong>Crutches</strong></td>
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<td><strong>Gait Trainers</strong></td>
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<tr>
<td>E0705</td>
<td>1 per 5 years</td>
</tr>
<tr>
<td></td>
<td>Lifts</td>
</tr>
</tbody>
</table>
The following mobility aids are not a benefit of Home Health Services:

- Items including but not limited to tire pumps, a color for a wheelchair, gloves, back packs, and flags are not considered medically necessary
- Mobile standers, power standing system on a wheeled mobility device
- Vehicle lifts and modifications
- Permanent ramps, vehicle ramps, and home modifications
- Stairwell lifts of any type
- Elevators or platform lifts of any type
- Chairs with incorporated seat lifts
- An attendant control, for safety, all power chairs are to include a stop switch
- Powered mobility device for use only outside the home

**Note:** For clients who are 21 years of age or older, requests for mobility aids that do not meet the criteria through Title XIX Home Health Services may be considered under the Texas Medicaid Home Health—Durable Medical Equipment (DME) Exceptional Circumstances process.

Texas Medicaid does not reimburse separately for associated DME charges, including battery disposal fees or state taxes. Reimbursement for associated charges is included in the reimbursement for the specific piece of equipment. White canes for the blind are considered self help adaptive aids and are not a benefit of Home Health Services.

### 2.2.18 Nutritional (Enteral) Products, Supplies, and Equipment

Medical nutritional products including enteral formulas and food thickener, may be approved for clients who have specialized nutritional requirements.

Medical nutritional products must be prescribed by a physician and be medically necessary.

Enteral nutritional products are those food products that are included in an enteral treatment protocol. They serve as a therapeutic agent for health maintenance and are required to treat an identified medical condition. Nutritional products, supplies, and equipment may be a benefit when provided in the home under Home Health Services.
For clients who are 20 years of age and younger and do not meet criteria through Home Health Services, products, supplies, and equipment can be considered through the Comprehensive Care Program (CCP).

**Note:** For clients who are 21 years of age or older, requests for nutritional (enteral) products, supplies and equipment that do not meet the criteria through Title XIX Home Health Services may be considered under the Texas Medicaid Home Health—Durable Medical Equipment (DME) Exceptional Circumstances process.

### 2.2.18.1 Enteral Nutritional Products, Feeding Pumps, and Feeding Supplies

Enteral nutritional products and related feeding supplies and equipment are a benefit through Home Health Services for clients who are 21 years of age and older and require tube feeding as their primary source of nutrition. The enteral product, supply, or equipment must be part of the medical POC outlined and maintained by the treating physician.

Enteral nutritional products may be reimbursed with the following procedure codes:

<table>
<thead>
<tr>
<th>Procedure Codes</th>
<th></th>
</tr>
</thead>
<tbody>
<tr>
<td>B4100</td>
<td>B4103</td>
</tr>
<tr>
<td>B4104</td>
<td>B4149</td>
</tr>
<tr>
<td>B4150</td>
<td>B4152</td>
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<tr>
<td>B4153</td>
<td>B4154</td>
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<td>B4155</td>
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<td>B4158</td>
<td>B4159</td>
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<tr>
<td>B4160</td>
<td>B4161</td>
</tr>
<tr>
<td>B4162</td>
<td></td>
</tr>
</tbody>
</table>

Enteral formulas consisting of semi-synthetic intact protein or protein isolates (procedure codes B4150 and B4152) are appropriate for the majority of clients requiring enteral nutrition.

Special enteral formulas or additives (procedure code B4104) may be considered for prior authorization with supporting documentation submitted by the client’s physician indicating the client’s medical needs for these special enteral formulas.

Special enteral formula may be reimbursed with the following procedure codes:

<table>
<thead>
<tr>
<th>Procedure Codes</th>
<th></th>
</tr>
</thead>
<tbody>
<tr>
<td>B4149</td>
<td>B4153</td>
</tr>
<tr>
<td>B4154</td>
<td>B4155</td>
</tr>
<tr>
<td>B4157</td>
<td>B4161</td>
</tr>
<tr>
<td>B4162</td>
<td></td>
</tr>
</tbody>
</table>

Pediatric nutritional products (procedure codes B4103, B4158, B4159, B4160, B4161, and B4162) are restricted to clients who are 20 years of age and younger.

Food thickener may be considered for clients with a swallowing disorder.

Enteral nutritional supplies and equipment may be reimbursed with the following procedure codes and limitations:

<table>
<thead>
<tr>
<th>Procedure Codes</th>
<th>Limitations</th>
</tr>
</thead>
<tbody>
<tr>
<td>A4322</td>
<td>30 per month</td>
</tr>
<tr>
<td>A5200</td>
<td>2 per month</td>
</tr>
<tr>
<td>B4034</td>
<td>Up to 31 per month</td>
</tr>
<tr>
<td>B4035</td>
<td>Up to 31 per month</td>
</tr>
<tr>
<td>B4036</td>
<td>Up to 31 per month</td>
</tr>
<tr>
<td>B4081</td>
<td>As needed</td>
</tr>
<tr>
<td>B4082</td>
<td>As needed</td>
</tr>
<tr>
<td>B4083</td>
<td>As needed</td>
</tr>
<tr>
<td>B4087</td>
<td>2 per rolling year</td>
</tr>
<tr>
<td>B4088</td>
<td>2 per rolling year</td>
</tr>
</tbody>
</table>

*Appropriate limitations for miscellaneous procedure code T1999 are determined on a case-by-case basis through prior authorization.*
Enteral nutritional supplies and equipment (Procedure code B9998) may be reimbursed as needed with the following modifiers and limitations:

**Procedure Codes** | **Limitations**
---|---
B4105 | Up to 62 per month
B9002 | 1 purchase every 5 years; 1 rental per month
T1999* | As needed*  
If procedure code T1999 is used for a needleless syringe, the allowed amount is 8 per month.

*Appropriate limitations for miscellaneous procedure code T1999 are determined on a case-by-case basis through prior authorization.

Enteral nutritional supplies and equipment (Procedure code B9998) may be reimbursed as needed with the following modifiers and limitations:

**Procedure Codes** | **Descriptions** | **Limitations**
---|---|---
B9998 with modifier U1 | Disposable G-tube adapter set | 4 per month
B9998 with modifier U2 | Nonobturated gastrostomy or jejunostomy tube with insertion supplies and extensions | 2 per rolling year
B9998 with modifier U3 | Low profile enteral extension set | 4 per month
B9998 with modifier U4 | Standard gastrostomy tube | 2 per rolling year
B9998 with modifier U5 | Standard enteral extension set | 4 per month

* Appropriate limitations for miscellaneous procedure codes B9998 are determined on a case-by-case basis through prior authorization.

### 2.2.18.2 Prior Authorization Requirements

Prior authorization is required for most enteral products, supplies, and equipment provided through Home Health Services. Requests are reviewed for medically necessary amounts based on caloric needs as indicated by the client’s physician.

**2.2.18.2.1 Clients who are 21 years of age and older**

Enteral nutrition and related supplies and equipment may be considered for prior authorization for clients who are 21 years of age and older when all or part of the client’s nutritional intake is received through a feeding tube, and the enteral formula is:

- The client’s sole source of nutrition
- The client’s primary source of nutrition
  - An enteral tube feeding is considered the primary source of nutrition when it comprises more than 70 percent of the caloric intake needed to maintain the client’s weight.
  - The percent of calories provided by an enteral formula may be calculated by dividing the client’s daily calories supplied by the enteral formula by the daily caloric intake ordered by the physician to maintain the client’s weight. The result is multiplied by 100 to determine the percentage of calories provided by the enteral formula.

Related supplies and equipment may be considered for prior authorization when criteria for nutritional products are met, and medical necessity is included for each item requested.

Renewal of the prior authorization will be considered based on medical necessity.

Prior authorization may be given for up to 6 months. Prior authorization may be recertified with documentation supporting ongoing medical necessity for the nutritional products requested.
2.2.18.2.2 Clients who are 20 years of age and younger

Prior authorization for nutritional products is not required for a client who is 20 years of age and younger and who meets at least one of the following criteria:

- Client receives all or part of their nutritional intake through a tube.
- Client has a metabolic disorder that has been documented with one of the following diagnosis codes:

<table>
<thead>
<tr>
<th>Diagnosis Codes</th>
</tr>
</thead>
<tbody>
<tr>
<td>C880 C965 C966 D472 D800 D801 D802 D803</td>
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<tr>
<td>D804 D805 D806 D807 D808 D809 D810 D811</td>
</tr>
<tr>
<td>D812 D814 D816 D817 D8189 D819 D820 D821</td>
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<tr>
<td>D822 D823 D824 D828 D829 D830 D831 D832</td>
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<td>D838 D839 D840 D841 D8481 D84821 D84822 D8489</td>
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<td>D849 D890 D891 D893 D8982 D8989 D899 E201</td>
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<td>E670 E671 E672 E673 E678 E68 E700 E701</td>
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<td>E710 E71110 E71111 E71118 E7112 E71120 E71121 E71128 E7119</td>
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<td>Z931</td>
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</table>
Prior authorization is required for nutritional products that are provided through CCP to clients who do not meet the criteria above.

A completed CCP Prior Authorization Request Form that prescribes the nutritional product and/or related services must be signed and dated by a prescribing physician who was familiar with the client before requesting prior authorization. The completed CCP Prior Authorization Request Form must include the procedure codes and numerical quantities for the services requested. A copy of the completed, signed, and dated CCP Prior Authorization Form must be maintained by the provider in the client’s medical record. The completed CCP Prior Authorization Request Form with the original dated signature must be maintained by the prescribing physician in the client’s medical record.

Requests for prior authorization must include the following documentation:

- Accurate diagnostic information pertaining to the underlying diagnosis or condition that resulted in the requirement for a nutritional product, as well as any other medical diagnoses or conditions, including:
  - The client’s overall health status.
  - Height and weight.
  - Growth history and growth charts.
  - Why the client cannot be maintained on an age-appropriate diet.
  - Other formulas tried and why they did not meet the client’s needs.
- Diagnosis or condition.
- The goals and timelines on the medical plan of care.
- Total caloric intake prescribed by the physician.
- Acknowledgement that the client has a feeding tube in place.

Related supplies and equipment for clients who require nutritional products may be considered for prior authorization when the criteria for nutritional products are met and medical necessity is included for each item requested.

Prior authorization may be given for up to 12 months. Prior authorization may be recertified with documentation that supports the ongoing medical necessity of the requested nutritional products.

A retrospective review may be performed to ensure that the documentation included in the client’s medical record supports the medical necessity of the requested service.

Requests for prior authorization, when required, must include the necessary product information.

Prior authorization of nutritional pudding products may be considered for children who have a documented oropharyngeal motor dysfunction and receive greater than 50 percent of their daily caloric intake from a nutritional pudding product.

Requests for electrolyte replacement products, such as Pedialyte or Oralyte, require documentation of medical necessity, including:

- The underlying acute or chronic medical diagnoses or conditions that indicate the need to replace fluid and electrolyte losses.
- The presence of mild to moderate dehydration due to the persistent mild to moderate diarrhea or vomiting.

Electrolyte replacement products are not indicated for clients with:

- Intractable vomiting
- Adynamic ileus
• Intestinal obstruction or perforated bowel
• Anuria, oliguria, or impaired homeostatic mechanism
• Severe, continuing diarrhea, when intended for use as the sole therapy

2.2.18.2.3 Enteral Formulas
Enteral formulas require prior authorization. Requests for prior authorization must include the necessary product information.

2.2.18.2.4 Nasogastric, Gastrostomy, or Jejunostomy Feeding Tubes
Feeding tubes require prior authorization.

Additional feeding tubes may be prior authorized if documentation submitted supports medical necessity, such as infection at gastrostomy site, leakage, or occlusion.

2.2.18.2.5 Enteral Feeding Pumps
Enteral feeding pumps, with and without alarms, require prior authorization.

Enteral feeding pumps may be considered for prior authorization for lease or purchase with documentation of medical necessity indicating that the client meets the following criteria:
• Gravity or syringe feedings are not medically indicated
• The client requires an administration rate of less than 100 ml/hr
• The client requires night-time feedings
• The client has one of the following medical conditions (this list is not all-inclusive):
  • Reflux or aspiration
  • Severe diarrhea
  • Dumping syndrome
  • Blood glucose fluctuations
  • Circulatory overload

2.2.18.2.6 Enteral Supplies
Enteral supplies require prior authorization, with the exception of irrigation syringes (procedure code A4322) and percutaneous catheter or tube anchoring devices (procedure code A5200) within the allowable limits.

Procedure code B4034 will not be prior authorized for use in place of procedure code A4322 for irrigation syringes when they are not part of a bolus administration kit.

Gravity bags and pump nutritional containers are included in the feeding supply kits and will not be prior authorized separately.

Procedure code B4105 will require prior authorization, and may be considered with documentation of medical necessity indicating that the client meets all the following criteria:
• The client has exocrine pancreatic insufficiency.
• The client utilizes an enteral feeding pump.
• The client utilizes a compatible formula and the amount of formula (mL) the client is receiving daily is documented.

Note: One cartridge can be used with up to 500mL of formula, with a maximum of two cartridges used per day. Procedure code B4105 will be limited to 62 per month.
For clients who are 5 through 20 years of age, procedure code B4105 will be considered through the Comprehensive Care Program (CCP).

Specific items may be considered for prior authorization using miscellaneous procedure code B9998 and modifiers U1, U2, U3, or U5.

Requests for a backpack or carrying case for a portable enteral feeding pump may be considered for prior authorization for purchase only, under miscellaneous code B9998, for clients who meet all of the following medical necessity criteria:

- The client requires enteral feedings lasting greater than eight hours continuously, or feeding intervals exceed the time that the client must be away from home to:
  - Attend school or work.
  - Participate in extensive, physician-ordered outpatient therapies.
  - Attend frequent, multiple medical appointments.
  - The client is ambulatory, or uses a wheelchair which will not support the use of a portable pump by other means, such as an IV pole.
  - The portable enteral feeding pump is client owned.

### 2.2.18.3 Documentation Requirements

To request prior authorization for nutritional formula, supplies, or equipment, the following documentation must be provided:

- Accurate diagnostic information pertaining to the underlying diagnosis or condition as well as any other medical diagnoses or conditions, to include the client’s overall health status
- Diagnosis or condition (including the appropriate International Classification of Diseases, Tenth Revision, Clinical Modification [ICD-10-CM] code)
- A statement from the ordering physician noting that enteral nutritional products for tube feedings are the client’s sole or primary source of nutrition
- The goals and timelines on the medical POC
- Total caloric intake prescribed by the physician
- Acknowledgement that the client has a feeding tube in place

### 2.2.18.4 Nutritional Counseling

Clients for whom nutritional products are being requested may benefit from nutritional counseling. Nutritional counseling is a benefit of CCP if it is provided to treat, prevent, or minimize the effects of illness, injury, or other impairment.

**Refer to:** Subsection 2.10, “Medical Nutrition Counseling Services (CCP)” in the *Children’s Services Handbook (Vol. 2, Provider Handbooks)* for information about nutritional counseling.

### 2.2.18.5 Women, Infants, and Children Program (WIC)

Generic nutritional products that have been approved by the United States Department of Agriculture (USDA) for use in the Women, Infants, and Children Program (WIC) may be approved for use by CCP clients.

While CCP does not require that a client access WIC, it is only recommended as another source of services for clients who are 4 years of age and younger, or clients who are pregnant or breast feeding. Nutritional products are not provided to infants who are 11 months of age and younger unless medical necessity is documented.
2.2.18.6 Managed Care Clients

Nutritional products that are provided to WIC clients are carved-out of the Medicaid Managed Care Program and must be billed to TMHP for payment consideration. Carved-out services are those that are rendered to Medicaid Managed Care clients but are administered by TMHP and not the client’s managed care organization (MCO).

Nutrition products that are provided to other Medicaid Managed Care Program clients (other than WIC clients) are not carved out and must be submitted to the managed care organization that administers the client’s Medicaid managed care benefits.

2.2.18.7 Noncovered Services

CCP will not cover the following:

- Nutritional products that are traditionally used for infant feeding.
- Nutritional products for the primary diagnosis of failure to thrive, failure to gain weight, or lack of growth. The underlying cause of failure to thrive, gain weight, and lack of growth is required.
- Nutritional bars.
- Nutritional products for clients who could be sustained on an age-appropriate diet.

2.2.19 Orthotic Services (CCP)

Orthoses, including orthopedic shoes, wedges, and lifts, are a benefit of Texas Medicaid when provided by a licensed orthotist or a licensed prosthetist/orthotist through CCP for clients who are birth through 20 years of age.

The following orthoses and related services may be reimbursed when medical necessity criteria are met:

- Spinal orthoses and additions to spinal orthoses, including those for scoliosis
- Lower-limb orthoses and additions to lower-limb orthoses, including fracture orthoses
- Foot orthoses, including inserts, orthopedic shoes, surgical boots, heel lifts, and wedges
- Upper-limb orthoses and additions to upper-limb orthoses, including fracture orthoses
- Other orthopedic devices, including protective helmets and dynamic splints
- Repairs, replacements, and modifications
- Orthotic device training

*Note:* Training in the use of an orthotic device for a client who has not worn one previously, has not worn one for a prolonged period, or is receiving a different type is a benefit when the training is provided by a physical or occupational therapist.

*Refer to:* The Physical Therapy, Occupational Therapy, and Speech Therapy Services Handbook (Vol. 2, Provider Handbooks) for additional information about physical and occupational therapy services.

The following definitions are used by Texas Medicaid:

- An orthosis is defined as: A custom-fabricated or custom-fitted medical device designed to provide for the support, alignment, prevention or correction of neuromuscular or musculoskeletal disease, injury, or deformity. The term does not include a fabric or elastic support, corset, arch support, low temperature plastic splint, a truss, elastic hose, cane, crutch, soft cervical collar, orthosis for diagnostic or evaluation purposes, dental appliance, or other similar device carried in stock and sold by a drugstore, department store or corset shop.
• A brace is defined as: An orthosis or orthopedic appliance that supports or holds in correct position any movable part of the body, and that allows for motion of that part. It must be a rigid or semirigid device used for the purpose of supporting a weak or deformed body part or restricting or eliminating motion in a diseased or injured body part.

To be considered for reimbursement, orthoses must be dispensed, fabricated, or modified by a licensed orthotist or licensed prosthetist/orthotist enrolled with Medicare and CCP. The following applies:

• Upper extremity customized splints made with low-temperature materials and inhibitive casting may be provided by occupational or physical therapists.

• Other orthopedic devices addressed in the orthotic section may be provided by a Medicaid-enrolled DME vendor.

• Orthopedic shoes must be provided by a shoe vendor enrolled as a DME provider.

The date of service for a custom-made or custom-fitted orthosis is the date the supplier places an order for the equipment and incurs liability for the equipment. The custom-made or custom-fitted orthosis will be eligible for reimbursement as long as the service is provided during a month the client is eligible for Medicaid.

The following items and services are included in the reimbursement for an orthotic device and not reimbursed separately:

• Client evaluation, measurement, casting, or fitting of the orthosis.

• Repairs due to normal wear and tear during the 90 days following delivery.

• Adjustments or modifications of the orthotic device made when fitting the orthosis and for 90 days from the date of delivery (adjustments and modifications during the first 90 days are considered part of the purchase of the initial device).

Orthopedic shoes that are attached to a brace must be billed by the vendor that bills for the brace.

Reimbursement for lifts and wedges may include the cost of the prescription shoe.

2.2.19.1 Noncovered Orthotic Services

The following circumstances are not a benefit of Texas Medicaid:

• Orthoses whose sole purpose is for restraint

• Orthoses provided solely for use during sports-related activities in the absence of an acute injury or other indicated medical condition

• Orthotic devices prescribed by a chiropractor

Diagnoses that are not considered medically necessary include, but are not limited to, the following:

• Tired feet

• Fatigued feet

• Nonsevere bow legs

• Valgus deformity of the foot, except as outlined in the orthotic section

• Pes planus (flat feet), except when there is a coexisting medical condition as outlined in the orthotic section

Orthopedic shoes with deluxe features, such as special colors, special leathers, and special styles, are not considered medically necessary, and the features do not contribute to the accommodative or therapeutic function of the shoe.
A foot-drop splint and recumbent positioning device and replacement interface are not considered medically necessary in a client with foot drop who is nonambulatory, because there are other more appropriate treatment modalities.

A static ankle-foot orthosis (AFO) or AFO component is not medically necessary if:

- The contracture is fixed.
- The client has foot drop without an ankle flexion contracture.
- The component is used to address knee or hip positioning, because the effectiveness of this type of component is not established.

A pneumatic thoracic-lumbar-sacral orthosis is considered experimental and investigational and is not a benefit of Texas Medicaid.

2.2.19.2 Prior Authorization and Documentation Requirements

Prior authorization is required for all orthoses and related services.

Before submitting a request for prior authorization for orthosis, the orthosis provider must have a completed CCP Prior Authorization Form requesting the orthosis or related services that has been signed and dated by a physician who is familiar with the client. The completed CCP Prior Authorization Form must include the procedure codes and quantities for requested services. A copy of the completed, signed, and dated form must be maintained by the orthosis provider in the client’s medical record. The completed CCP Prior Authorization Form with the original dated signature must be maintained by the prescribing physician in the client’s medical record.

- To complete the prior authorization process electronically, the orthosis provider must complete the prior authorization requirements through any approved electronic method and retain a copy of the signed and dated CCP Prior Authorization Request form in the client’s medical record at the provider’s place of business.
- To complete the prior authorization process by paper, the orthosis provider must fax or mail the completed CCP Prior Authorization Request Form to the CCP prior authorization unit and retain a copy of the signed and dated CCP form in the client’s medical record at the provider’s place of business.

To facilitate determination of medical necessity and avoid unnecessary denials, the physician must provide correct and complete information, including documentation for medical necessity of the equipment and supplies requested. The physician must maintain documentation of medical necessity in the client’s medical record. The provider may be asked for additional information to clarify or complete a request for the service or device.

All requests for prior authorization must include documentation of medical necessity including, but not limited to, documentation that the device is needed for one of the following general indications:

- To reduce pain by restricting mobility of the affected body part.
- To facilitate healing following an injury to the affected body part or related soft tissue.
- To facilitate healing following a surgical procedure on the affected body part or related soft tissue.
- To support weak muscles or a deformity of the affected body part.

Prior authorization requests for some types of orthosis require additional documentation. See the appropriate sections for additional documentation needed for each service.

The provider must keep the following written documentation in the client’s medical record:

- The prescription for the device.
• Orthotic devices must be prescribed by a physician (M.D. or D.O.) or a podiatrist. A podiatrist prescription is valid for conditions of the ankle and foot.

• The prescription must be dated on or before the initial date of the requested dates of service, which can be no longer than 90 days from the signature date on the prescription.

• Accurate diagnostic information that supports the medical necessity for the requested device. A retrospective review may be performed to ensure that the documentation included in the client’s medical record supports the medical necessity of the requested service or device.

A prior authorization is valid for a maximum period of six months from the prescription signature date. At the end of the six-month authorization period, a new prescription is required for prior authorization of additional services.

The actual date of service is the date the supplier has placed an order for the equipment and has incurred liability for the equipment.

2.2.19.2.1 Spinal Orthoses

Spinal orthoses include, but are not limited to, cervical orthoses, thoracic rib belts, thoracic-lumbar-sacral orthoses (TLSO), sacroiliac orthoses, lumbar orthoses, lumbar-sacral orthoses (LSO), cervicalthoracic-lumbar-sacral orthoses (CTLSO), halo procedures, spinal corset orthoses, and spinal orthoses for scoliosis.

Spinal orthoses will be considered for prior authorization with documentation of one of the general indications.

2.2.19.2.2 Lower-Limb Orthoses

Lower-limb orthoses include, but are not limited to, hip orthoses (HO), Legg Perthes orthoses, knee orthoses (KO), ankle-foot orthoses (AFO), knee-ankle-foot orthoses (KAFO), hip-knee-ankle-foot orthoses (HKAFO), fracture orthoses, and reciprocating gait orthoses (RGO).

In addition to the general indication requirements, lower-limb orthoses will be considered for prior authorization with documentation of the following criteria for specific orthotic devices:

Ankle-Foot Orthoses

AFOs used during ambulation will be considered for prior authorization for clients with documentation of all of the following:

• Weakness or deformity of the foot and ankle.
• A need for stabilization for medical reasons.
• Anticipated improvement in functioning during activities of daily living (ADLs) with use of the device.

AFOs not used during ambulation (static AFO) will be considered for prior authorization for clients with documentation of one of the following conditions:

• Plantar fasciitis.

• Plantar flexion contracture of the ankle, with additional documentation that includes all of the following:
• Dorsiflexion on pretreatment passive range of motion testing is at least ten degrees.
• The contracture is interfering or is expected to interfere significantly with the client’s functioning during ADLs.
• The AFO will be used as a component of a physician-prescribed therapy plan care, which includes active stretching of the involved muscles or tendons.
• There is reasonable expectation that the AFO will correct the contracture.

Knee-Ankle-Foot Orthoses
KAFOs used during ambulation will be considered for prior authorization for clients with documentation that supports medical necessity for additional knee stabilization.

KAFOs that are custom-fabricated (molded-to-patient model) for ambulation will be considered for prior authorization when at least one of the following criteria is met:

• The client cannot be fit with a prefabricated AFO/KAFO.
• The condition that necessitates the orthosis is expected to be permanent or of long-standing duration (more than six months).
• There is a need to control the knee, ankle, or foot in more than one plane.
• The client has a documented neurological, circulatory, or orthopedic status that requires custom fabrication to prevent tissue injury.
• The client has a healing fracture that lacks normal anatomical integrity or anthropometric proportions.

Reciprocating Gait Orthoses
Reciprocating gait orthoses will be considered for prior authorization for clients with spina bifida or similar functional disabilities.

The prior authorization request must include a statement from the prescribing physician that indicates medical necessity for the RGO, the PT treatment plan, and documentation that the client and family are willing to comply with the treatment plan.

2.2.19.2.3 Foot Orthoses
Foot orthoses include, but are not limited to, foot inserts, orthopedic shoes, wedges, and lifts.

Foot orthoses will be considered for prior authorization for clients with documentation of all of the following:

• The client has symptoms associated with the particular foot condition.
• The client has failed to respond to a course of appropriate, conservative treatment, including PT, injections, strapping, or anti-inflammatory medications.
• The client has at least one of the following:
  • Torsional conditions, such as metatarsus adductus, tibial torsion, or femoral torsion
  • Structural deformities
  • Hallux valgus deformities
  • In-toe or out-toe gait
  • Musculoskeletal weakness

In addition to the general indication requirements, foot orthoses will be considered for prior authorization with documentation of the following criteria for specific orthotic devices:

Foot Inserts
Removable foot inserts will be considered for prior authorization for clients with documentation of at least one of the following medical conditions:

• Diabetes mellitus.
• History of amputation of the opposite foot or part of either foot.
• History of foot ulceration or pre-ulcerative calluses of either foot.
• Peripheral neuropathy with evidence of callus formation of either foot.
• Deformity of either foot.
• Poor circulation of either foot.

Removable foot inserts may be covered independently of orthopedic shoes with documentation that the client has appropriate footwear into which the insert can be placed.

A University of California at Berkeley (UCB) removable foot insert will be considered for prior authorization with documentation that the device is required to correct or treat at least one of the following conditions:
• A valgus deformity and significant congenital pes planus with pain.
• A structural problem that results in significant pes planus, such as Down syndrome.
• Acute plantar fasciitis.

Orthopedic Shoes

Orthopedic shoes must be prescribed by a licensed physician (M.D. or D.O.) or a podiatrist. An orthopedic shoe is used by clients whose feet, although impaired, are essentially intact. An orthopedic shoe differs from a prosthetic shoe, which is used by clients who are missing all or most of the forefoot.

Orthopedic shoes will be considered for prior authorization when at least one of the following criteria is met:
• The shoe is permanently attached to a brace.
• The shoe is necessary to hold a surgical correction, postoperative casting, or serial or clubfoot casting.

An orthopedic shoe may be prior authorized up to one year from the date of the surgical procedure.

Only one pair of orthopedic shoes will be prior authorized every three months. Two pairs of shoes may be purchased at the same time; in such situations, however, additional requests for shoes will not be considered for another six months.

Requests for orthopedic shoes that do not meet the criteria listed above may be considered for prior authorization with documentation of medical necessity.

Wedges and Lifts

Wedges and lifts must be prescribed by a licensed physician (M.D. or D.O.) or a podiatrist and must be for treatment of unequal leg length greater than one-half inch.

2.2.19.2.4 Upper-Limb Orthoses

Upper-limb orthoses include, but are not limited to, shoulder orthoses (SO), elbow orthoses (EO), elbow-wrist-hand orthoses (EWHO), elbow-wrist-hand-finger orthoses (EWHFO), wrist-hand-finger orthoses (WHFO), wrist-hand orthoses (WHO), hand-finger orthoses (HFO), finger orthoses (FO), shoulder-elbow-wrist-hand orthoses (SEWHO), shoulder-elbow orthoses (SEO), and fracture orthoses.

In addition to the general indication requirements, upper-limb orthoses will be considered for prior authorization with documentation of the following criteria for specific orthotic devices.
2.2.19.2.5 Other Orthopedic Devices

Protective Helmets
Protective helmets will be considered for prior authorization for clients with a documented medical condition that makes the client susceptible to injury during ADLs. Covered medical conditions include the following:

- Neoplasm of the brain
- Subarachnoid hemorrhage
- Epilepsy
- Cerebral palsy

Requests for all conditions other than those listed above require submission of additional documentation that supports the medical necessity of the requested device.

Dynamic Splints
Static and dynamic mechanical stretching devices will be considered for prior authorization for a four month rental period when the request is submitted with the following documentation:

- Client’s condition
- Client’s current course of therapy
- Rationale for the use of the static or dynamic mechanical stretching device
- Agreement by the client or family that the client will comply with the prescribed use of the static or dynamic mechanical stretching device

After completion of the four-month rental period, the provider may submit a request for purchase of the static or dynamic mechanical stretching device. Requests for purchase of the static or dynamic mechanical stretching device must include documentation that the four-month rental period was successful and showed improvement in the client’s condition as measured by the following:

- Demonstrated increase in range of motion
- Demonstrated improvement in the ability to complete ADLs or perform activities outside the home

2.2.19.2.6 Related Services

Repairs, Replacements, and Modifications to Orthoses
Within the guarantee of the manufacturer, providers are responsible, without charge to the client or to Texas Medicaid, for replacement or repair of equipment or any part thereof that is found to be nonfunctional because of faulty material or workmanship.

Service and repairs must be handled under any warranty coverage an item may have. If there is no warranty, providers may request prior authorization for the necessary service and repairs.

A repair because of normal wear or a modification because of growth or change in medical status will be considered for prior authorization if it proves to be more cost effective than replacing the device.

The request for repairs must include a breakdown of charges for parts and the number of hours of labor required to complete the repairs. No charge is allowed for pickup or delivery of the item or for the assembly of Medicaid-reimbursed parts. The following information must be submitted with the request:

- The description and procedure code of the item being serviced or repaired.
- The age of the item.
- The number of times the item has been previously repaired.
• The replacement cost for the item.

The anticipated life expectancy of an orthotic device is six months. Requests for prior authorization for the replacement of a device before its usual life expectancy has ended must include documentation that explains the need for the replacement.

Replacement of orthotic equipment will be considered when the item is out of warranty and repairing the item is no longer cost-effective or when loss or irreparable damage has occurred. A copy of the police or fire report, when appropriate, and the measures to be taken to prevent reoccurrence must be submitted with the prior authorization request.

2.2.19.3 Cranial Molding Orthosis

2.2.19.3.1 Services, Benefits, and Limitations

Cranial molding orthosis (procedure code S1040) may be a benefit when all of the following criteria are met:

• The client is CCP eligible.
• The client is 3 through 18 months of age.
• The client requires a cranial molding orthosis as part of the treatment plan for a documented diagnosis of synostotic plagiocephaly.

The limitation for procedure code S1040 is one device per lifetime.

The definition for cosmetic, as it applies to cranial molding orthosis, includes surgery or other services used primarily to improve appearance and not to restore or correct significant deformity resulting from disease, trauma, congenital or developmental anomalies, or previous therapeutic process.

2.2.19.3.2 Noncovered Services

A cranial molding orthosis that is used for the treatment of positional plagiocephaly is considered cosmetic, and therefore is not a benefit of Texas Medicaid.

The effective use of a cranial molding orthosis for the treatment of brachycephaly, or a high cephalic index without cranial asymmetry has not been clearly documented, is not medically necessary, and therefore is not a benefit of Texas Medicaid.

2.2.19.3.3 Prior Authorization and Documentation Requirements

Cranial molding orthoses do not require prior authorization for clients with a diagnosis of synostotic plagiocephaly. Documentation of medical necessity must be maintained in the client’s medical record.

Prior authorization requests for a cranial molding orthosis for congenital conditions that are not outlined in this section may be considered by the Medical Director on a case-by-case basis with documentation of medical necessity. Additional devices beyond the once-per-lifetime benefit may be considered for prior authorization with documentation of all of the following:

• The initial device was obtained to treat synostotic plagiocephaly.
• Treatment with the device has been effective.
• The new device is needed due to growth.

To facilitate determination of medical necessity and avoid unnecessary denials, the physician must provide correct and complete information, including documentation of medical necessity for the equipment requested. The physician must maintain documentation of medical necessity in the client’s medical record. The requesting provider may be asked for additional information to clarify or complete a request for an additional cranial molding orthosis.
The completed CCP Prior Authorization Form, which includes the DME must be signed and dated by the prescribing physician familiar with the client’s condition. The completed CCP Prior Authorization Form must be maintained by the requesting provider and the prescribing physician. The original signature copy must be kept by the physician in the client’s medical record.

2.2.19.4 Thoracic-Hip-Knee-Ankle Orthoses (THKAO) (Vertical or Dynamic Standers, Standing Frames, Braces, and Parapodiums)

2.2.19.4.1 Services, Benefits, and Limitations

THKAO (vertical or dynamic standers, standing frames or braces, and parapodiums), including all accessories, require prior authorization. A THKAO may be considered if the client requires assistance to stand and remain standing.

Parapodium

A parapodium is used to help clients with neuromuscular diseases or conditions resulting in a lack of sufficient muscle power in the trunk and lower extremities to stand with their hands free. It helps develop a sense of balance and aids in learning functional movements such as standing with the hands free. A parapodium acts as an exoskeleton, providing side struts and chest, hip, knee, and foot bracing.

A parapodium may be considered for reimbursement for one of the following levels:

- **Level One:** Small Parapodium—The client has a maximum axillary height of 35 inches and a maximum weight of 55 pounds (normal age range is 1 through 10 years of age).
- **Level Two:** Medium parapodium—The client has a maximum axillary height of 41 inches and a maximum weight of 77 pounds (normal age range is 5 through 12 years of age).
- **Level Three:** Large parapodium—The client has a maximum axillary height of 45 inches and a maximum weight of 115 pounds (normal age range is 10 through 16 years of age).

Labor for parapodium assembly may be prior authorized.

Procedure code E0638 must be submitted with one of the following modifiers:

- **UA-Standing frame/table system, one position (e.g., upright, supine, or prone stander), any size, including pediatric, with or without wheels. Medicaid level of care 10, as defined by each state**
  
  **Note:** Use modifier UA to identify an upright or prone system stander.

- **UB-Standing frame/table system, one position (e.g., upright, supine, or prone stander), any size, including pediatric, with or without wheels. Medicaid level of care 11, as defined by each state**
  
  **Note:** Use modifier UB to identify a supine stander.

**Standing Frame or Brace**

A standing frame or brace is used to help very young clients, who are 12 months of age and older, who have good head control in the upright position and who have a neuromuscular disease or condition resulting in a lack of sufficient muscle power in the trunk and lower extremities to stand with their hands free.

Providers must use procedure code E0638 for a standing frame or brace.

**Vertical or Dynamic Stander**

A vertical stander or dynamic stander is used to initiate standing for clients who cannot maintain a good standing posture or may never be able to stand independently. A vertical stander is used to develop weight bearing through the legs in order to decrease demineralization and to promote better body awareness. Documentation for these standers must address medical necessity for the standers to be mobile.

Providers must use procedure code E0642 for the purchase of a dynamic stander.
2.2.19.4.2 Prior Authorization and Documentation Requirements

THKAO (vertical or dynamic standers, standing frames or braces, and parapodiums), including all accessories, requires prior authorization.

THKAO may be considered for clients who are CCP-eligible and who require assistance to stand and remain standing when documentation submitted clearly shows that it is medically necessary and will correct or ameliorate the client’s disability or physical or mental illness or condition.

Prior authorization may be considered for the THKAOs with the following documentation:

- Diagnoses relevant to the requested equipment, including functioning level and ambulatory status
- Anticipated benefits of the equipment
- Frequency and amount of time of a standing program
- Anticipated length of time the client will require this equipment
- Client’s height, weight, and age
- Anticipated changes in the client’s needs, anticipated modifications, or accessory needs, as well as the growth potential of the stander

2.2.20 Prosthetic Services

2.2.20.1 Services, Benefits, and Limitations

External prostheses are a benefit of Texas Medicaid when provided by a licensed prosthetist or licensed prosthetist/orthotist through CCP for clients who are birth through 20 years of age.

The following prostheses and related services may be reimbursed when medical necessity criteria are met:

- Lower limb
- Upper limb
- Craniofacial
- External breast
- Repair, replacements, and modifications
- Prosthetic training
- Accessories to prostheses

Prosthetic training by a physical or occupational therapist for a lower limb prosthesis or an upper extremity prosthesis is a benefit for clients who have not worn one previously or for a prolonged period or who are receiving a different type.

Refer to: Section 5, “Children’s Therapy Services Clients birth through 20 years of age” in the Physical Therapy, Occupational Therapy, and Speech Therapy Services Handbook (Vol. 2, Provider Handbooks).

To be considered for reimbursement, prostheses must be dispensed, fabricated, or modified by a licensed prosthetist or licensed prosthetist/orthotist enrolled with Medicare and CCP.

The date of service for a custom-made or custom-fitted prosthesis is the date the supplier places an order for the equipment and incurs a liability for the equipment. The custom-made or custom-fitted prosthesis will be eligible for reimbursement as long as the service is provided during a month the client is eligible for Medicaid.
The following items and services are included in the reimbursement for a prosthetic device and not reimbursed separately:

- Evaluation of the residual limb and gait
- Measurement, casting, or fitting of the prosthesis
- Cost of base component parts and labor contained in the base procedure code description
- Repairs due to normal wear and tear during the 90 days following delivery
- Adjustments or modifications of the prosthesis or the prosthetic component made when fitting the prosthesis or component and for 90 days from the date of delivery when the adjustments are not necessitated by changes in the residual limb or the client’s functional ability

In general, base codes do not represent a complete device. To include the additional components necessary for a complete device, providers may bill additional components with a code that is used in addition to a base code. Addition codes may also be used to indicate modifications to a device. The values assigned to the additional codes do not represent the actual value of the component or modification, but only the difference between the total value and the value of the base code. As a result, reimbursement of an addition does not involve subtraction of any amounts from the base code allowance.

### 2.2.20.1.1 Noncovered Prosthetic Services

Prosthetic devices prescribed by a chiropractor are not a benefit of Texas Medicaid.

A vacuum-assisted socket system (procedure code L5781 or L5782), which is a specialized vacuum pump, is considered experimental and investigational, and is not a benefit of Texas Medicaid.

Myoelectric hand prostheses for conditions other than the absence of forearm(s) and hand(s) are considered experimental and investigational and are not a benefit of Texas Medicaid.

A prosthetic device customized with enhanced features is not considered medically necessary if ADLs can be met with a standard prosthetic device.

Accessories that are not required for the effective use of a prosthetic device are not considered medically necessary.

### 2.2.20.2 Prior Authorization and Documentation Requirements

Prior authorization is required for all prosthetic devices.

A completed CCP Prior Authorization Form requesting the prosthesis must be signed and dated by a physician familiar with the client before requesting prior authorization for all prostheses. The completed CCP Prior Authorization Form must include the procedure codes and numerical quantities for services requested. A copy of the completed, signed, and dated form must be maintained by the prosthesis provider in the client’s medical record. The completed CCP Prior Authorization Form with the original dated signature must be maintained by the prescribing physician in the client’s medical record.

To complete the prior authorization process by paper, the prosthesis provider must fax or mail the completed CCP Prior Authorization Request Form to the CCP prior authorization unit and retain a copy of the signed and dated CCP form in the client’s medical record at the provider’s place of business.

To complete the prior authorization process electronically, the prosthesis provider must complete the prior authorization requirements through any approved electronic methods and retain a copy of the signed and dated CCP Prior Authorization Request form in the client’s medical record at the provider’s place of business.
To facilitate determination of medical necessity and avoid unnecessary denials, the physician must provide correct and complete information, including documentation for medical necessity of the equipment or supplies requested. The physician must maintain documentation of medical necessity in the client’s medical record. The provider may be asked for additional information to clarify or complete a request for the service or device.

All requests for prior authorization must include documentation of medical necessity including, but not limited to, documentation that the client meets the following general indications for the requested device:

- The prosthesis replaces all or part of the function of a permanently inoperative, absent, or malfunctioning part of the limb, and identification of the specific limb that is being replaced by the prosthesis.
- The prosthesis is required for ADLs or for rehabilitation purposes, and identification of the ADLs or rehabilitation purpose for which the prosthesis is required.

The provider must keep the following written documentation in the client’s medical record:

- The prescription for the device.
- Prosthetic devices must be prescribed by a physician (M.D. or D.O.).
- The prescription must be dated prior to or on the initial date of the requested dates of service, which can be no longer than 90 days from the signature date on the prescription.
- Accurate diagnostic information that supports the medical necessity for the requested device. (A retrospective review may be performed to ensure that the documentation included in the client’s medical record supports the medical necessity of the requested service or device.)
- The specific make, model, and serial number of the prosthetic components.
- The treatment plan outlining the therapy program prescribed by the treating physician, including expected goals with the use of the prosthesis.
- A statement submitted by the physician that indicates that the client or client’s family or caregiver demonstrates willingness to comply with the therapy program.

Prior authorization is valid for a maximum period of six months from the prescription signature date. At the end of the six-month authorization period, a new prescription is required for prior authorization of additional services.

The actual date of service is the date the supplier has placed an order for the equipment and has incurred liability for the equipment.

### 2.2.20.2.1 Lower-Limb Prostheses

Lower limb prostheses include, but are not limited to, the following:

- Partial foot, ankle, and knee disarticulation sockets
- Above-knee short prostheses
- Hip and knee disarticulation prostheses
- Postsurgical prostheses
- Preparatory prostheses
- Additions to lower extremity prostheses
- Replacement sockets
A basic lower limb prosthesis consists of the following:

- A socket or connection between the residual limb and the prosthesis
- A suspension mechanism attaching the socket to the prosthesis
- A knee joint that provides support during stance, smooth control during the swing phase, and unrestricted motion for sitting and kneeling
- An exoskeleton or endoskeleton pylon (tube or shell) that attaches the socket to the terminal device
- A terminal device (foot)

In addition to the general indication requirements, the following additional documentation is also required for all lower limb prostheses:

- Written documentation of the client’s current and potential functional levels. A functional level is defined as a measurement of the capacity and potential of the individuals to accomplish their expected post-rehabilitation daily function. The potential functional ability is based on reasonable expectations of the treating physician and the prosthetist and includes, but is not limited to, the following:
  - The client’s history, including prior use of a prosthesis if applicable
  - The client’s current condition, including the status of the residual limb and any coexisting medical conditions
  - The client’s motivation to ambulate and ability to achieve independent transfers or ambulation with the use of a lower limb prosthesis
  - The following functional modifiers and levels have been defined by the Centers for Medicare & Medicaid Services (CMS):

<table>
<thead>
<tr>
<th>Functional Level</th>
<th>Functional Modifier</th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td>Level 0</td>
<td>K0</td>
<td>Does not have the ability or potential to ambulate or transfer safely with or without assistance, and a prosthesis does not enhance quality of life or mobility.</td>
</tr>
<tr>
<td>Level 1</td>
<td>K1</td>
<td>Has the ability or potential to use a prosthesis for transfers or ambulation on level surfaces at fixed cadence. Typical of the limited and unlimited household ambulator</td>
</tr>
<tr>
<td>Level 2</td>
<td>K2</td>
<td>Has the ability or potential for ambulation with the ability to traverse low level environmental barriers such as curbs, stairs, or uneven surfaces. Typical of the limited community ambulator.</td>
</tr>
<tr>
<td>Level 3</td>
<td>K3</td>
<td>Has the ability or potential for ambulation with variable cadence. Typical of the community ambulator who has the ability to traverse most environmental barriers and may have vocational, therapeutic, or exercise activity that demands prosthetic utilization beyond simple locomotion.</td>
</tr>
<tr>
<td>Level 4</td>
<td>K4</td>
<td>Has the ability or potential for prosthetic ambulation that exceeds basic ambulation skills, exhibiting high-impact, stress, or energy levels. Typical of the prosthetic demands of the child, active adult, or athlete.</td>
</tr>
</tbody>
</table>

A client whose functional level is zero (0) is not a candidate for a prosthetic device; the device is not considered medically necessary.

Advanced knee, ankle, and foot prostheses procedure codes must be submitted with the appropriate functional modifier in the table above.
Microprocessor-Controlled Lower Limb Prostheses

Microprocessor-controlled lower limb prostheses (e.g., Otto Bock C-Leg, Intelligent Prosthesis, or Ossur Rheo Knee) will be considered for prior authorization for clients who have a transfemoral amputation from a nonvascular cause, such as trauma or tumor and a functional level of 3 or above, and who meet the following criteria:

- The individual has adequate cardiovascular reserve and cognitive learning ability to master the higher level of technology and to allow for faster than normal walking speed.
- The individual demonstrates the ability to ambulate at a faster than baseline rate using a standard prosthetic application with a swing and stance control knee.
- The individual has a demonstrated need for long-distance ambulation at variable rates (greater than 400 yards) on a daily basis. Use of the limb in the home or for basic community ambulation is not sufficient to justify provision of the computerized limb instead of standard limb applications.
- The individual has a demonstrated need for regular ambulation on uneven terrain or for regular use on stairs. Use of the limb for limited stair climbing in the home or employment environment is not sufficient evidence for prescription of this device over standard prosthetic application.

The licensed prosthetist or licensed prosthetist/orthotist providing the device must be trained in the fitting and programming of the microprocessor-controlled prosthetic device.

Foot Prostheses

The following foot prostheses will be considered for prior authorization for clients whose documented functional level is 1 or above:

- A solid ankle-cushion heel (SACH) foot
- An external keel SACH foot or single axis ankle/foot

A flexible-keel foot or multi-axial ankle/foot will be considered for prior authorization for clients whose documented functional level is 2 or above.

A flex foot system, energy storing foot, multiaxial ankle/foot, dynamic response, or flex-walk system or equivalent will be considered for prior authorization for clients whose documented functional level is 3 or above.

A prosthetic shoe will be considered for prior authorization if it is an integral part of a prosthesis for clients with a partial foot amputation.

Ankle Prosthesis

An axial rotation unit will be considered for prior authorization for clients whose documented functional level is 2 or above.

Knee Prosthesis

A single-axis, constant-friction knee and other basic knee systems will be considered for prior authorization for clients whose documented functional level is 1 or above. A fluid, pneumatic, or electronic knee prosthesis will be considered for prior authorization for clients whose documented functional level is 3 or above. A high-activity knee control frame will be considered for prior authorization for clients whose documented functional level is 4.

Prosthetic Substitutions or Additions for Below-Knee Prostheses

Prosthetic substitutions or additions (procedure codes L5629, L5638, L5639, L5646, L5647, L5704, L5785, L5962, and L5980) are not considered medically necessary when an initial below-knee prosthesis (procedure code L5500) or a preparatory below-knee prosthesis (procedure codes L5510, L5520, L5530, or L5540) is provided.
Prosthetic substitutions or additions (procedure codes L5620, L5629, L5645, L5646, L5670, L5676, L5704, and L5962) are not considered medically necessary when a below-knee preparatory, prefabricated prosthesis (procedure code L5535) is provided.

**Sockets**

Prior authorization for test (diagnostic) sockets for an individual prosthesis is limited to a quantity of two test sockets. Prior authorization for same-socket inserts for an individual prosthesis is also limited to a quantity of two. Requests for test sockets or same-socket inserts beyond these limitations must include documentation of medical necessity that supports the need for the additional sockets.

### 2.2.20.2.2 Upper-Limb Prostheses

Upper limb prostheses include, but are not limited to, the following:

- Partial hand prostheses
- Wrist and elbow disarticulation prostheses
- Shoulder and interscapular thoracic prostheses
- Immediate postsurgical or early fitting prostheses
- Preparatory prostheses
- Terminal devices
- Replacement sockets
- Inner sockets-externally powered
- Electric hand, wrist, and elbow prostheses

Upper limb prostheses will be considered for prior authorization with documentation of all of the general indication requirements. The additional criteria in the following sections apply for specific prosthetic devices.

**Myoelectric Upper Limb Prostheses**

A myoelectric upper limb prosthetic device is considered medically necessary when all of the following criteria have been met:

- The client has sufficient neurological, myocutaneous, and cognitive function to operate the prosthesis effectively.
- The client has an amputation or missing limb at the wrist or above (e.g., forearm, elbow, and so on).
- The client is free of comorbidities that could interfere with maintaining function of the prostheses (e.g., neuromuscular disease).
- The client retains sufficient microvolt threshold in the residual limb to allow proper function of the prostheses.
- Standard body-powered prosthetic devices cannot be used or are insufficient to meet the functional needs of the patient in performing ADLs.
- The client does not function in an environment that would inhibit function of the prosthesis (e.g., a wet environment or a situation involving electrical discharges that would affect the prosthesis).

### 2.2.20.2.3 External Breast Prostheses

External breast prostheses will be considered for prior authorization for clients who have congenital absence of a breast or who have had a mastectomy.
2.2.20.2.4 Craniofacial Prostheses
Craniofacial prostheses include, but are not limited to, external nasal, ear, and facial prostheses. Craniofacial prostheses will be considered for prior authorization with documentation that the device is necessary to correct an absence or deformity of the affected body part.

2.2.20.2.5 Related Services

Accessories to Prostheses
Accessories to prostheses, such as stump stockings and harnesses will be considered for prior authorization when they are essential to the effective use of the prosthetic device.

Repairs, Replacements, and Modifications to Prostheses
Repairs due to normal wear and tear will be considered for prior authorization after 90 days from the date of delivery of the initial prosthesis, when the repair is:

• Necessary to make the equipment functional.
• More cost-effective than the replacement of the prosthetic device.

Providers must include documentation that supports medical necessity when they request prior authorization.

Additional information from the provider may be requested to determine cost-effectiveness.

Replacement of prosthetic equipment will be considered for coverage when loss or irreparable damage has occurred. A copy of the police or fire report when appropriate and the measures to be taken to prevent re-occurrence must be submitted with the prior authorization request.

Socket replacements will be considered for prior authorization with documentation of functional or physiological need, including, but not limited to, changes in the residual limb, functional need changes, or irreparable damage or wear due to excessive weight or prosthetic demands of very active amputees.

Children typically require new prosthetic devices every 12 to 18 months, although the actual lifespan of a device depends on the child’s rate of skeletal growth. Prosthetic devices for children must accommodate growth and other physiological changes.

Components and systems that allow for growth or increase the lifespan of the prosthesis may include the following:

• Growth-oriented suspension systems and modifications
• Use of modular systems
• Use of flexible sockets
• Use of removable sockets (slip or triple-wall sockets)
• Use of distal pads
• Modification of socket liners
• Increasing or decreasing sock thickness

Modifications due to growth or change in medical status will be considered for prior authorization with documentation of medical necessity.

Medical necessity for requested components or additions to the prosthesis is based on the client’s current functional ability and the expected functional potential as defined by the prosthetist and the ordering physician.
2.2.21 Phototherapy Devices

Phototherapy devices are not a benefit of Title XIX Home Health Services. Phototherapy devices are a benefit of Texas Medicaid through CCP for clients who are birth through 20 years of age.

2.2.21.1 Services, Benefits, and Limitations

The rental of phototherapy devices (procedure code E0202) for use in the home are a benefit of Texas Medicaid for low-risk infants.

Low-risk infants are 35 or more weeks gestation at birth, without comorbidity, and with a total serum bilirubin (TSB) level within the following ranges:

<table>
<thead>
<tr>
<th>Infant’s Gestation at Birth</th>
<th>TSB for infant 0-24 hours of age*</th>
<th>TSB for infant 25-48 hours of age*</th>
<th>TSB for infant 49-72 hours of age*</th>
<th>TSB for infant older than 72 hours of age*</th>
</tr>
</thead>
<tbody>
<tr>
<td>38 weeks or greater</td>
<td>6–11</td>
<td>12–15</td>
<td>15–18</td>
<td>18–21</td>
</tr>
</tbody>
</table>

* Infant age when TSB level is drawn.
TSB levels are expressed in milligrams per deciliter (mg/dl).

The DME provider must perform routine maintenance and provide instructions to the parent or guardian on the safe use of the phototherapy device. Rental of a phototherapy device is reimbursed as a daily global fee and is limited to one per day, per client, any provider.

Providers may not bill for those days the phototherapy device is at the client’s home and is not in use.

Skilled nursing (SN) visits for clients requiring phototherapy services may be reimbursed separately through Title XIX Home Health Services for nonroutine clinical teaching and assessment. Routine laboratory specimens are obtained during the SN visit, and may only be considered when the alternative to obtaining the specimen is to transport the client by ambulance.

If a client who is receiving PDN services requires phototherapy, instructions in the use of the equipment must be part of the existing PDN authorized hours. SN visits will not be allowed on the same day as PDN services.

In accordance with American Academy of Pediatrics (AAP) guidelines, providers must conduct ongoing assessments for risk of severe hyperbilirubinemia for all infants who receive home phototherapy.

Initiation of home phototherapy for medium- and high-risk infants is not a benefit of Texas Medicaid. As defined by the AAP, medium- and high-risk infants should be considered for more extensive initial treatment in an inpatient setting. Medium- and high-risk infants include, but are not limited to, those who have one of the following known risk factors:

- Acidosis
- Albumin less than 3.0 g/dl
- Asphyxia
- Glucose-6-phosphate dehydrogenase (G6PD) deficiency
- Isoimmune hemolytic disease (blood group incompatibility)
- Jaundice within the first 24 hours
- Sepsis
- Significant lethargy
- Temperature instability
2.2.21.2 Prior Authorization and Documentation Requirements

Home phototherapy devices require prior authorization and are provided only for the days that are medically necessary.

For low-risk infants, prior authorization will be considered for phototherapy services that begin in the home.

For stabilized infants who began phototherapy treatment during their hospitalization and have been discharged from the hospital, prior authorization will be considered for the continuation of phototherapy services in the home. Initial prior authorization may be given for a maximum of seven days of home phototherapy. A new CCP Prior Authorization Request Form must be submitted to request more than seven days of home phototherapy.

The following documentation is required to support medical necessity when requesting home phototherapy services:

- A diagnostic evaluation, which must include, but is not limited to, a normal history and physical exam, and normal laboratory values for the following, as medically indicated:
  - Complete blood count with differential
  - Platelets
  - Blood smear for red blood cell morphology
  - Reticulocyte count
  - Urinalysis
  - Maternal and infant blood typing
  - Coombs test
- TSB level (in mg/dl)
- Gestational age
- Documentation of adequate infant hydration, as demonstrated by 4-6 wet diapers per day and 3-4 stools per day
- Documentation stating that infant weight loss does not exceed 10 percent of the infant’s birth weight
- Physician’s plan of care
- Anticipated number of days the client will need the phototherapy treatment
- Documentation of parental education regarding the importance of monitoring and follow-up

When requesting prior authorization for a hospitalized infant that requires continued home phototherapy, providers must submit documentation that indicates all pre-existing medium- or high-risk factors have resolved or stabilized.

Providers must submit the following additional documentation for prior authorization requests for previously hospitalized infants that require continued home phototherapy or for more than seven days of home phototherapy:

- TSB level greater than 13 mg/dl and trending downward. TSB levels less than 13 will require medical review to determine medical necessity.

**Note:** According to AAP guidelines, phototherapy may be discontinued when the TSB level falls below 13-14 mg/dl; however, exceptions to the guidelines may be considered. As a result, documentation must include the rationale for not discontinuing phototherapy when the TSB level drops below 13 mg/dl.
• Birth weight and current weight demonstrating weight gain.

  **Note:** According to AAP guidelines, breast-fed infants are expected to gain 15-30 grams per day (1/2-1 ounce per day) through the first 2-3 months of life.

### 2.2.21.2.1 Retroactive Eligibility

Newborn babies may not have a Medicaid number at the time that services are ordered by the physician and provided by the supplier. In these cases, prior authorization may be given retroactively for services rendered between the start date and the date that the client’s Medicaid number becomes available.

The provider is responsible for finding out the effective dates of client eligibility.

The provider has 95 days from the date on which the client’s Medicaid number becomes available (add date) to obtain prior authorization for services that were already rendered.

### 2.2.22 Prothrombin Time/International Normalized Ratio (PT/INR) Home Testing Monitor

PT/INR home testing monitors are a benefit of Title XIX Home Health Services for clients who require chronic oral anticoagulation due to one of the following:

• Mechanical heart valve
• Chronic atrial fibrillation
• Venous thromboembolism (including both deep vein thrombosis [DVT] and pulmonary embolism)
• Ventricular assist device (VAD) awaiting a heart transplant

The PT/INR home testing monitor is a portable, battery-operated instrument for the quantitative determination of PT/INR from whole blood obtained by finger-stick. This product is designed to aid in the management of high-risk clients who take oral anticoagulants.

  **Note:** For clients who are 20 years of age and younger and do not meet criteria for coverage through Title XIX Home Health Services, home PT/INR monitors and related testing supplies may be considered through CCP.

  **Note:** For clients who are 21 years of age or older, requests for PT/INR monitors and related testing supplies that do not meet the criteria through Title XIX Home Health Services may be considered under the Texas Medicaid Home Health—Durable Medical Equipment (DME) Exceptional Circumstances process.

The following procedure codes are included in this benefit:

• Procedure code E1399 may be reimbursed for the rental or purchase of the monitor.
• Procedure code A9900 may be reimbursed for the related testing supplies.

Procedure codes E1399 and A9900 may be reimbursed to DME providers for services rendered in the home setting.

### 2.2.22.1 Prior Authorization

Prior authorization is required for the home PT/INR monitors and related testing supplies.

Prior authorization requests must be submitted within three business days of the date of service and must include documentation of medical necessity and a completed Home Health Services (Title XIX) DME/Medical Supplies Physician Order Form.

The completed Title XIX Form must be maintained by the requesting provider and the prescribing provider. The original signature copy must be kept in the provider’s medical record for the client.
To avoid unnecessary denials, the prescribing provider must provide correct and complete information, including documentation for medical necessity of the equipment and/or supplies requested. The prescribing provider must maintain documentation of medical necessity in the client’s medical record. The requesting provider may be asked for additional information to clarify or complete a request for the PT/INR monitor.

Prior authorization for the rental or purchase of a home PT/INR monitor and related testing supplies will be considered for clients who meet all the following criteria:

- The client is on anticoagulation therapy and has a current prescription for Warfarin or other oral anticoagulant.
- The client has been on anticoagulation therapy for at least three months prior to the request for the home PT/INR monitor.
- The client is required to self-test at least every two weeks.

Additionally, the client must have at least one of the following conditions documented in the request for prior authorization:

- Fluctuations of INR or PT/PTT levels with titration greater than once per week in anticoagulation dosing with copies of laboratory reports and resultant medication changes.
- A medical condition that limits physical movement, places the client under medical restrictions for isolation, or requires non-emergency ambulance transport for the purpose of obtaining laboratory specimens.
- Limited venous access that compromises the ability to obtain laboratory specimens for the adequate monitoring of anticoagulation therapy.

The prior authorization request will be evaluated upon receipt to determine whether the equipment will be rented, purchased, repaired, or modified based on the client’s needs, duration of use, and age of equipment.

*Note:* Skilled nursing (SN) visits will not be approved for the sole purpose of instructing the client on the use of the PT/INR home testing monitor. Any necessary instruction must be performed as part of the office visit with the prescribing physician.

2.2.23 *Respiratory Equipment and Supplies*

Respiratory equipment and supplies may be provided in the home under Title XIX Home Health Services.

Respiratory equipment and supplies must be prescribed by a physician, be FDA approved for the medical condition, and have federal financial participation available to be considered a medically necessary benefit. An eligible client must have compromised health status without the requested equipment or supplies.

Equipment provided for rental may be new or used. Equipment provided for purchase must be new and unused.

HHSC or its designee will determine whether respiratory equipment will be rented, purchased, or repaired based on the client’s needs and expected duration of use.

*Note:* When new unused equipment is initially provided for rental and is subsequently authorized for purchase, the provider is not required to replace the equipment.
Rental of equipment includes all necessary supplies, adjustments, repairs, and replacement parts. Supplies needed for use with client-owned equipment may be purchased separately within the limitations.

**Note:** [Revised] For clients who are 20 years of age and younger, respiratory equipment and supplies that do not meet the criteria under Title XIX Home Health Services may be considered through the Texas Health Steps-Comprehensive Care Program (THSteps-CCP).

**Note:** [Revised] For clients who are 21 years of age and older, respiratory equipment and supplies that do not meet the criteria under Title XIX Home Health Services may be considered through the Texas Medicaid Home Health-Durable Medical Equipment (DME) Exceptional Circumstances provision. Respiratory equipment and supplies are available without prior authorization up to the stated quantity limitation, unless otherwise specified in this handbook. Prior authorization is required for quantities exceeding the limitation.

### 2.2.23.1 Prior Authorization

Unless otherwise indicated, prior authorization is required for rental or purchase of respiratory equipment provided through Home Health services.

All miscellaneous procedure codes listed in this handbook require prior authorization.

#### 2.2.23.1.1 Initial Request

A completed, signed, and dated prior authorization request form ordering the DME or medical supplies must include the procedure codes and numerical quantities for services requested and must be signed and dated by the ordering physician and the representative of the DME and medical supply provider before requesting prior authorization for all DME and supplies.

A Home Health Services (Title XIX) DME/Medical Supplies Physician Order Form must be submitted for all DME services and supplies, unless the ordering physician is requesting the following:

- A continuous positive airway pressure (CPAP) or Bi-level positive airway pressure (BiPAP) and respiratory assist devices (RADs) are to be requested using a Texas Medicaid Prior Authorization Request for CPAP or RAD (Bi-level PAP) form.

  **Note:** Home ventilators (procedure codes E0465, E0466, and E0467) requested with CPAP or RAD settings must also be submitted on the Texas Medicaid Prior Authorization Request for CPAP or RAD (Bi-level PAP) form with a letter of medical necessity explaining why CPAP or RAD is not medically appropriate for the client.

- Oxygen Therapy is to be requested using a Texas Medicaid Prior Authorization Request for Oxygen Therapy Devices and Supplies form.

- Secretion and mucus clearance device are to be requested using a Texas Medicaid Prior Authorization Request for Secretion and Mucus Clearance Devices - Initial Request form or a Texas Medicaid Prior Authorization Request for Secretion and Mucus Clearance Devices -Renewal Request form for secretion and mucus clearance devices.

  **Note:** It is not necessary to submit a Title XIX form if one of the prior authorization forms listed above are submitted.

The following completed, signed, and dated prior authorizations forms must be maintained by the DME provider and ordering physician in the client’s files. The ordering physician must retain the completed, signed, and dated original form. The DME provider must retain copies of the completed original prior authorization form that contains the ordering physician’s dated signature. The following forms will not be accepted beyond 90 days from the date of the ordering physician’s signature:

- Title XIX Form
- Texas Medicaid Prior Authorization Request for CPAP or RAD (Bi-level PAP) form
• Texas Medicaid Prior Authorization Request for Oxygen Therapy Devices and Supplies form
• Texas Medicaid Prior Authorization Request for Secretion and Mucus Clearance Devices - Initial Request form
• Texas Medicaid Prior Authorization Request for Secretion and Mucus Clearance Devices -Renewal Request form

Prior authorization requests may be submitted to the TMHP Prior Authorization Department via mail, fax, or the electronic portal. Prescribing or ordering providers, dispensing providers, clients’ responsible adults, and clients may sign prior authorization forms and supporting documentation using electronic or wet signatures.

Refer to: Subsection 5.5.1.2, “Document Requirements and Retention” in “Section 5: Fee-for-Service Prior Authorizations” (Vol. 1, General Information) for additional information about electronic signatures.

To complete the prior authorization process electronically, the provider must complete the prior authorization requirements through any approved electronic methods and retain a copy of the signed and dated prior authorization form in the client’s medical record.

To complete the prior authorization process by paper, the provider must fax or mail the completed prior authorization request form to the Home Health prior authorization unit and retain a copy of the signed and dated prior authorization form in the client’s medical record.

To facilitate determination of medical necessity and avoid unnecessary denials, the physician must provide correct and complete information, including documentation for medical necessity for the equipment or supplies requested. The physician must maintain documentation of medical necessity in the client’s medical record.

The requesting provider may be asked for additional information to clarify or complete a request.

Retrospective review may be performed to ensure documentation supports the medical necessity of the requested equipment or supplies.

A determination as to whether the equipment will be rented, purchased, repaired, or modified will be made by HHSC or its designee based on the client’s needs, duration of use, and age of the equipment.

Equipment that has been purchased may be considered for replacement when loss or irreparable damage has occurred outside the warranty terms, conditions, and limitations. A copy of the police or fire report when appropriate and the measures to be taken to prevent reoccurrence must be submitted with the prior authorization request.

2.2.23.1.2 * Respiratory Equipment for Clients not Meeting Required Criteria

Clients who are 20 years of age and younger

[Revised] For clients who are 20 years of age and younger, respiratory equipment and supplies that do not meet the criteria under Title XIX Home Health Services may be considered through the Texas Health Steps-Comprehensive Care Program (THSteps-CCP).

Clients who are 21 years of age and older

[Revised] For clients who are 21 years of age and older, respiratory equipment and supplies that do not meet the criteria under Title XIX Home Health Services may be considered through the Texas Medicaid Home Health-Durable Medical Equipment (DME) Exceptional Circumstances provision.

Refer to: [Revised] Section 2.2.3, “Home Health DME and Supplies Exceptional Circumstances Provision” in this handbook for additional information about the Exceptional Circumstances provision."
2.2.23.1.3 Renewal Requests for all Respiratory Equipment

Providers are expected to submit documentation for recertification requests as outlined in this handbook. If no specific documentation requirements are outlined, providers are to submit the following:

- A new prior authorization request form
- All of the initial request requirements
- A physician attestation that the treatment has been effective and the client has been compliant with treatment

2.2.23.1.4 Repair to Client-Owned Equipment

Repairs to client-owned equipment may be prior authorized as needed with documentation of medical necessity. Technician fees are considered part of the cost of the repair.

HHSC or its designee reserves the right to request additional documentation about the need for repairs when there is evidence of abuse or neglect to equipment by the client, client’s family, or caregiver. When there is documented proof of abuse or neglect, requests for repairs will not be prior authorized.

Providers are responsible for maintaining documentation in the client’s medical record specifying the repairs and supporting medical necessity and must include all the following:

- The date of purchase
- The serial number of the current equipment (as applicable)
- The cause of the damage or need for repairs
- What steps the client or caregiver will take to prevent further damage if repairs are due to an accident
- When requested, the cost of purchasing new equipment as opposed to repairing current equipment

Temporary replacement of client-owned respiratory equipment during the repair may be considered for prior authorization for one month using procedure code K0462.

Labor for repair of client-owned respiratory equipment may be considered for prior authorization using procedure code K0739 up to a maximum of two hours per day (maximum quantity of 8 units).

Routine maintenance of rental equipment is the provider’s responsibility.

2.2.23.2 Small Volume Nebulizers (SVN)

Small volume nebulizers (SVNs) (procedure code E0570) and related supplies (procedure codes A7003, A7004, and A7005) may be considered for purchase without prior authorization for the conditions listed below:

<table>
<thead>
<tr>
<th>Condition</th>
</tr>
</thead>
<tbody>
<tr>
<td>Bronchiectasis - Any type</td>
</tr>
<tr>
<td>Cystic Fibrosis (CF) with pulmonary manifestations</td>
</tr>
<tr>
<td>Pneumonia - Any type</td>
</tr>
<tr>
<td>Influenza</td>
</tr>
<tr>
<td>Bronchitis - Any type</td>
</tr>
<tr>
<td>Emphysema - Any type</td>
</tr>
<tr>
<td>Asthma - Any type</td>
</tr>
<tr>
<td>Chronic Obstructive Pulmonary Disease (COPD) - any type</td>
</tr>
<tr>
<td>Pneumoconiosis - Any type</td>
</tr>
</tbody>
</table>
Supporting documentation of medical necessity must be maintained in the client’s medical record, which must be available on request and is subject to retrospective review including but not limited to the following:

- Completed Title XIX Form signed and dated by the client’s prescribing physician
- Name of the medication(s) ordered for administration through the nebulizer treatments
- Frequency and duration of the need for the prescribed nebulizer treatments

SVN and related supplies may be considered for conditions not listed above with prior authorization when submitted with the following documentation of medical necessity:

- Completed Title XIX Form signed and dated by the client’s treating physician
- Justification supporting the use of an SVN to treat the client’s diagnosis
- Name of the medication(s) ordered for administration through the nebulizer
- Frequency and duration of need for the prescribed nebulizer treatments

### 2.2.23.3 Ultrasonic Nebulizers (USNs)

An ultrasonic nebulizer (USN) or electronic aerosol generator (procedure code E0574) is an electrically powered device that uses a piezoelectric crystal to generate aerosol. This crystal transducer converts radio waves into high-frequency mechanical vibrations (sound).

Ultrasonic nebulizers are a benefit when medically necessary and may be considered for purchase with prior authorization.

#### 2.2.23.3.1 Prior Authorization

Purchase of a USN may be considered medically necessary when submitted with all the following documentation:

- Client meets the criteria for a SVN
- Client requires equipment for delivery of one of the following:
  - Treprostinil to treat pulmonary arterial hypertension (PAH) when used to diminish symptoms associated with exercise
  - Tobramycin to treat cystic fibrosis (CF)

<table>
<thead>
<tr>
<th>Condition</th>
</tr>
</thead>
<tbody>
<tr>
<td>Acute, Sub-acute or Chronic Respiratory conditions</td>
</tr>
<tr>
<td>Respiratory conditions due to radiation, smoke, unspecified and specified external agents</td>
</tr>
<tr>
<td>Abnormal Sputum</td>
</tr>
<tr>
<td>Other diseases of the of the trachea and bronchus</td>
</tr>
<tr>
<td>Tracheostomy Status</td>
</tr>
<tr>
<td>Attention to tracheostomy</td>
</tr>
<tr>
<td>HIV with pulmonary manifestations</td>
</tr>
<tr>
<td>Pneumocystosis</td>
</tr>
<tr>
<td>Complications of a specified or unspecified transplanted organ, bone marrow or stem cells</td>
</tr>
<tr>
<td>Primary Pulmonary Hypertension</td>
</tr>
<tr>
<td>Other Chronic Pulmonary Heart Disease</td>
</tr>
</tbody>
</table>
USN may also be considered medically necessary for conditions other than those listed only when all of the following criteria have been met:

- Client meets the criteria for a SVN
- Client’s treating physician attests that the client has been compliant with other nebulizer and medication therapy
- Use of a SVN has failed to control the client’s disease, such as preventing the client from utilizing the hospital or emergency room

### 2.2.23.4 Humidification Therapy and Heating Elements

Humidification involves adding water vapor and sometimes heat to the inspired gas. Humidification therapy and heating elements are provided using the following procedure codes:

<table>
<thead>
<tr>
<th>Procedure Codes</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>Large Volume nebulizer jar (water jar)</strong></td>
</tr>
<tr>
<td>A7007</td>
</tr>
<tr>
<td><strong>Humidifiers</strong></td>
</tr>
<tr>
<td>E0550</td>
</tr>
<tr>
<td><strong>Other Supplies</strong></td>
</tr>
<tr>
<td>E0565</td>
</tr>
</tbody>
</table>

#### 2.2.23.4.1 Heating Elements

Heated humidifiers (procedure code E0562) and heated nebulizers (procedure code E0585) are used for clients with bypassed upper airways, clients receiving mechanical ventilatory support, and clients with high flow positive airway pressure devices. If heat is required for use with a large volume nebulizer (LVN), an immersion element (procedure code E1372) can be added.

#### 2.2.23.4.2 Prior Authorization

Humidification therapy and heating elements may be considered for rental or purchase with prior authorization with documentation of medical necessity.

Indications for humidification and warming of inspired gases are humidifying dry medical gases and overcoming humidity deficit created when upper airway is bypassed.

Clinical signs and symptoms that may be an indication that airway humidification is medically necessary include the following:

- Dry, nonproductive cough
- Increased airway resistance
- Increased incidence of infection
- Increased work of breathing
- Complaint of substernal pain and airway dryness
- Thick, dehydrated secretions

Providers must specify the site of room air or oxygen delivery (e.g., nose or mouth, hypopharynx, trachea) and state how much heat and humidity is needed to mitigate the cold, dry gas delivered through the site.

Providers must submit a Title XIX form and all of the following documentation to obtain prior authorization for the monthly rental or purchase of humidification therapy or heating elements:

- Evidence that the client has a tracheostomy or tracheobronchial stent
• Evidence that the client has thick tenacious secretions not responsive to normal levels of humidification provided with routine humidifiers used with regulators or flow meters
• Evidence that the client is not currently renting a ventilator
• Evidence that the client is not currently renting a compressor for the delivery of humidification

Providers must specify the site of room air or oxygen delivery (e.g., nose or mouth, hypopharynx, trachea) and state how much heat and humidity is needed to mitigate the cold, dry gas delivered through the site.

2.2.23.5 Large Volume Nebulizer Jars (Water Jars) and Compressors

Large volume nebulizer jars (procedure codes A7007 and A7017) used with compressors in humidification systems are a benefit when medically necessary and may be considered for purchase without prior authorization.

If heat is required, a heating element, such as an immersion element, can be added.

The autoclavable nebulizer (procedure code E0580 - glass or plastic) for use with a regulator or flow meter may be considered with prior authorization and documentation of medical necessity.

Equipment used with a large volume nebulizer to create a humidification system is a benefit when medically necessary with prior authorization.

A compressor (procedure code E0565) may be considered for rental or purchase with prior authorization. A compressor and heater (procedure code E0585), or large volume ultrasonic nebulizer (procedure code E0575) used to create the humidification systems when combined with a compressor are available for purchase and require prior authorization.

2.2.23.5.1 Prior Authorization

A large volume ultrasonic nebulizer and a nebulizer with compressor and heater may be considered for purchase with prior authorization when all the following criteria are met:

• The client has thick, tenacious secretions
• The client has one of the following medical conditions:
  • Cystic fibrosis
  • Bronchiectasis
  • A tracheostomy
  • A tracheobronchial stent

The compressor may also be considered when all of the following criteria are met:

• The compressor is needed for the administration of pentamidine using a filtered nebulizer
• The client has one of the following medical conditions:
  • HIV with pulmonary complications
  • Pneumocystosis
  • Complications of organ transplants

2.2.23.6 Intermittent Positive-Pressure Breathing (IPPB) Devices

IPPB is not the therapy of first choice for delivering aerosol or as a method of lung hyperinflation when other therapies can reliably meet the clinical objectives prescribed for the client. Prior authorization of an IPPB device (procedure code E0500) is available for rental or purchase and will be considered with documentation of ineffective response with use of other modalities such as treatment with a cough assist device.
Rental of the IPPB device includes all supplies, such as humidification and tubing.

In accordance with the American Association for Respiratory Care (AARC) recommendations, IPPB may be considered when one of the following indications is documented:

- The client has a need to improve lung expansion
- The need to deliver aerosol medication to a client when other methods of delivery have been unsuccessful

### 2.2.23.6.1 Prior Authorization

IPPB requires prior authorization and may be considered with documentation of ineffective response to treatment when other modalities, such as a cough assist device, have failed, when prescribed in accordance with the AARC recommendations, and there is medical necessity to improve lung expansion due to one of the following:

- The presence of clinically significant pulmonary atelectasis when other forms of therapy have been unsuccessful or the client cannot cooperate with the treatment
- The inability to clear secretions adequately due to pathology that severely limits the client’s ability to ventilate or cough effectively and failure to respond to other modes of treatment, including but not limited to:
  - Neuromuscular disorders or kyphoscoliosis with associated decreases in lung volumes and capacities
  - Presence of acute severe bronchospasm or exacerbated COPD that fails to respond to other therapy
  - Deliver aerosol medication when other methods of delivery have been unsuccessful including, but not limited to:
    - The client who has fatigue as a result of ventilatory muscle weakness such as neuromuscular disease, kyphoscoliosis, or spinal cord injury
    - The clients with severe hyperinflation where IPPB may decrease dyspnea and discomfort during nebulized therapy

### 2.2.23.7 Controlled Dose Inhalation Drug Delivery Systems

A controlled dose inhalation drug delivery system (procedure code K0730) is a benefit when it is medically necessary to deliver the drug iloprost (e.g. Ventavis) and is restricted to clients with pulmonary artery hypertension. The controlled dose inhalation drug delivery system is available for purchase with prior authorization when medically necessary.

#### 2.2.23.7.1 Prior Authorization

Prior authorization is required for purchase of a controlled dose inhalation drug delivery system when used with iloprost and may be considered when the client has a diagnosis of pulmonary artery hypertension and the pulmonary hypertension is not secondary to pulmonary venous hypertension or disorders of the respiratory system.

### 2.2.23.8 *Continuous Positive Airway Pressure (CPAP) and Respiratory Assist Devices (RAD) including Bi-Level PAP*

Continuous positive airway pressure (CPAP) (procedure code E0601) and respiratory assist devices (RADs) (procedure code E0470) without set back up rate are available for rental or purchase with prior authorization.

RADs with set back up rate (procedure codes E0471 and E0472) are a benefit when medically necessary and may be considered for rental with prior authorization for clients requiring:

- Treatment of obstructive sleep apnea
• Restrictive thoracic disorders
• Severe chronic obstructive pulmonary disease
• Central sleep apnea
• Complex sleep apnea
• Hypoventilation syndrome

Note: Other conditions may be considered based on medical necessity.

RADs with a set backup respiratory rate are available for rental, only when medically necessary.

Humidification devices (heated and non-heated) may be a benefit with prior authorization when medically necessary for rental or purchase for use with CPAP devices and RADs.

The following related supplies procedure codes are included in the CPAP and RAD rental and will not be reimbursed separately:

[Revised]

<table>
<thead>
<tr>
<th>Procedure Codes</th>
</tr>
</thead>
<tbody>
<tr>
<td>A7029</td>
</tr>
<tr>
<td>A7039</td>
</tr>
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</table>

The following procedure codes will deny if billed in the same month by any provider as procedure code E0472:

<table>
<thead>
<tr>
<th>Procedure Codes</th>
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<tbody>
<tr>
<td>A7029</td>
</tr>
<tr>
<td>A7039</td>
</tr>
</tbody>
</table>

2.2.23.8.1 Prior Authorization

CPAP devices (procedure code E0601) and RADs (procedure codes E0470, E0471, and E0472) require prior authorization. CPAP devices deliver a single, fixed pressure to the client during the night while sleeping. Some sleep breathing disorders do not benefit from CPAP and require treatment with RADs that are able to recognize the client’s breathing patterns and adjust pressure during the respiratory cycle during sleep.

Note: CPAP and RAD criteria are based on CMS coverage determinations.

CPAP and RAD accessories (headgear, chin straps, face masks, nasal pillows, cushions, nasal interfaces, tubing and filters), when used with the following procedure codes and within the maximum allowed limits, do not require prior authorization with a fee-for-service history of a client-owned CPAP and RAD device:

<table>
<thead>
<tr>
<th>Procedure Codes</th>
</tr>
</thead>
<tbody>
<tr>
<td>A7029</td>
</tr>
<tr>
<td>A7039</td>
</tr>
</tbody>
</table>

Note: RAD with backup rate used with an invasive interface (procedure codes E0471 and E0472) do require prior authorization. Supplies are included in the rental of procedure code E0472 and will not be authorized separately.
In the case of a client-owned RAD with backup rate that is used with invasive interface (procedure code E0472) that was purchased as a result of a rental or purchased through another payer source, proof of ownership of the device is required for consideration of reimbursement of associated supplies and accessories. A statement from the treating physician providing the make and model of the client-owned device, submitted with the claims appeal, will meet this requirement if the claims history is not available.

A CPAP device or a RAD without a set backup rate may be considered for an initial three month rental period with prior authorization. Following the initial three month rental period, if the CPAP or RAD without a set backup rate is effective, the device may be considered for purchase. Both devices may also be considered for continued rental with renewal at three month intervals up to 12 months.

A CPAP device and a RAD without a set backup rate may be considered purchased after 12 months of rental through the same provider and a request for purchase or further rental will not be considered.

A RAD with a set backup respiratory may be considered for an initial 3 month rental period with prior authorization and will be considered for rental only.

Humidification devices (heated or non-heated) for use with a CPAP or RAD device may be a benefit with prior authorization when medically necessary. Documentation submitted must support why humidification is medically necessary for use with positive pressure ventilation.

Prior authorization may be considered for initial and renewal requests for CPAP and RAD devices with submission of the Texas Medicaid Prior Authorization Request for CPAP or RAD (Bi-level PAP) form (new with each request) completed, signed and dated by the treating physician with the following sections completed:

- A and B for initial requests
- A and C for renewal requests

Additional documentation (e.g., titration sleep studies) as necessary to support the medical necessity of the service required as outlined below for the initial and renewal requests.

**Initial Request for a Continuous Positive Airway Pressure (CPAP) System**

The American Academy of Sleep Medicine (AASM) Guidelines state that it is clinically appropriate to treat clients who are 18 through 20 years of age using adult criteria.

A CPAP device (procedure code E0601) may be considered for an initial three-month rental period based on documentation supporting the medical necessity and appropriateness of the device. Documentation must include that the client has had a sleep study, lasting minimum of two hours, and at least one of the following criteria:

- For clients who are 17 years of age and younger, polysomnography results documenting an apnea-hypopnea index (AHI) greater than one event per hour may be used to establish medical necessity

- For clients who are 18 years of age and older, polysomnography results documenting an AHI or a respiratory disturbance index (RDI) greater than or equal to 15 events per hour

- For clients who are 18 years of age and older, an AHI or RDI greater than five events per hour with documentation of at least one of the following:
  - Excessive daytime sleepiness assessed by either the Epworth Sleepiness Scale (ESS) with a result greater than 10 or the Multiple Sleep Latency Test (MSLT) with a result less than 6
  - Symptoms of impaired cognition, mood disorders, or insomnia
  - Hypertension (systolic blood pressure greater than 140 mm Hg or diastolic blood pressure greater than 90 mm Hg)
  - Ischemic heart disease or previous myocardial infarction
  - History of stroke
• Greater than 20 episodes of oxygen desaturation to less than 85 percent during a full night sleep study
• Any one episode of oxygen desaturation of less than 70 percent
• Pulmonary hypertension

CPAP may be medically necessary for the treatment of obstructive sleep apnea (OSA) in clients younger than 18 years of age when one of the following criteria is documented:
• Adenoidectomy or tonsillectomy is contraindicated
• Adenoidectomy or tonsillectomy is delayed
• Adenoidectomy or tonsillectomy has been unsuccessful in relieving symptoms of OSA

Documentation must be maintained in the client’s medical record that the client or responsible caregiver has received instruction from the DME provider on the proper use and care of the device and supplies.

Renewal Request for a CPAP System

Prior authorization for purchase or an additional three month CPAP rental after the initial three-month rental period will be considered with all of the following documentation completed, signed, and dated by the client’s treating physician:
• A new Texas Medicaid Prior Authorization Request for CPAP or RAD (Bi-level PAP) form
• Documentation of medical necessity supporting:
  • The client’s continuous use of the equipment for a minimum of four hours per 24 hour period
  • The client’s symptoms as documented by the treating physician are improved with use of the CPAP

Continued rental of CPAP may be considered for up to 12 months with renewal at 3 month intervals. A CPAP device will be considered purchased after 12 months of continuous rental through the same provider.

Initial Request for Respiratory Assist Devices (RADs), including BiPAP - with and without a Set Backup Respiratory Rate

A RAD with or without a set back up rate may be considered for prior authorization when the client has one of the following medical conditions as documented by a sleep study and meets criteria for medical necessity for the specific medical condition:
• Obstructive sleep apnea (OSA)
• Restrictive thoracic disorders (i.e., neuromuscular diseases or severe thoracic cage abnormalities)
• Severe COPD
• Central sleep apnea (CSA), complex sleep apnea (CompSA)
• Hypoventilation syndrome

Initial Request for RAD for the Treatment of Obstructive Sleep Apnea (OSA)

A RAD without backup may be considered for an initial three month trial period for the treatment of OSA with prior authorization and submission of all of the following:
• All the required documentation delineated on the Texas Medicaid Prior Authorization Request for CPAP or RAD (Bi-level PAP) form outlined in the section for CPAP
• The client meets the criteria for the initial CPAP rental
• The documentation supports that CPAP has been tried and one of the following is documented:
• The client’s treating physician verifies that a therapeutic trial of CPAP was conducted in the home or a facility setting and failed to be effective in treating the client’s OSA

• A CPAP device was found to be ineffective during the initial facility based or sleep laboratory titration trial testing

If a CPAP device is tried and found ineffective during the initial facility-based titration or home trial, substitution of a RAD does not require a new face-to-face clinical evaluation or a new sleep test.

**Initial Request for RAD for the Treatment of Restrictive Thoracic Medical Conditions**

A RAD without a set backup rate requires prior authorization and may be considered for the treatment of thoracic medical conditions when all of the following are met:

• The client is diagnosed with a neuromuscular disorder (e.g., Duchenne muscular dystrophy, ALS, spinal cord injuries) or the client has a diagnosis of a severe thoracic cage abnormality (e.g., severe chest wall deformities) negatively impacting the client’s respiratory effort

• Significant respiratory insufficiency is documented by one of the following:
  • An arterial blood gas (ABG) PaCO2 greater than or equal to 45 mm Hg, obtained while awake and breathing the client’s routinely prescribed fraction of inspired oxygen concentration (FiO2)
  • Sleep oximetry demonstrates oxygen saturation less than or equal to 88 percent for 5 minutes or longer of continuous nocturnal recording time (minimum recording time of 2 hours), obtained while client is breathing his or her routinely prescribed FiO2

  **Note:** *FiO2 is the concentration of oxygen prescribed for routine use by the client. For example, if the client does not normally use supplemental oxygen, their prescribed oxygen is room air (FiO2 of 21 percent).*

For clients who have been diagnosed with a neuromuscular disorder only, documentation must support one of the following:

• Maximal inspiratory pressure less than 60 cm H2O

• Forced vital capacity less than 50 percent of predicted volume

A RAD with a set backup rate requires prior authorization and may be considered for the treatment of thoracic medical conditions when all of the following are met:

• The client meets the criteria for use of the RAD without a backup rate for the treatment of a thoracic medical condition

• The ordering physician certifies to all of the following:
  • Client has tried a RAD without a backup rate for at least 60 days
  • The client was compliant in the use of the device (using the device on average 4 or more hours in a 24 hour day)
  • The desired therapeutic respiratory response was not achieved with the RAD without a set backup rate

**Initial Request for RAD for the Treatment of Severe Chronic Obstructive Pulmonary Disease (COPD)**

A RAD without a backup rate may be considered for the treatment of severe COPD, with prior authorization, when all of the following criteria are met:

• An arterial blood gas PaCO2 less than 52 mm Hg, obtained while awake and when the client is either using 2 LPM of oxygen or the client’s prescribed FiO2 (the blood gas should be drawn while the client is using whichever concentration of oxygen is the higher of the two)
• Sleep oximetry demonstrates oxygen saturation less than or equal to 88 percent for 5 minutes or longer of continuous nocturnal recording time (minimum recording time of 2 hours), obtained while breathing oxygen at 2 LPM or the client’s prescribed FIO2 (whichever is higher)

• Prior to initiating therapy, documentation of sleep apnea and that treatment with CPAP has been considered with an explanation of why it was ruled out

To rule out the use of a CPAP, formal sleep testing is not required if there is sufficient information in the medical record submitted with the request to demonstrate that the client does not suffer from some form of sleep apnea (obstructive sleep apnea (OSA), CSA, or CompSA) as the predominant cause of awake hypercapnia or nocturnal arterial oxygen desaturation.

A RAD with a backup feature will be considered with prior authorization for severe COPD when the all of the following criteria are met:

• The client meets the criteria for use of the RAD without a backup rate for COPD

• The ordering physician certifies to all of the following:
  • Client has tried a RAD without a backup rate for at least 60 days
  • The client was compliant in the use of the device (using on average 4 or more hours in a 24 hour day)
  • The desired therapeutic respiratory response was not achieved with the RAD without a set backup rate

**Initial Request for RAD for the Treatment of Central Sleep Apnea (CSA) or Complex Sleep Apnea (CompSA)**

CSA or CompSA is characterized by the development of central apneas or central hypopneas during pressure titrations performed in a sleep lab titration study for CPAP or RAD without a backup rate. A RAD without a backup rate will be considered with prior authorization for the treatment of CSA or CompSA when a facility based polysomnogram is performed and supports all of the following:

• The client has a diagnosis of CSA or CompSA

• The sleep study documents one of the following:
  • The sum total of central hypopneas plus central apneas is greater than 50 percent of the total apneas and hypopneas rate
  • A central hypopnea/apnea rate index greater than 5 events per hour; and significant improvement of the sleep-associated hypoventilation while breathing the clients prescribed FiO2
  • Documentation ruling out CPAP as effective therapy if either OSA or CSA is a component of the initially observed sleep associated hypoventilation

A RAD with a backup rate will be considered with prior authorization for the treatment of CSA or CompSA when all of the following are met:

• The client meets the criteria for use of the RAD without a backup rate for the treatment of CSA or CompSA

• The ordering physician certifies to all of the following:
  • Client has tried a RAD without a backup rate for at least 60 days
  • The client was compliant in the use of the device (using on average 4 or more hours in a 24 hour day)
• The desired therapeutic respiratory response was not achieved with the RAD without a set backup rate

**Initial Request for RAD for the Treatment of Hypoventilation Syndrome**

A RAD without a backup rate may be considered for treatment of hypoventilation syndrome with prior authorization when all of the following criteria are met:

- An initial arterial blood gas PaCO$_2$, obtained while awake with the client breathing their prescribed FIO$_2$, greater than or equal to 45 mm Hg
- Spirometry shows a forced expired volume in 1 sec (FEV$_1$) or the forced vital capacity (FVC) greater than or equal to 70 percent
- A facility-based polysomnogram demonstrates oxygen saturation less than or equal to 88 percent for 5 minutes or longer of continuous nocturnal recording time (minimum recording time of 2 hours) not caused by obstructive upper airway events

A RAD with a set backup respiratory rate may be considered with prior authorization for the treatment of hypoventilation syndrome when one of the following are met:

- The client has hypoventilation syndrome as determined by a facility-based polysomnogram that demonstrates the desired respiratory therapeutic effects were not achieved with a RAD without a backup rate
- The client meets the criteria for RAD without a backup rate for hypoventilation syndrome, and the physician documents the desired respiratory therapeutic effects were not achieved with the RAD without a backup rate.

**Renewal Request for RAD with or without a Backup Rate**

Prior Authorization is required for renewal of a RAD with or without a backup rate.

Prior authorization for purchase of RAD without a set back up rate or continued rental of a RAD with or without a backup rate, after completion of the initial three-month rental period, may be considered with all of the following documentation completed, signed, and dated by the client’s treating physician:

- A new Texas Medicaid Prior Authorization Request for CPAP or RAD (Bi-level PAP) form
- Attestation from the treating physician that states the client is continuing to use the equipment at a minimum of four hours in a 24 hour period
- Client symptoms are improved as documented by the client’s treating physician

When recertifying a RAD with or without a set backup rate for significant respiratory insufficiency, documentation of a capillary blood gas (CBG) demonstrating a PaCO$_2$ greater than or equal to 45 mm Hg, obtained while awake and breathing the client’s routinely prescribed FiO$_2$ may be submitted in lieu of an ABG.

### 2.2.23.9 Mechanical Ventilation

Invasive and noninvasive ventilators (procedure codes E0465, E0466, and E0467) are considered for rental only with prior authorization and documentation of medical necessity indicating a clinical need for mechanical ventilation.

Mechanical ventilation may be considered for the treatment of, but not limited to:

- Neuromuscular or musculoskeletal diseases and conditions affecting the respiratory muscles
- Thoracic restrictive diseases
- Chronic respiratory failure
• The following procedure codes will deny if billed in the same month by any provider as procedure codes E0465 and E0466:

<table>
<thead>
<tr>
<th>Procedure Codes</th>
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</thead>
<tbody>
<tr>
<td>A4481</td>
</tr>
</tbody>
</table>

The following procedure codes will deny if billed in the same month by any provider as procedure code E0467:

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<td>A7013</td>
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<td>A7031</td>
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<tr>
<td>A7525</td>
</tr>
<tr>
<td>E0465</td>
</tr>
<tr>
<td>E0570</td>
</tr>
</tbody>
</table>

2.2.23.9.1 Prior Authorization

All ventilators and related equipment require prior authorization.

The DME provider is responsible for ensuring that there is a contingency plan to manage interruptions in the use of equipment such as emergency situations and mechanical failures that would be life threatening for the client. Acceptable plans include input from the client’s treating practitioner that takes into account the severity of the client’s medical condition and time constraints in providing emergency support.

A ventilator may be considered for an initial 3-month rental period. Following the initial 3-month rental period, if the ventilator was effective, it may be considered for ongoing 6 month rental periods.

A home ventilator with an invasive or noninvasive interface (procedure codes E0465, E0466, and E0467) is not intended for use as a CPAP or RAD; however, a home ventilator may be considered when medically necessary. A home ventilator requested with CPAP or RAD settings must be submitted on the Texas Medicaid Prior Authorization Request for CPAP or RAD (Bi-level PAP) form with a letter of medical necessity explaining why a CPAP or RAD is not medically appropriate for the client.

Prior authorization may be considered for initial and renewal requests for mechanical ventilators with submission of all of the following:

• A completed Title XIX Form signed and dated by the client’s treating physician (new with each request)

• Attestation from the treating physician that the mechanical ventilator is medically necessary and the client is compliant with the use of the equipment

The completed Title XIX Form must specify all ventilator settings and must be maintained by the DME provider and the treating physician in the client’s medical record.

The monthly ventilator rental includes all ventilator related supplies regardless of the client’s duration of use, whether 24 hours per day or less, including, but not limited to:

• Internal filters

• External filters

• Ventilator circuits with an exhalation valve
• High and low pressure alarms
• Humidification systems including supplies and solutions (e.g., sterile or distilled water)
• In-line compressors and related supplies
• Replacement ventilator
• Tracheostomy tube filters and humidification devices, such as heat moisture exchangers (procedure code A4483)

Note: A replacement ventilator is a replacement for the primary ventilator due to mechanical or other malfunction and is considered a part of the rental agreement and is not authorized separately from the primary ventilator request.

Oxygen rental is not considered a ventilator supply and may be considered for separate prior authorization.

Refer to: Subsection 2.2.23.12, “Oxygen Therapy” in this handbook for more information.

Heated or non-heated humidification requires a separate prior authorization for rental or purchase for client owned ventilators. Documentation submitted must support why it is medically necessary for the use with the ventilation.

Refer to: Subsection 2.2.23.4, “Humidification Therapy and Heating Elements” in this handbook for more information.

When prescribed by a treating physician and medically necessary for clients who require mechanical ventilation, the following devices will be prior authorized for rental:

• A home ventilator that uses an invasive interface, such as a tracheostomy tube (procedure code E0465)
• A home ventilator that uses a non-invasive interface, such as a mask or chest shell (procedure code E0466)
• A multifunction respiratory device (procedure code E0467)

Rental of a chest shell (cuirass or “clam shell”) (procedure code E0457) and chest wrap (procedure code E0459) for use with a mechanical ventilator is limited to up to three months.

Purchase of a chest shell (cuirass or “clam shell”) may be prior authorized for purchase following the initial three-month rental period of the mechanical ventilator. The prescribing physician must include the predicted length of treatment and the client’s compliance with the equipment.

2.2.23.10 Back-Up Mechanical Ventilator

A back-up ventilator is defined as an identical or similar device used to meet the same medical needs as the primary ventilator for the client, but provided in the home as a precaution in case of malfunction of the primary ventilator.

Requests for a back-up ventilator in the home must meet both the following criteria to be considered medically necessary:

• The client cannot maintain spontaneous ventilations for four or more consecutive hours.
• The client lives in an area where a replacement ventilator cannot be provided within two hours.

Prior authorization requests for a back-up ventilator must include all of the following documentation:

• The amount of time the client is able to maintain spontaneous ventilation.
• The distance and/or delivery time of a replacement ventilator to the clients home.

All requests for a back-up ventilator in the home must be sent to the TMHP for Medical Director for review.
2.2.23.11 Secretion and Mucus Clearance Devices

Secretion and mucus clearance devices are a benefit when medically necessary, and are typically needed by clients diagnosed with cystic fibrosis (CF), chronic bronchitis, bronchiectasis, and ciliary dyskinesia syndromes, some forms of asthma, neuromuscular degenerative disorders, post-operative atelectasis, or thoracic wall defects.

Secretion and mucus clearance devices may be considered when documentation clearly shows the client has one of the following indications for this form of therapy as described by the AARC in the Clinical Practices Guidelines for Postural Drainage Therapy:

- Evidence of retained secretions
- Evidence that the client is having difficulty with the secretion clearance
- Presence of atelectasis caused by mucus plugging

The following secretion and mucus clearance devices or procedures do not require prior authorization:

- Incentive spirometers (procedure code A9284)
- Mucous clearance valved chamber (oscillating positive expiratory pressure (PEP), such as the Flutter Valve) (procedure code S8185)
- Moisture exchangers (procedure code A4483) only when used for mechanically ventilated clients who own their ventilator
- Tracheostoma filters, such as Thermovent T (procedure code A4481) for clients with a tracheotomy who are not mechanically ventilated

The following secretion and mucus clearance devices require prior authorization:

- Insufflation-exsufflation devices (e.g. Cough Assist, Cofflator) (procedure code E0482)
- Electrical percussors (procedure code E0480)
- The high-frequency chest wall oscillation (HFCWO) system (procedure code E0483)
- Percussion cup (procedure code E1399)
- Intermittent positive pressure breathing (IPPB) devices (procedure code E0500)

Procedure codes A7025 and A7026 will deny if billed in the same month by any provider as rental of procedure code E0483.

2.2.23.11.1 Prior Authorization

Prior authorization requests for the rental or purchase of secretion and mucus clearance devices requires submission of a Texas Medicaid Prior Authorization Request for Secretion and Mucus Clearance Devices - Initial Request form or a Texas Medicaid Prior Authorization Request for Secretion and Mucus Clearance Devices -Renewal Request form completed, signed, and dated by the client’s treating physician unless otherwise specified.

Clients requiring more than one secretion and mucus clearance device must have a pulmonologist as the treating physician who submits a signed and dated letter of medical necessity (LMN) on the physician’s letterhead stating the need of two devices.

Percussion Cups

Percussion cups, used when performing chest physiotherapy, may be medically necessary to loosen thick, mucus secretions, assist respiration, and prevent infections and require prior authorization. Percussion cups should be requested using the miscellaneous DME procedure code E1399.
**Electrical Percussor**

An electrical percussor (procedure code E0480) may be considered for rental or purchase with documentation of medical necessity including a description of all previous courses of therapy (such as manual percussion and postural drainage (P&PD) or valved devices) and why they did not adequately assist the client in airway mucus clearance.

**Cough Augmentation Device (e.g., mechanical insufflation-exsufflation or cough assist machine)**

A cough augmentation device (procedure code E0482) may be considered for prior authorization for rental only for those clients who have chronic pulmonary disease or neuromuscular disorders (including spinal cord injury) that affect the respiratory musculature, causing a weak, ineffectual, or absent cough.

Prior authorization of a cough augmentation device may be considered for an initial three-month rental period with all of the following documentation completed, signed, and dated by the client’s treating physician:

- Diagnosis and background history including, as applicable, recent illnesses, complications, medications used, history of recent hospitalizations, and results of pulmonary function studies (if applicable) due to diagnosis related complications
- History of school, work, or extracurricular activity absences or other clinical evidence supporting natural deterioration to the level of requiring the use of a cough augmentation device to clear the airways, such as a weak, ineffective cough as demonstrated by pulmonary function studies (PFTs)
- Medical reasons why the client, parent, guardian, or caregiver cannot perform chest physiotherapy, or why such therapies were previously not effective

Requests for prior authorization recertification must include documentation by the client’s treating physician that the client is compliant with the use of the equipment and that the treatment is effective.

**High-Frequency Chest Wall Oscillation (HFCWO) System**

A high-frequency chest wall oscillation (HFCWO) system (procedure code E0483) will not be prior authorized as first line treatment. The client must have trialed other percussion and postural drainage therapy, such as electric percussor or cough augmentation device, for a minimum of three months before a request for a HFCWO system will be considered for prior authorization.

A request for a HFCWO system may be considered for prior authorization for rental or purchase when submitted with documentation addressing why prior therapy was ineffective and documentation of one of the following conditions.

- Bronchiectasis when it is confirmed by CT scan and characterized by either a continuous daily productive cough for 6 months or frequent exacerbations of pulmonary infections (i.e., more than 2 times per year) requiring antibiotic therapy
- Cystic fibrosis or other documented chronic suppurative endobronchitis
- Chronic neuromuscular disorder affecting the client’s ability to cough or clear respiratory secretions
- Weak ineffective or absent cough caused by chronic pulmonary disease or a neuromuscular disorder

**Rental of the HFCWO System**

An initial three-month rental may be prior authorized for the HFCWO system with hose and vest (procedure code E0483) when submitted with the following documentation of medical necessity that is completed, signed, and dated by the client’s treating physician:

- A Texas Medicaid Prior Authorization Request for Secretion and Mucus Clearance Devices form (Initial or Renewal Request) form
• Client history of a chronic respiratory illness with exacerbation or change in baseline respiratory condition in the past 6 months, including extra nebulizer treatments for respiratory secretions, I.V. antibiotics, and hospitalizations

• Client history of school, work, or extracurricular activity absences due to diagnosis related symptoms, or pulmonary function testing in (PFTs) in past 6 months, if applicable

• Other appropriate (age, ability, skill) modes of chest physiotherapy (such as percussion and postural drainage therapy or mechanical device) that have been trialed by the client, parent, guardian, or caregiver for a minimum of three months before the HFCWO request and the reasons the trialed therapy was ineffective or contraindicated

• Documentation that any previous use of an HFCWO device did not result in aspiration, exacerbation of a gastrointestinal or pulmonary issue, or an exacerbation of seizure activity

If at the end of the initial three-month rental a determination of purchase cannot be made, an additional three month-rental may be considered for prior authorization when the request is submitted with the above documentation and documentation of compliance with ordered therapy.

**Purchase of the HFCWO System**

If at the end of the initial three-month rental, the HFCWO system (procedure code E0483) is documented to be effective, purchase of the system may be considered for prior authorization when submitted with all the following required documentation:

• A physician’s statement of the HFCWO system trial in a clinic, hospital, or the home setting documenting:
  - The results of the HFCWO system therapy
  - The effectiveness and tolerance of the system that includes evidence of vest tolerance
  - An explanation of the trial outcome

• The treating physician’s description and assessment of the effectiveness and tolerance of the system that includes the client’s diagnosis and the following background history:
  - Respiratory related complications and evidence of a decrease in these complications
  - Medications used, including IV antibiotic therapy with dosage, frequency and duration, including evidence of decreased respiratory-related medication use
  - Recent hospitalizations related to the client’s respiratory condition and evidence of shorter hospital length(s) of stay
  - Evidence of decreased hospitalizations
  - Evidence of fewer school, work, or extracurricular activity absences due to a diagnostic related condition or other clinical evidence supporting natural deterioration to the level of requiring the use of a HFCWO system to clear the airways, such as a weak, ineffective cough as demonstrated by pulmonary function studies (PFTs)
  - Evidence of the frequency and compliance graphs for the 3-month period showing the frequency prescribed by the physician for each day and use of the system at least 50 percent of the time

• A statement from the treating physician that the previous use of the HFCWO device has not resulted in aspiration, exacerbation of a gastrointestinal or pulmonary issue, or exacerbation of seizure activity.

A HFCWO system purchase will be reimbursed only once per lifetime, due to the lifetime warranty provided by the manufacturer. Requests for a vest replacement (procedure code A7025) must include documentation that supports the client can no longer wear the vest due to changes in the client’s condition such as changes in height, weight, or skin abrasions.
2.2.23.12 Oxygen Therapy

Devices used for in-home oxygen therapy including stationary oxygen concentrators (procedure code E1390), compressed gas (procedure code E0424), liquid oxygen (procedure code E0439), portable compressed gas cylinder (procedure code E0431), or liquid oxygen reservoir (procedure code E0433) systems are a benefit when medically necessary and require prior authorization.

Oxygen system rental includes, but is not be limited to:

- Oxygen concentrator or oxygen tanks
- Regulator
- Flow meter
- Humidifier
- Cannula or mask
- Tubing

Oxygen system supplies, including but not limited to a cannula or mask, refills, and tubing do not require prior authorization for client owned equipment.

Devices used for in-home oxygen therapy may be considered for the treatment of hypoxemia which may be the result of, but not limited to:

- Bronchopulmonary dysplasia or other respiratory diagnoses due to prematurity
- Respiratory failure or insufficiency; musculoskeletal weakness, such as that caused by Duchenne’s muscular dystrophy or spinal muscle atrophy
- Diagnosis of cluster headaches
- Severe lung disease, such as COPD, diffuse interstitial lung disease, cystic fibrosis, bronchiectasis, or widespread pulmonary neoplasm

Humidification during oxygen therapy may be a benefit with prior authorization and is provided as a component of the oxygen therapy rental. Humidification during oxygen delivery with client-owned equipment may be a benefit for rental or purchase with prior authorization when medically necessary.

The following procedure codes will deny if billed in the same month by any provider as procedure codes E0424, E0431, E0433, E0434, E0439, E1390, and K0738:

| Procedure Codes |
|-----------------|-----------------|-----------------|-----------------|-----------------|-----------------|-----------------|-----------------|-----------------|-----------------|
| A4615 | A4616 | A4617 | A4619 | A4620 | E0441 | E0442 | E0443 | E0444 | E0447 |
| E0580 | E1353 | E1355 |

2.2.23.12.1 Prior Authorization

Oxygen therapy criteria are based on AARC, American Thoracic Society, and British Thoracic Society Treatment Guidelines. Oxygen therapy related supplies, other than humidification, do not require prior authorization for client owned equipment. Humidification during oxygen delivery with client-owned equipment may be a benefit for rental or purchase with prior authorization when medically necessary.

All oxygen therapy equipment requires prior authorization and prior authorization may be considered for monthly rental only. A completed Texas Medicaid Prior Authorization Request for Oxygen Therapy Devices and Supplies form completed, signed, and dated by the client’s treating physician must be submitted.

Prescribing providers must maintain an original, completed, signed, and dated Texas Medicaid Prior Authorization Request for Oxygen Therapy Devices and Supplies form in the client’s medical record. The DME provider must maintain a copy of the complete, signed, and dated in the client’s record.
Stationary oxygen concentrators are the preferred oxygen therapy home delivery system. If other types of oxygen therapy home delivery systems are required, documentation of medical necessity to support an exception, such as frequent interruptions in electrical service or medical necessity for a higher oxygen concentration than can be obtained with a concentrator, must be provided. The other types of delivery systems include:

- Compressed gas cylinder systems (nonportable tanks).
- Liquid oxygen reservoir systems.

Multiple oxygen types (e.g., liquid and gas) will not be authorized concurrently.

**Initial Oxygen Therapy Medical Necessity Certification**

Oxygen ordered for ‘as needed’ or ‘PRN’ use does not provide a basis to determine if intermittent oxygen is reasonable and medically necessary for the client. Documentation must support the need for intermittent use of oxygen.

Prior authorization of home oxygen therapy for an initial three month rental period may be considered with submission of all of the following documentation in addition to the request:

- Evidence from the client’s treating physician of a determination that the client has severe lung disease or hypoxia-related symptoms that are expected to improve with oxygen therapy and the client’s blood gas studies meet the criteria indicated below
- A Texas Medicaid Prior Authorization Request for Oxygen Therapy Devices and Supplies form with Sections A and B that is completed, signed, and dated by the client’s treating physician documenting the client’s specific medical needs and the testing performed to determine the need for oxygen therapy including all of the following criteria:
  - The client’s medical diagnosis requiring oxygen therapy
  - The oxygen flow rate
  - An estimate of the frequency, duration of use (e.g. 2 liters per minute, 12 hours per day), and duration of need (e.g., 3 months)
  - A qualifying blood gas assessment may be supported by the results of either pulse oximetry or an arterial blood gas and includes all of the following:
    - Date of the testing
    - Results of the testing
    - If the blood gas assessment occurred during the client’s inpatient hospital stay, a blood gas performed no more than two days before discharge is acceptable.
    - If a blood gas is obtained while the client is at home, the assessment must be performed while the client is in a stable chronic state (i.e., not during a period of acute illness or an exacerbation of their underlying disease) within the 30 day period prior to the request for service.

Oxygen therapy coverage is available under one of the three group categories outlined below, if the client has an eligible condition as described above.

**Group I Oxygen Therapy Category**

Group I - Prior authorization may be considered for clients of any age with significant hypoxemia with documentation of any of the following:

- An arterial pO2 (partial pressure of oxygen) equal to or less than 55 mm Hg or an arterial oxygen saturation equal to or less than 88 percent, taken at rest, breathing room air
• An arterial pO2 equal to or less than 55 mm Hg or arterial oxygen saturation at or below 88 percent, taken during sleep and lasting for at least 5 continuous minutes for clients who have a pO2 at or above 56 mm Hg or an arterial oxygen saturation at or above 89 percent while awake

• A decrease in arterial pO2 more than 10 mm Hg, or a decrease in arterial oxygen saturation of more than 5 percent, for at least 5 continuous minutes during sleep with symptoms or signs reasonably attributable to hypoxemia (e.g., impairment of cognitive processes and nocturnal restlessness or insomnia)

• An arterial pO2 equal to or less than 55 mm Hg or an arterial oxygen saturation at or below 88 percent, taken during exercise for a member who demonstrates a pO2 at or above 56 mm Hg or an arterial oxygen saturation at or above 89 percent during the day while at rest. In this case, supplemental oxygen may be provided for use during exercise if there is evidence the use of oxygen improves the hypoxemia that was demonstrated during exercise when the client was breathing room air

**Group I Oxygen Therapy for Clients who are 20 Years of Age and Younger**

Prior authorization may be considered for clients 20 years of age and younger when evidenced by any of the above or the following documentation:

• A neonate, and premature infant of any age who have not reached their 40th week of gestational maturity with an arterial pO2 of less than 60 mmHg or an arterial oxygen saturation level is less than 92 percent

• An infant with chronic neonatal lung disease with an arterial oxygen saturation equal to or less than 92 percent

• Other medical conditions that may be considered with supporting documentation include, but are not limited to:
  • Infants with bronchopulmonary dysplasia
  • Infants with apnea of prematurity, or recurrent cyanotic apneic episodes
  • Children with severe pulmonary hypotension
  • Children who have sickle cell anemia with respiratory conditions
  • Infants or children who have idiopathic pulmonary hypertension with sleep associated desaturations or a documented need for an emergent use of oxygen

<table>
<thead>
<tr>
<th>Oxygen Saturation</th>
<th>pO2 in mm HG</th>
<th>Required Parameters</th>
<th>Clients/Comments</th>
</tr>
</thead>
<tbody>
<tr>
<td>Less than or equal to 88 percent</td>
<td>Less than or equal to 55</td>
<td>At rest on room air</td>
<td>Infants, children and adults</td>
</tr>
<tr>
<td>Less than or equal to 88 percent</td>
<td>Less than or equal to 55</td>
<td>For greater than or equal to 5 minutes during sleep, when “at rest” criteria not met</td>
<td>Infants, children and adults</td>
</tr>
<tr>
<td>Less than or equal to 88 percent</td>
<td>Less than or equal to 55</td>
<td>During exercise, when criteria for “at rest” is not met</td>
<td>Infants, children and adults</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Tests provided must document the following results:</td>
<td>If only criteria - documentation from physician: client’s hypoxemia improved with use of O2 with exercise</td>
</tr>
<tr>
<td></td>
<td></td>
<td>At rest on room air</td>
<td></td>
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<tr>
<td></td>
<td></td>
<td>Exercising without oxygen</td>
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<tr>
<td></td>
<td></td>
<td>Exercising with oxygen</td>
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</tbody>
</table>
Intermittent acute oxygen therapy at home is not routinely recommended for seizures as there is no evidence that it reduces seizure duration, reduces harm from prolonged seizures, or improves quality of life for the child or family.

**Group II Oxygen Therapy Category**
Group II-Prior authorization may be considered for clients of any age whose arterial pO2 is 56-59 mm Hg or whose arterial blood oxygen saturation is 89 percent with documentation of any of the following:

- Dependent edema suggesting congestive heart failure (CHF)
- Cor pulmonale (pulmonary hypertension)
- Erythrocythemia with a hematocrit greater than 56 percent

**Group III Oxygen Therapy Category**
Group III-Prior authorization may be considered for clients with a diagnosis of cluster headaches with documentation of all of the following:

- Neurological evaluation with diagnosis of cluster headache
- Documentation of failed medication therapy

For clients whose only diagnosis is OSA, documentation must support the client’s oxygen sleep desaturation was not corrected with use of CPAP or other RADs.

**Oxygen Therapy Recertification**
Prior authorization of oxygen therapy rental after an initial three-month rental period may be considered for periods of six month at a time with the submission of all of the following documentation:

- A new Texas Medicaid Prior Authorization Request for Oxygen Therapy Devices and Supplies form with Sections A and B completed, signed, and dated by the client’s treating physician
- Evidence of a continued need for oxygen therapy
- Evidence from the client’s treating physician of the client’s compliance with the oxygen therapy

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<th>Required Parameters</th>
<th>Clients/Comments</th>
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</thead>
<tbody>
<tr>
<td>Decrease of greater than 5 percent</td>
<td>Decrease of greater than 10 mm Hg</td>
<td>For greater than 5 minutes taken during sleep and Client has signs or symptoms attributable to hypoxemia</td>
<td>Infants, children and adults</td>
</tr>
<tr>
<td>Less than 92 percent</td>
<td>Less than 60</td>
<td>At rest on room air</td>
<td>Neonates, premature infants who are &lt; than their 40 week gestational maturity</td>
</tr>
<tr>
<td>Less than or equal to 92 percent</td>
<td>At rest on room air Chronic neonatal lung disease</td>
<td>Infants and children</td>
<td></td>
</tr>
<tr>
<td>Greater than or equal to 89 percent</td>
<td>Greater than or equal to 56</td>
<td>Documentation from physician supporting the medical need for the oxygen therapy due to medical conditions requiring different parameters</td>
<td>Clients of any age</td>
</tr>
</tbody>
</table>

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<td>Documentation from physician supporting the medical need for the oxygen therapy due to medical conditions requiring different parameters</td>
<td>Clients of any age</td>
</tr>
</tbody>
</table>
• A new arterial blood gas assessment using either pulse oximetry or arterial blood gases
• Evidence that the client meets the criteria for any of the above Groups (I, II, or III) for oxygen therapy

If the above criteria for oxygen therapy are not met medical necessity for recertification of oxygen therapy will be considered by a medical director with documentation from the treating physician supporting the client’s need for oxygen therapy.

**Stationary Oxygen Systems**

Rental of a stationary oxygen system includes, but is not limited to, the nasal cannula or mask, tubing, and a basic bubble humidification system.

The types of covered stationary oxygen delivery systems include:

- Oxygen concentrators
- Compressed oxygen gas cylinder systems
- Liquid oxygen cylinder systems

**Portable Oxygen Systems**

Portable oxygen therapy may be considered for prior authorization when medical necessity documentation indicates that the client requires the use of oxygen in the home and would benefit from the use of a portable oxygen system when traveling outside of the home environment.

Portable oxygen systems will not be considered for prior authorization for travel outside of the home environment for clients who qualify for oxygen usage based solely on oxygen saturation levels during sleep.

The types of covered portable oxygen and portable oxygen related delivery systems include:

- Portable tanks for compressed oxygen gas cylinder systems
- Portable tanks for liquid oxygen cylinder systems
- Home compressor attachment used on an oxygen compressor to fill oxygen tanks
- Portable gaseous oxygen system home compressor

**2.2.23.13 Cardiorespiratory Monitor (CRM)**

A cardiorespiratory monitor (CRM) (procedure codes E0618 and E0619) for purchase or rental may be a benefit when medically necessary and may be considered for clients who require moment-to-moment cardiac and respiratory monitoring due to potential for sudden unexpected deterioration. For infants who are four months of age and younger, a CRM with recording feature (procedure code E0619) may be considered for rental without prior authorization for a maximum of two months. Prior authorization is required for clients who are 5 months of age and older.

*Note:* The American Academy of Pediatrics (AAP) recommends that infant monitoring using an infant CRM should not be used as a strategy to prevent sudden infant death syndrome (SIDS).

Procedure codes A4556 and A4557 will deny if billed in the same month by any provider as procedure codes E0618 and E0619.

**2.2.23.13.1 Prior Authorization**

For infants who are four months of age and younger, a CRM with recording feature (procedure code E0619) may be considered for rental without prior authorization for a maximum of two months with documentation of one of the following conditions:

- Central apnea (respiratory control disorders)
- Cardiac rhythm issues
A completed Title XIX Form signed and dated by the client’s treating physician must be maintained in the client’s medical record.

If a two-month rental has expired for infants who are 4 months of age and younger and continued CRM is medically necessary, a submitted prior authorization request on a completed Title XIX Form signed and dated by the client’s treating physician (new with each request) must include the following:

- The client has on-going, documented cardiorespiratory episodes (e.g. apnea or dysrhythmia)
- A physician interpretation, signed and dated by the physician, of the most recent two-month’s CRM data recorded downloads

A CRM with or without recording feature (procedure code E0618 or E0619) may be considered for prior authorization for rental or purchase for clients who are 5 months of age and older when submitted with a completed Title XIX Form signed and dated by the client’s treating physician (new with each request) for one of the following conditions:

- An episode of apparent life-threatening event (ALTE) in an infant who is 12 months of age or younger
- Symptomatic central apnea
- Technology dependence such as:
  - Mechanical ventilation
  - Tracheostomy with a critical airway obstruction
  - Assisted ventilation dependence
  - Cardiac dysrhythmia with significant risk of morbidity or mortality

### 2.2.23.14 Tracheostomy Tubes and Related Supplies

Tracheostomy tubes (procedure codes A7520, A7521, and A7522) are medically necessary for clients with a tracheostomy and are available for purchase with prior authorization.

Tracheostomy supplies, including inner cannulas, are available for purchase when medically necessary without prior authorization within the stated benefit limits.

A tracheostomy speaking valve is considered a medically necessary accessory that enhances the function of the tracheostomy and is available for purchase without prior authorization when requested within the stated benefit limits.

A tracheostomy speaking valve (procedure code L8501) is available for purchase and is limited to one per six months without prior authorization.

### 2.2.23.14.1 Prior Authorization

For the initial tracheostomy tube request, three tubes may be considered for prior authorization in the first month of service (two the same size and one smaller for emergencies).

For the next five months of the initial prior authorization period and for subsequent requests, one tracheostomy tube will be prior authorized per month.

More than one tracheostomy tube per month may be considered on a case-by-case basis with medical documentation supporting why the tracheostomy tube must be changed more frequently in order to meet the client’s medical needs.
Requesting tracheostomy supplies above the defined limitation

Tracheostomy supplies requests for clients 20 years of age and younger that exceed the defined limits require prior authorization with documentation supporting the medical need of the quantity requested and may be considered under the Comprehensive Care Program and must be requested on a CCP Prior Authorization Request Form.

Tracheostomy supplies requests for clients who are 21 years of age and older that exceed the defined limits require prior authorization with documentation supporting the medical need of the quantity requested and may be considered by a medical director with documentation of medical necessity and must be requested on a Title XIX Form.

Modifiers for Tracheostomy Tubes

When requesting prior authorization for non-customized or non-specialized tracheostomy tubes without specialized functions, providers must submit the most appropriate procedure code with no modifier.

When requesting prior authorization for specialized but non-customized tracheostomy tubes with specialized functions, providers submit the request with modifier U1.

When requesting prior authorization for customized tracheostomy tubes, providers must submit the request with modifier U2.

With the use of either modifier U1 or U2, the following documentation is required:

- A physician statement of the reason the client cannot use a standard tracheostomy tube
- The manufacturer’s information on the specialized functions of the tracheostomy tube or the order form describing the customization of the tracheostomy tube

Manufacturer’s retail or invoice pricing information is required when using modifier U2.

Tracheostomy Tube Inner Cannula and Required Modifier

Clients with a tracheostomy tube with a reusable inner cannula (procedure code A4623) are allowed one reusable inner cannula per month without prior authorization.

Requests for more than one reusable inner cannula per month require prior authorization and medical documentation from the client’s physician to support the need for more than one reusable inner cannula per month.

Clients with a tracheostomy tube with a disposable inner cannula (procedure code A4623 with modifier U3) are allowed 31 disposable inner cannulas per month without prior authorization. Request for more than 31 disposable inner cannulas per month require prior authorization and documentation from the client’s treating physician to support the medical need for more than 31 disposable inner cannulas per month.

Custom tracheostomy tubes are manufactured with reusable inner cannulas. The reusable inner cannulas are included in the prior authorization for any custom tracheostomy tube authorized.

2.2.23.15 Suction Machines and Related Supplies

A suction machine (procedure code E0600) may be considered for purchase with prior authorization if medically necessary for clients who have difficulty raising and clearing secretions. Suction supplies (suction canisters, suction tubing, tracheal suction catheters, and oropharyngeal suction catheters) (procedure codes A4605, A4624, A4628, A7000, and A7002) are medically necessary for use with a suction machine. These supplies are available, if medically necessary, for purchase without prior authorization, up to the stated benefit limits unless otherwise indicated. Only one type of tracheal suction catheter is allowed per month.
In most cases in the home setting, sterile suction catheters (procedure codes A4605 and A4624) and sterile saline for suctioning (procedure code A4216) are considered medically necessary only for tracheostomy suctioning. Sterile saline for tracheal suctioning (procedure code A4216) does not require prior authorization when requested within the stated benefit limits.

Procedure code A4605 will deny if billed in the same month by any provider as procedure code A4624.

2.2.23.15.1 Prior Authorization

Suction machines, suction canisters, suction tubing, tracheal suction tubes, and oropharyngeal suction catheters are a benefit with documented medical need to have oral, nasopharyngeal, or tracheal suctioning performed. Suction supplies do not require prior authorization within the stated benefit limits.

Suction canister filters are limited to one every two months with prior authorization and should be requested using the miscellaneous DME procedure code A9900. Suction equipment and supplies requests for clients who are 20 years of age and younger that exceed the allowed limits require prior authorization with documentation supporting the medical need of the quantity requested, may be considered under Comprehensive Care Program, and must be requested on a CCP Prior Authorization Request Form.

Suction equipment and supplies requests for clients who are 21 years of age and older that exceed the allowed limits require prior authorization with documentation supporting the medical need of the quantity requested and may be considered by a medical director with documentation of medical necessity supporting the medical need of the quantity requested and must be requested on a Title XIX Form.

2.2.23.16 Other Respiratory Supplies

Other respiratory supplies are a benefit when medically necessary and are available without prior authorization up to the stated quantity limitation unless otherwise indicated:

- Non-sterile (clean) respiratory supplies are considered standard of care and are clinically appropriate in the home setting
- Sterile respiratory supplies are a benefit with prior authorization when medically necessary and documentation clearly demonstrates that the client’s medical needs cannot be met with non-sterile (clean) supplies

2.2.23.16.1 Prior Authorization

Respiratory supplies are included in the rental of the respiratory equipment and will not be prior authorized separately, but may be prior authorized for equipment that is owned by the client.

To request prior authorization for respiratory supplies for use with client-owned equipment, the following documentation must be provided by the client’s treating physician:

- A completed Title XIX Form signed and dated by the client’s treating physician (new with each request)
- Accurate diagnostic information pertaining to the underlying diagnosis or condition as well as any other medical diagnoses or conditions, to include the client’s overall health status
- The prescribed respiratory care regimen, to include frequency, duration, and supplies needed
- Treatment for infection, if present
- Quantity of disposable supplies prescribed per month by the physician
- Medical justification for the quantity and type of supply requested
2.2.23.17 Bag Valve Mask (BVM) Resuscitator

A bag valve mask (BVM) resuscitator or handheld resuscitation bag for a ventilator-dependent client (procedure code S8999) is commonly used when the client is temporarily removed from mechanical ventilation. A BVM for a ventilator dependent client may be a benefit for purchase when medically necessary without prior authorization.

A BVM for a non-ventilator client with or without a tracheotomy and with an appropriately sized face mask when appropriate who requires manual respiratory assistance due to inadequate or no breathing may be a benefit when medically necessary and requires prior authorization using the miscellaneous DME procedure code E1399 and will be reviewed by a medical director.

2.2.23.18 Short Term Pulse Oximetry Services, Benefits, and Limitations

A pulse oximeter (procedure code E0445) may be rented for short term use for no more than one calendar month in a six calendar month period. The short term rental of a pulse oximeter may be a benefit when medically necessary and does not require prior authorization for clients with one of the following conditions:

- Weaning the client from home oxygen
- Change in the client’s condition that requires an adjustment in the liter flow of their home oxygen treatment
- To determine the client’s appropriate home oxygen liter flow for ambulation, exercise, or sleep
- To determine the client’s appropriate home oxygen liter flow for those who have neuromuscular disease involving the respiratory muscles, with chronic lung disease, or with severe cardiopulmonary disease.

2.2.23.19 Long Term Pulse Oximetry Services, Benefits, and Limitations

Long term pulse oximetry (procedure code E0445 with modifier U4) may be a benefit of Texas Medicaid through the Comprehensive Care Program (CCP) with prior authorization. Long-term rental and purchase of a pulse oximeter is a benefit for clients who are birth through 20 years of age. A pulse oximeter for long-term use is defined as equipment rented for more than one calendar month in a six-month period. Long-term pulse oximetry is limited to clients eligible for CCP who qualify for medically necessary services beyond the limits of the home health benefit.

A long-term pulse oximeter must be a device that:

- Is a bedside or tabletop device.
- Provides continuous oxygen saturation monitoring.
- Requires use of wired probes.
- Has battery and alternating current (A/C) capability.

2.2.23.19.1 Long Term Pulse Oximeter Prior Authorization

Prior Authorization for Clients 21 Years of Age and Older

Pulse oximetry that is medically necessary for more than one calendar month in a six calendar month period requires prior authorization and will be considered for clients 21 years of age and older by a medical director when submitted with a complete Title XIX form, signed and dated by the treating physician and with documentation of medical necessity, such as:

- When weaning from a ventilator or oxygen and an earlier weaning attempt was unsuccessful that includes why it was unsuccessful
- Documentation of changes in the client’s condition since the failed weaning attempt
Prior Authorization for Clients Birth through 20 Years of Age

Prior authorization is required for rental or purchase of a long-term pulse oximeter (procedure code E0445 with modifier U4) for clients who are birth through 20 years of age. HHSC or its designee will determine whether the long-term pulse oximeter equipment will be rented, purchased, or repaired based on the client’s needs and expected duration of use. Only new, unused equipment will be purchased.

The prior authorization request must include the following documentation:

- A completed CCP Prior Authorization Request form
- Documentation of the cause of the oxygen lability
- Documentation that a caregiver or medical healthcare provider is present who has been trained in use of the oximeter and how to respond to readings in a medically safe and appropriate manner, and the client meets one of the following criteria:
  - Client is oxygen or ventilator dependent and has respiratory system instability as noted in the documentation
  - Client experiences respiratory complications that require equipment that has oxygen saturation monitoring capabilities

A long-term pulse oximeter may be prior authorized for monthly rental up to a maximum of six months. Recertification for an additional six-month period may be considered for a maximum of six additional months.

A long-term pulse oximeter will be considered purchased and owned by the client when the total monthly payments equal the purchase cost for the equipment. Prior authorization for purchase or continued rental of the long-term pulse oximeter at the end of a 12-month rental period will not be considered. A long-term pulse oximeter may be prior authorized for purchase when a purchase is determined to be more cost effective than leasing the device with supplies.

Long-term pulse oximeter equipment that has been purchased is anticipated to last a minimum of five years. Replacement of equipment may also be considered for prior authorization when loss or irreparable damage has occurred outside the warranty terms, conditions, and limitations. A copy of the police or fire report, when appropriate, and the measures to be taken to prevent reoccurrence must be submitted with prior authorization request.

A long-term pulse oximeter used to monitor a client with a history of seizure activity is not routinely recommended for seizures as there is no evidence that it reduces seizure duration, reduces harm from prolonged seizures, or improves quality of life for the child or family.

Pulse Oximetry Supplies

Pulse oximeter probes for client-owned equipment do not require prior authorization and are limited as follows:

- Disposable pulse oximeter probes (procedure code A4606) are limited to four per month
- Reusable pulse oximeter sensor probes (procedure code A4606 with modifier U5) are limited to one every six months

Procedure code A4606 and procedure code A4606 with modifier U5 will deny if billed in the same month by any provider as procedure code E0445.

Prior authorization may be considered for quantities greater than four per month with documentation supporting medical necessity.
Pulse oximeter probes (procedure code A4606 [disposable] or A4606 [reusable] with modifier U5) are included in the pulse oximeter equipment rental. Pulse oximeter probes will be denied if billed with rental of pulse oximeter equipment (procedure code E0445 or E0445 with modifier U4) in the same month of service by any provider.

The rental of equipment includes all necessary supplies, adjustments, repairs, and replacement parts.

### 2.2.23.20 Procedure Codes and Limitations for Respiratory Equipment and Supplies

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<tr>
<th>Procedure Code</th>
<th>Maximum Limitations</th>
<th>Requires Prior Authorization</th>
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<td>A4483</td>
<td>15 per calendar month</td>
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<tr>
<td>A4556</td>
<td>15 per calendar month</td>
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<td>A4557</td>
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<tr>
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<td>10 per calendar month</td>
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<td>A4623</td>
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| A4624          | 85 per calendar month  
Only 1 type of tracheal catheter is allowed per month| No |
<p>| A4627          | 1 per 6 calendar months| No |
| A4628          | 2 per calendar month| No |
| A4629          | 30 per calendar month| No |
| A7000          | 10 per calendar month| No |
| A7002          | 8 per calendar month| No |
| A7003          | 2 per calendar month| No |
| A7004          | 2 per calendar month| No |
| A7005          | 1 per 6 calendar months| No |
| A7006          | 1 per calendar month| No |
| A7007          | 2 per calendar month| No |
| A7009          | 1 per calendar year| No |
| A7010          | 1 per 2 calendar months| No |
| A7012          | 2 per calendar month| No |
| A7013          | 2 per calendar month| No |
| A7014          | 1 per 3 calendar months| No |
| A7015          | 1 per calendar month| No |
| A7016          | 2 per calendar year| No |
| A7017          | 1 per 3 rolling years| No |
| A7018          | 4 per calendar month| No |
| A7025          | 1 per lifetime| Yes |
| A7026          | 1 per 6 calendar months| No |
| A7027          | 1 per 3 calendar months| No |
| A7028          | 1 per calendar month| No |
| A7029          | 2 per calendar month| Included in CPAP or RAD rental, No prior authorization for client-owned equipment if within policy limit |
| A7030          | 1 per 3 calendar months| Included in CPAP or RAD rental, No prior authorization for client-owned equipment if within policy limit |
| A7031          | 1 per calendar month| Included in CPAP or RAD rental, No prior authorization for client-owned equipment if within policy limit |</p>
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### 2.2.23.21 Services that are not a Benefit

The following are not a benefit of Title XIX Home Health Services:

- Respiratory equipment or supplies requested primarily for the convenience of the caregiver
- Rental of:
  - Mucus clearance valved chamber
  - Medication small volume nebulizer
  - Intrapulmonary percussive ventilation (IPV) system
  - Ultrasonic nebulizers
  - Oxygen supplies for rented equipment
  - Intermittent or spot-check pulse oximetry
- Purchase of:

<table>
<thead>
<tr>
<th>Procedure Code</th>
<th>Maximum Limitations</th>
<th>Requires Prior Authorization</th>
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<td>S8999 (Purchase)</td>
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• Bi-level PAP with set backup respiratory rate
• Intrapulmonary percussive ventilation (IPV)
• Ventilators
• Pulse oximeter as a Home Health benefit
• Intermittent or spot-check pulse oximetry
• Long term (greater than 1 calendar month) pulse oximeter is not a benefit of Home Health services.

Note: Clients 20 years of age and younger who qualify for medically necessary services beyond the limits of the short-term pulse oximeter benefit may request those long-term pulse oximeter services through the Comprehensive Care Program (CCP).

The following items are not a benefit of Title XIX Home Health Services because they either have no federal financial participation (FFP) available, are considered environmental equipment, or are considered ineffective or detrimental and as such are not considered medically necessary:

• Cool mist humidifiers
• Room air humidifiers
• Vaporizers
• Dehumidifiers
• Air conditioners
• Back-up generators

Sterile water is not medically necessary for humidifiers in the home setting and is therefore not a benefit.

2.2.24 Special Needs Car Seats and Travel Restraints

2.2.24.1 Special Needs Car Seats

A special needs car seat may be considered for reimbursement with prior authorization for a client who has outgrown an infant car seat and is unable to travel safely in a booster seat or seat belt.

A special needs car seat must have a top tether installed. The top tether is essential for proper use of the car seat. The installer is reimbursed for the installation by the manufacturer.

Providers must use procedure code E1399 for a special needs car seat.

Car seat accessories available from the manufacturer may be considered for reimbursement with prior authorization when medically necessary for correct positioning.

A stroller base for a special needs car seat is not a benefit of Texas Medicaid.

A special needs car seat may be considered for reimbursement with prior authorization for a client who has outgrown an infant car seat and is unable to travel safely in a booster seat or seat belt. Consideration must be given to the manufacturer’s weight and height limitations and must reflect allowances for at least 12 months of growth.

The provider must maintain a statement that has been signed and dated by the client’s parent or legal guardian in the client’s medical record that states the following:

• A top tether has been installed in the vehicle in which the client will be transported, by a manufacturer-trained vendor.
• Training in the correct use of the car seat has been provided by a manufacturer-trained vendor.
• The client’s parent or legal guardian has received instruction and has demonstrated the correct use of the car seat to a manufacturer-trained vendor.

To request prior authorization for a special needs car seat or accessories, the following documentation must be provided:

• The client’s weight must be at least 40 pounds, or the client’s height must be at least 40 inches.

• Supporting documentation must include the following and must be submitted for prior authorization:
  • Accurate diagnostic information pertaining to the underlying diagnosis or condition as well as any other medical diagnoses or conditions, to include the client’s overall health status.
  • A description of the client’s postural condition specifically including head and trunk control (or lack of control) and why a booster chair or seatbelt will not meet the client’s needs (the car seat must be able to support the head if head control is poor).
  • The expected long-term need for the special needs car seat.
  • A copy of the manufacturer’s certification for the installer’s training to insert the specified car seat.

A request for a client who does not meet the criteria may be considered on a case-by-case basis on review by HHSC or its designee.

2.2.24.2 Travel Safety Restraints

Providers must use procedure code E0700 for the purchase of travel safety restraints, such as ankle and wrist belts.

A travel safety restraint and ankle or wrist belts may be considered for reimbursement through CCP without prior authorization for clients with a medical condition requiring them to be transported in either a prone or supine position. The DME provider and the prescribing physician familiar with the client must maintain documentation in the client’s medical record supporting the medical necessity of the travel safety restraint.

2.2.25 Subcutaneous Injection Ports

A subcutaneous injection port is a sterile medication delivery device through which physician-prescribed medications can be injected directly into the subcutaneous tissue using a standard syringe and needle, an injection pen, or other manual injection device. The device can be used for multiple subcutaneous injections for a period of up to 72 hours, thereby avoiding repeated needle punctures of the skin. The device cannot be used with an injection pump.

A subcutaneous injection port, such as the I-Port or Insuflon, is a benefit of Texas Medicaid as a Title XIX Home Health service with prior authorization. Claims for a subcutaneous injection port must be submitted with procedure code A4211 and modifier U4.

Texas Medicaid may reimburse the device for clients who require multiple daily injections of a physician-prescribed medication and who meet the medical necessity criteria.
The subcutaneous injection port is not a benefit of Texas Medicaid as an item of convenience or for clients who are already receiving the medication through an ambulatory infusion pump. The device is considered an item of convenience if the client does not meet the criteria for medical necessity.

**Note:** For clients who are 20 years of age or younger and do not meet criteria for coverage through Title XIX Home Health Services may be considered through the Texas Health Steps Comprehensive Care Program (THSteps-CCP).

**Note:** For clients who are 21 years of age or older, requests that do not meet the criteria for coverage through Title XIX Home Health Services may be considered under the Texas Medicaid Home Health - Durable Medical Equipment (DME) Exceptional Circumstances process.

### 2.2.25.1 Prior Authorization

Prior authorization is required for a subcutaneous injection port. Initial prior authorizations will be issued for a trial period of up to 3 months. Prior authorizations that are issued after the successful completion of the initial trial period may be issued for a period of up to 6 months. Prior authorizations for subcutaneous injection ports are limited to a quantity of 10 individual ports per month. Additional ports will be considered for prior authorization with documentation of medical necessity.

#### 2.2.25.2 Documentation Requirements

The initial request for prior authorization must include documentation that indicates the client meets the following criteria for medical necessity:

- The client has a medical condition that requires multiple (i.e., 2 or more) subcutaneous, self-administered injections on a daily basis and has a current prescription for the injectable medication. Documentation must indicate the specific medical condition that is being treated, the name of the injectable medication, and the dosage and frequency of the injections.

  **Note:** “Self-administered” includes those injections administered by the client through a subcutaneous injection or by the caregiver to the client through a subcutaneous injection.

- The client or the caregiver has been unsuccessful with the self-administration of injections using a standard needle and syringe because the client demonstrates trypanophobia (i.e., severe needle phobia), as evidenced by documented physical or psychological symptoms. Documented symptoms may include, but are not limited to, the following:

<table>
<thead>
<tr>
<th>Condition</th>
<th>Possible Exhibited Symptoms</th>
</tr>
</thead>
<tbody>
<tr>
<td>Vaso-vagal trypanophobia</td>
<td>Physical symptoms such as changes in blood pressure, syncope, sweating, nausea, pallor, and tinnitus</td>
</tr>
<tr>
<td>Associate trypanophobia</td>
<td>Psychological symptoms such as extreme anxiety, insomnia, and panic attacks</td>
</tr>
<tr>
<td>Resistive trypanophobia</td>
<td>Signs and symptoms such as combativeness, elevated heart rate, high blood pressure, and violent resistance to procedures involving needles or injections</td>
</tr>
</tbody>
</table>

The prescribing physician must include with the prior authorization request a written statement of medical necessity that identifies the client as an appropriate candidate for the subcutaneous injection port device. The physician’s statement or medical record documentation that is submitted with the prior authorization request must indicate the following:

- The client or caregiver has received instruction during an office visit on the proper placement and use of the device, with successful return demonstration. (Prior authorization requests for skilled nursing visits for the sole purpose of client instruction on the use of the subcutaneous injection port device will not be approved. Necessary instruction must be performed as part of the office visit with the prescribing physician.)

- The client has no known allergies or sensitivities to adhesives, silicone, or similar materials.
• The client has no skin infection at potential injection sites.

• The client’s most recent lab results related to the medical condition requiring treatment with daily subcutaneous injections must also be submitted with the prior authorization request. Lab results may include, but are not limited to, hemoglobin A1c (HbA1c) levels for clients with insulin dependent diabetes mellitus (IDDM) and partial thromboplastin time (PTT) for clients who are receiving anticoagulant therapy.

Requests for the renewal of the prior authorization after the initial trial period has ended must include documentation of the following:

• Ongoing signs and symptoms associated with the client’s trypanophobia.

• Improved compliance with the physician-prescribed injection regimen.

• Successful use of the device with no persistent pattern of the client’s dislodging the device during the initial trial period.

• Results of relevant lab tests performed upon completion of the initial trial period, including, but not limited to, HbA1c levels for clients with IDDM and PTT for clients who are receiving anticoagulant therapy.

  Note: For clients with IDDM, if the HbA1c level has not declined with use of the subcutaneous injection port, additional documentation must be submitted by the physician who documents the clinical determination about the lack of significant improvement in the HbA1c level. The renewal of the prior authorization will not be approved without this information.

2.2.26 Total Parenteral Nutrition (TPN) Solutions

2.2.26.1 Services, Benefits, and Limitations for Clients Birth through 20 Years of Age

In-home TPN for clients who are birth through 20 years of age may be considered through CCP. Eligible clients may receive short-term or long-term nutritional support when oral or enteral intake are unable to maintain adequate nutrition. Covered services must be medically necessary and prescribed by the physician.

Parenteral nutrition solution, supplies, and infusion pumps services may be reimbursed with the following procedure codes:

<table>
<thead>
<tr>
<th>Procedure Codes</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>Solution Procedure Codes</strong></td>
</tr>
<tr>
<td>B4164</td>
</tr>
<tr>
<td>B4197</td>
</tr>
<tr>
<td><strong>Supply Procedure Codes</strong></td>
</tr>
<tr>
<td>B4220</td>
</tr>
<tr>
<td><strong>Infusion Pump Procedure Codes</strong></td>
</tr>
<tr>
<td>B9004</td>
</tr>
</tbody>
</table>

  Note: Procedure code B4187 is a benefit for clients who are birth through 18 years of age.

2.2.26.2 Prior Authorization and Documentation Requirements for Clients Birth through age 20

Prior authorization is required for TPN solutions, lipids, supply kits, and infusion pumps that are provided through CCP for clients birth through 20 years of age. Renewal of the prior authorization will be considered on the basis of medical necessity.
TPN solutions, lipids, supply kits, and infusion pumps will be considered with prior authorization when documentation submitted clearly shows that it is medically necessary and will correct or ameliorate the client’s disability or physical or mental illness or condition. Documentation must include the following:

- Conditions that result in a loss of function of the gastrointestinal (GI) tract and the inability to obtain adequate nutrition by the enteral route, such as:
  - Infections of the pancreas, intestines, or other body organs that result in a loss of GI function
  - Inflammatory bowel disease
  - Necrotizing enterocolitis
  - Malnutrition
  - Trauma
  - Overwhelming systemic infections
  - Serious burns

- Conditions that result in an inability of the bowel to absorb nutrition, such as:
  - Extensive bowel resection
  - Severe, advanced bowel disease. Examples include short bowel syndrome (SBS), chronic intestinal pseudo-obstruction (CIPS), Hirschprungs disease (HD), Crohn’s disease, and ulcerative colitis
  - Prematurity
  - Leukemias
  - Congenital gastrointestinal anomalies
  - Acquired immunodeficiency syndrome

To facilitate determination of medical necessity and avoid unnecessary denials, the physician must provide correct and complete information, including documentation of medical necessity for the equipment and supplies requested.

Prior authorization requests for TPN must include the following information:

- Medical condition for which TPN is necessary
- Documentation of any trials with oral and enteral feedings
- Percent of daily nutritional needs from TPN
- A copy of the TPN formula or prescription that includes amino acids and lipids and is signed and dated by the physician
- A copy of the most recent laboratory results that includes potassium, calcium, liver function studies, and albumin

**Note:** Conditions or durations of need that are not listed above may be considered by HHSC or its designee with documentation of medical necessity.

The requesting provider may be asked for additional information to clarify or complete a request for TPN services.

The physician must also maintain documentation of medical necessity in the client’s medical record. Retrospective review may be performed to ensure that the documentation supports the medical necessity of the TPN services.
2.2.26.3 Services, Benefits, and Limitations for Clients 21 years of age and older

In-home TPN for clients who are 21 years of age and older may be considered through home health services. Eligible clients may receive long-term nutritional support when oral or enteral intake are unable to maintain adequate nutrition. "Long-term nutritional support" refers to treatment lasting 30 days or longer. Covered services must be medically necessary and prescribed by the physician.

Conditions that may require TPN include, but are not limited to the following:

- Bowel disease or disorder
- Cancer
- AIDS
- Coma
- Burns
- Peritonitis

Note: Conditions or a duration of need not listed above may be considered by HHSC or its designee with documentation of medical necessity.

TPN services are not a benefit when oral or enteral intake will maintain adequate nutrition.

Parenteral nutrition solution services may be reimbursed using the following procedure codes:

<table>
<thead>
<tr>
<th>Procedure Codes</th>
</tr>
</thead>
<tbody>
<tr>
<td>B4164</td>
</tr>
<tr>
<td>B4199</td>
</tr>
</tbody>
</table>

2.2.26.4 Prior Authorization and Documentation Requirements for Clients who are 21 Years of Age and Older

TPN solutions, lipids, supply kits, and infusion pumps must be prior authorized. Renewal of the prior authorization will be considered on the basis of medical necessity.

The administration of intravenous fluids and electrolytes cannot be billed as in-home TPN.

Claims for TPN must contain the 9-character prior authorization number in Block 23. Providers must consult with their vendor for the location of this field in the electronic claims format. The prescribing physician name and provider identifier must be in Block 17 and 17a or in the appropriate field of the provider's electronic software.

Requests for prior authorization must include the following information:

- Medical condition necessitating the need for TPN and long-term nutritional support.
- Documentation of any trials with oral or enteral feedings.
- Percent of daily nutritional needs from TPN.
- A copy of the TPN formula or prescription, including amino acids and lipids, signed and dated by the physician.
- A copy of the most recent laboratory results (to include potassium, calcium, liver function studies and albumin).

The requesting provider may be asked for additional information to clarify or complete a request for TPN services.
Retrospective review may be performed to ensure that the documentation supports the medical necessity of the TPN services.

### 2.2.26.5 Infusion Pumps and Supplies (All Ages)

Parenteral nutrition supplies may be reimbursed using procedure codes B4220, B4222, B4224, and B9999.

Prior authorization requests for miscellaneous procedure code B9999 must include the following:

- A detailed description of the requested item or supply.
- Documentation supporting the medical necessity for the requested item or supply.

Prior authorization requests for a portable parenteral nutrition infusion pump (procedure code B9004) must also include documentation of medical necessity demonstrating that:

- The client requires continuous feedings
- Feeding intervals exceed the time that the client must be away from home to:
  - Attend school or work.
  - Participate in extensive, physician-ordered outpatient therapies.
  - Attend frequent, multiple medical appointments.

Prior authorization for parenteral nutrition infusion pumps will be limited to one portable pump (procedure code B9004) or one stationary pump (procedure code B9006) at any one time, unless medical necessity for two infusion pumps is established. Supporting documentation for the additional pump must be included with the prior authorization request.

The infusion pump may be rented once a month or purchased once every five years.

### 2.2.26.6 Backpack or Carrying Case (All Ages)

A backpack or carrying case for a portable parenteral nutrition infusion pump may be a benefit of Home Health Services, when medically necessary and prior authorized, using procedure code B9999.

Prior authorization requests for miscellaneous procedure code B9999 must include the following:

- A detailed description of the requested item or supply.
- Documentation supporting the medical necessity for the requested item or supply.

Requests for a carrying case or backpack for the portable infusion pump will be considered for prior authorization under miscellaneous code B9999, for clients who meet the medical necessity criteria for the portable pump as outlined above. The following additional criteria apply:

- The client is ambulatory, or uses a wheelchair which will not support the use of a portable pump by other means, such as an intravenous (IV) pole.
- The portable enteral feeding pump is client-owned.

### 2.2.26.7 Reimbursement and Claims (All Ages)

No more than a one-week supply of solutions and additives may be reimbursed if the solutions and additives are shipped and not used because of the client’s loss of eligibility, change in treatment, or inpatient hospitalization. Any days that the client is an inpatient in a hospital or other medical facility or institution must be excluded from the daily billing. Payment for partial months will be prorated based upon the actual days of administration. The administration of intravenous fluids and electrolytes cannot be billed as in-home TPN.
Claims for TPN must contain the 9-character prior authorization number in Block 23. Providers must consult with their vendor for the location of this field in the electronic claims format. The prescribing physician name and provider identifier must be in Block 17 and 17a or in the appropriate field of the provider’s electronic software.

Retrospective review may be performed to ensure that the documentation supports the medical necessity of the TPN services.

2.2.27 Vitamin and Mineral Products

Vitamin and mineral products prescribed or ordered by a physician to treat various conditions are a benefit of Texas Medicaid through CCP for clients who are 20 years of age and younger. Vitamin and mineral products must be submitted with procedure code A9152 or A9153, the appropriate modifier, and the corresponding National Drug Code. Units must be based on the quantity dispensed, for up to a 30-day supply.

Note: It is acceptable for providers to bill in excess of a 30-day supply when billing for liquid formulations due to variances in container size.

For purposes of billing, one unit is equal to one dose. The total billable units are equal to the total doses requested on the prior authorization.

Providers must dispense the most cost-effective product prescribed in accordance with a prescription from a licensed physician. Organic products will not be reimbursed unless medical documentation is provided to substantiate the need for that formulation.

The following vitamin and mineral products may be a benefit when submitted with the corresponding procedure code and state-identified modifier:

<table>
<thead>
<tr>
<th>Vitamin or Mineral</th>
<th>Procedure Code</th>
<th>State-Identified Modifier</th>
</tr>
</thead>
<tbody>
<tr>
<td>Beta-carotene</td>
<td>A9152</td>
<td>U1</td>
</tr>
<tr>
<td>Vitamin A (retinol)</td>
<td>A9152</td>
<td>U1</td>
</tr>
<tr>
<td>Biotin</td>
<td>A9152</td>
<td>U2</td>
</tr>
<tr>
<td>Boric acid</td>
<td>A9152</td>
<td>U3</td>
</tr>
<tr>
<td>Copper</td>
<td>A9152</td>
<td>U3</td>
</tr>
<tr>
<td>Iodine</td>
<td>A9152</td>
<td>U3</td>
</tr>
<tr>
<td>Phosphorus</td>
<td>A9152</td>
<td>U3</td>
</tr>
<tr>
<td>Zinc</td>
<td>A9152</td>
<td>U3</td>
</tr>
<tr>
<td>Calcium</td>
<td>A9152</td>
<td>U4</td>
</tr>
<tr>
<td>Chloride</td>
<td>A9152</td>
<td>U5</td>
</tr>
<tr>
<td>Iron</td>
<td>A9152</td>
<td>U6</td>
</tr>
<tr>
<td>Magnesium</td>
<td>A9152</td>
<td>U7</td>
</tr>
<tr>
<td>Vitamin B1 (thiamin)</td>
<td>A9152</td>
<td>U8</td>
</tr>
<tr>
<td>Vitamin B2 (riboflavin)</td>
<td>A9152</td>
<td>U8</td>
</tr>
<tr>
<td>Vitamin B3 (niacin)</td>
<td>A9152</td>
<td>U8</td>
</tr>
<tr>
<td>Vitamin B5 (pantothenic acid)</td>
<td>A9152</td>
<td>U8</td>
</tr>
<tr>
<td>Vitamin B6 (pyridoxine, pyridoxal 5-phosphate)</td>
<td>A9152</td>
<td>U8</td>
</tr>
<tr>
<td>Vitamin B9 (folic acid)</td>
<td>A9152</td>
<td>U8</td>
</tr>
<tr>
<td>Vitamin B12 (cyanocobalamin)</td>
<td>A9152</td>
<td>U8</td>
</tr>
<tr>
<td>Vitamin C (ascorbic acid)</td>
<td>A9152</td>
<td>U9</td>
</tr>
</tbody>
</table>
### Vitamin or Mineral | Procedure Code | State-Identified Modifier
--- | --- | ---
Vitamin D (ergocalciferol) | A9152 | UA
Vitamin E (tocopherols) | A9152 | UB
Vitamin K (phytonadione) | A9152 | UC
Multiminerals | A9153 | U1
Multivitamins | A9153 | U2
Trace elements | A9153 | U3
Miscellaneous | A9152 or A9153 | UD

**Note:** Claims for multivitamins with any combination of additives must be submitted with modifier U2.

Vitamin and mineral products may be indicated for, but are not limited to, treatment of the following conditions:

<table>
<thead>
<tr>
<th>Vitamin or Mineral</th>
<th>Condition</th>
</tr>
</thead>
</table>
| Beta-carotene | Vitamin A deficiency  
Cystic fibrosis  
Disorders of porphyrin metabolism  
Intestinal malabsorption |
| Biotin | Biotin deficiency  
Biotinidase deficiency  
Carnitine deficiency |
| Boric acid | Recalcitrant vulva vaginitis |
| Calcium | Calcium deficiency  
Disorders of calcium metabolism  
Chronic renal disease  
Hypocalcemia and hypomagnesaemia of the newborn  
Intestinal disaccharidase deficiencies and disaccharide malabsorption  
Allergic gastroenteritis and colitis  
Hypocalcemia due to use of Depo-Provera contraceptive injection |
| Chloride | Hypochloremia  
Hypercapnia with mixed acid-base disorder  
Bronchopulmonary dysplasia |
| Copper | Disorders of copper metabolism |
| Iodine | Iodine deficiency  
Simple and unspecified goiter and nontoxic nodular goiter |
| Iron | Disorders of iron metabolism  
Iron deficiency anemia  
Siderochrestic anemia |
<table>
<thead>
<tr>
<th>Vitamin or Mineral</th>
<th>Condition</th>
</tr>
</thead>
<tbody>
<tr>
<td>Magnesium</td>
<td>Magnesium deficiency</td>
</tr>
<tr>
<td></td>
<td>Hypoparathyroidism</td>
</tr>
<tr>
<td>Phosphorus</td>
<td>Disorders of phosphorus metabolism</td>
</tr>
<tr>
<td>Vitamin A (retinol)</td>
<td>Vitamin A deficiency</td>
</tr>
<tr>
<td></td>
<td>Intestinal malabsorption</td>
</tr>
<tr>
<td></td>
<td>Disorders of the biliary tract</td>
</tr>
<tr>
<td></td>
<td>Cystic fibrosis</td>
</tr>
<tr>
<td>Vitamin B1 (thiamin)</td>
<td>Vitamin B1 deficiency</td>
</tr>
<tr>
<td></td>
<td>Disturbances of branched-chain amino-acid metabolism (e.g., maple syrup urine disease)</td>
</tr>
<tr>
<td></td>
<td>Disorders of mitochondrial metabolism</td>
</tr>
<tr>
<td></td>
<td>Wernicke-Korsakoff syndrome</td>
</tr>
<tr>
<td>Vitamin B2 (riboflavin)</td>
<td>Vitamin B2 deficiency</td>
</tr>
<tr>
<td></td>
<td>Disorders of fatty acid oxidation</td>
</tr>
<tr>
<td></td>
<td>Riboflavin deficiency, ariboflavinosis</td>
</tr>
<tr>
<td></td>
<td>Disorders of mitochondrial metabolism</td>
</tr>
<tr>
<td>Vitamin B3 (niacin)</td>
<td>Vitamin B3 deficiency</td>
</tr>
<tr>
<td></td>
<td>Disorders of lipid metabolism, (e.g., pure hypercholesterolemia)</td>
</tr>
<tr>
<td>Vitamin B5 (pantothenic acid)</td>
<td>Vitamin B5 deficiency</td>
</tr>
<tr>
<td>Vitamin B6 (pyridoxine, pyridoxal 5-phosphate)</td>
<td>Vitamin B6 deficiency</td>
</tr>
<tr>
<td></td>
<td>Sideroblastic anemia</td>
</tr>
<tr>
<td>Vitamin B9 (folic acid)</td>
<td>Vitamin B9 deficiency</td>
</tr>
<tr>
<td></td>
<td>Folate-deficiency anemia</td>
</tr>
<tr>
<td></td>
<td>Combined B12 and folate-deficiency anemia</td>
</tr>
<tr>
<td></td>
<td>Disorders of mitochondrial metabolism</td>
</tr>
<tr>
<td></td>
<td>Sickle-cell disease</td>
</tr>
<tr>
<td></td>
<td>Pernicious anemia</td>
</tr>
<tr>
<td>Vitamin B12 (cyanocobalamin)</td>
<td>Vitamin B12 deficiency</td>
</tr>
<tr>
<td></td>
<td>Disturbances of sulphur-bearing amino-acid metabolism (e.g., homocystinuria and disturbances of metabolism of methionine)</td>
</tr>
<tr>
<td></td>
<td>Pernicious anemia</td>
</tr>
<tr>
<td></td>
<td>Combined B12 and folate-deficiency anemia</td>
</tr>
<tr>
<td>Vitamin C (ascorbic acid)</td>
<td>Vitamin C deficiency</td>
</tr>
<tr>
<td></td>
<td>Anemia due to disorders of glutathione metabolism</td>
</tr>
<tr>
<td></td>
<td>Disorders of mitochondrial metabolism</td>
</tr>
<tr>
<td>Vitamin or Mineral</td>
<td>Condition</td>
</tr>
<tr>
<td>-------------------</td>
<td>-----------</td>
</tr>
</tbody>
</table>
| Vitamin D (ergocalciferol) | Vitamin D deficiency  
Galactosemia  
Glycogenosis  
Disorders of magnesium metabolism  
Intestinal malabsorption  
Chronic renal disease  
Cystic fibrosis  
Disorders of phosphorus metabolism  
Hypocalcemia  
Disorders of the biliary tract  
Hypoparathyroidism  
Intestinal disaccharidase deficiencies and disaccharide malabsorption  
Allergic gastroenteritis and colitis |
| Vitamin E (tocopherols) | Vitamin E deficiency  
Inflammatory bowel disease (e.g., Crohn’s, granulomatous enteritis, and ulcerative colitis)  
Disorders of mitochondrial metabolism  
Chronic liver disease  
Intestinal malabsorption  
Disorders of the biliary tract  
Cystic fibrosis |
| Vitamin K (phytonadione) | Vitamin K deficiency  
Congenital deficiency of other clotting factors  
Hypoprothrombinemia of the newborn  
Hemorrhagic disease of the newborn  
Intestinal malabsorption  
Acquired coagulation factor deficiency  
Cystic fibrosis  
Disorders of the biliary tract  
Chronic liver disease |
| Zinc | Zinc deficiency  
Wilson’s disease  
Acrodermatitis enteropathica |
| Multi-minerals | Other and unspecified protein-calorie malnutrition |
| Multi-vitamins | Cystic fibrosis  
Other and unspecified protein-calorie malnutrition |
| Trace elements | Mineral deficiency |
Prior authorization for vitamin and mineral products must be requested using the CCP Prior Authorization Request Form. Requests for prior authorizations must be submitted and approved before the date of dispensing the vitamin or mineral products. Prior authorization requests for vitamin and mineral products that are initiated before the date of the physician’s order will not be approved.

The following documentation must be submitted with the prior authorization request:

- A physician’s prescription with the name of the vitamin or mineral product, dosage, frequency, duration, and route of administration
- The MSRP or average wholesale price (AWP), whichever is applicable, or the provider’s documented invoice price
- The calculated price per dose
- Documentation that supports the medical necessity of the requested vitamin or mineral

The following sample tables, taken from the CCP Prior Authorization Request Form, are examples of the information that is required to submit a request for vitamin and mineral products:

- **Example 1: Vitamin D**

<table>
<thead>
<tr>
<th>HCPCS Code</th>
<th>Brief Description of Requested Services</th>
<th>Retail Price</th>
</tr>
</thead>
<tbody>
<tr>
<td>A9152 UA</td>
<td>Vitamin D (ergocalciferol) 10 ml bottle (8000 units/ml)</td>
<td>$40.00/bottle</td>
</tr>
<tr>
<td></td>
<td>Dose: 400 units (0.05 ml)</td>
<td>$0.20/dose</td>
</tr>
<tr>
<td></td>
<td>Route: PO</td>
<td></td>
</tr>
<tr>
<td></td>
<td>Frequency: QD</td>
<td></td>
</tr>
</tbody>
</table>

*Note: HCPCS codes and descriptions must be provided.*

- **Example 2: Multivitamin Tablets**

<table>
<thead>
<tr>
<th>HCPCS Code</th>
<th>Brief Description of Requested Services</th>
<th>Retail Price</th>
</tr>
</thead>
<tbody>
<tr>
<td>A9153 U2</td>
<td>Centrum Kids (80 tablets/bottle)</td>
<td>$8.99/bottle</td>
</tr>
<tr>
<td></td>
<td>Dose: 1 tablet</td>
<td>$0.11/dose</td>
</tr>
<tr>
<td></td>
<td>Route: PO</td>
<td></td>
</tr>
<tr>
<td></td>
<td>Frequency: QD</td>
<td></td>
</tr>
</tbody>
</table>

*Note: HCPCS codes and descriptions must be provided.*

Prior authorization requests for products, conditions, or quantities other than those described in the “Benefits” section of this handbook will be considered on a case-by-case basis after review by the medical director. Providers must submit documentation that the prescribed products are for a medically accepted indication. Documentation must include one of the following:

- FDA approval
- The use is supported by one or more citations that are included or approved for inclusion in the following compendia:
  - The American Hospital Formulary Service Drug Information
  - The United States Pharmacopoeia-Drug Information (or its successor publications)
  - The DRUGDEX Information System
• Two articles from major medical peer-reviewed literature that demonstrate validated, untested data for the use of the agent in a specific medical condition that is safe and effective

Prior authorization of vitamin and mineral products may be granted for up to six months, and for a quantity up to a 30-day supply.

**Note:** Quantities in excess of these limitations may be considered when requesting liquid formulations due to variances in container size.

Requests for additional vitamin and mineral products must be submitted before the current authorized period expires, but no more than 30 days before the expiration.

### 2.2.28 Wound Care Supplies or Systems

Wound care equipment and supplies may be a benefit under Texas Medicaid Title XIX Home Health services.

To be considered a Home Health benefit, all of the following criteria must be met:

- The client must be eligible for home health benefits.
- The criteria listed in this handbook for the requested equipment and supplies must be met.
- The equipment and supplies requested must be medically necessary.
- Federal financial participation must be available.
- The requested equipment and supplies must be safe and appropriate for use in the home.

This handbook addresses wound care equipment and supplies provided to the client by a Durable Medical Equipment (DME) provider in the home and outpatient hospital setting.

Wound care equipment and supplies are designed to assist in healing of wounds in conjunction with an individualized wound care regimen prescribed by a provider familiar with the client.

Comprehensive wound care regimens include but are not limited to:

- Maintenance of a clean, moist bed of granulation tissue
- Debridement to remove devitalized tissue
- Any necessary treatment to resolve infection
- Optimization of nutrition, circulation, ambulation, and chronic disease management

Providers are to consider the clinical efficacy of the wound care product, the client’s functional status, as well as the measurable signs of effective wound management when ordering products to treat wounds.

Measurable signs of wound management include, but are not limited to:

- A decrease in wound size, either in surface area or volume
- A decrease in amount of exudate
- A decrease in amount of necrotic tissue
- Improved infection status

Wounds are defined as acute or chronic:

- Acute wounds progress through the normal stages of wound healing and show definite signs of healing within four weeks
- Chronic wounds do not progress normally through the stages of healing (often getting “stalled” in one phase) and do not show evidence of healing within four weeks

Skin ulcers represent the majority of chronic wounds. Skin ulcers include but are not limited to:

- Venous ulcers (also known as venous insufficiency ulcers or stasis ulcers)
• Arterial ulcers
• Diabetic ulcers
• Pressure injuries or pressure ulcers

2.2.28.1 Wound Care Supplies

Medically necessary wound care equipment and supplies include, but are not limited to, cleansing agents, wound packing, dressings and coverings, compression garments, and negative pressure wound therapy (NPWT) systems.

Cleansers

Wound cleansing helps create an optimal healing environment and decreases the potential for infection. Cleansing agents and methods vary based on effectiveness and individual client needs. Wound cleansing agents may include, but are not limited to:

• Normal Saline
• Commercial wound cleansers
• Povidone Iodine
• Hydrogen peroxide
• Sodium Hypochlorite

Dressings

A dressing is a wet or dry, sterile or non-sterile, pad or compress that is designed to be in direct contact with the wound. A dressing is applied to promote healing and protect the wound from further harm. Dressings and related supplies may include, but are not limited to:

• Wound packing and fillers
• Gauze, impregnated or non-impregnated, sterile or non-sterile
• Dry dressings
• Collagen dressings
• Alginate or other fiber gelling dressings
• Composite dressings
• Antimicrobials
• Foam dressings
• Contact layers and transparent films
• Hydrocolloid, Hydrofiber, and Hydrogel dressings
• Specialty absorptive dressings
• Compression dressings and wraps
• Tape to secure dressings

Compression

Compression dressings, wraps or stockings apply pressure to body parts to control edema and aid circulation by redirecting blood centrally. Below the knee and above the knee compression stockings may be benefits for Texas Medicaid clients. Compression dressings or stockings may be used, but not limited to the following indications:

• Edema in Pregnancy
• Postural Hypotension
• Lymphedema
• Treatment of any of the following complications of chronic venous insufficiency:
  • Venous edema
  • Stasis Ulcers
  • Varicose veins (not including spider veins)
  • Lipodermatosclerosis

Custom burn compression garments may be a benefit with prior authorization and documentation supporting medical necessity.

2.2.28.2 Negative-Pressure Wound Therapy (NPWT) System

An NPWT system provides and maintains a moist wound environment and protects the wound during the healing process.

An NPWT system consists of a cell foam dressing that is placed in the wound bed, a suction catheter tip, an adhesive drape to cover the wound, suction tubing, and a computerized electric vacuum pump. An NPWT system uses continuous or intermittent sub-atmospheric pressure to remove excess interstitial fluid and remove growth factor inhibitors. The removal of inhibitors allows the growth factor to stimulate cell proliferation and migration. Removal of excess fluid also helps decrease periwound induration.

Drainage from the wound is collected in a canister. Typically, each wound dressing is changed two to three times per week. NPWT is often referred to as a “wound-vac.”

NPWT may be considered for clients with wounds including but not limited to:

• Pressure
• Arterial
• Venous stasis
• Diabetic ulcers
• Post-surgical wound dehiscence; non-adhering skin grafts; or surgical flaps required for covering such wounds

Prior authorization is not required for the initial 90 days of NPWT. Prior authorization is required for continued therapy after the initial 90 days, with the submission of the Wound Care Equipment and Supplies Order Form.

Providers must submit the required documentation for the NPWT system for any prior authorization periods after the initial 90 days of treatment. All documentation required for prior authorization must be maintained in the client’s medical record with the prescribing provider.

2.2.28.3 Exclusions

The following services are not a benefit of Texas Medicaid:

• Wound care supplies for use in the office or outpatient setting
• Equipment and supplies for stand-by or back-up use
• Topical or portable hyperbaric oxygen chambers (procedure code A4575) that are placed directly over the wound and provide higher concentrations of oxygen to the damaged tissue
• Non-contact normothermic wound therapy (NNWT) systems and associated supplies (procedure codes A6000, E0231, E0232)
• Rental or purchase of an electrical stimulation or electromagnetic wound treatment device (procedure code E0769), for use by the client or caregiver in the home setting
• Contact or non-contact ultrasound treatment for wounds
• Electrochemical low-dose tissue oxygenation systems
• Metabolically and non-metabolically active skin equivalents used in wound care are not reimbursed in the home setting
• Procedure code A6545 billed without AW modifier

2.2.28.4 Authorization Requirements

Requests for medically necessary wound care equipment or supplies for any eligible client that does not meet criteria as outlined in this handbook for requests for supplies above the defined limitations will be considered through the comprehensive care program. These requests require prior authorization with documentation supporting the medical necessity of the equipment and the quantity requested.

Note: For clients who are 21 years of age or older, requests for wound care equipment and supplies that do not meet the criteria through Title XIX Home Health Services may be considered under the Texas Medicaid Home Health - Durable Medical Equipment (DME) Exceptional Circumstances process.

Prior authorization must be requested on a completed Wound Care Equipment and Supplies Order Form, signed and dated by the physician.

• The completed Wound Care Equipment and Supplies Order Form must include the procedure codes and numerical quantities for services requested. The completed, signed, and dated form must be maintained by both the DME provider and the prescribing physician in the client’s medical record. The completed Wound Care Equipment and Supplies Order Form with the original dated signature must be maintained by the prescribing physician.

• To complete the prior authorization process by paper, the provider must fax or mail the Wound Care Equipment and Supplies Order Form to the Home Health prior authorization unit and retain a copy of the signed and dated form in the client’s medical record at the durable medical equipment provider’s place of business.

• To complete the prior authorization process electronically, the provider must complete the prior authorization requirements through any approved electronic methods and retain a copy of the signed and dated Wound Care Equipment and Supplies Order Form in the client’s medical record at the durable medical equipment provider’s place of business.

• To facilitate determination of medical necessity and avoid unnecessary denials, the provider must include correct and complete information, specifically documentation for medical necessity of the equipment or supplies requested. The provider must maintain documentation of medical necessity in the client’s medical record, to include the medical condition necessitating the need for the wound care supplies or wound care system.

Unless otherwise specified, wound care equipment and supplies are available without prior authorization up to the stated quantity limitations. Prior authorization is required for any quantities exceeding the limitations.

Prior authorization or additional documentation is required for:

• Quantities of wound care supplies exceeding the quantity limits listed in the Procedure Codes and Limitations table.

• Disposable wound suction (procedure code A9272). Submitted documentation must include justification addressing why no other wound care equipment and supplies will meet the client’s need.
• Continued use of negative pressure wound therapy (NPWT) for more than 90 days requires documentation outlined in NPWT Greater than 90 Days section of this handbook.

• Compression burn garments require documentation of an appropriate diagnosis and evidence of medical necessity. These requests will be reviewed by the medical director.

Prior Authorization is required for the following procedure codes:

<table>
<thead>
<tr>
<th>Procedure Codes</th>
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<tbody>
<tr>
<td>A6215</td>
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<tr>
<td>A6509</td>
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</table>

2.2.28.5 Documentation Requirements

Information from the following four categories must be submitted anytime that prior authorization is required. If prior authorization is not required, this documentation must be maintained in the client’s medical record and is subject to retrospective review.

Category 1: Medical History and Compliance

A comprehensive treatment plan, including the prescribed wound care and management planned for the client including, but not limited to:

• Any medical diagnosis or chronic condition that effects wound healing

• History of previous wound care treatments and outcomes with dates, including therapies initiated in a hospital or Skilled Nursing Facility (SNF)

• Continued management of unresolved compliance issues (i.e., missed medical appointments, refusing dressing changes, repositioning, smoking, poor nutritional intake or choices)

• Whether a family member/friend/caregiver agrees to be available to assist the client

Category 2: Wound Care Interventions

Relevant information related to the current wound, including:

• Any mechanical, surgical, enzymatic or autolytic tissue debridement (if performed)

• Treatment for infection (if present)

Category 3: Wound Description & Details

Detailed description of the wound including:

• Dates of previous and current assessments

• The measurements at the initiation of wound care and the current measurements, including length, width, depth and any undermining or tunneling

• Wound color

• Amount, quality, quantity, and odor of drainage (if present)

• Presence of granulation or eschar (if appropriate)

• The currently prescribed wound care regimen, to include types of dressings, frequency of dressing changes and supplies needed for each dressing change

• Frequency client will be seen by a licensed medical professional to assess wound healing and current wound treatment regimen
Category 4: Contraindications

Absence of the following contraindications:

- Untreated osteomyelitis within the vicinity of the wound
- Wound ischemia
- Gangrene
- Presence in the wound of necrotic tissue with eschar (if debridement has not been attempted)
- Cancer present in the wound or around the margins
- Presence of a fistula to an organ or body cavity within the vicinity of the wound
- Documentation explaining the appropriateness of wound care is required if any of the above contraindications are present

The requesting DME provider may be asked for additional information to clarify or complete a request for the wound care equipment or supplies including but not limited to:

- Overall health status of clients whose wounds are not progressing through the normal stages of healing, including but not limited to:
  - Albumin or pre-albumin (within 30 days)
  - Hemoglobin A1C (within 30 days)
  - Use of pressure reducing surfaces, repositioning, and encouraged ambulation

All services outlined in this section are subject to retrospective review to ensure that the documentation in the client’s medical record supports the medical necessity of the service(s) provided.

Reauthorization will be considered based on medical necessity, with a new prior authorization request.

NPWT Greater than 90 Days

NPWT may be considered beyond the initial 90-day treatment period for additional 30 day treatment periods with prior authorization justifying continued use.

For NPWT of greater than 90 days, providers must submit justification for continued use of NPWT including:

- The measurements at the initiation of NPWT and the current measurements, including length, width, depth and any undermining or tunneling

Additionally, providers must make note if any of the following contraindications are present:

- No measurable improvement of wound status occurring over the prior 90-day period.
- The wound care equipment or supplies are no longer being used by the client as prescribed.

NPWT Supplies

NPWT supplies are limited to a maximum of:

- 15 dressing kits or supplies (procedure code A6550) per wound per month; unless documentation supports that the wound size requires more than one dressing kit for each dressing change or if the provider has ordered more frequent dressing changes.
- 10 disposable canisters (procedure code A7000) per month; unless documentation provided indicates medical necessity for additional canisters.
2.2.28.6 Reimbursement and Billing Guidelines

Wound care equipment and supplies addressed in this section are reimbursed only when provided as a Title XIX Home Health Benefit. Supplies provided in an outpatient setting, such as a wound care clinic, are part of the facility fee and are not reimbursed separately.

Providers should only bill for one month of supplies at a time, even though prior authorization may be granted for up to six months.

Procedure code A6545 must be billed with modifier AW, claims will deny if billed without the modifier.

Providers are reimbursed for items addressed in this section by the lesser of:

- The providers billed charges
- The published fee determined by HHSC

Manual price is determined by HHSC, which is based on:

- The manufacturer’s suggested retail price (MSRP) less 18 percent or average wholesale price (AWP) less 10.5 percent, whichever is applicable, or
- The provider’s documented invoice cost.

If manual pricing is used, the provider must request prior authorization and submit documentation of one of the following:

- The MSRP or AWP, whichever is applicable.
- The provider’s documented invoice cost.

DME Certification

The DME Certification and Receipt Form is required and must be completed before reimbursement can be made for any equipment or appliance delivered to a client. The certification form must include the name of the item, the date the client received the equipment or appliance, and the dated signatures of the provider and the client or primary caregiver. This signed and dated form must be maintained by the DME provider in the client’s medical record.

The DME Certification and Receipt Form must be submitted for DME claims and appeals when:

- A single item meets or exceeds a billed amount of $2,500.00
- Multiple items submitted on the same date of service meet or exceed a total billed amount of $2,500.00

Claims submitted without the DME Certification and Receipt Form will be denied.

Clients who receive DME meeting or exceeding a total billed amount of $2,500.00 may be contacted to verify receipt of the equipment. If receipt of the equipment cannot be verified, the claim payment is eligible for recoupment.

2.2.28.7 Wound Care Procedures and Limitations

The procedure codes listed in the following table do not require prior authorization when requested within allowable limits.

Note: Quantities that exceed the limitations identified in the table will require prior authorization with documentation supporting medical necessity.

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The procedure codes in the following table require prior authorization.

Note: Providers must provide appropriate documentation of medical necessity and documentation for the quantity requested.

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• Any services, equipment, or supplies furnished to a client who is a resident of a public institution or a client in a hospital, SN facility, or intermediate care facility
• Any services or supplies furnished to a client before the effective date of Medicaid eligibility as certified by HHSC or after the date of termination of Medicaid eligibility
• Any services or supplies furnished without prior approval by TMHP, except as listed
• Any supplies or equipment used in a physician’s office, or inserted by a physician (e.g., low profile gastrostomy tube)
• Apnea monitors
• Blood products (the administration or the supplies and equipment used to administer blood products)
• Cardiac telemetry monitoring
• Chemotherapy administration or the supplies and equipment used to administer chemotherapy
• Diapers and wipes for clients who are 3 years of age and younger
• Dynamic orthotic cranioplasty (DOC)
• Environmental equipment, supplies, or services, such as room dehumidifiers, air conditioners, heater/air conditioner filters, space heaters, fans, water purification systems, vacuum cleaners, treatments for dust mites, rodents, and insects
• Home whirlpool baths, spas, home exercisers/gym equipment, hemodialysis equipment, safety wall rails, toys/therapy equipment
• IPV
• Nutritional counseling
• Orthotics, braces, prosthetics including but not limited to voice prosthetic, and artificial larynx
• Parapodiums
• Personal protective equipment (such as gloves, masks, gowns, and sharps containers) for use by a health-care provider, including but not limited to an RN, LVN, or attendant in the home setting
• Pneumocardiograms
• Seat lift chairs
• Shipping, freight, delivery travel time
• Structural changes to homes, domiciles, or other living arrangements
• Vehicle mechanical or structural modifications, such as wheelchair lifts

Refer to: Subsection 1.12, “Texas Medicaid Limitations and Exclusions” in “Section 1: Provider Enrollment and Responsibilities” (Vol. 1, General Information).

2.2.30 Procedure Codes That Do Not Require Prior Authorization

The procedure codes listed in the following table do not require prior authorization for clients who are receiving services under Home Health Services. Although prior authorization is not required, providers must retain a completed Home Health Services (Title XIX) DME/Medical Supplies Physician Order Form for these clients. For medical supplies not requiring prior authorization, a completed Home Health Services (Title XIX) DME/Medical Supplies Physician Order Form may be valid for a maximum of six months unless the physician indicates the duration of need is less. If the physician indicates the
duration of need is less than six months, then a new Home Health Services (Title XIX) DME/Medical Supplies Physician Order Form is required at the end of the duration of need. It is expected that reasonable, medically necessary amounts will be provided.

The use of these services is subject to retrospective review. This is not an all inclusive list.

<table>
<thead>
<tr>
<th>Procedure Codes</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>Nebulizer Supplies/Equipment</strong>*</td>
</tr>
<tr>
<td>E0570</td>
</tr>
<tr>
<td><strong>Incontinence Supplies</strong>**</td>
</tr>
<tr>
<td>A4310</td>
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<tr>
<td>A4326</td>
</tr>
<tr>
<td>A4352</td>
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<td>A5105</td>
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<tr>
<td><strong>Inhaler Equipment</strong></td>
</tr>
<tr>
<td>A4614</td>
</tr>
</tbody>
</table>

* Prior authorization is required for certain diagnoses and if limitations are exceeded. Refer to subsection 2.2.23.2, “Small Volume Nebulizers (SVN)” in this handbook.

** Prior authorization is required for some procedure codes if the maximum limitation is exceeded. Refer to subsection 2.2.15.9, “Incontinence Procedure Codes with Limitations” in this handbook.

2.3 Other or Special Provisions

2.3.1 Medicaid Relationship to Medicare

2.3.1.1 Possible Medicare Clients

It is the provider’s responsibility to determine the type of coverage (Medicare, Medicaid, or private insurance) that the client is entitled to receive. Home health providers must follow these guidelines:

- Clients who are 64 years of age and younger without Medicare Part A or B:
  - If the agency erroneously submits an SOC notice to Medicare and does not contact TMHP for prior authorization, TMHP does not assume responsibility for any services provided before contacting TMHP. The SOC date is no more than three business days before the date the agency contacts TMHP. Visits made before this date are not considered a benefit of the Home Health Services Program.

- Clients who are 65 years of age and older without Medicare Part A or Part B and clients with Medicare Part A or B regardless of age:
  - In filing home health claims, home health providers may be required to obtain Medicare denials before TMHP can approve coverage. When TMHP receives a Medicare denial, the SOC is determined by the date the agency requested coverage from Medicare. If necessary, the 95-day claims filing deadline is waived for these claims, provided TMHP receives notice of the Medicare denial within 30 days of the date on the MRAN containing Medicare’s final disposition.
  - If the agency receives the MRAN and continues to visit the client without contacting TMHP by telephone, mail, or fax within 30 days from the date on the MRAN, TMHP will provide coverage only for services provided from the initial date of contact with TMHP. The SOC date is determined accordingly. TMHP must have the MRAN before considering the request for prior authorization.
2.3.1.2 Benefits for Medicare and Medicaid Clients

For eligible Medicare/Medicaid clients, Medicare is the primary payer and providers must bill Medicare before submitting a claim to Medicaid. Medicaid pays the Medicare deductible on Part B claims for qualified home health clients.

Home health service prior authorizations may be given for HHA services, certain medical supplies, equipment, or appliances suitable for use in the home in one of the following instances:

- When an eligible Medicaid client (enrolled in Medicare) who does not qualify for home health services under Medicare because SN care, PT, or OT are not a part of the client’s care.
- When the medical supplies, equipment, or appliances are denied by Medicare Part B and are a benefit of Home Health Services.

Federal and state laws require the use of Medicaid funds for the payment of most medical services only after all reasonable measures have been made to use a client’s third party resources or other insurance.

Note: If the client has Medicare Part B coverage, contact Medicare for prior authorization requirements and reimbursement. If the service is a Part B benefit, do not contact TMHP for prior authorization. Texas Medicaid will only pay the deductible and coinsurance according to current payment guidelines on the electronic crossover claim.

TMHP will not prior authorize or reimburse the difference between the Medicare payment and the retail price for Medicare Part B eligible clients.

Refer to: “Section 8: Third Party Liability (TPL)” (Vol. 1, General Information).

Subsection 2.7, “Medicare Crossover Claim Reimbursement” in “Section 2: Texas Medicaid Fee-for-Service Reimbursement” (Vol. 1, General Information).

2.3.1.3 Medicare and Medicaid Prior Authorization

Contact TMHP for prior authorization of Medicaid services (based on medical necessity and benefits of Home Health Services) within 30 days of the date on the MRAN.

Note: For MQMB clients, do not submit prior authorization requests to TMHP if the Medicare denial reason states “not medically necessary.” Medicaid only will consider prior authorization requests if the Medicare denial states “not a benefit” of Medicare.

Qualified Medicare Beneficiaries (QMB) are not eligible for Medicaid benefits. Texas Medicaid is only responsible for premiums, coinsurance, or deductibles on these clients according to payment guidelines. Providers should not submit prior authorization requests to the TMHP Home Health Services Prior Authorization Department for these clients.

To ensure Medicare benefits are used first in accordance with Texas Medicaid regulations, the following procedures apply when requesting Medicaid prior authorization and payment of home health services for clients.

Contact TMHP for prior authorization of Medicaid services (based on medical necessity and benefits of Home Health Services) within 30 days of the date on the MRAN. Fax a copy of the original Medicare MRAN and the Medicare appeal review letter to the TMHP Home Health Services Prior Authorization Department for prior authorization.

Note: Claims for STAR+PLUS MQMB clients (those with Medicare and Medicaid) must always be submitted to TMHP as noted on these pages. The STAR+PLUS health plan is not responsible for these services if Medicare denies the service as not a benefit.

When the client is 65 years of age and older or appears otherwise eligible for Medicare such as blind and disabled, but has no Part A or Part B Medicare, the TMHP Home Health Services Prior Authorization Department uses regular prior authorization procedures. In this situation, the claim is held for a midyear...
status determined by HHSC. The maximum length of time a claim may be held in a “pending status” for Medicare determination is 90 days. After the waiting period, the claim is paid or denied. If denied, the EOB code on the R&S report indicates that Medicare is to be billed.

Refer to: Section 3, “Home Health Skilled Nursing and Home Health Aide Services” in the Home Health Nursing and Private Duty Nursing Services Handbook (Vol. 2, Provider Handbooks).

2.4 Claims Filing and Reimbursement

2.4.1 Claims Information

Providers must use only type of bill (TOB) 321 in Form Locator (FL) 4 of the UB-04 CMS-1450. Other TOBs are invalid and result in claim denial.

Home Health services must be submitted to TMHP in an approved electronic format or on a CMS-1500 or a UB-04 CMS-1450 paper claim form. Submit home health DME and medical supplies to TMHP in an approved electronic format, or on a CMS-1500 or on a UB-04 CMS-1450 paper claim form. Providers may purchase UB-04 CMS-1450 and CMS-1500 paper claim forms from the vendor of their choice. TMHP does not supply them.

When completing a CMS-1500 or a UB-04 CMS 1450 paper claim form, providers must include all required information on the claim, as TMHP does not key information from attachments. Superbills, or itemized statements, are not accepted as claim supplements.

Refer to: “Section 3: TMHP Electronic Data Interchange (EDI)” (Vol. 1, General Information) for information on electronic claims submissions.

“Section 6: Claims Filing” (Vol. 1, General Information) for general information about claims filing.


Subsection 6.5, “CMS-1500 Paper Claim Filing Instructions” in “Section 6: Claims Filing” (Vol. 1, General Information) for instructions on completing paper claims.

Outpatient claims must have the appropriate revenue code and, if appropriate, the corresponding HCPCS code or narrative description. The prior authorization number must appear on the UB-04 CMS-1450 claim in Block 63 and in Block 23 of the CMS-1500 claim. The certification dates or the revised request date on the POC must coincide with the DOS on the claim. Prior authorization does not waive the 95-day filing deadline requirement.

Home health service claims should not be submitted for payment until Medicaid certification is received and a prior authorization number is assigned.

2.4.2 Reimbursement

DME and expendable medical supplies are reimbursed in accordance with 1 TAC §355.8023. Providers can refer to the Online Fee Lookup (OFL) or the applicable fee schedule on the TMHP website at www.tmhp.com. Providers may also request a hard copy of the fee schedule by contacting the TMHP Contact Center at 1-800-925-9126.

DME and expendable supplies, other than nutritional products, that have no established fee, are subject to manual pricing at the documented MSRP less 18 percent or the provider’s documented invoice cost.

Nutritional products that have no established fee are subject to manual pricing at the documented AWP less 10.5 percent or at the provider’s documented invoice cost.
For reimbursement, providers must note the following:

- Claims are approved or denied according to the eligibility, prior authorization status, and medical appropriateness.
- Claims must represent a numerical quantity of 1 month for supplies according to the billing requirements.
- DME/supplies must be provided by either a Medicaid enrolled home health agency’s Medicaid/DME supply provider or an independently-enrolled Medicaid/DME supply provider. Both must enroll and bill using the provider identifier enrolled as a DME supplier. File these services on a CMS-1500 claim form.

  **Note:** Medical social services and speech-language pathology services are available to clients who are 20 years of age and younger and are not a benefit of Home Health Services. These services may be considered a benefit for clients who qualify for CCP.

Texas Medicaid does not reimburse separately for associated DME charges, including but not limited to, battery disposal fees or state taxes. Reimbursement for any associated charges is included in the reimbursement for a specific piece of equipment.

**Refer to:** Subsection 2.2, “Fee-for-Service Reimbursement Methodology” in “Section 2: Texas Medicaid Fee-for-Service Reimbursement” (Vol. 1, General Information) for more information about reimbursement.

Texas Medicaid implemented mandated rate reductions for certain services. The Online Fee Lookup (OFL) and static fee schedules include a column titled “Adjusted Fee” to display the individual fees with all mandated percentage reductions applied. Additional information about rate changes is available on the TMHP website at www.tmhp.com/resources/rate-and-code-updates/rate-changes.

### 2.4.3 Home Health Agency Reimbursement for DME Services

Home health agencies are reimbursed for DME and medical supplies in accordance with 1 TAC §355.8023. Providers can refer to the OFL or the applicable fee schedule on the TMHP website at www.tmhp.com. Providers may also request a hard copy of the fee schedule by contacting the TMHP Contact Center at 1-800-925-9126. DME and medical supplies, other than nutritional products, that have no established fee are subject to manual pricing at the documented MSRP less 18 percent or the provider’s documented invoice cost.

### 2.4.4 Prohibition of Medicaid Payment to Home Health Agencies Based on Ownership

Medicaid denies home health services claims when TMHP records indicate that the physician ordering treatment has a significant ownership interest in, or a significant financial or contractual relationship with, the nongovernmental home health agency billing for the services. Federal regulation Title 42 CFR §424.22 (d) states that “a physician who has a significant financial or contractual relationship with, or a significant ownership in a nongovernmental home health agency may not certify or recertify the need for home health services care services and may not establish or review a plan of treatment.”

A physician is considered to have a significant ownership interest in a home health agency if either of the following conditions apply:

- The physician has a direct or indirect ownership of five percent or more in the capital, stock, or profits of the home health agency.
- The physician has an ownership of five percent or more of any mortgage, deed of trust, or other obligation that is secured by the agency, if that interest equals five percent or more of the agency’s assets.
A physician is considered to have a significant financial or contractual relationship with a home health agency if any of the following conditions apply:

- The physician receives any compensation as an officer or director of the home health agency.
- The physician has indirect business transactions, such as contracts, agreements, purchase orders, or leases to obtain services, supplies, equipment, space, and salaried employment with the home health agency.
- The physician has direct or indirect business transactions with the home health agency that, in any fiscal year, amount to more than $25,000 or 5 percent of the agency’s total operating expenses, whichever is less.

When providing CCP services and general home health services, the provider must file these on two separate UB-04 CMS-1450 paper claim forms with the appropriate prior authorization number, and must send them to the appropriate address.

Claims denied because of an ownership conflict will continue to be denied unless the home health agency submits documentation indicating that the ordering physician no longer has a significant ownership interest in, or a significant financial or contractual relationship with, the home health agency providing services. Documentation must be sent to TMHP Provider Enrollment at the address indicated in subsection A.11, “Written Communication With TMHP” in “Appendix A: State, Federal, and TMHP Contact Information” (Vol. 1, General Information).

### Claims Resources

Refer to the following sections or forms when filing claims:

<table>
<thead>
<tr>
<th>Resource</th>
<th>Location</th>
</tr>
</thead>
<tbody>
<tr>
<td>Acronym Dictionary</td>
<td>“Appendix C: Acronym Dictionary” (Vol. 1, General Information)</td>
</tr>
<tr>
<td>Automated Inquiry System (AIS)</td>
<td>Subsection A.10, “TMHP Telephone and Fax Communication” in “Appendix A: State, Federal, and TMHP Contact Information” (Vol. 1, General Information)</td>
</tr>
<tr>
<td>CMS-1500 Paper Claim Filing Instructions</td>
<td>Subsection 6.5, “CMS-1500 Paper Claim Filing Instructions” in “Section 6: Claims Filing” (Vol. 1, General Information)</td>
</tr>
<tr>
<td>State, federal, and TMHP contact information</td>
<td>“Appendix A: State, Federal, and TMHP Contact Information” (Vol. 1, General Information)</td>
</tr>
<tr>
<td>TMHP electronic claims submission information</td>
<td>Subsection 6.2, “TMHP Electronic Claims Submission” in “Section 6: Claims Filing” (Vol. 1, General Information)</td>
</tr>
<tr>
<td>TMHP Electronic Data Interchange (EDI)</td>
<td>“Section 3: TMHP Electronic Data Interchange (EDI)” (Vol. 1, General Information)</td>
</tr>
</tbody>
</table>

### Contact TMHP

The TMHP Contact Center at 1-800-925-9126 is available Monday through Friday from 7 a.m. to 7 p.m., Central Time.
5 Forms

The following linked forms can also be found on the Forms page of the Provider section of the TMHP website at www.tmhp.com:

<table>
<thead>
<tr>
<th>Forms</th>
</tr>
</thead>
<tbody>
<tr>
<td>Addendum to Home Health Services (Title XIX) DME/Medical Supplies Physician Order Form</td>
</tr>
<tr>
<td>CCP Prior Authorization Request Form</td>
</tr>
<tr>
<td>DME Certification and Receipt Form</td>
</tr>
<tr>
<td>External Insulin Pump Prior Authorization Form</td>
</tr>
<tr>
<td>Home Health Services (Title XIX) DME/Medical Supplies Physician Order Form Instructions</td>
</tr>
<tr>
<td>Home Health Services (Title XIX) DME/Medical Supplies Physician Order Form</td>
</tr>
<tr>
<td>Home Health Plan of Care (POC) Instructions</td>
</tr>
<tr>
<td>Home Health Plan of Care (POC)</td>
</tr>
<tr>
<td>Home Health Prior Authorization Checklist</td>
</tr>
<tr>
<td>Texas Medicaid Prior Authorization Request for CPAP or RAD (Bi-level PAP)</td>
</tr>
<tr>
<td>Texas Medicaid Prior Authorization Request for Oxygen Therapy Devices and Supplies form</td>
</tr>
<tr>
<td>Texas Medicaid Prior Authorization Request for Secretion and Mucus Clearance Devices—Initial Request form</td>
</tr>
<tr>
<td>Texas Medicaid Prior Authorization Request for Secretion and Mucus Clearance Devices –Renewal Request form</td>
</tr>
<tr>
<td>Texas Medicaid Provider Surety Bond and Instructions</td>
</tr>
<tr>
<td>Wheelchair/Scooter/Stroller Seating Assessment Form (CCP/Home Health Services)</td>
</tr>
<tr>
<td>Wound Care Equipment and Supplies Order Form</td>
</tr>
</tbody>
</table>

6 Claim Form Examples

The following linked claim form examples can also be found on the Claim Form Examples page of the Provider section of the TMHP website at www.tmhp.com:

<table>
<thead>
<tr>
<th>Claim Form Examples</th>
</tr>
</thead>
<tbody>
<tr>
<td>Home Health Services DME/Medical Supplies</td>
</tr>
</tbody>
</table>
GYNECOLOGICAL, OBSTETRICS, AND FAMILY PLANNING TITLE XIX SERVICES HANDBOOK

TEXAS MEDICAID PROVIDER PROCEDURES MANUAL: VOL. 2

MAY 2021
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1 General Information

The information in this handbook is intended for gynecological, obstetrics, and Texas Medicaid Title XIX family planning providers. The handbook provides information about Texas Medicaid’s benefits, policies, and procedures that are applicable to these service providers.

Important: All providers are required to read and comply with “Section 1: Provider Enrollment and Responsibilities” (Vol. 1, General Information). In addition to required compliance with all requirements specific to Texas Medicaid, it is a violation of Texas Medicaid rules when a provider fails to provide health-care services or items to Medicaid clients in accordance with accepted medical community standards and standards that govern occupations, as explained in 1 Texas Administrative Code (TAC) §371.1659. Accordingly, in addition to being subject to sanctions for failure to comply with the requirements that are specific to Texas Medicaid, providers can also be subject to Texas Medicaid sanctions for failure, at all times, to deliver health-care items and services to Medicaid clients in full accordance with all applicable licensure and certification requirements including, without limitation, those related to documentation and record maintenance.

Refer to: The Children’s Services Handbook (Vol. 2, Provider Handbooks) for more information about providing services to Texas Medicaid/Texas Health Steps (THSteps) clients.

“Section 1: Provider Enrollment and Responsibilities” (Vol. 1, General Information).

“Texas Medicaid Administration” in the Preliminary Information (Vol. 1, General Information).

The Healthy Texas Women Program Handbook (Vol. 2, Provider Handbooks) for information about women’s health services.

The Health and Human Services Commission Family Planning Program Services Handbook (Vol. 2, Provider Handbooks) for information about Health and Human Services Commission (HHSC) programs that provide family planning services.

This handbook contains information about Texas Medicaid fee-for-service benefits. For information about managed care benefits, providers can refer to the Medicaid Managed Care Handbook (Vol. 2, Provider Handbooks).

Managed care carve-out services are administered as fee-for-service benefits. A list of all carve-out services is available in the Medicaid Managed Care Handbook.

Refer to: Section 8, “Carve-Out Services” in the Medicaid Managed Care Handbook (Vol. 2, Provider Handbooks).

1.1 Family Planning Overview

TMHP processes family planning claims and encounters for two different funding sources:

- The HHSC Family Planning program funding for HHSC-contracted providers
- The Title XIX family planning funding for Texas Medicaid providers

HHSC awards contracts to agencies across Texas to provide services to low-income individuals who may not qualify for Texas Medicaid services. These awards are granted through a competitive procurement process. HHSC contracts with a variety of providers, including local health departments, universities, medical schools, private nonprofit agencies, FQHCs, RHCs, and hospital districts. All HHSC-contracted providers must first be enrolled in Title XIX Texas Medicaid.
Client eligibility requirements, reimbursement methodologies, client copayment guidelines, and covered services may differ for each funding source. Family planning funding cannot be used for elective abortion services.

- Title XIX funds are available for family planning services provided to Texas Medicaid clients. TMHP processes Title XIX claims and reimburses eligible services on a fee-for-service basis for family planning providers and a prospective payment system basis for FQHC and RHC providers.
- HHSC Family Planning Program contracts annually with family planning providers. TMHP processes claims and reimburses providers for services to eligible clients according to the individually granted funds.
- Funds are also available for women’s health and family planning services provided to Healthy Texas Women (HTW) clients. TMHP processes HTW claims and reimburses eligible services on a fee-for-service basis for family planning providers and a prospective payment system basis for FQHC and RHC providers.

1.1.1 Guidelines for Family Planning Providers

The following guidelines apply for all family planning services:

- Family planning services may be provided by a physician or under the direction of a physician, not necessarily personal supervision. A physician provides direction for family planning services through written standing delegation orders and medical protocols. The physician is not required to be on the premises for the provision of family planning services by a registered nurse (RN), physicians assistant (PA), nurse practitioner (NP), clinical nurse specialist (CNS), or certified nurse midwife (CNM).
- Services must be provided without regard to age, marital status, sex, race, ethnicity, parenthood, handicap, religion, national origin, or contraceptive preference.
- Texas Medicaid clients, including limited care clients, are allowed to choose any enrolled family planning service provider.
- Family planning clients must be allowed freedom of choice in the selection of contraceptive methods as medically appropriate.
- Family planning clients must be allowed the freedom to accept or reject services without coercion.
- Only family planning clients may consent to the provision of family planning services. Counseling should be offered to adolescents that encourages them to discuss their family planning needs with a parent, an adult family member, or other trusted adult.
- Sterilization services cannot be provided to any person who is 20 years of age or younger. For more information, HHSC providers may refer to the HHSC website at www.healthytexaswomen.org.

1.2 Payment Window Reimbursement Guidelines for Services Preceding an Inpatient Admission

According to the three-day and one-day payment window reimbursement guidelines, most professional and outpatient diagnostic and nondiagnostic services that are rendered within the designated timeframe of an inpatient hospital stay and are related to the inpatient hospital admission will not be reimbursed separately from the inpatient hospital stay if the services are rendered by the hospital or an entity that is wholly owned or operated by the hospital.

These reimbursement guidelines do not apply in the following circumstances:

- The professional services are rendered in the inpatient hospital setting.
- The hospital and the physician office or other entity are both owned by a third party, such as a health system.
• The hospital is not the sole or 100-percent owner of the entity.

Refer to: Subsection 3.7.4.14, “Payment Window Reimbursement Guidelines” in the Inpatient and Outpatient Hospital Services Handbook (Vol. 2, Provider Handbooks) for additional information about the payment window reimbursement guidelines.

2 Medicaid Title XIX Family Planning Services

2.1 Title XIX Provider Enrollment

Physician, FQHC, and RHC providers may provide Title XIX family planning services for Texas Medicaid clients under the provider’s Texas Medicaid provider number. No additional enrollment is required to provide Title XIX family planning services.

Refer to: Subsection 7.1, “Provider Enrollment” in the Medical and Nursing Specialists, Physicians, and Physician Assistants Handbook (Vol. 2, Provider Handbooks) for information about physician provider enrollment.

Subsection 4.1, “Enrollment” in the Clinics and Other Outpatient Facility Services Handbook (Vol. 2, Provider Handbooks) for information about FQHC provider enrollment.

Subsection 7.1, “Enrollment” in the Clinics and Other Outpatient Facility Services Handbook (Vol. 2, Provider Handbooks) for information about RHC provider enrollment.

Family planning agencies must apply for enrollment with TMHP to receive an agency provider identifier. To be enrolled in Texas Medicaid, family planning agencies must meet the following requirements:

• Complete an agency enrollment application.

• Ensure that all services are furnished by, prescribed by, or provided under the direction of a licensed physician in accordance with the Texas Medical Board or Texas BON.

• Have a medical director who is a physician currently licensed to practice medicine in Texas, and submit a current copy of the medical director’s physician license.

• Have an established record of performance in the provision of both medical and educational counseling of family planning services as verified through client records, established clinic hours, and clinic site locations.

• Provide family planning services in accordance with HHSC standards of client care for family planning agencies.

• Be approved for family planning services by the HHSC Family Planning Program.

Note: An RHC can also apply for enrollment as a family planning agency.

The effective date for participation is the date an approved provider agreement with Medicaid is established and the provider is assigned a Medicaid provider identifier.

Providers cannot be enrolled if their license is due to expire within 30 days. A current license must be submitted.

Refer to: “Section 1: Provider Enrollment and Responsibilities” (Vol. 1, General Information) for more information about enrollment procedures.

Subsection 6.3.6, “Benefit Code” in the Section 6: Claims Filing (Vol. 2, Provider Handbooks) for more information about benefit codes.
2.2 Services, Benefits, Limitations, and Prior Authorization

This section includes information on family planning services funded through Title XIX Medicaid.

Family planning services are preventive health, medical, counseling, and educational services that assist individuals in managing their fertility and achieving optimal reproductive and general health. Title XIX services include:

- Family planning annual exams
- Other family planning office or outpatient visits
- Laboratory services
- Radiology services
- Contraceptive devices and related procedures
- Drugs and supplies
- Medical counseling and education
- Sterilization and sterilization-related procedures (i.e., tubal implants, tubal ligation, vasectomy, and anesthesia for sterilization)

Providers must use one of the following diagnosis codes in conjunction with all family planning procedures and services:

<table>
<thead>
<tr>
<th>Diagnosis Codes</th>
</tr>
</thead>
<tbody>
<tr>
<td>Z30011</td>
</tr>
<tr>
<td>Z3009</td>
</tr>
<tr>
<td>Z30433</td>
</tr>
<tr>
<td>Z9852</td>
</tr>
</tbody>
</table>

One of the diagnosis codes in this table must be included in Block 24 E of the CMS-1500 claim form referencing the appropriate procedure code. The choice of diagnosis code must be based on the type of family planning service performed.

*Note:* Title XIX family planning services are exempt from the limited program and rules.

2.2.1 Family Planning Annual Exams

An annual family planning exam consists of a comprehensive health history and physical examination, which includes the following:

- Medical laboratory evaluations as indicated
- An assessment of the client’s problems and needs
- The implementation of an appropriate contraceptive management plan
Family planning providers must bill the most appropriate evaluation and management (E/M) visit procedure code for the complexity of the annual family planning examination provided. To bill an annual family planning examination, one of the following procedure codes must be billed with modifier FP and a family planning diagnosis code:

<table>
<thead>
<tr>
<th>Procedure Codes</th>
</tr>
</thead>
<tbody>
<tr>
<td>99202</td>
</tr>
</tbody>
</table>

**Important:** Only the annual family planning examination requires modifier FP. All other family planning office visits do not. One annual family planning examination is allowed per year. Claims filed incorrectly may be denied.

The following table summarizes the uses for the E/M procedure codes and the corresponding billing requirements for the annual examination:

<table>
<thead>
<tr>
<th>Billing Criteria</th>
<th>Frequency</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>New patient:</strong> Most appropriate E/M procedure code with modifier FP and a family planning diagnosis code</td>
<td>One new patient E/M code every 3 years following the last E/M visit provided the client by that provider or a provider of the same specialty in the same group</td>
</tr>
<tr>
<td><strong>Established patient:</strong> Most appropriate E/M procedure code with modifier FP and a family planning diagnosis code</td>
<td>Once a year*</td>
</tr>
</tbody>
</table>

* The established patient procedure code will be denied if a new patient procedure code has been billed in the same year.

An annual family planning examination (billed with modifier FP) will not be reimbursed when submitted with the same date of service as a surgical procedure or an additional E/M visit.

If another condition requiring an E/M office visit beyond the required components for the annual examination is discovered, the provider may submit a claim for the additional visit using modifier 25 to indicate that the client’s condition required a significant, separately identifiable E/M service. Documentation supporting the provision of a significant, separately identifiable E/M service must be maintained in the client’s medical record and made available to Texas Medicaid upon request.

### 2.2.1.1 FQHC Reimbursement for Family Planning Annual Exams

To receive their encounter rate for the annual family planning examination, FQHCs must use the most appropriate E/M procedure code for the complexity of service provided as indicated in the previous table in subsection 2.2.1, “Family Planning Annual Exams” in this handbook.

The annual exam is allowed once per fiscal year, per client, per provider. Two additional family planning office or outpatient visits may be reimbursed to the FQHC within the same year for the same client.

A new patient visit for the annual exam may be reimbursed once every three years following the last E/M visit provided to the client by that provider or a provider of the same specialty in the same group. The annual examination must be billed as an established patient visit if E/M services have been provided to the client within the last three years.

Reimbursement for services payable to an FQHC is based on an all-inclusive rate per visit.

### 2.2.2 Other Family Planning Office or Outpatient Visits

Other family planning E/M visits are allowed for routine contraceptive surveillance, family planning counseling and education, contraceptive problems, suspicion of pregnancy, genitourinary infections, and evaluation of other reproductive system symptoms.
During any visit for a medical problem or follow-up visit, the following must occur:

- An update of the client’s relevant history
- Physical exam, if indicated
- Laboratory tests, if indicated
- Treatment or referral, if indicated
- Education and counseling, or referral, if indicated
- Scheduling of office or clinic visit, if indicated

Title XIX family planning providers must use one of the following procedure codes based on the complexity of the visit with a family planning diagnosis for other family planning office or outpatient visits:

<table>
<thead>
<tr>
<th>Procedure Codes</th>
<th>99202</th>
<th>99203</th>
<th>99204</th>
<th>99205</th>
<th>99211</th>
<th>99212</th>
<th>99213</th>
<th>99214</th>
<th>99215</th>
</tr>
</thead>
</table>

**Important:** Family planning E/M office and outpatient visits should not be billed with modifier FP. Claims filed incorrectly may be denied.

The following table summarizes the uses for the E/M procedure codes and the corresponding billing requirements for each type of visit:

<table>
<thead>
<tr>
<th>Billing Criteria</th>
<th>Frequency</th>
</tr>
</thead>
<tbody>
<tr>
<td><em>New patient:</em> Most appropriate E/M procedure code with a family planning diagnosis code</td>
<td>One new patient E/M code every 3 years following the last E/M visit provided the client by that provider or a provider of the same specialty in the same group</td>
</tr>
<tr>
<td><em>Established patient:</em> Most appropriate E/M procedure code with a family planning diagnosis code</td>
<td>As needed*</td>
</tr>
<tr>
<td><em>The established patient procedure code will be denied if a new patient procedure code has been billed in the same year.</em></td>
<td></td>
</tr>
</tbody>
</table>

**Refer to:** Subsection 2.2, “Services, Benefits, Limitations, and Prior Authorization” in this handbook for the list of family planning diagnosis codes.

A general family planning office or outpatient visit (billed without modifier FP) will not be reimbursed when submitted with the same date of service as a surgical procedure or an additional E/M visit. If another condition requiring an E/M office visit beyond the required components for an office visit, family planning visit, or surgical procedure is discovered, the provider may submit a claim for the additional visit using modifier 25 to indicate that the client’s condition required a significant, separately identifiable E/M service. Documentation supporting the provision of a significant, separately identifiable E/M service must be maintained in the client’s medical record and made available to Texas Medicaid upon request.

**2.2.2.1 FQHC Reimbursement for Other Family Planning Office or Outpatient Visits**

FQHCs may be reimbursed for three family planning encounters per year, per client, regardless of the reason for the encounter. The three encounters may include any combination of general family planning, annual family planning exams, or services. Procedure codes J7296, J7297, J7298, J7300, J7301, and J7307 may be reimbursed in addition to the FQHC encounter payment. When seeking reimbursement for an intrauterine device (IUD) or implantable contraceptive capsule, providers must submit on the same claim the procedure code for the family planning service provided and the
procedure code for the contraceptive device. The contraceptive device is not subject to FQHC limitations. Providers must use modifier U8 when submitting claims for a contraceptive device purchased through the 340B Drug Pricing Program.

A family planning diagnosis code must be billed along with the most appropriate informational procedure codes for the services that were rendered. Reimbursement for services payable to an FQHC is based on an all-inclusive rate per visit.

Refer to: Section 4, “Federally Qualified Health Center (FQHC)” in the Clinics and Other Outpatient Facility Services Handbook (Vol. 2, Provider Handbooks) for more information about FQHC services.

2.2.2.2 RHC Reimbursement for Other Family Planning Office or Outpatient Visits

RHC’s may receive an encounter rate when submitting claims with procedure code T1015, in addition to a flat “add on” fee for a LARC device (procedure codes J7296, J7297, J7298, J7300, J7301, and J7307).

Refer to: Subsection 7.2.1.4, “Family Planning Services” in the Clinics and Other Outpatient Facility Services Handbook (Vol. 2, Provider Handbooks) for more information about family planning services performed by an RHC.

2.2.3 Laboratory Procedures

All family planning laboratory services must be billed with a family planning diagnosis code.

2.2.3.1 Clinical Laboratory Improvement Amendments (CLIA) Requirement

All providers of laboratory services must comply with the rules and regulations of the CLIA. Providers who are not in compliance with CLIA will not be reimbursed for laboratory services. Only the office or lab that holds the appropriate CLIA certificate and that actually performs the laboratory test procedure may be reimbursed for the procedure.

Refer to: Subsection 2.1.1, “Clinical Laboratory Improvement Amendments (CLIA)” in the Radiology and Laboratory Services Handbook (Vol. 2, Provider Handbooks).

2.2.3.2 Medical Record Documentation

Medicaid family planning service providers must document in the client’s medical record the medical necessity of all ordered laboratory services. The medical record documentation must also reference an appropriate diagnosis.

2.2.3.3 Lab Specimen Handling and Testing

Any test specimen sent to a laboratory may be reimbursed to the laboratory that performs the test and not to the referring family planning provider.

If the provider that obtains the specimen does not perform the laboratory procedure, the provider that obtains the specimen may be reimbursed one lab handling fee per day, per client, using procedure code 99000 and a family planning diagnosis code for the handling or conveyance of the specimen from the provider’s office to a laboratory. More than one lab handling fee may be reimbursed per day if multiple specimens are obtained and sent to different laboratories.

Handling fees are not paid for Pap smears or cultures. The appropriate procedure code may be reimbursed for Pap smear interpretations when billed with modifier SU indicating that the screening and interpretation were actually performed in the office.

2.2.3.4 Providing Information to the Reference Laboratory

When sending any specimen, including Pap smears, to the reference laboratory, the family planning provider must provide the reference laboratory with the client’s name, address, Texas Medicaid number, and a family planning diagnosis so the laboratory may bill Texas Medicaid for its family planning lab services.
2.2.4 Radiology Services

Procedure codes 74000, 74010, and 76830 may be reimbursed for services performed for the purpose of localization of an IUD.

2.2.5 External Contraceptives, Long Acting Reversible Contraceptives (LARCs), and Related Procedures

2.2.5.1 External Contraceptives

Procedure codes A4261 (cervical cap) and A4266 (diaphragm) may be reimbursed separately from the fitting and instruction (procedure code 57170).

Procedure codes A4261 and A4266 may be reimbursed when they are billed with one of the following diagnosis codes:

<table>
<thead>
<tr>
<th>Diagnosis Codes</th>
</tr>
</thead>
<tbody>
<tr>
<td>Z30011</td>
</tr>
<tr>
<td>Z30431</td>
</tr>
</tbody>
</table>

2.2.5.2 Long Acting Reversible Contraception: Intrauterine Devices

2.2.5.2.1 Insertion of the LARC IUD

The IUD and the insertion of the IUD may be reimbursed using procedure code J7296, J7297, J7298, J7300, J7301, and 58300.

Procedure codes J7296, J7297, J7298, J7300, and J7301 may be reimbursed when they are billed with one of the following diagnosis codes:

<table>
<thead>
<tr>
<th>Diagnosis codes</th>
</tr>
</thead>
<tbody>
<tr>
<td>Z30011</td>
</tr>
<tr>
<td>Z3041</td>
</tr>
<tr>
<td>Z309</td>
</tr>
</tbody>
</table>

2.2.5.2.2 Removal of the LARC IUD

Procedure code 58301 may be reimbursed when an IUD is extracted from the uterine cavity. An office visit will not be reimbursed when billed on the same date of service as procedure code 58301.

When a vaginal, cervical, or uterine surgery procedure code is submitted with the same date of service as the IUD removal procedure code or the IUD replacement procedure code, the following reimbursement may apply:

- The other vaginal, cervical, or uterine surgical procedure may be reimbursed at full allowance.
- The removal or the replacement of the IUD will be denied.

2.2.5.3 Long Acting Reversible Contraception: Implantable Contraceptive Capsules

The contraceptive capsule and the implantation of the contraceptive capsule may be reimbursed using procedure code J7307. Providers must use modifier U8 when submitting claims for a contraceptive device purchased through the 340B Drug Pricing Program.

Procedure code 11981 may be reimbursed for the insertion of the contraceptive capsule when it is billed with a family planning diagnosis code.

Procedure code 11983 may be reimbursed for the removal with reinsertion of the contraceptive capsule when it is billed with a family planning diagnosis code. Progesterone-containing subdermal contraceptive capsules (Norplant) were previously used for birth control. Although subdermal contraceptive
capsules are no longer approved by the Food and Drug Administration (FDA), the removal of the implanted contraceptive capsule may be considered for reimbursement with procedure code 11976 or 11982.

2.2.5.4 Immediate Postpartum Insertion of LARCs: IUDs and Implantable Contraceptive Capsules

Procedure codes for LARCs may be reimbursed in addition to the hospital Diagnosis related group (DRG) payment when insertion is performed immediately postpartum. "Immediately postpartum" refers to the following:

- Insertion within 10–15 minutes of placental delivery for IUDs
- Insertion prior to discharge for implantable contraceptive capsules.

Medicaid MCOs must adopt claim processing procedures to reimburse hospital and facility providers for immediate postpartum LARC devices in addition to the rate for delivery services.

For claims submitted to TMHP, hospital and facility providers must submit an outpatient claim with the appropriate procedure code for the contraceptive device in addition to the inpatient claim for the delivery services.

For claims submitted to a Texas Medicaid managed care organization (MCO), providers must follow the MCO’s claim processing procedures for reimbursement of immediate postpartum LARC devices in addition to the rate for delivery services.

2.2.6 Contraceptive Drugs, Supplies, and Prophylactics

The following procedure codes may be reimbursed for contraceptive drugs, supplies, and prophylactics:

<table>
<thead>
<tr>
<th>Procedure Codes</th>
</tr>
</thead>
<tbody>
<tr>
<td>A4267</td>
</tr>
</tbody>
</table>

* Procedure code J3490 may be reimbursed when a prescription medication to treat a genital infection is provided to the client.

Procedure code J1050 with modifier U1 may be reimbursed for services rendered to female clients as medically appropriate for the purpose of contraception. A quantity of 1 must be billed.

Procedure code J1050 (no modifier) may be reimbursed for services rendered to male and female clients of any age for other indications as appropriate. Providers must bill the appropriate quantity based on the amount used in milligrams (mg).

For Texas Medicaid Title XIX services, procedure code J1050 is not diagnosis-restricted. For Title XIX family planning services, procedure code J1050 must be billed with a valid family planning diagnosis code.

Procedure codes A4268, A4269, and S4993 may be reimbursed when they are billed with one of the following diagnosis codes:

<table>
<thead>
<tr>
<th>Diagnosis Codes</th>
</tr>
</thead>
<tbody>
<tr>
<td>Z30011</td>
</tr>
<tr>
<td>Z30431</td>
</tr>
</tbody>
</table>

Procedure code A9150 is not reimbursed through Title XIX Medicaid for the medication to treat a monilia infection. The drug is available through the Medicaid Vendor Drug Program with a prescription.

Providers must use modifier U8 when submitting claims for a contraceptive device purchased through the 340B Drug Pricing Program.
Refer to: Subsection 1.1, “About the Vendor Drug Program” in the Outpatient Drug Services Handbook (Vol. 2, Provider Handbooks) for information about this program.

2.2.6.1 Prescriptions and Dispensing Medication

Family planning agencies may do one or both of the following:

- Dispense family planning drugs and supplies directly to the client and bill accordingly.
- Write a prescription for the client to take to a pharmacy.

Family planning drugs and supplies that are dispensed directly to the client must be billed to TMHP for Texas Medicaid fee-for-service clients. Only family planning agencies may be reimbursed for dispensing family planning drugs and supplies. Family planning agencies may be reimbursed for dispensing up to a 1-year supply of contraceptives in a 12-month period using procedure code J7303, J7304, or S4993. The appropriate family planning diagnosis code must be included on the claim.


Title XIX clients may have prescriptions filled at the clinic pharmacy or at another pharmacy. Pharmacies under the Vendor Drug Program are allowed to fill all prescriptions as prescribed. Family planning drugs and supplies are exempt from the three prescriptions-per-month rule for up to a six-month supply.

2.2.6.2 Pharmacy Benefit for Long-Acting Reversible Contraception Products

Certain LARC products are available as a pharmacy benefit of Family Planning and are available through a limited number of specialty pharmacies that work with LARC manufacturers. Providers can refer to the Texas Medicaid/CHIP Vendor Drug Program website at www.txvendordrug.com/formulary/larc.shtml for additional information, including a list of covered products and participating specialty pharmacies.

2.2.6.3 Medroxyprogesterone Acetate (Depo-Provera)

Medroxyprogesterone acetate injectable suspension (Depo-Provera) has been approved by the FDA as a method of contraception. Intramuscular injections of medroxyprogesterone acetate given at 90-day intervals has been proven to be a long-term method of preventing pregnancy. Medroxyprogesterone acetate injectable suspension is reimbursed by Texas Medicaid to providers of family planning services. Medroxyprogesterone acetate must be billed using procedure code J1050 with modifier U1 and a valid family planning diagnosis codes.

2.2.6.4 Injection Administration

Injection administration billed by a provider is reimbursed separately from the medication. If billed without procedure code J1050 and modifier U1, procedure code 96372 must be billed with a family planning diagnosis and a description of the medication in the Remarks field of the claim. Injection administration is not payable to outpatient hospitals.

Refer to: Subsection 2.2, “Services, Benefits, Limitations, and Prior Authorization” in this handbook for a list of family planning diagnosis codes.

2.2.7 Medical Counseling and Education

Procedure code H1010 for the instruction in natural family planning methods may be reimbursed once per day, per person or per couple, when billed by any provider with one of the following diagnosis codes:

<table>
<thead>
<tr>
<th>Diagnosis Codes</th>
</tr>
</thead>
<tbody>
<tr>
<td>Z30011 Z30018 Z3002 Z3009 Z302 Z3040 Z3041 Z3042</td>
</tr>
<tr>
<td>Z30430 Z30431 Z30432 Z30433 Z3049 Z308 Z309 Z9851</td>
</tr>
<tr>
<td>Z9852</td>
</tr>
</tbody>
</table>
Procedure code H1010 is intended to instruct a couple or an individual in methods of natural family planning. Two sessions (one per client) may be billed for separate, individual sessions, or one session may be billed for counseling and education if provided in a joint session. Each session may be billed separately or the two sessions may be billed together with a total charge for both sessions.

### 2.2.8 Sterilization and Sterilization-Related Procedures

For a complete list of Title XIX sterilization procedures, providers can refer to the Texas Medicaid fee schedules located on the TMHP website at [http://public.tmhp.com/FeeSchedules/Default.aspx](http://public.tmhp.com/FeeSchedules/Default.aspx).

#### 2.2.8.1 Sterilization Consent

Per federal regulation 42 CFR 50, Subpart B, all sterilization procedures require an approved Sterilization Consent Form.

**Note:** The Texas Medicaid - Title XIX Acknowledgment of Hysterectomy Information form is not sterilization consent.

**Refer to:**
- Sterilization Consent Form (English) on the TMHP website at [www.tmhp.com](http://www.tmhp.com).
- Sterilization Consent Form (Spanish) on the TMHP website at [www.tmhp.com](http://www.tmhp.com).
- Sterilization Consent Form Instructions on the TMHP website at [www.tmhp.com](http://www.tmhp.com).

#### 2.2.8.2 Anesthesia for Sterilization

Procedure codes 00840, 00851, and 00940 may be reimbursed for anesthesia for sterilization services in accordance with standard anesthesia billing requirements. Providers must include a valid family planning diagnosis code on the claim.

**Refer to:** Subsection 6.2.5.2, “Anesthesia” in “Section 6: Claims Filing” (Vol. 1, General Information) for more information about anesthesia modifiers.

#### 2.2.8.3 Occlusive Sterilization Device

Procedure code A4264 may be reimbursed for the occlusive sterilization system (micro-insert), and may be reimbursed separately from the surgery (procedure code 58565) to place the device.

Providers must bill procedure code A4264 on the same date of service by the same provider as the occlusive sterilization system (micro-insert).

Procedure code 58565 is considered bilateral and is limited to once per lifetime, any provider.

#### 2.2.8.4 Tubal Ligation

Procedure code 58600, 58615, 58670, or 58671 may be reimbursed for tubal ligations.

#### 2.2.8.5 Vasectomy

Procedure code 55250 may be reimbursed for any sterilization procedure that is performed on a male by a family planning agency. This procedure code may be reimbursed as a global fee to include preoperative, intra-operative, and postoperative services by all parties involved. Vasectomies are considered to be permanent, once-per-lifetime procedures. If a vasectomy has previously been reimbursed for the client, providers may appeal with documentation that supports the medical necessity for the repeat sterilization.

#### 2.2.8.6 Facility Fees for Sterilization

Hospital-based and freestanding ambulatory surgical centers (HASCs/ASCs) may be reimbursed for procedure code 55250, 58260, 58262, 58565, 58600, 58615, 58670, or 58671. An appropriate family planning diagnosis code must be billed when reporting facility fees for procedure codes 58565 or 58670.
Refer to: Ambulatory Surgical Center on the TMHP website at www.tmhp.com for a claim form example.

Subsection 5.2.13, “Gynecological and Reproductive Health and Family Planning Services” in the Inpatient and Outpatient Hospital Services Handbook (Vol. 2, Provider Handbooks) for additional gynecological, reproductive health, and family planning services procedure codes that may be reimbursed to ASC and HASC providers.

2.2.9 Prior Authorization
Prior authorization is not required for family planning services, including sterilization and sterilization-related procedures.

2.2.10 Non-covered Services

2.2.10.1 Family Planning Services for Undocumented Aliens
Undocumented aliens are identified on the client eligibility card as having limited Medicaid eligibility by the classification of Type Program (TP) 30, 31, 34, and 35. Under Texas Medicaid, these clients are only eligible for emergency services, including emergency labor and delivery. Texas Medicaid emergency-only services do not cover Title XIX family planning services.

2.3 Documentation Requirements
All services require documentation to support the medical necessity of the service rendered, including gynecological and reproductive health services, and family planning services. Gynecological and reproductive health services, and family planning services are subject to retrospective review and recoupment if documentation does not support the service billed.

2.4 Claims Filing and Reimbursement

2.4.1 Claims Information
Providers may use the following claim forms to submit claims to TMHP:

<table>
<thead>
<tr>
<th>Providers</th>
<th>Forms</th>
</tr>
</thead>
<tbody>
<tr>
<td>Medicaid Fee-For-Service Claims Submitted to TMHP</td>
<td></td>
</tr>
<tr>
<td>All family planning services provided by physicians, PAs, NPs, CNSs, CNMs, and family planning agencies who also contract with HHSC</td>
<td>2017 claim form or approved electronic format</td>
</tr>
<tr>
<td>Medicaid family planning providers who do not contract with HHSC</td>
<td>2017 claim form, CMS-1500 claim form, or approved electronic format of either form</td>
</tr>
<tr>
<td>Hospitals</td>
<td>UB-04 CMS-1450 claim form or approved electronic format</td>
</tr>
<tr>
<td>FQHCs not contracted with HHSC</td>
<td>UB-04 CMS-1450, 2017 claim form, or approved electronic format of either form</td>
</tr>
<tr>
<td>FQHC also contracts with HHSC</td>
<td>2017 claim form or approved electronic format</td>
</tr>
</tbody>
</table>

The following applies when filing claims:
- All claims and Sterilization Consent Forms submitted by family planning agencies must be submitted with benefit code FP3.
- Family planning services billed by RHCs must include modifier AJ, AM, SA, or U7. These services must be billed using the appropriate national place of service (72) for an RHC setting.
When completing a 2017, CMS-1500, or UB-04 CMS-1450 claim form, all required information must be included on the claim, as TMHP does not key any information from claim attachments. Superbills, or itemized statements, are not accepted as claim supplements.

All claims must be filed within approved filing deadlines.

Denied claims may be appealed.

Providers may copy 2017 Claim Form on the TMHP website at www.tmhp.com.

Providers may purchase CMS-1500 and UB-04 CMS-1450 claim forms from the vendor of their choice. TMHP does not supply the forms.

Refer to: “Section 3: TMHP Electronic Data Interchange (EDI)” (Vol. 1, General Information) for information on electronic claims submissions.

“Section 6: Claims Filing” (Vol. 1, General Information) for general information about claims filing.


Subsection 6.5.4, “CMS-1500 Instruction Table” in “Section 6: Claims Filing” (Vol. 1, General Information).


“Section 7: Appeals” (Vol. 1, General Information) for information about appealing claims.

Blocks that are not referenced are not required for processing by TMHP and may be left blank.

RHCs must use their National Provider Identifier (NPI), the appropriate benefit code as applicable, and the appropriate modifier and place of service as outlined in this section.

2.4.1.1 Family Planning and Third Party Liability

Federal and state regulations mandate that family planning client information be kept confidential. Because seeking information from third party insurance may jeopardize the client’s confidentiality, prior insurance billing is not a requirement for billing family planning for any title program.

2.4.2 Billing Procedures for Non-Family-Planning Services Provided During a Family Planning Visit (Title XIX Only)

When a non-family-planning service is provided during a family planning visit or the client is offered family planning services during a medical visit, the following billing process must be used:

A family planning clinic must bill for non-family-planning services using the performing provider’s identifier. The clinic provider identifier is used to bill family planning services only.

The performing provider must bill both family planning services and non-family-planning services, using the correct provider identifier.

The FQHC must bill both family planning services and non-family-planning services, using the correct provider identifier.
• An RHC may bill a rural health encounter for a non-family-planning medical condition or use the physician’s or NP’s provider identifier to bill for family planning services. If the RHC also is enrolled as a family planning agency, the family planning services may be billed using the agency’s family planning provider identifier and the appropriate national place of service (72) for an RHC setting.

2.4.3 National Drug Code

*Refer to:* Subsection 6.3.4, “National Drug Code (NDC)” in “Section 6: Claims Filing” (Vol. 1, General Information).

2.4.4 National Correct Coding Initiative (NCCI) and Medically Unlikely Edit (MUE) Guidelines

The Healthcare Common Procedure Coding System (HCPCS) and Current Procedural Terminology (CPT) codes included in the Texas Medicaid Provider Procedures Manual are subject to NCCI relationships, which supersede any exceptions to NCCI code relationships that may be noted in the manuals and bulletins. Providers should refer to the CMS NCCI web page for correct coding guidelines and specific applicable code combinations.

In instances when Texas Medicaid limitations are more restrictive than NCCI MUE guidance, Texas Medicaid limitations prevail.

3 Breastfeeding Support Services

Breast pumps facilitate an infant’s ability to receive their mother’s own breast milk when it would be difficult to do so without equipment. Breast pump equipment is a benefit when provided by durable medical equipment (DME) suppliers and medical supply companies in the home. DME suppliers may deliver breast pump equipment to a client who is still in the hospital, but for claims purposes, the place of service should indicate the home setting.

A breast pump may be obtained under an eligible mother’s Medicaid client number; however, if a mother is no longer eligible for Texas Medicaid and there is a need for a breast pump or parts, then breast pump equipment must be obtained under the infant’s Medicaid client number.

Any provider who is familiar with the mother’s or infant’s health may order a breast pump. The ordering provider may include, but is not limited to, obstetricians, gynecologists, neonatologists, and pediatricians. Providers in the hospital setting or the community setting may write the order for breast pump equipment.

DME providers are not precluded from releasing equipment to infants with a Medicaid pending status when they have an order from the direct care provider. A Medicaid identification number is typically assigned to an infant within 24 hours, but it may take longer. During this time, DME providers are expected to release equipment to the client as ordered by the direct care provider who is familiar with the infant.

*Note:* For clients who are 21 years of age or older, requests for breast pumps that do not meet the criteria through Title XIX Home Health Services may be considered under the Texas Medicaid Home Health—Durable Medical Equipment (DME) Exceptional Circumstances process.

*Note:* Medicaid providers should supply breast pump equipment to qualifying clients. If clients do not meet criteria as outlined, mothers should also be informed that breast pumps may be available through the Special Supplemental Nutrition Program for Women, Infant, and Children (WIC), contingent upon WIC’s issuance criteria.

The American Academy of Pediatrics (AAP) recommends that infants exclusively breastfeed for the first six months and continue to breastfeed through 12 months of age or longer.
Breastfeeding refers to both the infant’s feeding of breast milk directly from the breast as well as the feeding of mother’s own expressed breast milk to the infant.

Breast milk provides unsurpassed nutrition and immune protections, influencing the growth and development of infants, and is a significant primary prevention strategy for improving infant health outcomes.

Breastfeeding is encouraged as a means to prevent various illnesses and conditions and to promote the health and wellness of mothers and infants. In some circumstances, breast pumps may be necessary for breastfeeding.

Breastfeeding reduces the infant’s risk for illness, including ear infections, gastrointestinal and bacterial infections, respiratory infections, asthma, diabetes, obesity, and leukemia; and reduces the premature infant’s risk of necrotizing enterocolitis. Breastfeeding also reduces the mother’s risk of developing conditions including breast and ovarian cancers, diabetes, and cardiovascular disease. With proper breastfeeding support and management, breastfeeding may also be protective against post-partum depression.

Infant health can be directly impacted by the ability of the infant’s mother to provide her own breast milk. Therefore, it is beneficial to enhance the opportunities for infants to receive their mother’s milk through the use of equipment when needed.

The American Academy of Pediatrics (AAP) and the Centers for Disease Control and Prevention (CDC) advise that mothers who are infected with human immunodeficiency virus (HIV), Human T-lymphotropic virus (HTLV)-1, and HTLV-2 infection should not breastfeed as the virus may be passed to their infant.

All breast pumps must meet the following specifications:

- Comply, be registered, and be cleared with the Federal Drug Administration (FDA)
- Allow for pumping sessions to be efficiently completed within 30 minutes
- Be adaptable for several sizes of breast shields (flanges), including larger sizes, so as to accommodate different sizes of breasts and nipples
- Have an adjustable and wide-range of suction pressure at the breast shield during use, typically from 30 millimeters of mercury up to 250 millimeters of mercury (mm Hg)
- Have a mechanism or written guidelines to prevent or instruct the user from achieving a vacuum level over 250 mm Hg
- Be portable

The following breast pump procedure codes are a benefit of Texas Medicaid with the listed limitations:

<table>
<thead>
<tr>
<th>Procedure Codes</th>
<th>Additional Information</th>
<th>Limitations</th>
</tr>
</thead>
<tbody>
<tr>
<td>A4281, A4282, A4283, A4284, A4285, A4286</td>
<td>Breast pump parts for use with a pump that has been purchased. All parts must be submitted with modifier U8.</td>
<td>Each part - up to 2 times within 12 months from the breast pump date of purchase</td>
</tr>
<tr>
<td>E0602*</td>
<td>Purchase of a personal-use, manual breast pump</td>
<td>Once within 12 months from the date of birth</td>
</tr>
<tr>
<td>E0603*</td>
<td>Purchase of a personal-use, electric breast pump</td>
<td></td>
</tr>
</tbody>
</table>

*Only one of these procedure codes may be reimbursed when submitted for the same date of service by any provider
The rental or purchase of a breast pump, as well as any replacements or parts, must be billed using the mother’s Medicaid identification number, or if she is no longer eligible, using the infant’s Medicaid identification number.

Personal-use breast pump is for use by only one individual. Personal-use pumps are not to be shared, or used by anyone other than the original owner. Use by more than one individual may pose a risk of cross-contamination, may result in infection or illness of mother and infant, and may void the equipment warranty.

A manual or electrical (AC and/or DC) personal-use breast pump may be considered for purchase only.

A multiple-user electric breast pump is also known as a hospital-grade electric breast pump. Hospital-grade electric breast pumps are designed for repeated uses throughout the pump’s lifetime, by more than one woman and infant. The risk of cross-contamination is eliminated via a closed system. Pumped breast milk must not reach the motor. Hospital grade pumps are durable, heavy-duty breast pumps.

A hospital-grade breast pump may be considered for rental only.

Pump kits, which are specific to each breast pump manufacturer’s requirements, provide the necessary supplies and accessories to allow expression of breast milk.

Procedure codes E0602 and E0603 will be denied when submitted within the same calendar month as procedure code E0604.

Procedure code E0602 will be denied when submitted within one year from procedure code E0603, any provider.

Claims for parts will not be reimbursed when billed for the same day as the purchase of breast pump equipment. Reimbursement is for purchase or rental, with documented medical necessity and prior authorization when appropriate, as outlined in this handbook.

Refer to: Subsection 2.3.2.3, “Breast Pump Claims Filing for MCO Services” in the Medicaid Managed Care Handbook (Vol. 2, Provider Handbooks) for more information related to breast pump coverage for newborns.

### 3.1 Breast Pump Kit Specifications

A breast pump kit is included in the purchase or rental of a breast pump, and is not separately reimbursed. Kits should include the following:

- Breast shields (flanges) that are adjustable and flexible, or flanges that are available in several different sizes if rigid, including larger sizes
- All accessories necessary for pumping two breasts simultaneously for electric pumps, or at least one breast manually for manual handle-squeeze pumps
- All parts necessary to easily convert an electric pump to a manual pump
- At least one extra set of membranes and valve replacements
• At least one extra diaphragm replacement for closed-system pumps
• At least two collection bottles with spill-proof standard size caps, that are bisphenol-A (BPA) and DEHP-free
• Accessories and supplies must be compatible with the pump provided. Materials must be of durable quality for withstanding repeated boiling, washing, and pumping use.

3.2 Replacement Parts
Breast pump replacement parts (procedure codes A4281, A4282, A4283, A4284, and A4285) must be billed with modifier U8.

Replacement parts will be denied when submitted for the same date of service as a breast pump.

If the breast pump was not purchased by Texas Medicaid and requires replacement parts, the following documentation of a client-owned device must be submitted:

• Purchase date
• Serial number
• Purchasing entity of the device
• Copy of the receipt, if available

Note: Parts for a hospital-grade electric breast pump (procedure code E0604), and routine servicing and all necessary repairs to ensure the unit remains functional for the client’s needs, are included in the rental of the pump and is the responsibility of the DME supplier.

Refer to: Subsection 6.4.1, “National Correct Coding Initiative (NCCI) Guidelines” in “Section 6: Claims Filing” (Vol. 1, General Information) for information about NCCI MUE guidelines.

3.3 Personal-Use Manual Breast Pump Medical Necessity Criteria
Manual breast pumps (procedure code E0602) are personal-use, hand-operated, handle-squeeze pumps that are appropriate for short-term or occasional uses related to, but not limited to, any of the following:

• Infrequent separation from infants; such as mothers who work or go to school part-time for less than 10 hours per week, and who do not meet criteria for electric or hospital-grade pumps.
• Resolving brief uncomplicated periods of plugged duct
• Short-term concerns of mild engorgement
• Flat, retracted, or inverted nipples, and the mother does not meet the criteria for electric or hospital-grade pumps
• Cracked or fissured nipples, and the mother does not meet the criteria for electric or hospital-grade pumps

Note: Manual breast pumps are not recommended for pumping on a regular basis, or for attempting to establish a milk supply.

3.3.1 Specifications
Manual breast pumps must include an independent milk collection bottle. The pump cylinder must not be the milk-collecting container.

3.4 Personal-Use Electric Breast Pump Medical Necessity
Electric Breast Pumps (procedure code E0603) have a motor and are electrically (AC and/or DC) operated. Personal-use double-collection electric breast pumps are recommended for their functionality and efficiency; they allow expression of breast milk from both breasts simultaneously.
Personal use double-collection electric breast pumps are for mothers and infants who are breastfeeding with limited, minor, or no complications. Personal-use double-collection electric breast pumps are recommended for pumping and maintaining a milk supply related to, but not limited to, any of the following:

- Regular separation from infants; such as mothers returning to work or school for 10 or more hours per week
- Infants detained in the hospital, who do not meet the criteria for a multiple-user electric breast pump
- Significant breast engorgement
- Breast abscess
- Mastitis
- If the mother is to receive short-term treatment with medication or therapies that may be transmitted through breast milk, but she wishes to maintain her milk supply by pumping and discarding her milk in the interim

### 3.4.1 Specifications

Electric breast pumps must meet the following specifications:

- Be adaptable for simultaneous pumping of both breasts (double-collection)
- Have an adjustable suction pressure range necessary for preventing nipple trauma
- Automatically cycle with an adjustable variable cycling rate, typically 30 to 60 or more cycles per minute
- Include a battery option or adapter to be used as an alternate power source when electricity is not immediately available

**Note:** Personal-use, single-collection electric pumps cannot simultaneously pump both breasts. Single-collection pumps are not recommended, as they are neither effective in maintaining a long-term milk supply nor efficient when pumping during short periods, such as work breaks. Double-collection breast pumps are the standard personal-use electric pump recommended by Texas Medicaid for breastfeeding infants and mothers.

### 3.5 Hospital-Grade Electric Breast Pump Medical Necessity Criteria

Multiple-user, hospital-grade electric breast pumps (procedure code E0604) are heavy-duty, durable, closed-system pumps designed to be loaned multiple times throughout the pump’s lifespan.

Rental of a multiple-user, hospital-grade electric breast pump is recommended for moderate to significant breastfeeding complications. Hospital-grade electric breast pumps are recommended for pumping related to, but not limited to, any of the following:

- Infants who cannot suck well, or have an uncoordinated swallow/suck reflex, due to respiratory disease or congenital disorder
- Infants diagnosed with failure to thrive, cardiac problems, or other special needs
- Infants who are chronically ill
- Infants of low birth-weight with increased nutritional needs
- Infants with severe feeding or digestive problems, as described by the provider in documentation
- Prematurity (less than 37 weeks gestation)
- Multiple births (e.g., twins, triplets, etc.)
• Long-term separation of mother and infant due to hospitalization
• Mothers experiencing conditions affecting their milk production, or low-milk supply, as described in documentation by the prescribing provider familiar with the client
• Mothers needing to induce lactation for establishing their milk supply, but are unable to do so without a hospital-grade breast pump

  Note: A closed-system pump requires a personal-use milk collection pump kit, included in the rental, but to be kept by the individual and not for return with the pump.

3.5.1 Specifications
Use of a hospital-grade breast pump may be covered when the use of a hospital-grade breast pump is determined to be medically necessary and appropriate, as documented by the provider.

A hospital-grade electric breast pump must meet the following specifications:
• Be adaptable for simultaneous pumping of both breasts with an adjustable suction for preventing nipple trauma
• Automatically cycle with adjustable or variable cycling that closely mimics the suckling action of an infant
• Electrical (AC and/or DC)
• Include an adapter to be used as an alternate power source when electricity is not immediately available
• Must not allow milk to contact the housing unit or internal pump-motor at any time when the multiple-user pump is used per manufacturer’s instructions

3.6 Prior Authorization
Prior authorization requests must include the following:
• A completed Home Health Services (Title XIX) DME/Medical Supplies Physician Order Form prescribing the durable medical equipment, signed and dated by the prescribing provider familiar with the client
• The prescribing provider must provide correct and complete information on the form, including accurate medical necessity of the equipment requested.

To complete the prior authorization process, the DME provider must submit the completed Home Health Services (Title XIX) DME/Medical Supplies Physician Order Form to the THMP Home Health Unit.

3.6.1 Replacement Parts
Prior authorization is not required for up to 2 replacements of each part within 12 months from the breast pump’s date of purchase.

Prior authorization is required for parts that exceed the limitations outlined in this handbook.

Requests must be submitted with appropriate documentation to support the need for additional replacement parts. The following documentation must be included under “If applicable” in section B of the Home Health Services (Title XIX) DME/Medical Supplies Physician Order Form:
• The provider must attest that the mother continues to use the equipment for breastfeeding.
• The provider must indicate that the requested part is required for improved pumping efficiency (e.g., larger flanges), or is damaged or lost and affecting the function of the pump.
3.6.2 Personal-Use Manual or Electric Breast Pumps

Prior authorization is not required for the purchase of a manual or electric personal-use breast pump, within 12 months from the date of birth. Prior authorization is required for the replacement of a manual or electric personal-use breast pump due to damage or loss, within 12 months from the purchase date.

Requests for the replacement of a Texas Medicaid purchased personal-use breast pump (procedure code E0602 or E0603) must include the following documentation:

- A statement from the provider describing the loss or damage and what measures will be taken to prevent reoccurrence
- A copy of the police or fire report, when appropriate. The report must also be maintained in the client’s medical record.

Note: HHSC or its designee reserves the right to request additional documentation about the need for replacement when there is evidence of abuse or neglect to equipment by the client, client’s family, or caregiver. Requests for replacement when there is documented proof of abuse or neglect will not be approved.

3.6.3 Hospital-Grade Breast Pump

The initial 60-day rental of a hospital-grade pump does not require prior authorization.

The subsequent rental of a hospital-grade breast pump does require prior authorization. Subsequent rental requests may be considered for 90-day increments only. A maximum of 3 prior-authorized 90-day increments will be allowed within the 12 months following birth.

If an infant with medical necessity requires the extended rental of a hospital-grade electric breast pump, beyond these limitations, the claim may be considered for reimbursement upon appeal with documentation.

The prescribing provider familiar with the client must identify pumping as the mother’s primary method for expressing her breast milk, and must include a statement that clearly describes the infant’s medical necessity. Medically necessary conditions may include the following:

- Short-bowel syndrome
- Severe malabsorption syndromes
- Severe feeding intolerances or immunological deficiencies

Clients who no longer qualify for the continued rental of a hospital-grade breast pump may still qualify for the purchase of a breast pump as outlined in this handbook.

Purchase of a manual breast pump will not be reimbursed during the rental of a hospital-grade pump or if a personal-use electric breast pump was already purchased within the frequency limitations outlined in this handbook.

Purchase of a personal-use electric breast pump will not be reimbursed during the rental of a hospital-grade breast pump.

A hospital-grade electric breast pump will be considered purchased and owned by the client when the monthly payments for rental, through the same provider, equals the purchase cost for the equipment.

The following is required:

- The DME provider must notify the client when the rental equipment is considered purchased due to an extended rental. Proof of ownership must be provided to the client by the DME provider.
- Proof of client ownership of the device is required for reimbursement of replacement parts
• A statement from the DME provider indicating the make and model of the client-owned device, along with proof of client ownership, must be submitted with claim appeals for reimbursement of parts.

A hospital-grade breast pump that has been purchased due to extended rental is anticipated to last the minimum time frame indicated by the manufacturer’s warranty.

3.7 Documentation Requirements
Direct care providers must maintain the following documentation in the client’s medical record:
• Client’s specific medical necessity regarding the specific type of breast pump equipment ordered
• Anticipated duration of need regarding the circumstances or conditions related to the type of equipment ordered
• Infant's age (or gestational age, if premature)
• Documentation of the mother’s intent to breastfeed

3.7.1 DME Certification Form
The DME Certification and Receipt Form must be submitted by DME claims and appeals when:
• A single item meets or exceeds a billed amount of $2,500
• Multiple items submitted on the same date of service meet or exceed a total billed amount of $2,500

The DME Certification and Receipt Form is required and must be completed before reimbursement can be made for any DME delivered to a client. The certification form must include the following:
• Name of the item
• Date the client received the DME
• Date signature of the DME provider and the client or client’s primary caregiver

The signed and dated form must be maintained by the DME provider in the client’s record. Claims submitted without the DME Certification and Receipt Form will be denied.

Clients who receive DME meeting or exceeding a total billed amount of $2,500 may be contacted to verify receipt of the equipment. If receipt of the equipment cannot be verified, the claim payment is eligible for recoupment.

3.8 Services that are not a Benefit
The following breastfeeding support services are not benefits of Texas Medicaid:
• Personal-use electric breast pumps that are only capable of single-collection pumping, one breast at a time
• Breastfeeding support services in the preconception or prenatal period
• Breastfeeding support services for infants who are not breastfeeding and the mother has no intent to breastfeed

4 Obstetric Services

4.1 Services, Benefits, Limitations, and Prior Authorization
Antepartum care, antenatal surveillance, perinatal procedures, infant deliveries, and postpartum care are a benefit of Texas Medicaid.
Medicaid reimburses prenatal care, deliveries, and postpartum care as individual services. Providers may choose one of the following options for billing maternity services:

- Providers may itemize each service individually on one claim form and file at the time of delivery. The filing deadline is applied to the date of delivery.
- Providers may itemize each service individually and submit claims as the services are rendered. The filing deadline is applied to each individual date of service.

Providers who only provide prenatal care and choose to submit prenatal visit charges on one claim form have the filing deadline applied to the estimated date of confinement (EDC) that must be stated in Block 24D of the CMS-1500 claim form.

Laboratory (including pregnancy tests) and radiology services provided during pregnancy must be billed separately and claims must be received by TMHP within 95 days of the date of service.

Medicaid may reimburse only one delivery or Cesarean section procedure code per client in a seven-month period; reimbursement includes multiple births. Delivering physicians who perform regional anesthesia or nerve block do not receive additional reimbursement because these charges are included in the reimbursement for the delivery except as outlined.

Refer to:

Procedure code 99140 is not considered for reimbursement when submitted with diagnosis code O80 for a normal delivery or with diagnosis code O82 for a Cesarean delivery when one of these diagnosis codes is documented on the claim as the referenced diagnosis. The referenced diagnosis must indicate the complicating condition. An emergency is defined as a situation when delay in treatment of the client poses a significant health threat to a client’s life, bodily organ, or body part.

Hospital admissions resulting from conditions or comorbidities complicating labor should be billed using the appropriate E/M procedure codes. These codes are not subject to the three-day pre-care period but are not payable on the date of delivery or the following six-week post-care period.

The procedure codes listed in the tables below may be reimbursed by Texas Medicaid. Providers can refer to the Texas Medicaid Static Fee Schedules and the Online Fee Look-up for rate and coverage information about specific procedure codes.

Prior authorization requests may be submitted to the TMHP Prior Authorization Department via mail, fax, or the electronic portal. Prescribing or ordering providers, dispensing providers, clients’ responsible adults, and clients may sign prior authorization forms and supporting documentation using electronic or wet signatures. Medical record documentation must include assessment findings that substantiate the medical necessity for each diagnostic test performed.

### 4.1.1 Antepartum and Fetal Invasive Procedures

The following procedure codes may be submitted for antepartum and fetal invasive procedures:

<table>
<thead>
<tr>
<th>Procedure Codes</th>
</tr>
</thead>
<tbody>
<tr>
<td>36460 59000 59001 59012 59015 59020 59025 59030 59050 59051</td>
</tr>
<tr>
<td>59070 59074 59076 82731 84112</td>
</tr>
</tbody>
</table>

Antenatal surveillance includes fetal contraction stress test (procedure code 59020), fetal nonstress test (procedure code 59025), and fetal biophysical profile with or without nonstress testing (procedure code 76818 or 76819).
The American Congress of Obstetricians and Gynecologists (ACOG) states “because antepartum fetal surveillance results have not been definitively demonstrated to improve perinatal outcome, all indications for antepartum testing must be considered somewhat relative. In general, antepartum fetal surveillance has been employed in pregnancies in which the risk of antepartum fetal demise is increased.” Accordingly, some of the conditions under which testing may be appropriate, include but are not limited to the following:

- Maternal conditions:
  - Antiphospholipid syndrome
  - Hyperthyroidism (poorly controlled)
  - Hemoglobinopathies (hemoglobin SS, SC, or S-thalassemia)
  - Cyanotic heart disease
  - Systemic lupus erythematosus
  - Chronic renal disease
  - Type I diabetes mellitus
  - Hypertensive disorders
- Pregnancy-related conditions:
  - Pregnancy-induced hypertension
  - Decreased fetal movement
  - Oligohydramnios
  - Polyhydramnios
  - Intrauterine growth restriction
  - Postterm pregnancy
  - Isoimmunization (moderate to severe)
  - Previous fetal demise (unexplained or recurrent risk)
  - Multiple gestation (with significant growth discrepancy)

Procedure codes 59020 and 59025 billed with revenue code 729 for outpatient facilities may be reimbursed on the same day by a different provider without appeal; however, if billed more than once per day by the same provider, it will be denied. The provider may appeal with documentation supporting the performance of the test more than once on the same day by the same provider.

A fetal fibronectin (fFN) enzyme immunoassay (procedure code 82731) may be considered for reimbursement through Texas Medicaid when performed between 22 0/7 and 34 6/7 weeks for women with risk factors for preterm labor with or without symptoms of preterm labor.

Fetal intrauterine transfusion (procedure code 36460) and cordocentesis (procedure code 59012) are restricted to the diagnoses listed in the following table:

<table>
<thead>
<tr>
<th>Diagnosis Codes</th>
</tr>
</thead>
<tbody>
<tr>
<td>O353XX0</td>
</tr>
<tr>
<td>O360110</td>
</tr>
<tr>
<td>O360121</td>
</tr>
<tr>
<td>O360132</td>
</tr>
<tr>
<td>O360193</td>
</tr>
</tbody>
</table>
FIUT (procedure code 36460) is reimbursed as a global fee and, therefore, includes all other services provided by the same physician, including umbilical blood sampling or cordocentesis (procedure code 59012).

Appeals for cordocentesis performed for a diagnosis other than the ones listed in the appropriate table in the policy will be reviewed on a case by case basis.

In addition to the physician performing the FIUT (procedure code 36460), another physician may assist with echography control.

Therapeutic amniocentesis (procedure code 59001) is restricted to the following diagnosis codes:

<table>
<thead>
<tr>
<th>Diagnosis Codes</th>
</tr>
</thead>
<tbody>
<tr>
<td>O360914</td>
</tr>
<tr>
<td>O360925</td>
</tr>
<tr>
<td>O360939</td>
</tr>
<tr>
<td>O361110</td>
</tr>
<tr>
<td>O361121</td>
</tr>
<tr>
<td>O361132</td>
</tr>
<tr>
<td>O361193</td>
</tr>
<tr>
<td>O361914</td>
</tr>
<tr>
<td>O361925</td>
</tr>
<tr>
<td>O361939</td>
</tr>
<tr>
<td>O368210</td>
</tr>
<tr>
<td>O368221</td>
</tr>
<tr>
<td>O368232</td>
</tr>
<tr>
<td>O368293</td>
</tr>
</tbody>
</table>

Transabdominal amnioinfusion (procedure code 59070), fetal fluid drainage (e.g., vesicocentesis, thoracocentesis, paracentesis), including ultrasound guidance (procedure code 59074), and fetal shunt placement, including ultrasound guidance (procedure code 59076) are restricted to one of the following diagnosis codes:

<table>
<thead>
<tr>
<th>Diagnosis Codes</th>
</tr>
</thead>
<tbody>
<tr>
<td>O360910</td>
</tr>
<tr>
<td>O360925</td>
</tr>
<tr>
<td>O360939</td>
</tr>
</tbody>
</table>
## 4.1.2 Vaginal and Cesarean Deliveries

The following procedure codes submitted with the appropriate modifier may be a benefit for vaginal or Cesarean deliveries:

### Procedure Codes

<table>
<thead>
<tr>
<th>Code</th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td>59409</td>
<td>Prior to 39 Weeks and Medically Necessary</td>
</tr>
<tr>
<td>59410</td>
<td>39 Weeks or Later</td>
</tr>
<tr>
<td>59514</td>
<td>Prior to 39 Weeks and Not Medically Necessary</td>
</tr>
<tr>
<td>59515</td>
<td></td>
</tr>
<tr>
<td>59612</td>
<td></td>
</tr>
<tr>
<td>59614</td>
<td></td>
</tr>
<tr>
<td>59620</td>
<td></td>
</tr>
<tr>
<td>59622</td>
<td></td>
</tr>
<tr>
<td>S8415</td>
<td>Procedure code S8415 is for home delivery supplies</td>
</tr>
</tbody>
</table>

One of the following modifiers must be billed with the procedure codes indicated above for vaginal and cesarean deliveries:

### Modifiers

<table>
<thead>
<tr>
<th>Modifier</th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td>U1</td>
<td>Prior to 39 Weeks and Medically Necessary</td>
</tr>
<tr>
<td>U2</td>
<td>39 Weeks or Later</td>
</tr>
<tr>
<td>U3</td>
<td>Prior to 39 Weeks and Not Medically Necessary</td>
</tr>
</tbody>
</table>

Claims will deny if submitted for a delivery prior to 39 weeks of gestation and not medically necessary, or for a delivery service with no modifier.

Claims will deny or recoupment will occur for associated claims for deliveries that are performed prior to 39 weeks and are determined to be not medically necessary including:

- Claims for the provider performing the vaginal or Cesarean delivery
- Inpatient and outpatient hospital claims inclusive of the delivery, planned Cesarean section, induction with vaginal delivery or failed induction with subsequent Cesarean section
- Birthing center claims inclusive of induction with vaginal delivery
- Claims for medical or surgical admission, including ICU, due to the complications of the delivery for the mother

Home deliveries must be billed with procedure code 59409 or 59410; including postpartum care.

Licensed midwives will not be reimbursed for home deliveries.

### 4.1.2.1 Home Deliveries

Home deliveries and the home supplies for the delivery (S8415) require submitting a written prior authorization request during the client’s third trimester of pregnancy. Home deliveries will not be prior authorized to a licensed midwife.
Documentation must include:

- A statement signed by a licensed physician who examined the client during the third trimester and determined at the time of examination the client is not at high risk for complications and is suitable for a home delivery
- A plan for access to emergency transport for mother and neonate, if needed

### 4.1.3 Elective Deliveries Prior to 39 Weeks

Texas Medicaid restricts any Cesarean section, labor induction, or any delivery following labor induction to one of the following criteria:

- Gestational age of the fetus should be determined to be at least 39 weeks
- When the delivery occurs prior to 39 weeks, maternal and/or fetal conditions must dictate medical necessity for the delivery

**Note:** Records are subject to retrospective review. Payments made for Cesarean section, labor induction, or any delivery following labor induction that fail to meet these criteria (as determined by review of medical documentation), are subject to recoupment. Recoupment may apply to all services related to the delivery, including additional physician fees, birthing center, and inpatient and outpatient hospital fees.

### 4.1.4 Other Vaginal and Cesarean Delivery Procedures

The following vaginal and Cesarean delivery procedures do not require vaginal and Cesarean delivery modifiers:

<table>
<thead>
<tr>
<th>Procedure Codes</th>
</tr>
</thead>
<tbody>
<tr>
<td>59525</td>
</tr>
</tbody>
</table>

**Refer to:** Subsection 4.1.8, “Obstetric Ultrasound” in this handbook for information about ultrasound limitations.

### 4.1.5 Surgical Treatment of Early Intrauterine Failed Pregnancy

A provider must conduct an evaluation to make a definitive diagnosis of early intrauterine failed pregnancy; this may include a comprehensive medical history and examination, serum human chorionic gonadotropin (hCG) testing, other lab tests, and ultrasound examination.

The following procedure codes may be submitted for surgical management of early intrauterine failed pregnancy. The provider must choose the most appropriate code:

<table>
<thead>
<tr>
<th>Procedure Codes</th>
</tr>
</thead>
<tbody>
<tr>
<td>59812 59820 59821</td>
</tr>
</tbody>
</table>

Performing providers must maintain appropriate documentation in the medical record of the client’s early intrauterine failed pregnancy and the surgical treatment provided for the individual circumstances.

Surgical treatment of early intrauterine failed pregnancy (procedure codes 59812, 59820, and 59821) must include one of the following diagnosis codes as the reference or primary diagnosis on the claim. The provider must choose the most appropriate code:

<table>
<thead>
<tr>
<th>Diagnosis Codes</th>
</tr>
</thead>
<tbody>
<tr>
<td>O021 O0339 O034 O071 O0730 O0739</td>
</tr>
</tbody>
</table>
4.1.6 Abortion

The following procedure codes may be submitted for abortion services:

<table>
<thead>
<tr>
<th>Procedure Codes</th>
<th>Modifier</th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td>59830 59840 59841 59850 59851 59852 59855 59856 59857</td>
<td>G7</td>
<td>Pregnancy resulted from rape or incest or pregnancy certified by physician as life threatening</td>
</tr>
</tbody>
</table>

Abortion services are benefits of Texas Medicaid if submitted with the following modifier:

In accordance with federal directives, abortions may be reimbursed when performed to save the life of the mother or for pregnancies resulting from rape or incest.

In accordance with federal law, providers are required to use specific language regarding the reason the mother’s condition is life-threatening. An abortion for a life-threatening condition must be due to a physical disorder, injury, or illness, including a life-endangering physical condition caused by or arising from the pregnancy itself that would place the woman in danger of death unless an abortion was performed.

Reimbursement of an abortion is based on the physician’s certification that the abortion was performed to save the life of the mother, to terminate pregnancy resulting from rape, or to terminate pregnancy resulting from incest.

One of the following statements, signed by the physician is mandatory for any abortion performed. Substitute wording will not be accepted. One of these statements must accompany any claim for abortion in order for reimbursement to be made:

- “I, (physician’s name), certify that on the basis of my professional judgment, an abortion procedure is necessary because (client’s full name, Medicaid number, and complete address) suffers from a physical disorder, injury, or illness, including a life-endangering physical condition caused by or arising from the pregnancy itself that would place her in danger of death unless an abortion is performed.”

  Signature

- “I, (physician’s name), certify that on the basis of my professional judgment, an abortion procedure for (client’s full name, Medicaid number, and complete address) is necessary to terminate a pregnancy that was the result of rape. I have counseled the client concerning the availability of health and social support services and the importance of reporting the rape to the appropriate law enforcement authorities.”

  Signature

- “I, (physician’s name), certify that on the basis of my professional judgment, an abortion procedure for (client’s full name Medicaid number, and complete address) is necessary to terminate a pregnancy that was the result of incest. I have counseled the client concerning the availability of health and social support services and the importance of reporting the incest to the appropriate law enforcement authorities.”

  Signature

A stamped or typed physician signature on the original certification statement is not acceptable. The physician signature must be an original signature. A copy of the signed certification statement must be submitted with each claim for reimbursement. Faxes are not acceptable at this time. The physician must maintain the original certification statement in the client’s file.
Abortion services must be billed with modifier G7.

Performing physicians, facilities, anesthesiologists, and certified respiratory nurse anesthetist (CRNA) providers must submit modifier G7 with the appropriate procedure code when requesting reimbursement for abortion procedures that are within the scope of the rules and regulations of Texas Medicaid. Modifier G7 must be submitted with the procedure code that identifies abortion services.

**Important:** To bill a Texas Medicaid client for a service that TMHP denies as not medically necessary, the billing provider must ensure that the client or client’s guardian has signed an acknowledgment statement. The statement must be obtained by a provider who has contact with the client.

### 4.1.6.1 Services Related to Abortion Procedures

An anesthesia service that is provided for an abortion procedure may be reimbursed if the abortion procedure meets medical necessity criteria and complies with the Texas Medicaid guidelines in the section above.

All other services that are related to abortion services are also subject to medical necessity review. All services that are related to a non-covered abortion procedure will be denied or recouped.

### 4.1.7 Other Maternity Care and Delivery Services

The following procedure codes may be submitted for other maternity care and delivery services:

<table>
<thead>
<tr>
<th>Procedure Codes</th>
</tr>
</thead>
<tbody>
<tr>
<td>59100</td>
</tr>
<tr>
<td>59897</td>
</tr>
</tbody>
</table>

### 4.1.8 Obstetric Ultrasound

The following procedure codes may be submitted for obstetric ultrasound services:

<table>
<thead>
<tr>
<th>Procedure Codes</th>
</tr>
</thead>
<tbody>
<tr>
<td>76801</td>
</tr>
<tr>
<td>76817</td>
</tr>
</tbody>
</table>

Texas Medicaid requires providers to follow the documentation requirements as set forth in the Diagnostic Ultrasound section of the Current Procedural Terminology (CPT) manual for the diagnostic studies of the fetus, including when ultrasound is used to guide a procedure.

Documentation requirements set forth in the CPT manual include, but are not limited to:

- Permanently recorded images with measurements, when measurements are clinically indicated
- Final written report included in the client’s medical record (includes written interpretation)
- Report must include description of elements that comprised a “complete” or “limited” exam and the reasons an element could not be visualized
- Permanently recorded images are also required for ultrasound guidance procedures of the site to be localized. In addition, description of the localization process, either separately or within the report of the procedure when the guidance is utilized.

Permanently recorded images must be available on request by the Texas Health and Human Services Commission (HHSC).

Prior authorization is required for greater than three obstetrical ultrasounds per pregnancy. Requests for additional obstetric ultrasounds may be considered when submitted with documentation of medical necessity on the Obstetric Ultrasound Prior Authorization Request Form.
Refer to: Obstetric Ultrasound Prior Authorization Request Form on the TMHP website at www.tmhp.com.

Authorization is not required for obstetric ultrasounds performed in the emergency department, outpatient observation, or inpatient hospital setting.

Texas Medicaid follows the American Congress of Obstetricians and Gynecologists (ACOG) indications for sonography.

First trimester ultrasounds may be medically necessary for, but not limited to, the following conditions:
- Confirm the presence of an intrauterine pregnancy
- Evaluation of a suspected ectopic pregnancy
- Evaluation of vaginal bleeding
- Evaluation of pelvic pain
- Estimation of gestational age
- Diagnosis or evaluation of multiple gestations
- Confirmation of cardiac activity
- Adjunct to chorionic villus sampling or localization and removal of an intrauterine device
- Assessment of certain fetal anomalies, such as anencephaly, in clients at high risk
- Evaluation of maternal pelvic or adnexal masses or uterine abnormalities
- Screening for fetal aneuploidy
- Evaluation of suspected hydatidiform mole

Second- and third-trimester ultrasounds may be medically necessary for the following conditions:
- Estimation of fetal age
- Evaluation of fetal growth
- Evaluation of vaginal bleeding
- Evaluation of cervical insufficiency
- Evaluation of abdominal and pelvic pain
- Determination of fetal presentation
- Adjunct to amniocentesis or other procedure
- Evaluation of suspected multiple gestation
- Evaluation of a significant discrepancy between uterine size and clinical dates
- Evaluation of pelvic mass
- Evaluation of suspected hydatidiform mole
- Adjunct to cervical cerclage placement
- Evaluation of a suspected ectopic pregnancy
- Evaluation of suspected fetal death
- Evaluation of suspected uterine abnormality
- Evaluation for fetal well-being
- Evaluation of suspected amniotic fluid abnormalities
• Evaluation of suspected placental abruption
• Adjunct to external cephalic version
• Evaluation for premature rupture of membranes or premature labor
• Evaluation for abnormal biochemical markers
• Follow-up evaluation of a fetal anomaly
• Follow-up evaluation of placental location for suspected placenta previa
• Evaluation for clients with a history of previous congenital anomaly
• Evaluation of fetal condition in late registrants for prenatal care
• Assessment of findings that may increase the risk of aneuploidy
• Screening for fetal anomalies

A request for retroactive authorization must be submitted no later than 14 calendar days, beginning the day after the study is completed.

Requests for prior authorization or retroactive authorization may be submitted by mail or an approved electronic method.

The Obstetric Ultrasound Prior Authorization Request Form must be completed, signed, dated, and maintained in the client’s medical record by the provider requesting the test. The form must include information related to medical necessity of the test including all of the following:

• Procedure requested (CPT code) and quantity requested
• The trimester(s) during which the requested ultrasounds will be performed
• The date range during which the procedure(s) will be performed
• Client’s estimated date of confinement (EDC) at the time the request is submitted
• Diagnosis

Additional documentation to support medical necessity may include any of the following:

• Treatment history
• Treatment plan
• Medications
• Previous imaging results

The Obstetric Ultrasound Prior Authorization Request Form must be completed, signed, and dated by the ordering provider (physician, nurse practitioner/clinical nurse specialist, certified nurse midwife [CNM], or physician assistant) when requesting prior authorization for obstetric ultrasounds, regardless of the method of request for authorization.

Residents may order obstetric ultrasounds; however, the attending physician must sign the authorization form and provide the group or supervising provider’s provider identifier.

Providers may be requested to provide additional documentation.

Obstetric ultrasounds provided in the emergency department or outpatient observation must be submitted with Modifier U6 when submitted on the professional claim form in order to be considered for reimbursement.
Obstetric ultrasounds provided in the emergency department or hospital observation must be submitted with the appropriate corresponding emergency services or hospital observation revenue code in order to be considered for reimbursement.

**Note:** *Any obstetric ultrasound performed in the emergency department or hospital observation will not count toward the limit of three per pregnancy.*

Any obstetric ultrasound claims submitted with or without prior authorization for the initial three will count toward the limit of three per pregnancy.

For transvaginal obstetric ultrasound performed in addition to one of the transabdominal examinations, documentation is required to substantiate the need to perform both tests on the same day.

Reimbursement for obstetric ultrasounds may be considered on appeal when submitted with documentation of any one of the following:

- Ultrasound was performed for a different pregnancy
- The provider was unable to obtain the previous ultrasound records from a different provider
- The provider was new to treating the client and was not aware the client had received three obstetric ultrasounds

Only one appeal will be considered per client for the same provider. Providers must obtain prior authorization for additional obstetric ultrasounds performed after the appealed service.

Add-on procedure codes (76802, 76810, 76812, and 76814) when billed with the primary procedure code for obstetric ultrasounds do not count toward the limit of three per pregnancy.

Claims for add-on codes for multiple fetuses should be billed with Modifier 76 if greater than one additional fetus. Claims for multiple fetuses greater than two will be considered on appeal with documentation indicating number of fetuses.

Three dimensional (3-D) rendering of obstetric ultrasound (procedure code 76376 or 76377) is not a benefit of Texas Medicaid.

Procedure code 76810 must be billed in conjunction with primary procedure code 76805, by the same provider.

Procedure code 76812 must be billed in conjunction with primary procedure code 76811, by the same provider.

Procedure code 76814 must be billed in conjunction with primary procedure code 76813, by the same provider.

### 4.1.9 Diagnostic Ultrasound and Ultrasonic Guidance

The following procedure codes may be submitted for diagnostic ultrasound and ultrasonic guidance services:

<table>
<thead>
<tr>
<th>Procedure Codes</th>
<th>76818</th>
<th>76819</th>
<th>76941</th>
<th>76946</th>
</tr>
</thead>
</table>

Ultrasonic guidance (procedure code 76941) is restricted to the diagnoses listed in the following table:

<table>
<thead>
<tr>
<th>Diagnosis Codes</th>
</tr>
</thead>
<tbody>
<tr>
<td>O353XX0</td>
</tr>
<tr>
<td>O360110</td>
</tr>
<tr>
<td>O360121</td>
</tr>
<tr>
<td>O360132</td>
</tr>
</tbody>
</table>
Ultrasonic guidance for intrauterine fetal transfusion (procedure code 76941) will be reimbursed separately when billed by a different physician.

Fetal biophysical profile (procedure code 76818 or 76819), when billed with 76805, 76810, 76811, 76812, 76813, 76814, 76815, or 76816, will be reimbursed separately.

### 4.1.10 Doppler Studies

The following procedure codes may be submitted for doppler study services:

<table>
<thead>
<tr>
<th>Procedure Codes</th>
</tr>
</thead>
<tbody>
<tr>
<td>76820</td>
</tr>
<tr>
<td>76821</td>
</tr>
</tbody>
</table>

As supported by ACOG, umbilical artery doppler (procedure code 76820) is limited to suspected intrauterine growth restriction (IUGR), post-term gestation, diabetes mellitus, systemic lupus erythematosus, or antiphospholipid antibody syndrome.

Middle cerebral artery doppler (procedure code 76821) is indicated, but not limited to fetuses who are alloimmunized.

### 4.1.11 Echocardiography

The following procedure codes may be submitted for echocardiography services:

<table>
<thead>
<tr>
<th>Procedure Codes</th>
</tr>
</thead>
<tbody>
<tr>
<td>76825</td>
</tr>
<tr>
<td>76826</td>
</tr>
<tr>
<td>76827</td>
</tr>
<tr>
<td>76828</td>
</tr>
</tbody>
</table>

Fetal echocardiography (procedure codes 76825, 76826, 76827, and 76828) may be reimbursed for the following factors/syndromes:

- Fetal Risk Factors
  - Extracardiac anomalies
  - Chromosomal
- Anatomic
- Fetal cardiac dysrhythmia
  - Irregular rhythm
  - Tachycardia
  - Bradycardia
- Nonimmune hydrops fetalis
- Suspected cardiac anomaly on ultrasound
- Abnormal fetal situs

- Maternal Risk Factors
  - Congenital heart disease
  - Cardiac teratogen exposure
    - Lithium
    - Alcohol
    - Phenytoin
    - Trimethadione
    - Isoretinoin
  - Maternal Metabolic Disorders
    - Diabetes mellitus
    - Phenylketonuria
- Familial Risk Factors for congenital heart disease
  - Previous sibling
  - Paternal
- Syndromes
  - Marfan’s
  - Noonan’s
  - Tuberous sclerosis

### 4.1.12 Hydroxyprogesterone Caproate

The following procedure codes may be submitted for hydroxyprogesterone caproate:

<table>
<thead>
<tr>
<th>Procedure Code</th>
<th></th>
</tr>
</thead>
<tbody>
<tr>
<td>J1726</td>
<td>J1729</td>
</tr>
</tbody>
</table>

The following documentation supporting medical necessity for administration of a hydroxyprogesterone caproate injection must be maintained in the client’s medical record:

- The client’s treatment is initiated between 16 weeks, 0 days and 20 weeks, 6 days gestation
- The client’s treatment continues, as medically indicated, through 36 weeks, 6 days gestation or delivery, whichever occurs first
- The client has a singleton pregnancy
The client has had a prior, singleton spontaneous preterm delivery before 37 weeks gestation. Hydroxyprogesterone caproate is limited to a maximum of 21 doses per pregnancy.

Prior authorization is not required for either the compounded or trademarked versions of hydroxyprogesterone caproate for injection (procedure codes J1726 and J1729).

Requests for initiation of the client’s treatment after 20 weeks, 6 days gestation but not beyond 24 weeks gestation will be considered on a case-by-case basis. A prior authorization request must be submitted by the physician to the Special Medical Prior Authorization (SMPA) department with documentation to support the medical necessity of starting treatment at this stage of gestation, and must be approved by the Medical Director.

Procedure code J1726 and J1729 are restricted to the following payable diagnosis:

<table>
<thead>
<tr>
<th>Diagnosis Codes</th>
</tr>
</thead>
<tbody>
<tr>
<td>O09211</td>
</tr>
<tr>
<td>O09212</td>
</tr>
<tr>
<td>O09213</td>
</tr>
<tr>
<td>O09219</td>
</tr>
</tbody>
</table>

Hydroxyprogesterone caproate must be submitted with an NDC. Hydroxyprogesterone caproate is administered once weekly (every 7 days) by injection.

### 4.1.13 Fetal Surgery

The following procedure codes may be submitted for fetal surgery:

<table>
<thead>
<tr>
<th>Procedure Codes</th>
</tr>
</thead>
<tbody>
<tr>
<td>59072</td>
</tr>
<tr>
<td>S2401</td>
</tr>
<tr>
<td>S2402</td>
</tr>
<tr>
<td>S2403</td>
</tr>
<tr>
<td>S2405</td>
</tr>
<tr>
<td>S2409</td>
</tr>
<tr>
<td>S2411</td>
</tr>
</tbody>
</table>

Fetal surgery procedures require prior authorization.

Prior authorization requests must be submitted on a Special Medical Prior Authorization (SMPA) Request Form to the SMPA department.

Refer to: Special Medical Prior Authorization (SMPA) Request Form on the TMHP website at www.tmhp.com.

Procedure codes S2401, S2402, S2403, S2405 and S2409 may be authorized for Texas Medicaid only when the hospital is a member of the North American Fetal Therapy Network (NAFTNet).

The pediatric surgeon for procedure codes S2401, S2402, S2403, S2405, or S2409 must submit documentation which includes:

- A clear description of the fetal malformation(s). The malformation(s) must interfere with the intrauterine organ development and fetal survival and have potential fatal consequences before or after birth
- Evidence that in utero correction of the fetal congenital malformation(s) results in a clinical outcome that is better than that which would be seen in expectant management

**Note:** Services and procedures that are investigational or experimental are not a benefit of Texas Medicaid.

Umbilical cord occlusion (procedure code 59072) may be considered when all the following is documented:

- Diagnosis of monoamniotic-monochorionic twins is present
- Spontaneous fetal death of one of the twins with the presence of hydrops
- The ratio of the acardia twin weight to the pump twin weight is greater than 50 percent

Diagnosis Codes

<table>
<thead>
<tr>
<th>O09211</th>
<th>O09212</th>
<th>O09213</th>
<th>O09219</th>
</tr>
</thead>
</table>

Procedure Codes

| 59072 | S2401 | S2402 | S2403 | S2405 | S2409 | S2411 |
• The abdominal circumference of the twin with reversed arterial perfusion fetus is greater than or equal to the abdominal circumference of the pump twin

  **Note:** Elective abortions are not benefits of Texas Medicaid

Fetoscopic laser therapy for treatment of twin-to-twin transfusion syndrome (procedure code S2411) may be considered when all the following are documented:

• Fetal gestational age of less than 26 weeks
• Evidence of polyhydramnios in the recipient fetus
• Donor fetus is oligohydramniotic
• Evidence of abnormal blood flow documented by Doppler studies in one or both fetuses

4.1.14 **Antenatal and Postnatal Care Visits**

The following procedure codes may be submitted for antenatal and postnatal care visits:

<table>
<thead>
<tr>
<th>Procedure Codes</th>
</tr>
</thead>
<tbody>
<tr>
<td>59430 99202 99203 99204 99205 99211 99212 99213 99214 99215</td>
</tr>
<tr>
<td>99341 99342 99343 99344 99345 99347 99348 99349 99350</td>
</tr>
</tbody>
</table>

Texas Medicaid reimburses prenatal care, deliveries, and postpartum care as individual procedures. Prenatal and postpartum care visits billed in an inpatient hospital (POS 3), will be denied as part of another procedure when billed within the three days before delivery or the six weeks after delivery. The inpatient intrapartum and postpartum care are included in the fee for the delivery or Cesarean section and should not be billed separately.

Physicians (obstetricians, family practice physicians, and maternal-fetal medicine specialists), CNMs, and maternity service clinics (MSCs) are limited to 20 prenatal care visits per pregnancy and one postpartum care visit after discharge from the hospital.

Licensed midwives (LMs) are limited to 20 outpatient antepartum care visits per pregnancy to be performed in a birthing center; postpartum visits are not separately reimbursed. Routine pregnancies are anticipated to require around 11 visits per pregnancy and high-risk pregnancies are anticipated to require around 20 visits per pregnancy. More frequent visits may be necessary for high-risk pregnancies. High-risk obstetrical visits are not limited to 20 visits per pregnancy. The provider can appeal with documentation supporting a complication of pregnancy. Documentation reflecting the need for increased visits must be maintained in the physician’s files and is subject to retrospective review.

Providers must bill the most appropriate new or established patient prenatal or postnatal visit procedure code. New patient codes may be used when the client has not received any professional services from the same physician, or another physician of the same specialty who belongs to the same group, within the last three years (36 months).

When billing for prenatal services, use modifier TH with the appropriate E/M procedure code to the highest level of specificity.

  **Note:** Failure to use the TH modifier may result in recoupment of payment rendered.

LMs are not reimbursed for postpartum visits.

One postpartum care procedure code may be reimbursed per pregnancy. The claim for the postpartum visit may be submitted with either procedure code 59430 or with a delivery procedure code (59410, 59515, 59614, or 59622) that includes postpartum care. The reimbursement amount for the submitted procedure code covers all postpartum care per pregnancy regardless of the number of postpartum visits provided.
Postpartum depression screening is a benefit at the infant’s Texas Health Steps medical checkup or follow-up visit, as a separately reimbursed service in the 12 months following the infant’s birth.

Refer to: Section 5.3.11.1.4, “Postpartum Depression Screening” in the Children’s Services Handbook (Vol. 2, Provider Handbooks) for more information regarding postpartum depression screening for mothers during the infant’s Texas Health Steps medical checkup or follow-up visit.

Any other E/M office visit will not be reimbursed when billed date of service, by the same provider, as any antenatal or postpartum office visit. Modifier 25 may be used to identify a significant separately identifiable E/M service by the same physician on the same date of service as the procedure or other service. Documentation that supports the provision of a significant, separately identifiable E/M service must be maintained in the client’s medical record and made available to Texas Medicaid upon request.

Note: All unlisted surgical procedure codes have a 42 global period assigned by Texas Medicaid.

4.1.14.1 Maternity Service Clinic (MSC)

MSCs are limited provider clinics, unrelated to a hospital, that only provide maternity services. An MSC will be reimbursed for antepartum and/or postpartum care visits only. Hemoglobin, hematocrit, and urinalysis procedures are included in the charge for antepartum care and not separately reimbursed. Services other than antepartum and postpartum care visits will be denied.

4.1.15 Birthing Centers—Professional Services

The following procedures may be performed by professionals in the birthing center setting:

<table>
<thead>
<tr>
<th>Procedure Codes</th>
<th>59409</th>
<th>59410***</th>
<th>99218</th>
<th>99219</th>
<th>99220</th>
</tr>
</thead>
<tbody>
<tr>
<td>***Licensed Midwives may not use code 59410.</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

The following table includes procedure codes that may be benefits for licensed midwife (LM) services rendered in the birthing center setting:

<table>
<thead>
<tr>
<th>Procedure Codes</th>
<th>59409</th>
<th>99202</th>
<th>99211</th>
<th>99212</th>
<th>99218</th>
<th>99219</th>
<th>99220</th>
</tr>
</thead>
</table>

Note: Licensed Midwives may not use code 59410.

Antenatal services provided by LMs may be a benefit when billed with modifier TH. If the client is discharged prior to delivery, procedure codes 99218, 99219, or 99220 may be billed by the professional for labor services only. Clinical documentation that clearly demonstrates level of medical decision making (i.e., moderate or complex) must be included in the client’s medical record. All medical documentation is subject to retrospective review. Those services not supported by the documentation in the client’s medical record are subject to recoupment.

Refer to: Subsection 9.2.44, “Newborn Services” in the Medical and Nursing Specialists, Physicians, and Physician Assistants Handbook (Vol. 2, Provider Handbooks) for more information related to the care of the newborn.

4.1.16 Birthing Centers—Facility Services

The following procedures may be performed by birthing center facilities:

| Procedure Codes | 59409 | S4005 |
Deliveries at a facility licensed as a birthing center by the Department of State Health Services (DSHS) must be billed with procedure code 59409.

If the client is discharged prior to delivery, procedure code S4005 may be billed by the facility for labor services only.

### 4.1.17 Tobacco Use Cessation

Counseling for cessation of the habit of using tobacco products by pregnant women is a benefit of Texas Medicaid.

The following procedure codes listed may be billed for tobacco use cessation counseling using an applicable diagnosis code:

<table>
<thead>
<tr>
<th>Procedure Codes</th>
</tr>
</thead>
<tbody>
<tr>
<td>99406</td>
</tr>
</tbody>
</table>

One of the following diagnosis codes must be submitted for tobacco use cessation with the appropriate procedure code:

<table>
<thead>
<tr>
<th>Diagnosis Codes</th>
</tr>
</thead>
<tbody>
<tr>
<td>O99330</td>
</tr>
</tbody>
</table>

Refer to: Subsection 9.2.56.3.3, “Tobacco Use Cessation” in the Medical and Nursing Specialists, Physicians, and Physician Assistants Handbook (Vol. 2, Provider Handbooks) for additional information related to tobacco use cessation counseling and additional covered diagnoses.

Procedure codes 99406 and 99407 are limited to once per day, same or different procedure code, any provider.

Tobacco use cessation services delivered by group modality are limited to a maximum of 8 individuals and must be billed with modifier HQ.

Procedure codes 99406 and 99407 may be billed in any combination, individual or group, by the same or different provider, and are limited to eight services per rolling year. Additional services require documentation of medical necessity to exceed the established limit.

### 4.1.18 Zika Virus Testing

Refer to: Subsection 2.2.13.1, “Zika Virus Testing” in the Radiology and Laboratory Services Handbook (Vol. 2, Provider Handbooks) for information about Zika virus testing.

### 5 Noninvasive Prenatal Screening (NIPS)

Noninvasive prenatal screening (NIPS) is a benefit of Texas Medicaid when medically necessary for the advanced screening of fetal chromosomal abnormalities in pregnant women who meet specific screening criteria. Genetic screening results, when informative, may influence clinical management decisions.

NIPS may be performed as early as ten weeks gestation for specific fetal aneuploidy screening, restricted to Trisomy 13, Trisomy 18, Trisomy 21, and fetal sex chromosome aneuploidy (SCA). To determine NIPS appropriateness, a baseline ultrasound, if not previously performed, is strongly recommended to confirm viability, the number of fetuses, and gestational dating.

If NIPS provides an abnormal screening result, invasive prenatal confirmatory diagnostic testing is strongly encouraged due to the potential risk of a false-positive result from NIPS. Confirmatory diagnostic tests include chorionic villus sampling (CVS) or amniocentesis.
It is recommended that clients who receive an indeterminate result be offered further genetic counseling, comprehensive evaluation with ultrasound, and diagnostic testing due to the increased risk of aneuploidy. Higher body mass index (BMI) may affect NIPS results. Clients weighing more than 250 pounds are at risk of having an inconclusive result from NIPS.

NIPS does not assess the risk for fetal anomalies such as neural tube defects or ventral wall defects. Ultrasound evaluation and maternal serum alpha-fetoprotein screening should be offered for these risk assessments.

If a fetal structural anomaly (e.g. hydrops, cystic hygroma, cardiac malformations, abdominal wall defects, or skeletal abnormalities) is identified upon ultrasound examination, it is recommended that diagnostic testing be offered rather than NIPS.

NIPS must be ordered by the medical provider rendering direct care to the client. The provider must order the most appropriate screening based on the client’s medical history and the results of previous screenings, if available. The provider must clarify for the client the option to decline, and the provider must document that the option to decline was clearly provided in the client’s medical record.

Note: Some noninvasive prenatal screenings include an extended panel that screens for microdeletions and additional trisomies, such as T16 and T22. However, this use has not been validated, and the “opt-out” box on the requisition form should be checked.

5.1 Screening for Fetal Sex Chromosome Aneuploidy

In addition to trisomy (e.g., T13, T18, T21), NIPS may also screen for fetal SCA (e.g., 45, X; 47, XXX; 47, XXY; 47, XYY).

Note: Currently, clinical evidence is unclear for concluding the net health benefits when using cell-free fetal DNA to screen for fetal sex chromosome aneuploidy (SCA). The potential benefit of early detection for management must be balanced against potential harms, stigmatization, and distorted perceptions of the child.

Sex chromosome aneuploidy of maternal origin should be considered when NIPS results suggest fetal sex chromosome aneuploidy (e.g., 45, X; 47, XXX, 47, XXY; 47, XYY). Other considerations include the risk for incidental findings with NIPS. Appropriate client counseling is encouraged.

5.1.1 Screening Criteria

NIPS is a benefit for singleton pregnancies. At least one of the following criteria must be met for a client to be eligible for NIPS:

- Fetal ultrasound indicates risk of aneuploidy
- Fetal ultrasound indicates structural anomalies associated with aneuploidy, and the mother wishes to postpone invasive diagnostic testing
- History of pregnancy with aneuploidy
- Maternal age of 35 years or older at time of delivery
- Parental balanced Robertsonian translocation of chromosome 13 or 21
- Abnormal serum screening results for the current pregnancy:
  - First trimester screen
  - Sequential screen
  - Integrated screen
  - Quadruple screen
5.2 Genetic Counseling Requirement

Genetic counseling must be provided by a trained genetic counselor, nurse specialist in genetics, maternal-fetal medicine specialist, or other medical provider (e.g., obstetrician) possessing expertise in genetic counseling who is not affiliated with the genetic screening laboratory. Both pre- and post-screening counseling must provide the depth of content and time for the client to make an informed decision.

The client must be provided with information about the purpose and nature of the screenings. Documentation in the medical record must reflect that the client has been given information on the benefits, risks, and limitations of advanced screening; as well as the nature, inheritance, and implications of genetic disorders. Documentation requirements include all of the following:

- Pre-screening genetic counseling:
  - The date that formal pre-screening counseling was provided, with the name and qualifications of the counseling professional
  - The explanation of risks, benefits, and limitations that was discussed with the client
  - The client’s ability to understand the risks, benefits, and limitations and the client’s informed choice to proceed with NIPS as evidenced by the client’s signature on a consent form specific to the NIPS to be performed
  - The client’s other prenatal radiological or lab results, if available, to support medical necessity of NIPS

- Post-screening genetic counseling:
  - The client’s NIPS results
  - The date that formal post-screening counseling was provided, with the name and qualifications of the counseling professional
  - The clear, non-directive explanation provided to the client concerning the findings and implications of the NIPS results
  - The client’s ability to understand the results and explanation provided

The genetic counseling must be nondirective. The purpose of the provider’s information is not to direct the client, but to allow the client to make informed medical and personal decisions.

Clients should be informed that a negative NIPS result does not ensure an unaffected pregnancy.

5.3 Prior Authorization

Prior authorization is required for NIPS procedure codes 81420 and 81507. The prior authorization request must be submitted on the Special Medical Prior Authorization (SMPA) Request Form completed, signed, and dated by the provider rendering direct care to the client, and include the performing laboratory’s TPI in section D of the form. The ordering provider must share the authorization number with the laboratory provider submitting the claim. Requests from laboratories will not be processed.

The expected dates of service requested in Section B of the Special Medical Prior Authorization (SMPA) Request Form must not exceed 60 days. Prior authorizations will only be approved for 60 days, during which time the client must obtain the screening.

**Note:** For prior authorization requests submitted before the client’s 10th week of gestation, the expected dates of service must begin no sooner than the 10th week of gestation. Approved prior authorizations will expire 60 days from the start of service date indicated on the SMPA form.
The provider must indicate on the prior authorization request form that the client meets required criteria (as noted above in Screening Criteria).

The request for prior authorization should document that the client was provided counseling regarding potential outcomes of aneuploidy screening, as well as potential outcomes of fetal sex chromosome aneuploidy screening when included, and that she understands the implications associated with each possible aneuploidy result.

Prior authorization requests may be submitted to the TMHP Special Medical Prior Authorization Department via mail, fax, or the electronic portal. Ordering providers may sign prior authorization forms and supporting documentation using electronic or wet signatures.

Medical documentation submitted by the ordering provider must verify any indications the provider included on the form, such as the client’s age, history of affected pregnancy or family history, anomalous ultrasound findings, or abnormal maternal serum results. Requisition forms from the laboratory are not sufficient for verification of genetic history.

A no-call or inconclusive result is possible and further diagnostic testing is strongly recommended in these cases.

NIPS procedure codes 81420 and 81507 are limited to once per pregnancy. Additional tests will not be authorized.

**Note:** Providers may appeal denied claims with documentation of a new pregnancy.

### 5.3.1 Additional Documentation Requirements

In addition to the documentation of pre and post genetic counseling, and the option to decline NIPS provided to the client, the following NIPS documentation must also be maintained in the client’s medical record and is subject to retrospective review:

- The appropriateness and benefit of NIPS specific to the client
- The client’s specific high-risk criteria

### 5.4 NIPS Limitations

Procedure codes 81507 and 81420 are restricted to female clients only who are 10 through 55 years of age. Procedure code 81420 will be denied when billed during the same pregnancy as procedure code 81507, by any provider. Claims that have been paid for procedure code 81420 are subject to recoupment if procedure code 81507 is submitted later for the same pregnancy.

### 5.5 Non-Covered Services

The following NIPS services are not a benefit of Texas Medicaid:

- NIPS as part of a routine prenatal laboratory assessment
- NIPS if performed without informed patient choice and pre- and post-screen genetic counseling from a qualified professional
- NIPS for women who do not meet the criteria outlined above
- NIPS for women with multiple gestations (e.g., twins, triplets, etc.)
- NIPS for screening of chromosomal microdeletion syndromes
- NIPS for screening of trisomy other than T13, T18, or T21
- NIPS for sex determination, paternity determination, or non-medical reasons
- NIPS is not reimbursed with procedure code 81599
6  Gynecological Health Services

6.1  Services, Benefits, Limitations, and Prior Authorization
Gynecological examinations, surgical procedures, and treatments are benefits of Texas Medicaid. The following gynecological procedures and services may be benefits of Texas Medicaid:

- Gynecological and family planning examinations
- Contraceptives
- Diagnostic tests
- Surgical procedures
- Gynecological treatments

Refer to: Section 2, "Medicaid Title XIX Family Planning Services" in this handbook for information about contraception, sterilizations, and family planning annual examinations.

6.2  Surgical and Laparoscopic Treatment of Ectopic Pregnancy
Surgical and laparoscopic treatment of ectopic pregnancy (procedure codes 59120, 59121, 59130, 59135, 59136, 59140, 59150, and 59151) is a benefit of Texas Medicaid.

6.3  Laparoscopic Procedures
Laparoscopic procedures (procedure codes 58545, 58546, 58578, and 58674) are a benefit of Texas Medicaid.

6.4  Endometrial Cryoablation
Endometrial cryoablation (procedure codes 58353 and 58356) is a benefit of Texas Medicaid.

6.5  Uterine Suspension
Uterine suspension (procedure codes 58400 and 58410) is a benefit of Texas Medicaid.

6.6  Vulvectomy
Vulvectomy (procedure code 56620) is a benefit of Texas Medicaid.

6.6.1  Prior Authorization for Vulvectomy
Prior Authorization is required for vulvectomy. The prior authorization request must include documentation of one of the following conditions:

- Vulvar intraepithelial neoplasia (VIN)
- Labial enlargement that results in abrasion, irritation, or intractable skin infection

Note: A vulvectomy will not be considered for cosmetic reasons.

6.7  Salpingostomy
Salpingostomy (procedure codes 58673 and 58770) is a benefit of Texas Medicaid.

6.7.1  Prior Authorization for Salpingostomy
Prior authorization is required for salpingostomy.
The prior authorization request must include documentation of one or more of the following conditions:

- Ectopic pregnancy
- Hydrosalpinx unrelated to infertility
- Salpingitis unrelated to infertility
- Torsion of the fallopian tube
- Abscess of the fallopian tube
- Peritubal adhesions unrelated to infertility
- Cyst or tumor of the fallopian tube unrelated to infertility
- Hematosalpinx

6.8 **Ovarian Wedge Resection**

Ovarian wedge resection (procedure code 58920) is a benefit of Texas Medicaid.

6.8.1 **Prior Authorization for Ovarian Wedge Resection**

Prior Authorization is required for ovarian wedge resection.

The prior authorization request must include documentation of polycystic ovarian syndrome (PCOS).

*Note:* Ovarian wedge resection will not be considered to improve chances of conceiving if the PCOS lead to infertility.

6.9 **Assays for the Diagnosis of Vaginitis**

Vaginitis assay procedure codes 87480, 87510, 87660, 87661, 87797, 87798, and 87800 are benefits of Texas Medicaid.

Providers are reminded to code to the highest level of specificity with a diagnosis to support medical necessity when submitting procedure code 87797.

Claims may be subject to retrospective review if they are submitted with diagnosis codes that do not support medical necessity.

If a positive test result was not treated, documentation must be present indicating why treatment was not rendered.

6.10 **Hysteroscopy**

Hysteroscopy (procedure codes 58555, 58558, 58559, 58560, 58561, 58562, 58563, and 58579) is a benefit of Texas Medicaid.

6.11 **Examination Under Anesthesia**

Pelvic examination under anesthesia (procedure code 57410) is considered part of another gynecological surgery performed the same day.

If the examination is performed as an independent procedure or at the time of a nongynecological surgery, the procedure may be reimbursed.

6.12 **Laminaria Insertion**

Insertion of a laminaria or dilatation (procedure code 59200) is a benefit of Texas Medicaid.
6.13 Hysterectomy Services

Texas Medicaid reimburses hysterectomies when they are medically necessary. Texas Medicaid does not reimburse hysterectomies performed for the sole purpose of sterilization.

Providers can use any of the following procedure codes to submit claims for hysterectomy procedures:

<table>
<thead>
<tr>
<th>Procedure Codes</th>
</tr>
</thead>
<tbody>
<tr>
<td>51925 58150 58152 58180 58200 58210 58240 58260 58262 58263</td>
</tr>
<tr>
<td>58267 58270 58275 58280 58285 58290 58291 58292 58294 58295</td>
</tr>
<tr>
<td>58542 58543 58544 58548 58550 58552 58553 58554 58570 58571</td>
</tr>
<tr>
<td>58572 58573 58575 59135 59525</td>
</tr>
</tbody>
</table>

Providers can refer to the Texas Medicaid fee schedules on the TMHP website at www.tmhp.com for components and fees that may be reimbursed.

6.13.1 Hysterectomy Acknowledgment

Hysterectomy services are considered for reimbursement when a signed Texas Medicaid - Title XIX Acknowledgment of Hysterectomy Information form is faxed to TMHP, the claim is filed with a signed Texas Medicaid - Title XIX Acknowledgment of Hysterectomy Information form, or documentation supporting that the Texas Medicaid - Title XIX Acknowledgment of Hysterectomy Information form could not be obtained or was not necessary.

All Texas Medicaid clients (including those in a STAR or STAR+PLUS Program health plan) receiving hysterectomy services must sign a Texas Medicaid - Title XIX Acknowledgment of Hysterectomy Information form. The acknowledgment must be submitted to TMHP with the claim or to the client’s health plan.

The Texas Medicaid - Title XIX Acknowledgment of Hysterectomy Information form must be signed and dated by the client. The statement must indicate that the client was informed both orally and in writing before the surgery that the hysterectomy would leave her permanently incapable of bearing children.

**Note:** A client representative’s signature will be required for mentally incompetent clients.

The client’s eligibility file is updated upon receipt of the signed Texas Medicaid - Title XIX Acknowledgment of Hysterectomy Information form. Claims for services related to the hysterectomy cannot be reimbursed unless the signed Texas Medicaid - Title XIX Acknowledgment of Hysterectomy Information form is on file; therefore to avoid claim denials, each individual provider involved in the hysterectomy procedure is encouraged to submit a copy of the valid Texas Medicaid - Title XIX Acknowledgment of Hysterectomy Information form rather than relying on another provider to do so.

The provider is responsible for maintaining the original, signed copy of the Texas Medicaid - Title XIX Acknowledgment of Hysterectomy Information form in the client’s medical record when a claim is submitted for consideration of payment. These records are subject to retrospective review.

When a hysterectomy, whether abdominal or vaginal, is performed without a client’s acknowledgement form:

- The hysterectomy procedure code is denied.
- The other surgical procedures are evaluated for their clinical relevance.
- Multiple procedures are processed according to the multiple surgery guidelines.
A Texas Medicaid - Title XIX Acknowledgment of Hysterectomy Information form is not required if the performing physician certifies that at least one of the following circumstances existed before the surgery:

- The patient was already sterile before the hysterectomy, and the cause of the sterility is stated (e.g., congenital disorder, sterilized previously, or postmenopausal). Providers must use a post menopause or sterilization diagnosis code on the claim form. If the provider submits a claim and does not attach the acknowledgment, the provider must maintain the signed statement in the client’s records, and the physician’s signature will not be required on the claim form. These records are subject to retrospective review.

- The patient requires a hysterectomy on an emergency basis because of a life-threatening situation. The physician must state the nature of the emergency and certify that it was determined that prior acknowledgment was not possible. Because the acknowledgment may be signed the day of or an hour before surgery, an emergency situation requires that the patient be unconscious or under sedation and unable to sign the acknowledgment.

Although the Texas Medicaid - Title XIX Acknowledgment of Hysterectomy Information form is not required if the criteria previously listed are met, the performing physician must certify that one or more of the circumstances existed prior to the surgery. This certification may be submitted before the claim is submitted or attached to the claim and signed by the performing provider.

Refer to: Title 42 of CFR 441.255 and 25 TAC Part 1, Chapter 29, Subchapter F, section 25.501 for more information.

Refer to: Texas Medicaid - Title XIX Acknowledgment of Hysterectomy Information on the TMHP website at www.tmhp.com.

For clients with retroactive Medicaid coverage, one of the following must be submitted with the Texas Medicaid - Title XIX Acknowledgment of Hysterectomy Information form:

- A copy of the client’s Your Texas Benefits Medicaid card, which covers the date of the hysterectomy.

- A copy of the retroactive approval notice for Medicaid coverage.

Faxing Forms

All Medicaid providers may fax the Texas Medicaid - Title XIX Acknowledgment of Hysterectomy Information form to 1-512-514-4218. The form must include the client's Texas Medicaid number. All consent forms should be faxed with a cover sheet that identifies the provider and includes the telephone number and address. If the fax is incomplete or the consent form is invalid, the form is returned by mail or fax for correction. Completed consent forms that are faxed for adjustments or appeals are validated in the TMHP system. However, claims associated with the consent forms must be appealed through the mail to Appeals/Adjustments at the following address:

Texas Medicaid & Healthcare Partnership
Attn: Appeals/Adjustments
PO Box 200645
Austin, TX 78720-0645

6.14 Pap Smear (Cytopathology Studies)

Pap smears are benefits of Texas Medicaid for early detection of cancer. Family planning clients are eligible for annual Pap smears. Procurement and handling of the Pap smear are considered part of the E/M of the client and are not reimbursed separately.
The following procedure codes are reimbursed only to pathologists and CLIA-certified laboratories (whose directors providing technical supervision of cytopathology services are pathologists):

<table>
<thead>
<tr>
<th>Procedure Codes</th>
</tr>
</thead>
<tbody>
<tr>
<td>88142</td>
</tr>
<tr>
<td>88166</td>
</tr>
</tbody>
</table>

These procedure codes must be billed with the place of service where the Pap smear is interpreted.

Procedure code 88141 is reimbursed in addition to and when billed with the cytopathology procedure codes in the table above.

Procedure code 88155 will only be reimbursed when billed in conjunction with one of the following procedure codes on the same date of service by the same provider:

<table>
<thead>
<tr>
<th>Procedure Codes</th>
</tr>
</thead>
<tbody>
<tr>
<td>88142</td>
</tr>
<tr>
<td>88167</td>
</tr>
</tbody>
</table>

**Note:** Pap smear procedures will not be reimbursed separately to either the physician or a laboratory when billed on the same day as a THSteps medical check-up visit.

### 6.15 Clitoroplasty and Vaginoplasty

Clitoroplasty and vaginoplasty are performed for clients who possess ovaries and are female by genetic sex, but the external genitalia are not those of a normal female. Surgical correction of abnormalities of the external genitalia is the only indicated treatment for this disorder. Clitoroplasty and vaginoplasty procedure codes 56805 and 57335 may be considered for reimbursement for female clients who are 20 years of age and younger when submitted for reimbursement with one of the following diagnosis codes:

<table>
<thead>
<tr>
<th>Diagnosis Codes</th>
</tr>
</thead>
<tbody>
<tr>
<td>E250</td>
</tr>
<tr>
<td>E3451</td>
</tr>
</tbody>
</table>

### 6.16 Documentation Requirements

All services require documentation to support the medical necessity of the service rendered, including gynecological services.

Gynecological health services are subject to retrospective review and recoupment if documentation does not support the service billed.

### 6.17 Claims Filing and Reimbursement

Gynecological services must be submitted to TMHP in an approved electronic format or on the CMS-1500 claim form. Providers may purchase CMS-1500 claim forms from the vendor of their choice. TMHP does not supply the forms. When completing a CMS-1500 claim form, all required information must be included on the claim, as TMHP does not key any information from claim attachments. Super-bills, or itemized statements, are not accepted as claim supplements.
Texas Medicaid rates for physicians and other practitioners are calculated in accordance with TAC §355.8085. Providers can refer to the Online Fee Lookup (OFL) or the applicable fee schedule on the TMHP website at www.tmhp.com.

Refer to: Subsection 2.2.1.1, “Non-emergent and Non-urgent Evaluation and Management (E/M) Emergency Department Visits” in “Section 2: Texas Medicaid Fee-For-Service Reimbursement” (Vol. 1, General Information).

Section 104 of the Tax Equity and Fiscal Responsibility Act (TEFRA) of 1982 requires that Medicare and Medicaid limit reimbursement for those physician services furnished in outpatient hospital settings (e.g., clinics and emergency situations) that are ordinarily furnished in physician offices.

### 6.17.1 NCCI and MUE Guidelines

The HCPCS and CPT codes included in the Texas Medicaid Provider Procedures Manual are subject to NCCI relationships, which supersede any exceptions to NCCI code relationships that may be noted in the manual. Providers should refer to the CMS NCCI web page or correct coding guidelines and specific applicable code combinations.

In instances when Texas Medicaid limitations are more restrictive than NCCI MUE guidance, Texas Medicaid limitations prevail.

### 6.18 National Drug Code

Refer to: Subsection 6.3.4, “National Drug Code (NDC)” in “Section 6: Claims Filing” (Vol. 1, General Information).

### 7 Claims Resources

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<thead>
<tr>
<th>Resource</th>
<th>Location</th>
</tr>
</thead>
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<td>Subsection 6.8, “Family Planning Claim Filing Instructions” in “Section 6: Claims Filing” (Vol. 1, General Information)</td>
</tr>
<tr>
<td>Acronym Dictionary</td>
<td>“Appendix C: Acronym Dictionary” (Vol. 1, General Information)</td>
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<tr>
<td>Automated Inquiry System (AIS)</td>
<td>Subsection A.10, “TMHP Telephone and Fax Communication” in “Appendix A: State, Federal, and TMHP Contact Information” (Vol. 1, General Information)</td>
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<tr>
<td>CMS-1500 Paper Claim Filing Instructions</td>
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</tr>
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<tr>
<td>TMHP electronic claims submission information</td>
<td>Subsection 6.2, “TMHP Electronic Claims Submission” in “Section 6: Claims Filing” (Vol. 1, General Information)</td>
</tr>
</tbody>
</table>
8 Contact TMHP

The TMHP Contact Center at 1-800-925-9126 is available Monday – Friday from 7 a.m. to 7 p.m., Central Time.

9 Forms

The following linked forms can also be found on the Forms page of the Provider section of the TMHP website at www.tmhp.com:

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<tr>
<td>Texas Medicaid - Title XIX Acknowledgment of Hysterectomy Information</td>
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<td>2017 Claim Form</td>
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10 Claim Form Examples

The following linked claim form examples can also be found on the Claim Form Examples page of the Provider section of the TMHP website at www.tmhp.com:

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<th>Claim Form Examples</th>
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<td>Nurse Practitioner/Clinical Nurse Specialist (Family Planning)</td>
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# HEALTH AND HUMAN SERVICES COMMISSION FAMILY PLANNING PROGRAM SERVICES HANDBOOK

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1 Provider Enrollment for HHSC Family Planning Program Contractors

Agencies that submit claims for HHSC Family Planning Program Services must have a contract with HHSC. The HHSC Family Planning Program determines client eligibility and benefits. Refer to the HHSC Family Planning Program Policy Manual for specific eligibility, services, and policy information at https://hhs.texas.gov/laws-regulations/handbooks/family-planning-program-policy-manual.

Refer to: “Section 1: Provider Enrollment and Responsibilities” (Vol. 1, General Information) for more information about enrollment procedures.


Subsection 1.1, “Family Planning Overview” in the Gynecological, Obstetrics, and Family Planning Title XIX Services Handbook (Vol. 2, Provider Handbooks) for more information about family planning funding sources, guidelines for family planning providers, and family planning services for undocumented aliens and legalized aliens.

2 Services, Benefits, Limitations, and Prior Authorization

This section contains information about family planning services funded through the HHSC Family Planning Program funding source, including:

- Family planning annual exams
- Other family planning office or outpatient visits
- Laboratory procedures
- Radiology services
- Contraceptive devices and related procedures
- Drugs and supplies
- Medical counseling and education
- Immunizations
- Breast and cervical cancer screening and diagnostic services
- Prenatal services
- Sterilization and sterilization-related procedures (i.e., tubal ligation, vasectomy, and anesthesia for sterilization)

Providers are encouraged to include the appropriate diagnosis codes on the claim in conjunction with all family planning procedures and services.

Refer to: The HHSC Family Planning Program Policy Manual.

The choice of diagnosis code must be based on the type of family planning service performed.

2.1 Family Planning Annual Exams

An annual family planning exam consists of a comprehensive health history and physical examination, including medical laboratory evaluations as indicated, an assessment of the client’s problems and needs, and the implementation of an appropriate contraceptive management plan.
HHSC family planning program providers must bill the most appropriate evaluation and management (E/M) with modifier FP visit procedure code for the complexity of the annual family planning examination provided. To bill an annual family planning examination, providers must include the appropriate E/M procedure codes and must be billed with modifier FP on the claim in conjunction with all family planning procedures and services.

Refer to: The HHSC Family Planning Program Policy Manual.

The following table summarizes the uses for the E/M procedure codes and the corresponding billing requirements for the annual examination:

<table>
<thead>
<tr>
<th>Billing Criteria</th>
<th>Frequency</th>
</tr>
</thead>
<tbody>
<tr>
<td>New patient: Appropriate E/M procedure code with modifier FP</td>
<td>One new patient E/M code every three years following the last E/M visit provided to the client by that provider or a provider of the same specialty in the same group</td>
</tr>
<tr>
<td>Established patient: Appropriate E/M procedure code with modifier FP</td>
<td>Once per state fiscal year*</td>
</tr>
</tbody>
</table>

* The established patient procedure code will be denied if a new patient procedure code has been billed for the annual examination in the same year.

For appropriate claims processing, providers are encouraged to use a family planning diagnosis code to bill the annual family planning exam.

Refer to: Subsection 2, “Services, Benefits, Limitations, and Prior Authorization” in this handbook for the list of family planning diagnosis codes.

An annual family planning examination (billed with modifier FP) will not be reimbursed when submitted with the same date of service as an additional E/M visit. If another condition requiring an E/M office visit beyond the required components for an office visit, family planning visit, or surgical procedure is discovered, the provider may submit a claim for the additional visit using Modifier 25 to indicate that the client’s condition required a significant, separately identifiable E/M service. Documentation supporting the provision of a significant, separately identifiable E/M service must be maintained in the client’s medical record and made available to Texas Medicaid upon request.

2.1.1 FQHC Reimbursement for Family Planning Annual Exams

FQHCs must use the most appropriate E/M procedure code for the complexity of service provided as indicated in the HHSC Family Planning Program Policy Manual.

The annual exam is allowed once per fiscal year, per client, per provider. Other family planning office or outpatient visits may be billed within the same year.

A new patient visit for the annual exam may be reimbursed once every three years following the last E/M visit provided to the client by that provider or a provider of the same specialty in the same group. The annual examination must be billed as an established patient visit if E/M services have been provided to the client within the last three years.

Refer to: The HHSC Family Planning Program Policy Manual.

2.2 Family Planning Office or Outpatient Visits

Other family planning E/M visits are allowed for routine contraceptive surveillance, family planning counseling and education, contraceptive problems, suspicion of pregnancy, genitourinary infections, and evaluation of other reproductive system symptoms.

During any visit for a medical problem or follow-up visit, the following must occur:

- An update of the client’s relevant history
• Physical exam, if indicated
• Laboratory tests, if indicated
• Treatment or referral, if indicated
• Education and counseling, or referral, if indicated
• Scheduling of office or clinic visit, if indicated

Refer to: The HHSC Family Planning Program Policy Manual for more information about general family planning office or outpatient visits.

The following table summarizes the uses for the E/M procedure codes and the corresponding billing requirements for general family planning office or outpatient visits:

<table>
<thead>
<tr>
<th>Billing Criteria</th>
<th>Frequency</th>
</tr>
</thead>
<tbody>
<tr>
<td>New patient: Appropriate E/M procedure code</td>
<td>One new patient E/M code every three years following the last E/M visit provided to the client by that provider or a provider of the same specialty in the same group</td>
</tr>
<tr>
<td>Established patient: Appropriate E/M procedure code</td>
<td>As needed*</td>
</tr>
</tbody>
</table>

* The established patient procedure code will be denied if a new patient procedure code has been billed for the annual examination in the same year.

For appropriate claims processing, providers are encouraged to use a family planning diagnosis code to bill the annual family planning exam.

Refer to: Subsection 2, “Services, Benefits, Limitations, and Prior Authorization” in this handbook for the list of family planning diagnosis codes.

2.2.1 FQHC Reimbursement for Family Planning Office or Outpatient Visits

FQHCs must use the most appropriate E/M procedure code for the complexity of service provided as indicated previously in the tables in the HHSC Family Planning Program Policy Manual.

The new patient procedure codes will be limited to one new patient E/M procedure code three years following the last E/M visit provided to the client by that provider or a provider of the same specialty in the same group. The annual examination must be billed as an established patient visit if E/M services have been provided to the client within the last three years.

A general family planning office or outpatient visit (billed without modifier FP) will not be reimbursed when submitted with the same date of service as an additional E/M visit. If another condition requiring an E/M office visit beyond the required components for an office visit, family planning visit, or surgical procedure is discovered, the provider may submit a claim for the additional visit using modifier 25 to indicate that the client’s condition required a significant, separately identifiable E/M service. Documentation supporting the provision of a significant, separately identifiable E/M service must be maintained in the client’s medical record and made available to Texas Medicaid upon request.

Refer to: The HHSC Family Planning Program Policy Manual.

Section 4, “Federally Qualified Health Center (FQHC)” in the Clinics and Other Outpatient Facility Services Handbook (Vol. 2, Provider Handbooks) for more information about FQHC services.

2.2.1.1 Laboratory Procedures

Refer to: The HHSC Family Planning Program Policy Manual for more information about laboratory procedures.
Appropriate documentation must be maintained in the client’s record.

**Refer to:** Subsection 2.1.1, “Clinical Laboratory Improvement Amendments (CLIA)” in the Radiology and Laboratory Services Handbook (Vol. 2, Provider Handbooks).

Texas Medicaid follows the Medicare categorization of tests for CLIA certificate holders.


For waived tests, providers must use modifier QW as indicated on the CMS website.

### 2.3 Immunization Administration

**Refer to:** The HHSC Family Planning Program Policy Manual for specific procedure codes that may be reimbursed for medications, immunizations, and vaccines.

#### 2.3.1 Human Papilloma Virus (HPV) Vaccine

**Refer to:** The HHSC Family Planning Program Policy Manual for specific procedure codes that may be reimbursed for medications, immunizations, and vaccines for HPV.

### 2.4 Radiology

**Refer to:** The HHSC Family Planning Program Policy Manual for specific procedure codes that may be reimbursed for radiology services performed for the purpose of localization of an IUD.

### 2.5 Contraceptive Devices and Related Procedures

#### 2.5.1 Barrier Contraceptives

**Refer to:** The HHSC Family Planning Program Policy Manual for specific procedure codes that may be reimbursed for barrier contraceptives separately from fitting and instruction.

#### 2.5.2 IUD

**Refer to:** The HHSC Family Planning Program Policy Manual for specific procedure codes that may be reimbursed for IUDs and the insertion of IUDs.

##### 2.5.2.1 Removal of the IUD

**Refer to:** The HHSC Family Planning Program Policy Manual for specific procedure codes that may be reimbursed for the removal of an IUD.

When a vaginal, cervical, or uterine surgery procedure code is submitted with the same date of service as the IUD removal procedure code or the IUD replacement procedure code, the following reimbursement may apply:

- The other vaginal, cervical, or uterine surgical procedure may be reimbursed at full allowance.
- The removal or the replacement of the IUD will be denied.

#### 2.5.3 Contraceptive Implants

The contraceptive implant, procedure code J7307, and the implantation of the contraceptive implant, procedure code 11981, may be reimbursed.

Progesterone-containing subdermal contraceptive implants (Norplant) were previously used for birth control. Although subdermal contraceptive implants are no longer approved by the FDA, the removal of the implanted contraceptive implant may be considered for reimbursement.
Refer to: The HHSC Family Planning Program Policy Manual for the appropriate contraceptive implant removal procedure code.

### 2.6 Drugs and Supplies

Refer to: The HHSC Family Planning Program Policy Manual for specific procedure codes that may be reimbursed for providing contraceptive methods.

#### 2.6.1 Prescriptions and Dispensing Medication

Providers may do one or both of the following:

- Dispense family planning drugs and supplies directly to the client and bill the HHSC Family Planning Program.
- Write a prescription for the client to take to a pharmacy.

Family planning drugs and supplies that are dispensed directly to the client must be billed to the HHSC Family Planning Program. Only providers with an appropriate pharmacy license may be reimbursed for dispensing family planning drugs and supplies. Provider types with an appropriate pharmacy license may be reimbursed for dispensing up to a one-year supply of contraceptives in a 12-month period.

Refer to: The HHSC Family Planning Program Policy Manual for more information about dispensing contraceptives.

HHSC Family Planning Program clients may have their prescriptions filled at the clinic pharmacy. HHSC Family Planning Providers can refer to the HHSC Family Planning Policy and Procedure Manual for additional guidance on dispensing medication.

#### 2.6.2 Oral Medication Reimbursement

Refer to: The HHSC Family Planning Program Policy Manual for more information about oral medication.

### 2.7 Family Planning Education

Refer to: The HHSC Family Planning Program Policy Manual for the procedure codes that may be reimbursed for providing Contraceptive Method Instruction.

#### 2.7.1 Medical Nutrition Therapy

For clients requiring intensive nutritional guidance, medical nutritional therapy can be provided as an allowable and billable service. Medical nutritional therapy, however, must be provided by a registered dietician in order to be reimbursed.

Refer to: The HHSC Family Planning Program Policy Manual for more information about medical nutritional therapy.

#### 2.7.2 Instruction in Natural Family Planning Methods

Counseling with the intent to instruct a couple or an individual in methods of natural family planning may be reimbursed twice a year.

Refer to: The HHSC Family Planning Program Policy Manual for more information about natural family planning.

### 2.8 Sterilization and Sterilization-Related Procedures

#### 2.8.1 Sterilization Consent

Per federal regulation 42 CFR 50, Subpart B, all sterilization procedures require an approved Sterilization Consent Form.
2.8.2 Incomplete Sterilizations
Sterilizations are considered to be permanent, once per lifetime procedures. If the claim is denied indicating a sterilization procedure has already been reimbursed for the client, the provider may appeal with documentation that supports the medical necessity for the repeat sterilization.

2.8.3 Tubal Ligation and Hysteroscopic Occlusion
Refer to: The HHSC Family Planning Program Policy Manual for more information about tubal ligation and hysteroscopic occlusion.

2.8.4 Vasectomy
Refer to: The HHSC Family Planning Program Policy Manual for more information about vasectomies.

Vasectomies are considered to be permanent, once-per-lifetime procedures. If the claim is denied indicating a vasectomy procedure has already been reimbursed for the client, the provider may appeal with documentation that supports the medical necessity for the repeat sterilization.

2.9 Prior Authorization
Prior authorization is not required for sterilization and sterilization-related procedures.

3 Documentation Requirements
All services require documentation to support the medical necessity of the service rendered, including gynecological and reproductive health services and family planning services.

Gynecological and reproductive health services and family planning services are subject to retrospective review and recoupment if documentation does not support the service billed.

4 Claims Filing and Reimbursement

4.1 Claims Information
Providers must use the appropriate claim form to submit HHSC Family Planning Program claims to TMHP. Claims for dates of service that span multiple contract periods must be submitted on separate claims for services performed within each contract period.

Note: To submit HHSC Family Planning Program claims using TexMedConnect, providers must choose Family Planning Program "Title X-DFPP" on the electronic version of the 2017 claim form.
4.1.1 Filing Deadlines

The following table summarizes the filing deadlines for HHSC Family Planning Program claims:

<table>
<thead>
<tr>
<th>Deadline</th>
<th>Appeals</th>
</tr>
</thead>
<tbody>
<tr>
<td>95 days from the date of service on the claim or date of any third party insurance explanation of benefits (EOB)</td>
<td>120 days from the date of the Remittance and Status (R&amp;S) Report on which the claim reached a finalized status</td>
</tr>
</tbody>
</table>

*Note:* As stated in the HHSC Family Planning Policy and Procedure Manual, all claims and appeals must be submitted and processed within 60 days after the end of the contract period.

4.1.2 Third Party Liability

Federal and state regulations mandate that family planning client information be kept confidential. Because seeking information from third party insurance may jeopardize the client’s confidentiality, prior insurance billing is not a requirement for billing family planning for any title program.

4.2 Reimbursement

Reimbursement for family planning procedures is available in the TMHP Online Fee Lookup (OFL) on the TMHP website at www.tmhp.com.

4.2.1 Funds Gone

HHSC family planning providers are contracted to provide services for a specific time period, either the state fiscal year or a contract period within the fiscal year. The providers receive a specific budget amount for their contract period. When their claims payments have reached their budget allowance, providers must continue to submit claims. The amount of funds that they would have received had the funds been available will be tracked as “funds gone.”

Providers may receive additional funds for a contract period at a later time. Claims identified as “funds gone” may be reimbursed at that time.

On the R&S Report, “Claims Paid” is the dollar amount of claims paid during this financial transaction period. “Approved to Pay/Not Funds Gone” is the dollar amount that has been processed and approved to pay, but the payment has not been issued yet. “Funds Gone” is the dollar amount that has been submitted after the provider’s budget allowance has been reached. The amount in “Approved to Pay/Not Funds Gone” added to the amount in “Funds Gone” will equal the amount in the “Approved to Pay - New Claims” section.

4.3 NCCI and MUE Guidelines

The Healthcare Common Procedural Coding System (HCPCS) and Current Procedural Terminology (CPT) codes included in the Texas Medicaid Provider Procedures Manual are subject to National Correct Coding Initiative (NCCI) relationships, which supersede any exceptions to NCCI code relationships that may be noted in the manual. Providers should refer to the CMS NCCI web page for correct coding guidelines and specific applicable code combinations.

In instances when Texas Medicaid limitations are more restrictive than NCCI Medically Unlikely Edit (MUE) guidance, Texas Medicaid limitations prevail.

4.4 National Drug Code

*Refer to:* Subsection 6.3.4, “National Drug Code (NDC)” in “Section 6: Claims Filing” (Vol. 1, General Information).
HEALTHY TEXAS WOMEN PROGRAM HANDBOOK

TEXAS MEDICAID PROVIDER PROCEDURES MANUAL: VOL. 2

MAY 2021
# HEALTHY TEXAS WOMEN PROGRAM HANDBOOK

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1 General Information

The information in this handbook is intended for Healthy Texas Women (HTW) program providers. The handbook provides information about Texas Medicaid’s HTW benefits, policies, and procedures that are applicable to these service providers.

**Important:** All providers are required to read and comply with Section 1: Provider Enrollment and Responsibilities. In addition to required compliance with all requirements specific to Texas Medicaid, it is a violation of Texas Medicaid rules when a provider fails to provide healthcare services or items to Medicaid clients, including HTW clients, in accordance with accepted medical community standards and standards that govern occupations, as explained in Title 1 Texas Administrative Code (TAC) §371.1659. Accordingly, in addition to being subject to sanctions for failure to comply with the requirements that are specific to Texas Medicaid, providers can also be subject to Texas Medicaid sanctions for failure, at all times, to deliver healthcare items and services to Medicaid clients in full accordance with all applicable licensure and certification requirements including, without limitation, those related to documentation and record maintenance.

**Refer to:** The Children’s Services Handbook (Vol. 2, Provider Handbooks) for more information about providing services to Texas Medicaid and Texas Health Steps (THSteps) clients.

“Section 1: Provider Enrollment and Responsibilities” (Vol. 1, General Information).

“Texas Medicaid Administration” in the Preliminary Information (Vol. 1, General Information).

The Healthy Texas Women website at [www.healthytexaswomen.org](http://www.healthytexaswomen.org) for information about family planning and the locations of clinics receiving family planning funding from HHSC.

The Gynecological, Obstetrics, and Family Planning Title XIX Services Handbook (Vol. 2, Provider Handbooks) for information about Texas Medicaid fee-for service and Title XIX family planning benefits for gynecological and reproductive health services.

The Clinics and Other Outpatient Facility Services Handbook (Vol. 2, Provider Handbooks) for information about services provided in a Federally Qualified Health Centers (FQHCs) and Rural Health Clinics (RHCs).

2 Healthy Texas Women (HTW) Program Overview

The goal of HTW is to expand access to women’s health and family planning services to reduce unintended pregnancies, positively affect the outcome of future pregnancies, and positively impact the health and wellbeing of women and their families in the eligible population.

HTW is established to achieve the following objectives:

- Implement the state policy to favor childbirth and family planning services that do not include elective abortions or the promotion of elective abortions.

- Ensure the efficient and effective use of state funds in support of these objectives and to avoid the direct or indirect use of state funds to promote or support elective abortions.

- Reduce the overall cost of publicly-funded healthcare (including federally-funded healthcare) by providing low-income Texans access to safe, effective services that are consistent with these objectives.

- Enforce Human Resources Code §32.024(c-1).
Healthy Texas Women Plus (HTW Plus) is a set of additional physical health, mental health, and substance use disorder benefits that are available to HTW clients during the first 12 months of their eligibility following a pregnancy. These outpatient services target major health conditions that contribute to maternal mortality and severe morbidity in the extended postpartum period and provide continuity of care for chronic conditions treated during the pregnancy period.

Health-care providers who want to provide HTW Plus services must be enrolled as Texas Medicaid providers and have completed the HTW provider certification process. There are no additional requirements for HTW providers to provide HTW Plus benefits within their scope of education and training, because HTW Plus is not a separate program from HTW.

2.1 Guidelines for HTW Providers

HTW provides family planning services, related preventive health services that are beneficial to reproductive health, and other preventive health services that positively affect maternal health and future pregnancies for women who:

- Are 15 through 44 years of age.
  
  \textit{Note:} Women who are 15 through 17 years of age must have a parent or legal guardian apply on their behalf.

- Are a United States citizen or eligible immigrant.

- Are a resident of Texas.

- Do not currently receive benefits through another Medicaid program (including Medicaid for Pregnant Women), Children’s Health Insurance Program (CHIP), or Medicare Part A or B.

- Have a household income at or below 200 percent of the federal poverty level.

- Are not pregnant.

- Do not have other insurance that covers the services that HTW provides.

\textbf{Exception:} A client who has other private health insurance may be eligible to receive HTW services if a spouse, parent, or other person would cause physical, emotional, or other harm to the client because the client filed a claim on the health insurance.

HTW services are provided by a physician or by another qualified health-care professional operating under physician direction. A physician provides direction for family planning services through written standing delegation orders and medical protocols. The physician is not required to be on the premises for the provision of family planning services by an RN, PA, NP, or CNS. HTW participants may receive services from any provider that participates in HTW.

HTW clients must be allowed freedom of choice in the selection of contraceptive methods as medically appropriate. They must also be allowed the freedom to accept or reject services without coercion. All HTW-covered methods of contraception must be made available to the client, either directly or by referral to another provider of contraceptive services. Services must be provided without regard to age, marital status, race, ethnicity, parenthood, disability, religion, national origin, or contraceptive preference.

Client eligibility can be verified by:

- Using TexMedConnect.
• Accessing the Medicaid Client Portal for Providers.
• Checking an electronic or printed copy of Your Texas Benefits Healthy Texas Women card.
• Calling the Automated Inquiry System at 1-800-925-9126.

Refer to: Subsection 4.4.3, “Client Eligibility Verification” in “Section 4: Client Eligibility” (Vol. 1, General Information).

HTW clients will have the following identifiers on the feedback received from the stated source:
• Medicaid Coverage: W - MA - TWHP
• Program Type:
  • 68 - MEDICAL ASSISTANCE - WOMEN'S HEALTH PROGRAM (HTW)
  • 69 - MEDICAL ASSISTANCE - WOMEN'S HEALTH PROGRAM (HTW PLUS)
• Program: 100 - MEDICAID
• Benefit Plan: 100 - Traditional Medicaid

HTW clients will receive 12 months of continuous eligibility unless:
• The client dies.
• The client voluntarily withdraws from HTW.
• The client no longer satisfies the HTW eligibility criteria.
• The client is certified for another Medicaid program, such as Medicaid for Pregnant Women, or CHIP.
• State law no longer allows the woman to be covered.
• HHSC or its designee determines the client provided information affecting her eligibility that was false at the time of application.

If a provider suspects that a HTW client has committed fraud on the application, the provider should report the client to the HHSC Office of Inspector General (OIG) at 1-800-436-6184.

2.1.1 Referrals
If a provider identifies a health problem that is not within their scope of practice, the provider must refer the HTW client to another provider or clinic that can treat her. As mandated by Texas Human Resources Code §32.024(c-1), HTW does not reimburse office visits during which clients are referred for elective abortions.

HHSC prefers that clients be referred to local indigent care services. However, the toll-free Information and Referral hotline 2-1-1, can assist clients and providers with locating low-cost health services for clients in need.

2.1.2 Referrals for Clients Diagnosed with Breast or Cervical Cancer
Medicaid for Breast and Cervical Cancer (MBCC) provides access to cancer treatment through full Medicaid benefits for qualified women diagnosed with breast or cervical cancer. Health facilities that contract with the Breast and Cervical Cancer Screening (BCCS) program are responsible for helping women with the MBCC application.

To find a BCCS provider, call 2-1-1. For questions about the BCCS program, contact the state office at 512-458-7796, or visit www.healthytexaswomen.org/bccs-program.

2.1.3 Abortions
Elective and non-elective abortions are not covered by HTW.
Texas Human Resources Code Section 32.024(c-1) and Title 1 Texas Administrative Code, §382.17 prohibit the participation of a provider that performs or promotes elective abortions or affiliates with an entity that performs or promotes elective abortions.

A provider that performs elective abortions (through either surgical or medical methods) or that is affiliated with an entity that performs or promotes elective abortions for any patient is ineligible to serve HTW clients and cannot be reimbursed for any services rendered to a HTW client. This prohibition only applies to providers delivering services to HTW clients.

“Elective abortion” means the intentional termination of a pregnancy by an attending physician who knows that the female is pregnant, using any means that is reasonably likely to cause the death of the fetus. The term does not include the use of any such means: (A) to terminate a pregnancy that resulted from an act of rape or incest; (B) in a case in which a woman suffers from a physical disorder, physical disability, or physical illness, including a life-endangering physical condition caused by or arising from the pregnancy, that would, as certified by a physician, place the woman in danger of death or risk of substantial impairment of a major bodily function unless an abortion is performed; or (C) in a case in which a fetus has a severe fetal abnormality, meaning a life-threatening physical condition that, in reasonable medical judgment, regardless of the provision of life-saving treatment, is incompatible with life outside the womb.

Certain providers that want to participate in HTW must certify that they do not perform or promote elective abortions and do not affiliate with any entity that does, as directed by HHSC.

Refer to: Subsection 2.2, “HTW Provider Enrollment” in this handbook for more information about certification regarding elective abortions.

2.2 HTW Provider Enrollment

Certain providers who have completed the Medicaid enrollment process through TMHP, and have certified that they do not perform elective abortions or affiliate with providers that perform elective abortion are eligible to participate.

Refer to: “Section 1: Provider Enrollment and Responsibilities” (Vol. 1, General Information) for more information about enrollment procedures.

Certain providers that want to participate in HTW must certify that they do not perform or promote elective abortions and do not affiliate with any entity that does, as directed by HHSC. Providers may complete the Healthy Texas Women Certification and disclose the required information as part of the Medicaid enrollment process, or at any time after completing the Medicaid enrollment process. New providers may use the TMHP website to submit the Healthy Texas Women Certification through the Provider Enrollment Portal (PEP). Medicaid-only providers may use the TMHP website to submit the Healthy Texas Women Certification through the Provider Information Management System (PIMS).

The following provider types are required to certify:

- Physician or physician group with a general surgery, family practice/general practice, gynecology, OB/GYN, internal medicine, pediatric specialty or other specialty whose services are covered under HTW Plus, or a clinic/group practice
- Federally Qualified Health Center (FQHC)
- Physician Assistant
- Nurse practitioner/clinical nurse specialist
- Certified nurse midwife/registered nurse/licensed midwife
- Maternity Services Clinic
- Family Planning Clinic
• Rural Health Clinic—Freestanding/Independent
• Rural Health Clinic—Hospital Based
• Ambulatory Surgical Center—Freestanding/Independent
• Chemical Dependency Treatment Facility—Clinic/group practice
• Opioid Treatment Provider (OTP)—Clinic/group practice
• Licensed Professional Counselor
• Licensed Clinical Social Worker (LCSW)
• Psychologist – Psychology group
• Medical Supplier (DME)—In-home hyperalimentation supplies, Respiratory therapists, DME/Pharmacy (CCP)

Information that providers submit through PIMS can be searched by clients who use the Find a Doctor feature on the HTW website at www.healthytexaswomen.org.

2.3 Services, Benefits, Limitations, and Prior Authorization
This section includes information on women’s health and family planning services funded through HTW. HTW benefits include:

• Contraceptive services
• Pregnancy testing and counseling
• Preconception health screenings (e.g., screening for obesity, hypertension, diabetes, cholesterol, smoking, and mental health)
• Sexually transmitted infection (STI) services
• Treatment for the following chronic conditions:
  • Hypertension
  • Diabetes
  • High cholesterol
• Breast and cervical cancer screening and diagnostic services:
  • Radiological procedures including mammograms
  • Screening and diagnosis of breast cancer
  • Diagnosis and treatment of cervical dysplasia
• Immunizations
• Treatment of postpartum depression

Refer to: Subsection 2.3.10, “HTW Plus Services, Benefits, and Limitations” in this handbook for HTW Plus benefit information.

The following procedure codes are benefits for HTW:

<table>
<thead>
<tr>
<th>Procedure Codes</th>
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<tbody>
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<td><strong>Contraceptive and STI Services</strong></td>
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<tr>
<td>00851</td>
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* CLIA waived test
^ QW Modifier
### Procedure Codes

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<th>Description</th>
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### Behavioral Health Services

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### Supplies and Services

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### Evaluation and Management

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### Breast Cancer Screening

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### Cervical Cancer Screening

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### Problem-Focused Gynecological Services

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### Immunizations and Vaccinations

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### Other Preventative Services

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<tr>
<td>82681</td>
<td>82465^</td>
</tr>
<tr>
<td>84479</td>
<td>85007</td>
</tr>
</tbody>
</table>

* CLIA waived test
^ QW Modifier
Procedure code G0433 will deny if billed on the same day by the same provider as procedure code 86703.

Procedure codes 96156, 96158, 96159, 96167, and 96168 are a benefit for clients who are 20 years of age and younger.

Procedure code 99473 is limited to one service per year, by any provider. Procedure code 99473 may be considered for reimbursement more than once per year when documentation of medical necessity is submitted with the claim.

Procedure code 99474 is limited to four services per year, by any provider, and it may be reimbursed only if a claim for procedure code 99473 has been submitted within the past 12 rolling months.

Self-measured blood pressure monitoring is a benefit when it is used as a diagnostic tool to help a physician diagnose hypertension in individuals whose blood pressure is either elevated or inconclusive when it is evaluated in the office alone.

Refer to: Subsection 2.1.1, “Clinical Laboratory Improvement Amendments (CLIA)” in the Radiology and Laboratory Services Handbook (Vol. 2, Provider Handbooks).

### 2.3.1 Family Planning History Check

HTW clients must receive family planning services annually, but no later than the third visit as an established client. These services must include family planning counseling and education, including natural family planning and abstinence. In order to receive reimbursement, all existing HTW clients must have received family planning services and/or counseling within the past rolling year.

The following HTW clients do not require a family planning history check:

- New clients
- Women who are sterilized
- Women who have a long-acting reversible contraception (LARC)

### 2.3.2 Family Planning Annual Exams

Family planning providers must bill one of the following E/M visit procedure code based on the complexity of the annual family planning examination provided:

<table>
<thead>
<tr>
<th>Procedure Codes</th>
</tr>
</thead>
<tbody>
<tr>
<td>86885 87270 87512 87529 87530 87661 88155 88160 88161 88165</td>
</tr>
<tr>
<td>88167 88172 94760 J0558 J0561 J0690 J2010</td>
</tr>
</tbody>
</table>

* CLIA waived test
^ QW Modifier

The following table summarizes the uses for the E/M procedure codes and the corresponding billing requirements for the annual examination:

<table>
<thead>
<tr>
<th>Billing Criteria</th>
<th>Frequency</th>
</tr>
</thead>
<tbody>
<tr>
<td>New patient: Most appropriate E/M procedure code</td>
<td>One new patient E/M code every 3 years following the last E/M visit provided to the client by that provider or a provider of the same specialty in the same group</td>
</tr>
</tbody>
</table>
Refer to: The Family Planning section of the HHSC website for information about the HHSC Family Planning Program.

2.3.2.1 FQHC Reimbursement for Family Planning Annual Exams

To receive their encounter rate for the annual family planning examination for HTW clients, FQHCs must use the most appropriate E/M procedure code for the complexity of service provided as indicated in the previous tables in Subsection 2.3.2, “Family Planning Annual Exams” in this handbook.

A new patient visit for the annual exam may be reimbursed once every three years following the last E/M visit provided to the client by that provider or a provider of the same specialty in the same group. The annual examination must be billed as an established patient visit if E/M services have been provided to the client within the last three years.

Reimbursement for services payable to an FQHC is based on an all-inclusive rate per visit.

2.3.3 Other Family Planning Office or Outpatient Visits

HTW does not cover office visits during which clients are referred for elective abortions.

A provider is allowed to bill clients for services that are not a benefit of HTW.

Refer to: Subsection 1.7.11.1, “Client Acknowledgment Statement” in “Section 1: Provider Enrollment and Responsibilities” (Vol. 1, General Information) for the requirements for billing clients.

2.3.3.1 FQHC Reimbursement for Other Family Planning Office or Outpatient Visits

FQHCs may be reimbursed for three family planning encounters per HTW client, per year. Procedure codes J7296, J7297, J7298, J7300, J7301, and J7307 may be reimbursed in addition to the FQHC encounter payment. When seeking reimbursement for an IUD or implantable contraceptive implant, providers must submit on the same claim the procedure code for the contraceptive device along with the procedure code for the encounter. The contraceptive device is not subject to FQHC limitations. Providers must use modifier U8 when submitting claims for a contraceptive device purchased through the 340B Drug Pricing Program.

Outpatient visits for non-family planning-related encounters that are provided by FQHCs for HTW or HTW Plus covered physical and behavioral health services may be reimbursed when medically necessary.

Reimbursement for services payable to an FQHC is based on an all-inclusive rate per visit.

Refer to: Section 4, “Federally Qualified Health Center (FQHC)” in the Clinics and Other Outpatient Facility Services Handbook (Vol. 2, Provider Handbooks) for more information about FQHC services.

2.3.4 Laboratory Procedures

The fee for the handling or conveyance of the specimen for transfer from the provider’s office to a laboratory may be reimbursed using procedure code 99000.

More than one lab handling fee may be reimbursed per day if multiple specimens are obtained and sent to different laboratories.

Note: When a provider who renders HTW laboratory services obtains a specimen but does not perform the laboratory procedure, the provider who obtains the specimen may be reimbursed one lab handling fee per day, per client.

Handling fees are not paid for Pap smears or cultures. When billing for Pap smear interpretations, the claim must indicate that the screening and interpretation were actually performed in the office by using the modifier SU (procedure performed in physician’s office).
If more than one of procedure codes 87480, 87510, 87660, 87661, or 87800 is submitted by the same provider for the same client with the same date of service, all of the procedure codes are denied. Only one procedure code (87480, 87510, 87660, 87661, or 87800) may be submitted for reimbursement, and providers must submit the most appropriate procedure code for the test provided.

**Note:** Providers must code to the highest level of specificity with a diagnosis to support medical necessity when submitting procedure code 87797.

**Refereto:** Subsection 2.1.1, “Clinical Laboratory Improvement Amendments (CLIA)” in the Radiology and Laboratory Services Handbook (Vol. 2, Provider Handbooks).

Appropriate documentation must be kept in the client’s record.

Claims may be subject to retrospective review if they are submitted with diagnosis codes that do not support medical necessity.

HTW follows the Medicare categorization of tests for CLIA certificate holders.


For waived tests, providers must use modifier QW as indicated on the CMS website.

**2.3.5 Contraceptive Devices**

Providers must use modifier U8 when submitting claims for a contraceptive device purchased through the 340B Drug Pricing Program.

An E/M procedure code will not be reimbursed when it is billed with the same date of service as procedure code 58301, unless the E/M visit is a significant, separately identifiable service from the removal of the IUD. If the E/M visit occurs on the same date of service as the removal of the IUD, modifier 25 may be used to indicate that the E/M visit was a significant, separately identifiable service from the procedure.

**Note:** HTW does not reimburse for counseling for, or provision of, emergency contraception.

**2.3.6 Drugs and Supplies**

**2.3.6.1 Prescriptions and Dispensing Medication**

Drugs and supplies that are dispensed directly to the client must be billed to HTW. Only providers with an appropriate pharmacy license may be reimbursed for dispensing family planning drugs and supplies. Provider types with an appropriate pharmacy license may be reimbursed for dispensing up to a one-year supply of contraceptives in a 12-month period using procedure code J7303, J7304, or S4993.

Pharmacies under the Vendor Drug Program are allowed to fill all prescriptions as prescribed. Family planning drugs and supplies are exempt from the three prescriptions-per-month rule for up to a six-month supply.

**Refereto:** Subsection 1.1, “About the Vendor Drug Program” in the Outpatient Drug Services Handbook (Vol. 2, Provider Handbooks) for information about this program.

**2.3.6.1.1 Long-Acting Reversible Contraception Products**

Certain LARC products are available as a pharmacy benefit of HTW and are available through a limited number of specialty pharmacies that work with LARC manufacturers. Providers can refer to the Texas Medicaid/CHIP Vendor Drug Program website at [www.txvendordrug.com/formulary/larc.shtml](http://www.txvendordrug.com/formulary/larc.shtml) for additional information, including a list of covered products and participating specialty pharmacies.

**2.3.7 Sterilization and Sterilization-Related Procedures**

Sterilizations are considered to be permanent, once per lifetime procedures. Denied claims may be appealed with documentation that supports the medical necessity for a repeat sterilization.
The sterilization services that are available to HTW clients include surgical or nonsurgical sterilization, follow-up office visits related to confirming the sterilization, and any necessary short-term contraception.

HTW covers sterilization as a form of birth control. To be eligible for a sterilization procedure through HTW, the client must be 21 years of age or older and must complete and sign a Sterilization Consent Form within at least 30 days of the date of the surgery but no more than 180 days. In the case of an emergency, there must be at least 72 hours between the date on which the consent form is signed and the date of the surgery. Operative reports that detail the need for emergency surgery are required.

2.3.7.1 Sterilization Consent
Per federal regulation 42 Code of Federal Regulations (CFR) 50, Subpart B, all sterilization procedures require an approved Sterilization Consent Form.

Note: The Texas Medicaid - Title XIX Acknowledgment of Hysterectomy Information form is not sterilization consent.


2.3.8 Treatment for Sexually Transmitted Infections (STIs)
HTW covers treatment for the following conditions:

- Gardnerella
- Trichomoniasis
- Candida
- Chlamydia
- Gonorrhea
- Herpes
- Syphilis

2.3.9 Immunizations and Vaccinations
HTW covers the following immunizations and vaccinations:

- HPV
- Hep A
- Hep B
- Chicken pox
- MMR
- Tdap
- Flu

2.3.10 HTW Plus Services, Benefits, and Limitations
HTW Plus provides enhanced postpartum services to HTW Plus clients for the following targeted health conditions and services:

- Behavioral health conditions
  - Individual, family, and group psychotherapy services
• Peer specialist services
• Cardiovascular and coronary conditions
  • Cardiovascular evaluation imaging and laboratory studies
  • Blood pressure monitoring equipment
  • Anticoagulant, antiplatelet, and antihypertensive medications
• Substance use disorders
  • Screening, brief intervention, and referral for treatment
  • Outpatient substance use counseling
  • Smoking cessation services
  • Medication-assisted treatment
  • Peer specialist services
• Diabetes
  • Laboratory studies
  • Additional injectable insulin options
  • Blood glucose testing supplies
  • Voice-integrated glucometers for women with diabetes who are visually impaired
  • Glucose monitoring supplies
• Asthma
  • Medications and supplies

The following procedure codes are benefits of HTW Plus:

<table>
<thead>
<tr>
<th>Procedure Codes</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>Behavioral Health Services</strong></td>
</tr>
<tr>
<td>90832 90833 90834 90836 90837 90838 90847 90853 90870 96130</td>
</tr>
<tr>
<td>96131 96136 96137 99354 99355 H0038</td>
</tr>
<tr>
<td><strong>Cardiovascular and Coronary Services</strong></td>
</tr>
<tr>
<td>37187 37211 37212 70498 70547 70548 71275 73706 74174 74175</td>
</tr>
<tr>
<td>75571 75574 75635 75716 75736 93005 93010 93015 93016 93017</td>
</tr>
<tr>
<td>93018 93041 93042 93224 93225 93226 93227 93241 93242 93243</td>
</tr>
<tr>
<td>93244 93245 93246 93247 93248 93306 93307 93308 93312 93320</td>
</tr>
<tr>
<td>93321 93325 93350 93351 93355 93360 93389 93923 93970 93971 94619</td>
</tr>
<tr>
<td>A4663 A4670</td>
</tr>
<tr>
<td><strong>Substance Use Disorder Services</strong></td>
</tr>
<tr>
<td>99408 H0001 H0004 H0005 H0020 H0038 H0049 J2310</td>
</tr>
<tr>
<td><strong>Diabetes Services</strong></td>
</tr>
<tr>
<td>83037 A4253 A4258 A4259 E2100 J1610 S5550 S5552</td>
</tr>
<tr>
<td><strong>Asthma Services</strong></td>
</tr>
<tr>
<td>94150 94617 A4614 A4627 E0570 J1720 J3301 J7611 J7613 J7614</td>
</tr>
<tr>
<td>J7620 J7626 J7644 S8101</td>
</tr>
</tbody>
</table>
HTW Plus benefits are subject to the same restrictions and limitations that are applied to Texas Medicaid’s coverage for the same procedure codes.

Referred: The relevant handbooks for detailed coverage information of HTW Plus benefits.

2.3.11 Prior Authorization
Prior authorization is not required for HTW services.

2.4 Documentation Requirements
All services require documentation to support the medical necessity of the service rendered, including HTW services.

HTW services are subject to retrospective review and recoupment if documentation does not support the service billed.

2.5 HTW Claims Filing and Reimbursement

2.5.1 Claims Information
Providers must use the appropriate claim form to submit HTW claims to TMHP.

Referred: Subsection 2.4, “Claims Filing and Reimbursement” in the Gynecological, Obstetrics, and Family Planning Title XIX Services Handbook (Vol. 2, Provider Handbooks) for more information about filing family planning claims.

2.5.1.1 HTW and Third Party Liability
Federal and state regulations mandate that family planning client information be kept confidential.

Because seeking information from third party insurance may jeopardize the client’s confidentiality, third party billing for HTW is not allowed.

2.5.2 Reimbursement
Providers can refer to the Online Fee Lookup (OFL) or the applicable fee schedule on the TMHP website at www.tmhp.com.

2.5.3 National Drug Code
2.5.4 NCCI and MUE Guidelines

The Health Care Procedure Coding System (HCPCS) and Current Procedural Terminology (CPT) codes included in the Texas Medicaid Provider Procedures Manual are subject to NCCI relationships, which supersede any exceptions to NCCI code relationships that may be noted in the manual. Providers should refer to the CMS NCCI web page for correct coding guidelines and specific applicable code combinations.

In instances when Texas Medicaid limitations are more restrictive than NCCI MUE guidance, Texas Medicaid limitations prevail.

3 Claims Resources

<table>
<thead>
<tr>
<th>Resource</th>
<th>Location</th>
</tr>
</thead>
<tbody>
<tr>
<td>Acronym Dictionary</td>
<td>“Appendix C: Acronym Dictionary” (Vol. 1, General Information)</td>
</tr>
<tr>
<td>Automated Inquiry System (AIS)</td>
<td>Subsection A.10, “TMHP Telephone and Fax Communication” in “Appendix A: State, Federal, and TMHP Contact Information” (Vol. 1, General Information)</td>
</tr>
<tr>
<td>2017 Claim Form Instructions</td>
<td>Subsection 6.8, “Family Planning Claim Filing Instructions” in “Section 6: Claims Filing” (Vol. 1, General Information)</td>
</tr>
</tbody>
</table>

4 Contact TMHP

The TMHP Contact Center at 1-800-925-9126 is available Monday-Friday from 7 a.m. to 7 p.m., Central Time.

5 Forms

The following linked forms can also be found on the Forms page of the Provider section of the TMHP website at www.tmhp.com:

<table>
<thead>
<tr>
<th>Forms</th>
</tr>
</thead>
<tbody>
<tr>
<td>Sterilization Consent Form Instructions</td>
</tr>
<tr>
<td>Sterilization Consent Form (English)</td>
</tr>
<tr>
<td>Sterilization Consent Form (Spanish)</td>
</tr>
<tr>
<td>2017 Claim Form</td>
</tr>
<tr>
<td>Healthy Texas Women Certification</td>
</tr>
</tbody>
</table>
HOME HEALTH NURSING AND PRIVATE DUTY NURSING SERVICES HANDBOOK

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1 General Information

This handbook contains information about Texas Medicaid fee-for-service benefits. The information in this handbook is intended for home health nursing services. Nursing services include home health skilled nursing visits, home health aide services, and private duty nursing services. The Handbook provides information about Texas Medicaid’s benefits, policies, and procedures applicable to these therapies.

For information about managed care services, refer to the Medicaid Managed Care Handbook. Managed care carve-out services are administered as fee-for-service benefits. A list of all carve-out services is available in Section 8, “Carve-Out Services” in the Medicaid Managed Care Handbook (Vol. 2, Provider Handbooks).

Important: All providers are required to read and comply with Section 1: Provider Enrollment and Responsibilities. In addition to required compliance with all requirements specific to Texas Medicaid, it is a violation of Texas Medicaid rules when a provider fails to provide healthcare services or items to Medicaid clients in accordance with accepted medical community standards and standards that govern occupations, as explained in Title 1 Texas Administrative Code (TAC) §371.1659. Accordingly, in addition to being subject to sanctions for failure to comply with the requirements that are specific to Texas Medicaid, providers can also be subject to Texas Medicaid sanctions for failure, at all times, to deliver health-care items and services to Medicaid clients in full accordance with all applicable licensure and certification requirements including, without limitation, those related to documentation and record maintenance.

Refer to: “Section 1: Provider Enrollment and Responsibilities” (Vol. 1, General Information) for more information about enrollment procedures.

1.1 Client Eligibility for Home Health Nursing and Aide Services

It is the provider’s responsibility to determine the type of coverage (Medicare, Medicaid, or private insurance) that the client is eligible to receive. To verify client Medicaid eligibility and retroactive eligibility, the home health agency, durable medical equipment (DME), or medical supplier must verify eligibility by accessing the Medicaid Client Portal for Providers. Additionally, eligibility can be verified by calling the Automated Inquiry System (AIS) at 1-800-925-9126 or the Texas Medicaid & Healthcare Partnership (TMHP) Electronic Data Interchange (EDI) Help Desk at 1-888-863-3638.

Home health clients do not need to be homebound to qualify for services.

The Medicaid client must be eligible on the date of service (DOS) and must meet all of the following requirements to qualify for home health services:

- Have a medical need for home health professional services, DME, or medical supplies that is documented in the client’s plan of care (POC) and considered a benefit under home health services
- Receive services that meet the client’s existing medical needs and can be safely provided in the client’s home
- Receive prior authorization from TMHP for most home health professional services, DME, or medical supplies


Note: Texas Health Steps (THSteps)-eligible clients who qualify for medically necessary services beyond the limits of this home health services benefit may receive those services through the Comprehensive Care Program (CCP).
1.1.1 Prior Authorization Requests for Clients with Retroactive Eligibility

Retroactive eligibility occurs when the effective date of a client’s Medicaid coverage is before the date the client’s Medicaid eligibility is added to TMHP’s eligibility file, which is called the “add date.”

For clients with retroactive eligibility, prior authorization requests must be submitted after the client’s add date and before a claim is submitted to TMHP.

For services provided to fee-for-service Medicaid clients during the client’s retroactive eligibility period, i.e., the period from the effective date to the add date, prior authorization must be obtained within 95 days from the client’s add date and before a claim for those services is submitted to TMHP. For services provided on or after the client’s add date, the provider must obtain prior authorization within three business days of the date of service.

The provider is responsible for verifying eligibility. The provider is strongly encouraged to access the Medicaid Client Portal for Providers, TexMedConnect, or AIS to verify eligibility frequently while providing services to the client. If services are discontinued before the client’s add date, the provider must still obtain prior authorization within 95 days of the add date to be able to submit claims.

Refer to: “Section 4: Client Eligibility” (Vol. 1, General Information).

1.1.2 Client Evaluation

When a home health agency receives a referral to provide home health nursing and therapy services for a client who is eligible for Texas Medicaid, the agency-employed registered nurse (RN) must evaluate the client in the home before calling TMHP for prior authorization. A home evaluation by the agency-employed RN is required for SN, home health aide (HHA), occupational therapist (OT), physical therapist (PT), DME, or medical supplies requested on a home health services POC. It is expected that appropriate referrals will be made between home health agencies and DME suppliers for care. It is recommended that DME suppliers keep open communication with the client’s physician to ensure the client’s medical record is current.

This evaluation must include assessment of the following:

- Medical necessity for home health services, DME, or medical supplies requested.
- Client safety.
- Appropriateness of care in the home setting.
- Capable caregiver available if clients are unable to perform their own care or monitor their own medical condition.

Following the RN’s assessment or evaluation of the client in the home setting for home health services needs, the agency-employed RN who completed the home evaluation must contact TMHP for prior authorization within three business days of the start of care (SOC).

1.2 Client Eligibility for PDN Services

PDN is considered medically necessary when a client has a disability, physical, or mental illness, or chronic condition and requires continuous, skillful observations, judgments, and interventions to correct or ameliorate his or her health status.

To be eligible for PDN services, a client must meet all the following criteria:

- Be birth through 20 years of age and eligible for Medicaid and THSteps
- Meet medical necessity criteria for PDN
- Have a primary physician who must:
  - Provide a prescription for PDN.
• Establish a POC.
• Provide documentation to support the medical necessity of PDN services.
• Provide continuing medical care and supervision of the client, including, but not limited to, examination or treatment within 30 calendar days prior to the start of PDN services, or examination or treatment that complies with the THSteps periodicity schedule, or is within six months of the PDN extension SOC date, whichever is more frequent (for extensions of PDN services). This requirement may be waived based on review of the client’s specific circumstances.

Note: The physician visit may be waived when a diagnosis has already been established by the physician, and the client is under the continuing care and medical supervision of the physician. A waiver is valid for no more than 365 days, and the client must be seen by his or her physician at least once every 365 days. The waiver must be based on the physician’s written statement that an additional evaluation visit is not medically necessary. This documentation must be maintained by the physician and the provider in the client’s medical record.

• Provide specific written, dated orders for the client who is receiving continuing or ongoing PDN services.
• Require care beyond the level of services provided under Texas Medicaid (Title XIX) home health services.

Clients who are birth through 17 years of age must reside with a responsible adult who is either trained to provide nursing care or is capable of initiating an identified contingency plan when the scheduled private duty nurse is unexpectedly unavailable.

PDN is based on the need for skilled care in the client’s home; however, these services may follow the client and may be provided in accordance with 42 CFR §440.80. The POS must be able to support the client’s health and safety needs. It must be adequate to accommodate the use, maintenance, and cleaning of all medical devices, equipment, and supplies required by the client. Necessary primary and backup utilities, communication, fire, and safety systems must be available at all times.

The amount and duration of PDN must always be commensurate with the client’s medical needs. Requests for services must reflect changes in the client’s condition that affect the amount and duration of PDN.

2 Enrollment

Refer to: Subsection 1.7.15, “Private Duty Nursing (PDN) Providers” in “Section 1: Provider Enrollment and Responsibilities” (Vol. 1, General Information) for enrollment information.

3 Home Health Skilled Nursing and Home Health Aide Services

3.1 Services, Benefits, Limitations, and Prior Authorization

Home health skilled nursing (SN) and HHA visits are a benefit of Texas Medicaid Title XIX home health services when a client requires home nursing services for an acute condition or an acute exacerbation of a chronic condition that can be met on an intermittent or part-time basis. For clients who are 20 years of age or younger, SN and HHA visits are a benefit of Texas Medicaid Title XIX home health services when a client requires nursing services for an acute condition, a chronic condition, or an acute exacerbation of a chronic condition that can be met on an intermittent or part-time basis. SN visits are
intended to provide SN care to promote independence and support the client living at home. HHA visits are intended to provide personal care services under the supervision of an RN, PT, or OT employed by the home health agency to promote independence and support the client living at home.

The following codes are a benefit of Texas Medicaid:

<table>
<thead>
<tr>
<th>Procedure Codes</th>
</tr>
</thead>
<tbody>
<tr>
<td>G0156</td>
</tr>
<tr>
<td>G0299</td>
</tr>
<tr>
<td>G0300</td>
</tr>
</tbody>
</table>

Title XIX home health services must be provided by a licensed and certified home health agency enrolled in Texas Medicaid.

When the client’s needs are beyond the benefit of Title XIX home health services, additional benefits may be accessed through the following:

- Services for clients who are 20 years of age or younger may include, but are not limited to, private duty nursing (PDN) or personal care services (PCS).
- Services for clients who are 21 years of age or older may include, but are not limited to, long-term care assistance.

Refer to: Section 4, “Private Duty Nursing (PDN) Services - CCP” in this handbook for information about PDN services. Subsection 2.11, “Personal Care Services (PCS) (CCP)” in the Children’s Services Handbook (Vol. 2, Provider Handbooks) for information about PCS services.

### 3.1.1 Medical Necessity

SN and HHA visits are considered medically necessary for a client who:

- Requires skillful observations and judgement to improve health status, skilled assessment, or skilled treatments or procedures.
- Requires individualized, intermittent, acute skilled care.
- Requires skilled interventions to improve health status, and if skilled intervention is delayed, it is expected to result in:
  - Deterioration of a chronic condition
  - Loss of function
  - Imminent risk to health status due to medical fragility, or risk of death
- Requires general supervision of nursing care provided by an HHA over whom the RN is administratively or professionally responsible.

Note: When documentation does not support medical necessity for home health SN or HHA visits, providers may be directed to possible alternative services based on the client’s age and needs.

### 3.2 Skilled Nursing and Home Health Aide Services

All SN and HHA services must be prior authorized.

The following definitions apply to Title XIX Home Health SN and HHA visits:

- Acute is defined as a condition or exacerbation that is anticipated to improve and reach resolution within 60 days.
- Part-time is defined as SN or HHA visits provided less than eight hours per day for any number of days per week. Part-time visits may be continuous up to 7.5 hours per day (not to exceed a combined total of three 2.5 hour visits).
• Intermittent is defined as SN or HHA visits provided for less than eight hours per visit and less frequently than daily. Intermittent visits may be delivered in interval visits up to 2.5 hours per visit, not to exceed a combined total of three visits per day.

SN visits are nursing services ordered by a physician, included in the Texas Medicaid home health services Plan of Care (POC), and provided by an RN or a licensed vocational nurse (LVN) currently licensed by the Board of Nurse Examiners of the State of Texas (BNE). SN visits may be considered when a client requires nursing services for an acute condition or an acute exacerbation of a chronic condition that can be met on an intermittent or part-time basis and typically has an end-point. SN visits may be provided on consecutive days.

HHA visits are services ordered by the physician, included in the nursing Texas Medicaid home health services POC, and are services the HHA is permitted to perform under State law. HHA visits may be considered when a client requires services for an acute condition or an acute exacerbation of a chronic condition that can be met on an intermittent or part-time basis and typically has an end-point. HHA visits will not be considered unless the client also requires SN or therapy services. HHA visits may be provided on consecutive days.

Refer to: The Physical Therapy, Occupational Therapy, and Speech Therapy Services Handbook (Vol. 2, Provider Handbooks) for information about home health PT, OT, and ST services.

3.2.1 Skilled Nursing Visits

SN visits are limited to SN procedures performed by an RN or LVN licensed to perform these services under the Texas Nursing Practice Act and include the following:

• Direct SN care, and parent, guardian, or caregiver training and education
• SN observation, assessment, and evaluation by an RN, provided a physician specifically requests that a nurse visit the client for this purpose, and the physician’s order reflects the medical necessity for the visit
• Supervision of delegated services provided by an HHA or others over whom the RN is administratively or professionally responsible

SN care consists of those services that must, under State law, be performed by an RN or LVN, and meet the criteria for SN services specified in the Code of Federal Regulations (42 CFR §§ 409.32, 409.33, and 409.44):

• In determining whether a service requires the skill of a licensed nurse, consideration must be given to the inherent complexity of the service, the condition of the client, and the accepted standards of medical and nursing practice.
• The fact that the SN service can be, or is, taught to the client or to the client’s family or friends does not negate the skilled aspect of the service when the service is performed by a nurse.
• If the service could be performed by the average nonmedical person, the absence of a competent person to perform it does not cause it to be a SN service.
• If the nature of a service is such that it can safely and effectively be performed by the average nonmedical person without direct supervision of a licensed nurse, the service cannot be regarded as a SN service.
• Some services are classified as SN services on the basis of complexity alone (e.g., intravenous and intramuscular injections or insertion of catheters), and if reasonable and necessary to the treatment of the client’s illness or injury, would be a benefit on that basis. However, in some cases, the client’s condition may cause a service that would ordinarily be considered unskilled to be considered an SN service. This would occur when the client’s condition is such that the service can be safely and effectively provided only by a nurse.
• A service, which, by its nature, requires the skills of a nurse to be provided safely and effectively continues to be a skilled service even if it is taught to the client, the client’s family, or other caregivers. Where the client needs the SN care and there is no one trained, able and willing to provide it, the services of a nurse would be reasonable and necessary to the treatment of the illness or injury.

• The SN services must be reasonable and necessary to the diagnosis and treatment of the client’s illness or injury within the context of the client’s unique medical condition. To be considered reasonable and necessary for the diagnosis or treatment of the client’s illness or injury, the services must be consistent with the nature and severity of the illness or injury, the client’s particular medical needs, and within accepted standards of medical and nursing practice. A client’s overall medical condition is a valid factor in deciding whether skilled services are needed. A client’s diagnosis should never be the sole factor in deciding whether the service the client needs is either skilled or not skilled.

• The determination of whether the services are reasonable and necessary should be made in consideration of the physician’s determination that the services ordered are reasonable and necessary. The services must, therefore, be viewed from the perspective of the condition of the client when the services were ordered, and what was, at that time, reasonably expected to be appropriate treatment for the illness or injury throughout the certification period.

• The SN care must be provided on a part-time or intermittent basis.

Professional nursing provided by an RN, as defined in the Texas Nursing Practice Act, means the performance of an act that requires substantial specialized judgment and skill, the proper performance of which is based on knowledge and application of the principles of biological, physical, and social science as acquired by a completed course in an approved school of professional nursing. The term does not include acts of medical diagnosis or the prescription of therapeutic or corrective measures. Professional nursing involves:

• The observation, assessment, intervention, evaluation, rehabilitation, care and counsel, or health teachings of a person who is ill, injured, infirm, or experiencing a change in normal health processes.

• The maintenance of health or prevention of illness.

• The administration of a medication or treatment as ordered by a physician, podiatrist, or dentist.

• The supervision of delegated nursing tasks or teachings of nursing.

• The administration, supervision, and evaluation of nursing practices, policies, and procedures.

• The performance of an act delegated by a physician.

• Development of the nursing care plan.

Vocational nursing, as defined in the Texas Nursing Practice Act, means a directed scope of nursing practice, including the performance of an act that requires specialized judgment and skill, the proper performance of which is based on knowledge and application of the principles of biological, physical, and social science as acquired by a completed course in an approved school of vocational nursing. The term does not include acts of medical diagnosis or the prescription of therapeutic or corrective measures. Vocational nursing involves:

• Collecting data and performing focused nursing assessments of the health status of an individual.

• Participating in the planning of the nursing care needs of an individual.

• Participating in the development and modification of the nursing care plan.

• Participating in health teaching and counseling to promote, attain, and maintain the optimum health level of an individual.

• Assisting in the evaluation of an individual’s response to a nursing intervention and the identification of an individual’s needs.
• Engaging in other acts that require education and training, as prescribed by board rules and policies, commensurate with the nurse’s experience, continuing education, and demonstrated competency.

### 3.2.2 Home Health Aide Visits

HHA visits are intended to provide hands-on personal care, performance of simple procedures as an extension of therapy or nursing services, assistance in ambulation or exercises, and assistance in administering medications that are ordinarily self-administered.

Any HHA services offered by a home health agency must be provided by a qualified HHA under the supervision of a qualified licensed individual (e.g., RN, PT, or OT) employed by the home health agency.

The duties of an HHA during a visit include, but are not limited to:

- Obtaining and recording the client’s vital signs (temperature, pulse, respirations, blood pressure)
- Observation, reporting and documentation of the client’s status, and the care or service furnished
- Personal care (hygiene and grooming) including, but not limited to:
  - Sponge, tub, or shower bath
  - Shampoo, sink, tub or bed bath
  - Nail and skin care
  - Oral hygiene
- Toileting and elimination care
- Ambulation
- Exercise
- Range of motion
- Safe transfer
- Positioning
- Assisting with nutrition and fluid intake
- Household services essential to the client’s health care at home
- Assistance with medications that are ordinarily self-administered
- Reporting changes in the client's condition and needs
- Completing appropriate documentation

### 3.2.2.1 Supervision of Home Health Aides

Supervision, as defined by the Texas Nursing Practice Act, is the process of directing, guiding, and influencing the outcome of an individual’s performance of an activity.

An RN or therapist (PT or OT) must provide the HHA written instructions for all the tasks delegated to the HHA. A therapist may prepare the written instructions if the client is receiving only HHA visits, which do not include delegated SN tasks, in addition to the therapy services.

The requirements for HHA supervision are as follows:

- When only HHA visits are provided, an RN must make a supervisory visit to the client’s residence at least once every 60 days. The supervisory visit must occur when the HHA is providing care to the client.
• When SN, PT, or OT visits are provided in addition to an HHA visit, an RN must make a supervisory visit to the client’s residence at least every two weeks. The supervisory visit must occur when the HHA is providing care to the client.

• When only PT or OT visits are provided in addition to HHA visits, the appropriate therapist may make the supervisory visit in place of an RN. The supervisory visit must occur when the HHA is providing care to the client.

• Documentation of HHA supervision must be maintained in the client’s medical record.

3.3 Home Health Skilled Nursing and Home Health Aide Services Providers

Providers must be a licensed and certified home health agency enrolled in Texas Medicaid and must comply with all applicable federal, state, and local laws and regulations, and Texas Medicaid’s policies and procedures.

All providers must maintain written policies and procedures for:

• Obtaining consent for medical treatment for clients in the absence of the primary caregiver that meets the standards of the Texas Family Code, Chapter 32.

• Obtaining physician signatures for all telephone orders within 14 calendar days of receipt of the order.

Providers must only accept clients on the basis of a reasonable expectation that the client’s needs can be adequately met in the place of service. The essential elements of safe and effective home health SN and HHA services include a trained parent, guardian, or caregiver, a primary physician, competent providers, and an environment that supports the client’s health and safety needs.

• The place of service must be able to support the health and safety needs of the client and must be adequate to accommodate the use, maintenance, and cleaning of all medical devices, equipment, and supplies required by the client.

• Necessary primary and back-up utility, communication, and fire safety systems must be available.

A parent or guardian, primary caregiver, or alternate care giver may not provide SN or HHA services even if he or she is an enrolled provider or employed by an enrolled provider.

3.4 Authorization Requirements

Home Health SN and HHA visits require prior authorization.

Prior authorization of SN or HHA visits, requires that a client’s primary physician:

• Provides an order for SN or HHA visits or recertification, identifying that the prescribed SN or HHA visits are medically necessary as defined in the Statement of Benefits

• The physician’s documentation in the client’s medical record must support the prescribed SN or HHA visits are medically necessary as defined in the Statement of Benefits

• The physician’s documentation in the client’s medical record must support that the client’s medical condition is sufficiently stable to permit safe delivery of SN or HHA visits as described in the home health services POC

• Establishes a medical POC, which is maintained in the client’s medical record

• Provides continuing care and medical supervision

• Provides specific written, dated orders for clients receiving SN or HHA visits

• Reviews and approves the home health services POC at least every 60 days, or more frequently as the physician determines necessary, including but not limited to when the client’s condition changes
SN visits requested primarily to provide the following will not be prior authorized:

- Respite care
- Child care
- Activities of daily living for the client
- Housekeeping services
- Routine post-operative disease, treatment, or medication teaching after a physician visit
- Routine disease, treatment, or medication teaching after a physician visit
- Individualized, comprehensive case management beyond the service coordination required by the Texas Nursing Practice Act

**Note:** Clients who are 20 years of age or younger may be eligible for private duty nursing services and personal care services through Texas Medicaid Private Duty Nursing Services and Personal Care Services Policies.

HHA visits requested for the following will not be prior authorized when:

- Primarily requested to perform housekeeping services
- Provided to a client residing in a hospital, SN facility, or intermediate care facility

**Note:** Clients who are 20 years of age or younger may be eligible for private duty nursing services and personal care services through Private Duty Nursing (PDN) Services - THSteps-CCP or Personal Care Services (PCS) - THSteps-CCP.

Certain facilities are required by licensure to meet all the medical needs of the client. SN or HHA visits will not be authorized for clients receiving care in any of the following facilities:

- Hospitals
- SN facilities
- Intermediate care facilities for the individuals with intellectual disability (ICF-IID)
- Special care facilities, including but not limited to, sub-acute units or facilities for the treatment of acquired immune deficiency syndrome (AIDS)
- Prescribed pediatric extended care centers, unless the SN and/or HHA services are provided before or after PPECC services, when rendered on the same day.

When a client, client’s responsible adult, or client’s physician notifies the SN and/or HHA service provider that the client also receives services from a PPECC, the SN and/or HHA service provider must coordinate services with the PPECC provider to prevent duplication of services.

**Note:** It is anticipated that the provision of SN and/or HHA services, in addition to PPECC would be uncommon.

### 3.4.1 Initial Assessment and Reassessments

When a provider has received a referral and has physician orders for SN or HHA services, the provider must have an RN perform an initial client assessment in the client’s home.

A client can be referred to a home health agency for SN or HHA services by:

- The client
- The client’s physician
- The client’s family
The client assessment or reassessment should include, but is not limited to, the following:

- A nursing assessment of medical necessity for the requested visits, which includes:
  - Complexity and intensity of the client’s care
  - Stability and predictability of the client’s condition
  - Frequency of the client’s need for SN care
  - Identified medical needs and goals
  - Description of wounds, if present
  - Cardiac status
- Whether the setting can support the health and safety needs of the client and is adequate to accommodate the use, maintenance, and cleaning of all medical devices, equipment, and supplies required by the client.
- Comprehension level of parent, guardian, caregiver, or client.
- Receptivity to training and ability level of the parent, guardian, caregiver, or client.

The initial assessment and any reassessments performed by an RN are required when changes in the client’s condition occur during the course of the authorization period. If there is no change in the client’s condition, the reassessment must document medical necessity, as defined in the Statement of Benefits, to support continued and ongoing SN or HHA visits beyond the initial 60 day authorization period.

A reassessment is required when the SN and/or HHA provider is notified by the client, client’s responsible adult, or the client’s physician that PPECC services have been initiated.

### 3.4.2 Home Health Services Plan of Care Requirements

The initial assessment or reassessments are used to establish and revise the home health services POC and must support the client’s medical necessity for SN services, HHA services, PT services, and OT services.

**Note:** Providers must use the Texas Medicaid home health services POC located on the TMHP Prior Authorization Texas Medicaid Forms web page; the Centers of Medicare and Medicaid (CMS) Form 485 will not be accepted.

The POC must be initiated and written in a clear and legible format by the RN and include the following:

- The client’s Medicaid number, the physician’s license number, and the provider’s Medicaid number
- Date the client was last seen by the physician
- The start of care (SOC) date for home health services
- All pertinent diagnoses
- The client’s mental status
- The prognosis
- The types of service requested, including the number of visits and amount, duration, and frequency
- The equipment or supplies required
- Rehabilitation potential
- Prior and current functional limitations
- Activities permitted
- Nutritional requirements
- Medications, including the dose, route and frequency
- Treatments, including amount and frequency
- Wound care orders and measurements
- Safety measures to protect against injury
- Available caregiver
- List all community or state agency services the client receives in the home including, but not limited to, primary home care (PHC), community based alternative (CBA), medically dependent children’s program (MDCP)
- Instructions for timely discharge or referral
- Documentation of coordination with PPECC, when a client receives ongoing skilled nursing in a PPECC setting. When a client receives PPECC, the SN and/or HHA provider must provide a medical rationale to support the need for SN and/or HHA services, when PPECC services are provided on the same day.

The POC must be accompanied by the physician’s signed and dated orders or must be signed and dated by the physician. The POC must include the SOC date (when the services will begin) and must be signed and dated by the assessing RN.

When a provider has received signed physician orders for SN or HHA visits, the POC does not require a physician signature before the provider contacts the claims administrator for prior authorization of services. The POC must be signed and dated by a physician familiar with the client prior to submitting a claim for services, and no later than 30 days from the SOC date.

The type and frequency of visits, supplies, or DME must appear in the POC before the physician signs the POC and may not be added after the physician has signed the POC. If any change in the POC occurs during a prior authorization period (additional visits, supplies, or DME), the provider must update the POC, have the physician sign the updated POC, and contact the claims administrator for prior authorization.

**Note:** Verbal physician orders may only be given to people authorized to receive them under state and federal law. They must be reduced to writing, signed and dated by the RN or qualified therapist responsible for furnishing or supervising the ordered service. The physician must sign the written copy of the verbal order within two weeks, or per agency policy if less than two weeks. A copy of the written verbal order must be maintained in the client’s medical record prior to and after being signed by the physician.

**Note:** All documentation, including all written and verbal orders, and all physician-signed POCs, must be maintained by the physician, and the home health agency must keep the original, signed copy of the POC in the client’s medical record.

The client must be seen by a physician within 30 days of the initial SOC, and at least once every six months thereafter unless the client’s condition changes.

A revised POC is required for every request for any change in SN or HHA visits. The revised POC must include all continuing and new orders. The revised POC must be updated to document any changes in the client’s condition or diagnosis.

A new POC is required with every request for recertification. The new POC must include all continuing and new orders. The new POC must document all changes in the client’s condition or diagnosis and reflect the need for continued SN or HHA services in relation to the original need for care. The physician must certify he or she has provided continuing care and medical supervision including, but not limited to, examination or treatment of the client within six months or when the client’s condition has changed.
3.4.2.1 Written Plan of Care (POC)

A home health services POC is required for SN, HHA, OT, or PT services. The POC is not required as an attachment with the claim, but a signed and dated POC must be maintained by the provider and primary physician in the client’s medical record. The client’s primary physician must recommend, sign, and date a POC. The POC must be initiated by the RN in a clear and legible format.

Referred to: Subsection 4.5, “Frequency and Duration Criteria for PT, OT, and ST Services” in the Physical Therapy, Occupational Therapy, and Speech Therapy Services Handbook (Vol. 2, Provider Handbooks) for additional information about PT, OT, and ST services.

Services billed in excess of those authorized for the prior authorization week or month are subject to recoupment.

For the home health services POC to be valid, the primary physician must sign and date it, and indicate when the services will begin. The home health agency must update and maintain the POC at least every 60 days or as necessitated by a change in the client’s condition.

3.4.2.2 DME and Medical Supplies Submitted with a Plan of Care (POC)

The cost of incidental medical supplies used during an SN or HHA visit are included in the rate for G0299 and G0300. Medical supplies left at the home for the client to use must be billed with the provider identifier enrolled as a DME supplier after prior authorization has been granted by the TMHP Home Health Services Prior Authorization Department.


When the home health services POC is used to submit a prior authorization for DME or medical supplies that will be used in conjunction with the professional services provided by the agency, such as SN, HHA, OT, or PT, the home health agency’s DME provider identifier must be submitted on the POC, and all of the requested DME and medical supplies must be listed in the “Supplies” section of the POC.

The POC does not require a physician’s signature before prior authorization of professional services, DME, or medical supplies is requested but does require the assessing RNs dated signature. The POC must be signed and dated by a primary physician familiar with the client prior to submitting a claim for services and no later than 30 days from the SOC date.

If the home health agency uses the Home Health Services (Title XIX) DME/Medical Supplies Physician Order Form, the agency must complete Section A. A primary physician familiar with the client must complete, sign, and date Section B prior to submission to TMHP for prior authorization of the requested DME or medical supplies.

The following information is required to consider these medical supplies for prior authorization:

- Item description
- Procedure code
- Quantity of each medical supply requested
- Manufacturer’s suggested retail price (MSRP) for items that do not have a maximum fee assigned

3.4.3 Prior Authorization of SN and HHA Services

Prior authorization requests may be submitted to the TMHP Prior Authorization Department by mail, fax, or the electronic portal. Prescribing or ordering providers, dispensing providers, clients’ responsible adults, and clients may sign prior authorization forms and supporting documentation using electronic or wet signatures.

Refer to: Subsection 5.5.1.2, “Document Requirements and Retention” in “Section 5: Fee-for-Service Prior Authorizations” (Vol. 1, General Information) for additional information about electronic signatures.

Home Health SN and HHA services require prior authorization. Providers must obtain authorization within three business days of the SOC date for an initial authorization. For recertifications, providers must obtain authorization within seven business days of the new SOC date. During the authorization process, providers are required to deliver the requested services from the SOC date, which is the date agreed to by the physician, the RN, the Home Health Agency, and the client, parent, guardian, or caregiver. The SOC must be documented on the POC.

A provider requesting prior authorization for SN or HHA Services must submit the following documentation:

- A completed client assessment
- A completed Texas Medicaid home health services POC that must:
  - Be signed and dated by the assessing RN
  - Signed and dated by the physician or submitted with the signed and dated physician’s orders.

Note: To complete the prior authorization process by paper, the SN or HHA provider must fax or mail the completed documentation to the Home Health prior authorization unit and retain a copy of the signed and dated documentation in the client’s medical record at the provider’s place of business.

Note: To complete the prior authorization process electronically, the SN or HHA provider must complete the prior authorization requirements through any approved electronic methods and retain a copy of the signed and dated documentation in the client’s medical record at the provider’s place of business.

Note: All documentation, including all written and verbal orders, and all physician-signed POCs, must be maintained by the ordering physician, and the home health agency must keep the original, signed copy of the POC in the client’s medical record.

Requests must be based on the medical needs of the client. Documentation must support the quantity and frequency of intermittent or part-time SN or HHA visits that will safely meet the client’s needs. The amount and duration of SN or HHA visits requested will be evaluated by the claims administrator. The home health agency must ensure the requested services are supported by the client assessment, POC, and the physician’s orders.

The length of the authorization is determined on an individual basis and is based on the goals and timelines identified by the physician, home health agency, RN, and client, parent, guardian, or caregiver. SN and HHA visits will be prior authorized for no more than 60 days at a time. As a client’s problems are resolved and goals are met, a client’s condition is expected to become more stable, and the client’s needs for SN and HHA services may decrease.

3.4.3.1 Routine Laboratory Specimens

SN visits to obtain routine laboratory specimens may be considered when the only alternative to obtain the specimen is to transport the client by ambulance.
3.4.3.2 **Home Phototherapy**
SN visits to address hyperbilirubinemia will not be considered for prior authorization if the client has an open authorization for home phototherapy. Home phototherapy is reimbursed as a daily global fee and includes coverage of SN visits for client, parent, or caregiver teaching and monitoring, and customary and routine laboratory work.

3.4.3.3 **Prothrombin Time/International Normalized Ration (TP/INR) Home Testing Device**
SN visits will not be authorized for the set-up or teaching of the Prothrombin Time/Internationalized Normalized Ration (TP/INR) home testing device.

3.4.3.4 **Total Parenteral Nutrition (TPN)**
SN visits to address total parenteral nutrition (TPN) must:
- Be provided by an RN appropriately trained in the administration of TPN.
- Include education of the client or caregiver regarding the in-home administration of TPN before administration initially begins.
- Include the use and maintenance of required supplies and equipment.
- Occur at least once every month to monitor the client’s status and to provide ongoing education to the client or caregivers regarding the administration of TPN.

For clients receiving PDN who also require TPN administration education, intermittent SN visits may be considered for separate prior authorization when:
- The PDN provider is not an RN appropriately trained in the administration of TPN, and the PDN provider is not able to perform the function.
- There is documentation to support the medical need for an additional skilled nurse to perform TPN.

The SN services may be prior authorized only for the client or caregiver training in TPN administration.

3.4.3.5 **Instruction in the Self-administration of Prescribed Injections**
For clients receiving SN visits, instruction to the client or caregiver in the self-administration of prescribed injections (IM, SQ, or IV), including but not limited to Factor 8 and IVIg are considered part of the existing authorized skilled nursing home visits. Additional nursing visits for teaching and (initial) supervision to the client or caregiver will not be allowed.

Instruction and initial supervision must be provided by an RN appropriately trained in the administration of the drug or product being administered, and the client and caregiver must be involved in the decision to self-administer the medication.

The client or caregiver administering the injectable medication (IM, SQ, or IV), including but not limited to Factor 8 product or IVIg, must:
- Be medically stable.
- Have a history of compliance with other medications.
- Have a simple drug regimen.
- Have the ability to read and understand directions on the medication label.
- Demonstrate knowledge of the administration technique, maintenance of the required supplies and equipment, and storage requirements.

SN visits will not be approved for the sole purpose of instructing the client on the use of the subcutaneous injection port device. Any necessary instruction must be performed as part of the office visit with the prescribing physician.
3.4.3.6 Prior Authorization Status and Limitations

The claims administrator will notify the provider of the authorization or other action taken on the request for services.

Up to a maximum combined total of three SN and HHA visits may be prior authorized per day. One visit may last up to a maximum of 2.5 hours. SN or HHA visits may be provided on consecutive days.

Note: When documentation does not support medical necessity for home health SN or HHA visits, providers may be directed to possible alternative services based on the client’s age and needs.

A nurse or HHA may be authorized to provide services to more than one client over the span of the day as long as:

• Each client’s care is based on an individualized POC; and
• Each client’s needs and POC do not overlap with another client’s needs and POC.

Settings in which a nurse or HHA provider may provide services in a provider-client ratio greater than 1:1 include, but are not limited to, homes with more than one client receiving home health services, foster homes, and independent living arrangements.

A prior authorization for SN or HHA visits is no longer valid when:

• The client is no longer eligible for Medicaid;
• The client no longer meets the medical necessity criteria for SN or HHA services;
• The place of service cannot provide for the health and safety of the client;
• The client, parent, guardian, or caregiver refuses to comply with the attending physician’s plan of treatment and compliance is necessary to assure the health and safety of the client; or
• The client changes providers and the change of notification is submitted to the claims administrator in writing with a prior authorization request from the new provider.

3.4.3.7 Canceling a Prior Authorization

The client has the right to choose their home health agency provider and to change providers. If the client changes providers, TMHP must receive a change of provider letter with a new POC or Home Health Services (Title XIX) DME/Medical Supplies Physician Order Form. The client must sign and date the letter, which must include the name of the previous provider, the current provider, and the effective date for the change.

The client is responsible for notifying the original provider of the change and the effective date. Prior authorization for the new provider can only be issued up to three business days before the date TMHP receives the change of provider letter and the new Home Health Services (Title XIX) DME/Medical Supplies Physician Order Form.

3.4.4 Medicare and Medicaid Prior Authorization

Qualified Medicare Beneficiaries (QMB) are not eligible for Medicaid benefits. Providers should not submit prior authorization requests to the TMHP Home Health Services Prior Authorization Department for these clients.

For eligible Medicare and Medicaid clients, Medicare is the primary insurance and providers must contact Medicare first for prior authorization and reimbursement. Home health service prior authorizations may be given for HHA services, certain medical supplies, or DME suitable for use in the home in one of the following instances:

• When an eligible Medicaid client (enrolled in Medicare) does not qualify for home health services under Medicare because SN care, OT, or PT are not a part of the client’s care.
When the medical supplies and DME are not a benefit of Medicare Part B and are a home health services benefit.

Federal and state laws require the use of Medicaid funds for the payment of most medical services only after all reasonable measures have been made to use a client’s third party resources or other insurance.

**Note:** If the client has Medicare Part B coverage, contact Medicare for prior authorization requirements and reimbursement. If the service is a Part B benefit, do not contact TMHP for prior authorization.

To ensure that Medicare benefits are used first in accordance with Texas Medicaid regulations, the following procedures apply when requesting Medicaid prior authorization and payment of home health services for clients:

- Contact TMHP for prior authorization of Medicaid services (based on medical necessity and home health services benefits) within 30 days of the date on the MRAN. Fax a copy of the original MRAN and the Medicare appeal review letter to the TMHP Home Health Services Prior Authorization Department for prior authorization.

- An MRAN is not required when a client is eligible for Medicare or Medicaid and needs HHA visits only. However, a skilled supervisory nursing visit must be made on the same day as the initial HHA visit and at least every 60 days (on the same day an HHA visit is made) thereafter as long as no skilled need exists. An SN supervisory visit is reimbursable, but an SN visit made for the primary purpose of assessing a client’s nursing care is not. The SOC date will be the date of the first requested Medicare home health services visit as listed on the original MRAN.

  **Note:** Claims for State of Texas Access Reform (STAR)+PLUS MQMB clients (those with Medicare and Medicaid) should always be submitted to TMHP as noted on these pages. The STAR+PLUS health plan is not responsible for these services if Medicare denies the service as not a benefit.

For Medicaid qualified Medicare beneficiary (MQMB) clients, do not submit prior authorization requests to TMHP if the Medicare denial reason states “not medically necessary.”

Medicaid will only consider prior authorization requests if the Medicare denial states “not a benefit” of Medicare.

When the client is 65 years of age or older or appears otherwise eligible for Medicare (e.g., a person who is blind or disabled), but has no Part A or Part B Medicare, the TMHP Home Health Services Prior Authorization Department uses regular prior authorization procedures. In this situation, the claim is held for a midyear status determined by HHSC. The maximum length of time a claim may be held in a “pending status” for Medicare determination is 90 days. After the waiting period, the claim is paid or denied. If denied, the EOB code on the R&S report indicates that Medicare is to be billed.

**Refer to:** Subsection 3.2.1, “Skilled Nursing Visits” in this handbook.

Home health providers should follow these guidelines:

- Clients who are 64 years of age and younger without Medicare Part A or B:
  - If the agency erroneously submits an SOC notice to Medicare and does not contact TMHP for prior authorization, TMHP does not assume responsibility for any services provided before contacting TMHP. The SOC date is no more than three business days before the date the agency contacts TMHP. Visits made before this date are not considered a benefit of Texas Medicaid.
  
- Clients who are 65 years of age and older without Medicare Part A or Part B and clients with Medicare Part A or B regardless of age:
• In filing home health claims, home health providers may be required to obtain Medicare denials before TMHP can approve coverage. When TMHP receives a Medicare denial, the SOC is determined by the date the agency requested coverage from Medicare. If necessary, the 95-day claims filing deadline is waived for these claims, provided TMHP receives notice of the Medicare denial within 30 days of the date on the MRAN containing Medicare’s final disposition.

• If the agency receives the MRAN and continues to visit the client without contacting TMHP by telephone, mail, or fax within 30 days of the date on the MRAN, TMHP will provide coverage only for services provided from the initial date of contact with TMHP. The SOC date is determined accordingly. TMHP must have the MRAN before considering the request for prior authorization.

TMHP will not prior authorize or reimburse the difference between the Medicare payment and the retail price for Medicare Part B eligible clients.

Refer to: “Section 8: Third Party Liability (TPL)” (Vol. 1, General Information).

3.5 Home Health SN and HHA Procedure Billing and Limitations

Home Health SN or HHA visits provided by home health agencies enrolled in Texas Medicaid must be billed using procedure codes G0299 (SN), G0300 (SN), and G0156 (HHA) and will be reimbursed per visit of up to 2.5 hours; not to exceed a combined total of three visits per day (7.5 hours total).

The reimbursement methodology for professional services delivered by home health agencies is a statewide visit rate calculated in accordance with 1 TAC §355.8021.

When services are provided to more than one client in the same setting, only the units directly provided to each client at distinct, separate time periods will be reimbursed. Provider documentation must support the services were delivered at distinct, separate time periods. Total home health services billed for all clients cannot exceed the individual provider’s total number of hours spent at the place of service.

One as needed (PRN) SN visit may be reimbursed every 30 days outside of the prior authorized visits when SN visits have been authorized for the particular client.

For reimbursement purposes, Home Health SN or HHA services are always billed as place of service 2 (home) regardless of the setting in which the services are actually provided. SN or HHA services provided in the day care or school setting will not be reimbursed.

For all clients, SN visits may be provided in the following locations:

• Home of the client, parent, guardian, or caregiver
• Foster homes
• Independent living arrangements

For all clients, HHA visits may be provided in the following locations:

• Home of the client, parent, guardian, or caregiver
• Foster homes
• Independent living arrangements

An immediate relative, parent or guardian, primary caregiver, or alternate care giver may not be reimbursed for HHA services even if he or she is an enrolled provider or employed by an enrolled provider.
SN and/or HHA services may be billed on the same day as PPECC services, but they may not be billed simultaneously with PPECC services. SN and/or HHA services may be billed before or after PPECC services.

Note: SN and/or HHA services are subject to retrospective review and possible recoupment when the medical record does not document the provision of SN and/or HHA services are medically necessary based on the client’s situation and needs. The service provider’s record must explain all discrepancies between the service hours approved and the service hours provided. For example: the parents released the provider from all responsibility for the service hours or the agency was not able to staff the service hours. The release of provider responsibility does not indicate the client does not have a medical need for the services during those time periods.

3.5.1 Skilled Nursing Visit for TPN Education
The nurse providing the intermittent SN visit for TPN services will only be reimbursed for time spent delivering client or family instruction and for direct client TPN services. The services delivered must be documented in the client’s record.

PDN and SN should not be routinely performed on the same date during the same time period.

PDN and SN will not be considered for reimbursement when the services are performed on the same date during the same time period without prior authorization approval.

If the SN visit for TPN education occurs during a time period when the PDN provider is caring for the client, both the PDN provider and the nurse educator must document in the client’s medical record the skilled services individually provided, including but not limited to:

- The start and stop time of each nursing provider’s specialized task(s)
- The client condition that requires the performance of skilled PDN tasks during the SN visit for TPN education
- The skilled services that each provided during that time period

Both the intermittent SN visit and the PDN services provided during the same time period may be recouped if the documentation does not support the medical necessity of each service provided.

A skilled nursing visit for TPN education is not allowable in a PPECC setting.

3.5.2 Medication Administration Limitations
Nursing visits for the purpose of administering medications are not a benefit if one of the following conditions exists:

- The medication is not considered medically necessary to the treatment of the individual’s illness or is not approved by the Food and Drug Administration (FDA) or is being used for indications not approved by the FDA.
- The administration of medication exceeds the therapeutic frequency or duration by accepted standards of medical practice.
- A medical reason does not prohibit the administration of the medication by mouth.
- The client, a primary caregiver, a family member, or neighbor have been taught or can be taught to administer subcutaneous (SQ/SC), intramuscular (IM), and intravenous (IV) injections and has demonstrated competency.
- The medication is a chemotherapeutic agent or blood product SQ/SC, IM, and IV injections.
4 Private Duty Nursing (PDN) Services - CCP

4.1 Services, Benefits, Limitations, and Prior Authorization

PDN services are a benefit of the Texas Health Steps-Comprehensive Care Program (THSteps-CCP) for Medicaid clients who are 20 years of age or younger. PDN services are nursing services, as described by the Texas Nursing Practice Act and its implementing regulations, for clients who meet the medical necessity criteria, and who require individualized, continuous, skilled care beyond the level of SN visits normally authorized under Texas Medicaid Home Health SN and Home Health Aide (HHA) Services. PDN services may be provided by a registered nurse (RN) or a licensed vocational nurse (LVN).

Note: Texas defines a licensed practical nurse (LPN) as a licensed vocational nurse (LVN).

The following procedure code is a benefit of Texas Medicaid when PDN services are provided by a home health agency or an independently enrolled RN or LVN. Appropriate modifiers from the Modifier table must be submitted for reimbursement purposes, but are not required for prior authorization. Independently enrolled RNs or LVNs must include modifier U3 along with TD or TE for reimbursement purposes. An appropriate diagnosis from the Diagnosis Codes table must be submitted when modifier UA is used to obtain additional reimbursement for clients with a tracheostomy or who are ventilator dependent.

<table>
<thead>
<tr>
<th>Procedure Code</th>
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<tbody>
<tr>
<td>T1000</td>
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<table>
<thead>
<tr>
<th>Modifiers</th>
<th>Description</th>
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<tbody>
<tr>
<td>TD</td>
<td>Registered nurse (RN)</td>
</tr>
<tr>
<td>TE</td>
<td>Licensed vocational nurse (LVN)</td>
</tr>
<tr>
<td>U3</td>
<td>Independently enrolled provider</td>
</tr>
<tr>
<td>UA</td>
<td>Specialized services</td>
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</tbody>
</table>

<table>
<thead>
<tr>
<th>Diagnosis Codes for use with Modifier UA only</th>
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</thead>
<tbody>
<tr>
<td>J9500</td>
</tr>
<tr>
<td>Z930</td>
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</table>

Example: Procedure code T1000 would be submitted with modifiers TD, U3, and UA for reimbursement purposes for PDN in increments of up to 15 minutes when provided by an independently enrolled RN to a client who has a tracheostomy or is ventilator dependent.

Because of the nature of the service being provided, some billing situations are unique to PDN. These billing requirements are as follows:

- All hours worked on one day must be billed together, on one detail, even if they involve two shifts. For example, if Nurse A works 7 a.m. to 11 a.m. and then returns and works 7 p.m. to 11 p.m., services must be billed for 8 hours (32 15-minute units) on one detail for that date of service.

- An individually-enrolled nurse will not be reimbursed for more than 16 hours of PDN services in one day.

PDN may be delivered in a provider to client ratio other than one-on-one. An RN or LVN may provide PDN services to more than one client over the span of the day as long as each client’s care is based on an individualized POC, and each client’s needs and POC do not overlap with another client’s needs and POC. Only the time spent on direct PDN for each client is reimbursed. Total PDN billed for all clients cannot exceed an individual provider’s total number of hours at the POS.
A single nurse may be reimbursed for services to more than one client in a single setting when the following conditions are met:

- The hours for PDN for each client have been authorized through the TMHP Prior Authorization Department.
- Only the actual “hands-on” time spent with each client is billed for that client.
- The hours billed for each client do not exceed the total hours approved for that client and do not exceed the actual number of hours for which services were provided.

**Example:** If the prior authorized PDN hours for Client A is four hours, Client B is six hours, and the actual time spent with both clients is eight hours, the provider must bill for the actual one-on-one time spent with each client, not to exceed the client’s prior authorized hours or total hours worked. It would be acceptable to bill four hours for Client A and four hours for Client B, or three hours for Client A and five hours for Client B. It would not be acceptable to bill five hours for Client A and three hours for Client B. It would be acceptable to bill ten hours if the nurse actually spent ten hours onsite providing prior authorized PDN services split as four hours for Client A and six hours for Client B. A total of ten hours cannot be billed if the nurse worked only eight hours.

For reimbursement purposes, PDN must always be submitted with POS 2 (home) regardless of the setting in which services are actually provided. PDN may be provided in any of the following settings:

- Client’s home
- Nurse provider’s home
- Client’s school
- Client’s daycare facility

A parent or guardian of a minor client, or the client’s spouse may not be reimbursed for PDN services even if he or she is an enrolled provider or employed by an enrolled provider.

PDN services are subject to retrospective review and possible recoupment when the medical record does not document the provision of PDN services are medically necessary based on the client’s situation and needs. The PDN services provider’s record must explain all discrepancies between the service hours approved and the service hours provided. For example: the parents released the provider from all responsibility for the service hours or the agency was not able to staff the service hours. The release of provider responsibility does not indicate the client does not have a medical need for the services during those time periods.

### 4.1.1 Medical Necessity

Texas Medicaid defines medically necessary EPSDT services as health care, diagnostic services, treatments, and other measures necessary to correct or ameliorate any disability, physical or mental illness, or chronic conditions.

Medicaid clients who are 20 years of age or younger, and who are eligible for THSteps, are entitled to all medically necessary PDN services to promote independence and support the client living at home.

PDN services are considered medically necessary when a client has a disability, physical or mental illness, or chronic condition, and he or she requires continuous, skillful observations, judgments, and interventions to correct or ameliorate his or her health status.

The following elements should always be addressed in documentation submitted with a request for PDN services:

- Dependent on technology to sustain life.
• Requires ongoing and frequent skilled interventions to maintain or improve health status; and delayed skilled intervention is expected to result in:
  • Deterioration of a chronic condition;
  • Loss of function;
  • Imminent risk to health status due to medical fragility; or
  • Risk of death.

4.1.2 PDN Services

All PDN services must be prior authorized.

PDN services provide nursing care and parent, guardian, or responsible adult training and education intended to:

• Optimize client health status and outcomes; and
• Promote family-centered, community-based care as a component of an array of service options by:
  • Preventing prolonged or frequent hospitalizations or institutionalization.
  • Providing cost-effective and quality care in the most appropriate, least restrictive environment.

PDN services are nursing services ordered by a physician, included in the nursing plan of care (POC), and provided by an RN or LVN.

Note: An advanced practice registered nurse (APRN) or a physician assistant (PA) may sign all documentation related to the provision of private duty nursing services on behalf of the client’s physician when the physician delegates this authority to the APRN or PA. The APRN or PA provider’s signature and license number must appear on the forms where the physician signature and license number blocks are required.

Professional nursing provided by an RN, as defined in the Texas Nursing Practice Act, means that the performance of an act that requires substantial specialized judgment and skill, the proper performance of which is based on knowledge and application of the principles of biological, physical, and social science as acquired by a completed course in an approved school of professional nursing. The term does not include acts of medical diagnosis or the prescription of therapeutic or corrective measures. Professional nursing involves:

• The observation, assessment, intervention, evaluation, rehabilitation, care and counsel, or health teachings of a person who is ill, injured, infirm, or experiencing a change in normal health processes.
• The maintenance of health or prevention of illness.
• The administration of a medication or treatment as ordered by a physician, podiatrist, or dentist.
• The supervision of delegated nursing tasks or teaching of nursing.
• The administration, supervision, and evaluation of nursing practices, policies, and procedures.
• The performance of an act delegated by a physician.
• Development of the nursing care plan.

Vocational nursing, as defined in the Texas Nursing Practice Act, means a directed scope of nursing practice, including the performance of an act that requires specialized judgment and skill, the proper performance of which is based on knowledge and application of the principles of biological, physical, and social science as acquired by a completed course in an approved school of vocational nursing. The term does not include acts of medical diagnosis or the prescription of therapeutic or corrective measures. Vocational nursing involves:

• Collecting data and performing focused nursing assessments of the health status of an individual.
• Participating in the planning of the nursing care needs of an individual.

• Participating in the development and modification of the nursing care plan.

• Participating in health teaching and counseling to promote, attain, and maintain the optimum health level of an individual.

• Assisting in the evaluation of an individual’s response to a nursing intervention and the identification of an individual’s needs.

• Engaging in other acts that require education and training, as prescribed by board rules and policies, commensurate with the nurse’s experience, continuing education, and demonstrated competency.

Professional and vocational nursing care consists of those services that must, under state law, be performed by an RN or LVN, and are further defined as nursing services in the Code of Federal Regulations (42 CFR §§ 409.32, 409.33, and 409.44).

• In determining whether a service requires the skill of a licensed nurse, consideration must be given to the inherent complexity of the service, the condition of the beneficiary, and the accepted standards of medical and nursing practice.

• The fact that the nursing care can be, or is, taught to the client or to the client’s family or friends does not negate the skilled aspect of the service when the service is performed by a nurse.

• If the service could be performed by the average nonmedical person, the absence of a competent person to perform it does not cause it to be a nursing service.

• If the nature of a service is such that it can safely and effectively be performed by the average nonmedical person without direct supervision of a licensed nurse, the services cannot be regarded as nursing care.

• Some services are classified as a nursing care on the basis of complexity alone (e.g., intravenous and intramuscular injections or insertion of catheters), and if medically necessary for the treatment of the client’s illness or injury, would be covered on that basis. However, in some cases, the client’s condition may cause a service that would ordinarily be considered unskilled to be considered nursing care. This would occur when the client’s condition is such that the service can be safely and effectively provided only by a nurse.

• A service that, by its nature, requires the skills of a nurse to be provided safely and effectively continues to be a skilled service even if it is taught to the client, the client’s family, or other responsible adults.

Because Texas Medicaid is obligated to provide all medically necessary PDN services, a parent or guardian is not obligated to provide PDN services even if the parent or guardian has received the appropriate training. Medically necessary PDN services will not be denied to clients based on the parent or guardian’s ability to provide the necessary PDN services.

PDN services that are intended to provide mainly respite care; child care; or do not directly relate to the client’s medical needs or disability are not a benefit of Texas Medicaid.

The delivery of PDN services may inherently result in the relief of the parent, guardian, or responsible adult, child care, or some nonmedical, nonskilled activities in the course of providing nursing care.

4.1.3 PDN Providers

PDN services must be provided by a licensed home health services agency or a licensed and certified home health services agency enrolled in Texas Medicaid or by an RN or LVN enrolled independently with Texas Medicaid. PDN providers must comply with all applicable federal, state, and local laws and regulations and Texas Medicaid policies and procedures.
All providers must maintain written policies and procedures for:

- Obtaining consent for medical treatment for clients in the absence of the parent or guardian that meet the standards of the Texas Family Code, Chapter 32.
- Obtaining physician signatures for all telephone orders within 14 calendar days of receipt of the order.

PDN providers must only accept clients on the basis of a reasonable expectation that the client’s needs can be adequately met in the place of service. The essential elements of safe and effective PDN services include a responsible adult when the client is a minor child, a contingency plan, a primary physician, competent providers, and an environment that supports the client’s health and safety needs.

The place of service must be able to support the health and safety needs of the client and must be adequate to accommodate the use, maintenance, and cleaning of all medical devices, equipment, and supplies required by the client. Necessary primary and back-up utilities, communications and fire safety systems must be available.

Clients who are 17 years of age or younger must reside with an identified responsible adult who is either trained to provide nursing care or is capable of initiating an identified contingency plan when the scheduled private duty nurse is unexpectedly unavailable.

An identified responsible adult is an individual 18 years of age or older who has agreed to accept the responsibility for a client’s provision of food, shelter, clothing, education, nurturing, and supervision. Responsible adults include, but are not limited to: biological parents, adoptive parents, foster parents, guardians, individuals court-appointed as managing conservators, and other family members by birth or marriage.

An identified contingency plan is a structured process, designed by the responsible adult and the PDN provider, by which a client will receive care when a scheduled private duty nurse is unexpectedly unavailable, and the responsible adult is unavailable, or is not trained, to provide the nursing care. The identified responsible adult must be able to initiate the contingency plan.

The responsible adult’s signature must be on the form acknowledging:

- Information about PDN has been discussed and received.
- PDN may change or end based on a client’s need for nursing care.
- PDN is not authorized for the primary purpose of providing respite, childcare, ADLs, or housekeeping.
- All requirements have been met before seeking prior authorization for PDN.
- The responsible adult has participated in the development of the POC and the nursing care plan for the client.
- Emergency plans have been made and are part of the client’s care plan.
- The client or responsible adult agrees to follow the physician’s POC.

**Note:** A responsible adult of a minor client or a client’s spouse may not be reimbursed for PDN even if the responsible adult is an enrolled provider or employed by an enrolled provider.

PDN services may be delivered in a provider or client ratio other than 1:1.

**4.1.4 Authorization Requirements**

PDN services require prior authorization.

All requests for PDN services must be based on the current medical needs of the client.
PDN services will not be prior authorized when:

- The client does not meet medical necessity criteria as defined in the Statement of Benefits.
- The client does not have a primary physician;
- The client is not 20 years of age or younger;
- When the client’s needs are not beyond the scope of services available through Medicaid Title XIX Home Health SN or HHA Services because the needs can be met on a part-time or intermittent basis.

Requests for PDN must be based on the current medical needs of the client.

The following criteria are considered for PDN prior authorization:

- The documentation submitted with the request is complete.
- The requested services are nursing services as defined by the Texas Nursing Practice Act and its implementing regulations.
- The explanation of the client’s medical needs is sufficient to support a determination that the requested services correct or ameliorate the client’s disability, physical or mental illness, or chronic condition.
- The client’s nursing needs cannot be met on an intermittent or part-time basis through Texas Medicaid (Title XIX) home health services skilled nursing services.
- There is no TPR financially responsible for the services.

Only those services that meet the medical necessity criteria for PDN are reimbursed. Before the TMHP Prior Authorization Department or MCO determines the requested nursing services do not meet the criteria, the Medical Director contacts the treating physician to determine whether additional information or clarification can be provided that would allow for the prior authorization of the requested PDN. If the Medical Director is not successful in contacting the treating physician or cannot obtain additional information or clarification, the Medical Director makes a decision based on the available information.

Providers must obtain prior authorization within three business days of the SOC for services that have not been prior authorized. During the prior authorization process, providers are required to deliver the requested services from the SOC date. The SOC date is the date agreed to by the physician, the PDN provider, and the client or responsible adult and is indicated on the submitted POC as the SOC date.

Note: The TMHP Prior Authorization Department does not prior authorize an SOC date earlier than seven calendar days before contact with TMHP.

Requests for nursing services must be submitted on the required Medicaid authorization forms and include supporting documentation. The supporting documentation must:

- Clearly and consistently describe the client’s current diagnosis, functional status, and condition.
- Consistently describe the treatment throughout the documentation.
- Provide a sufficient explanation as to how the requested nursing services correct or ameliorate the client’s disability, physical or mental illness, or condition.

4.1.4.1 Authorization Forms

The CCP Prior Authorization Request Form must be completed, signed, and dated by the physician. When PDN services are ordered, by signing the form the physician attests and certifies the client’s medical condition is sufficiently stable to permit safe delivery of PDN as described in the plan of care. All requested dates of service must be included.
The POC must be recommended, signed, and dated by the client’s primary physician. A POC must meet the standards outlined in the 42 CFR §484.18 related to the written POC. The primary physician must review and revise the POC, in consultation with the provider and the responsible adult, for each prior authorization, or more frequently as the physician deems necessary or the client’s situation changes.

The Nursing Addendum to Plan of Care for Private Duty Nursing and/or Prescribed Pediatric Extended Care Centers form addresses PDN eligibility criteria, nursing care plan summary, health history summary, 24-hour schedule, and the rationale for the hours of PDN requested.

The following is a description of the nursing care plan summary:

- The nursing care summary is not a complete nursing care plan.
- Information must be client-focused and detailed.
- The problem list must reflect the reasons that nursing services are needed. The problem list is not the nursing care plan. Providers must identify two-to-four current priority problems from their nursing care plan. The problem does not need to be stated as a nursing diagnosis. The problems listed must focus on the primary reasons that a licensed nurse is required to care for the client. Other attached documents are not accepted in lieu of this section.
- The Goals must relate directly to the problems listed and be client-specific and measurable. Goals may be short- or long-term; however, for many clients who receive PDN, the goals generally are long-term.
- The Outcomes are the effects of the provider’s nursing interventions and must be measurable. Generally, these are more short-term than goals. For initial requests, list expected outcomes. Extension requests should note the results of nursing interventions.
- The Progress must be viewed as a “yardstick” or continuum on which progress toward goals is marked. Initial requests must state expected progress for the authorization period. Extension requests must list the progress noted during the previous authorization period. It is recognized that all progress may not be positive.
- The addendum must summarize the client’s health problems relating to the medical necessity for PDN.
- The addendum must clearly communicate a picture of the client’s overall condition and nursing care needs.
- The summary of recent health history is imperative in determining whether the client’s condition is stable or if new nursing care needs have been identified. This section gives the PDN provider an opportunity to describe the client’s recent health problems, including acute episodes of illness, hospitalizations, injuries, and so on. The summary should create a complete picture of the client’s condition and nursing care needs. The summary may cover the previous 90 days, even though the authorization period is 60 days; however, the objective of the summary is to capture the client’s recent health problems and current health priorities. This section should not be merely a list of events. This section is the place to indicate the frequency of nursing interventions if they are different from the physician’s order on the POC, such as, the order may be for a procedure to be PRN (Pro Re Nata “As Needed”), but it is actually being performed every two hours.
- The addendum must include the rationale for increasing, decreasing, or maintaining the level of PDN and must relate to the client’s health problems and goals.
- The addendum must include the provider’s plan to decrease hours or discharge from service (if appropriate).
All direct-care services must be identified in the client’s 24-hour daily schedule. It is understood that the schedule may change, as the client’s needs change. The TMHP Prior Authorization Department does not have to be notified of changes in the schedule except as they occur when a PDN recertification is requested.

4.1.4.2 Primary Physician Requirements

The client must have a primary physician who provides continuing care and medical supervision, including, but not limited to, examination or treatment within 30 calendar days prior to the start of PDN services.

The physician visit may be waived when a diagnosis has already been established by the physician, and the client is under the continuing care and medical supervision of the physician. A waiver is valid for no more than 365 days, and the client must be seen by his or her physician at least once every 365 days. The waiver must be based on the physician’s written statement that an additional evaluation visit is not medically necessary. This documentation must be maintained by the physician and the provider in the client’s medical record.

Note: An advanced practice registered nurse (APRN) or a physician assistant (PA) may sign all documentation related to the provision of private duty nursing services on behalf of the client’s physician when the physician delegates this authority to the APRN or PA.

The primary physician must:

- Provide a prescription for PDN services
- Recommend, sign, and date a plan of care (POC)
- Sign a statement of need that PDN services are medically necessary
- Maintain documentation that the client’s medical condition will allow safe delivery of PDN services as described in the POC

The physician recommended POC must include the following:

- The client’s Medicaid number; the physician’s license number; and the provider’s Medicaid number
- Date the client was last seen by the physician
- The start of care (SOC) date for PDN services
- All pertinent diagnoses
- The client’s mental status
- The prognosis
- The types of service requested, including the amount, duration, and frequency
- The equipment or supplies required
- Rehabilitation potential
- Prior and current functional limitations
- Activities permitted
- Nutritional requirements
- Medications, including the dose, route, and frequency
- Treatments, including amount and frequency
- Wound care orders and measurements
- Safety measures to protect against injury
• Responsible adult when the client is a minor child
• Contingency plan
• List all community or state agency services the client receives in the home (including, but not limited to, PCS, Community First Choice (CFC), MDCP
• Instructions for timely discharge or referral
• Client specific goals, including if receiving PPECC, the goal of ensuring coordination of ongoing skilled nursing services with the PPECC provider.
• If the client also receives PPECC services, documentation that the client or client’s responsible adult has been involved in the POC development, and description of how ongoing skilled nursing services will be coordinated between PDN and PPECC providers.

Refer to: 42 CFR §484.18 for additional information about POC requirements.

Physician Recertification of PDN services:
• The primary physician’s medical care must comply with the THSteps periodicity schedule.
• The primary physician must provide specific, written, dated orders for clients who are receiving continuing or ongoing PDN services.

4.1.4.3 PDN Provider Requirements
When a provider receives a referral for PDN services, the provider must have an RN perform a nursing assessment of the client within the client’s home environment. This assessment must be performed before seeking prior authorization for PDN services, with any request for PDN services recertification, or any request to modify PDN service hours. The assessment includes, but is not limited to, the determination of:
• Medical necessity for PDN services
• Safety of providing care in the proposed setting;
• Appropriateness of care in the place of service
• Receptivity to training and ability level of the parent, guardian, or responsible adult
• The existing level of care and any additional health-care services to include, but not limited to, School Health and Related Services (SHARS), MDCP, PT, OT, ST, PCS, CFC, or case management services. Services provided under these programs will not prevent a client from obtaining medically necessary services. Certain school services are provided to meet education needs, not medical needs.

When an RN completes a client assessment and identifies a medical necessity for ADLs or health-related functions to be provided by a nurse, the scope of PDN services may include these ADLs or health-related functions.

Note: The TMHP Prior Authorization Department does not review or authorize PDN based on partial or incomplete documentation.

4.1.4.4 Prior Authorization of PDN Services
Prior authorization requests may be submitted to the TMHP Prior Authorization Department by mail, fax, or the electronic portal. Prescribing or ordering providers, dispensing providers, clients’ responsible adults, and clients may sign prior authorization forms and supporting documentation using electronic or wet signatures.

Refer to: Subsection 5.5.1.2, “Document Requirements and Retention” in “Section 5: Fee-for-Service Prior Authorizations” (Vol. 1, General Information) for additional information about electronic signatures.
PDN services require prior authorization. Providers must obtain authorization within three business days of the SOC for services that have not been prior authorized. During the authorization process, providers are required to deliver the requested services from the SOC date. The SOC date is the date agreed upon by the physician, the PDN provider, and the client, parent, or guardian and is indicated on the submitted POC as the SOC date.

**Note:** Not including PDN services provided during the authorization process, coverage periods may not coincide with calendar weeks or months. A prior authorized week coverage period begins from the day of the week the prior authorization period begins on and continues for seven days. For example, if the prior authorization starts on a Thursday, the prior authorization week runs Thursday through Wednesday. The number of nursing hours authorized for a week must be contained in that prior authorization week. Hours billed in excess of those authorized for the PAN week are subject to recoupment.

A PDN provider requesting prior authorization for PDN services must submit all of the following documentation:

- A completed THSteps-CCP Prior Authorization Request form signed and dated by the primary physician within 30 calendar days prior to the SOC date.
- A completed POC form, signed and dated by the primary physician within 30 calendar days prior to the SOC date.
- A completed Nursing Addendum to Plan of Care for Private Duty Nursing and/or Prescribed Pediatric Extended Care Centers form signed and dated by the primary physician, RN completing the assessment, and client, parent, guardian, or responsible adult within 30 calendar days prior to the SOC date. The Nursing Addendum form must include:
  - An updated problem list
  - An updated rationale or summary page
  - A contingency plan
  - A 24-hour daily care flowsheet
  - A signed Acknowledgement

All documentation must be maintained by the requesting PDN provider. The PDN provider may be asked to submit additional documentation including, but not limited to, nurse’s notes, medication administration records, seizure logs, and ventilator logs to support medical necessity as defined in this handbook.

The request can be submitted as follows:

- To complete the prior authorization process by paper, the PDN provider must fax or mail the completed PDN Request Documentation to the TMHP Prior Authorization Department and retain a copy of the signed and dated documentation in the client’s medical record at the provider’s place of business.
- To complete the prior authorization process electronically, the PDN provider must complete the prior authorization requirements through any approved electronic methods and retain a copy of the signed and dated documentation in the client’s medical record at the provider’s place of business.

Requests for authorizations of PDN services should always be commensurate with the client’s medical needs. Requests for services should reflect changes in the client’s condition that affect the amount and duration of PDN.

For clients who are receiving PDN services who also require phototherapy, parent or guardian education, instructional use of the phototherapy equipment, and obtaining laboratory specimens collection are included in the PDN care provided.
Authorizations for more than 16 hours per day will not be issued to a single, independently enrolled nurse.

The length of the authorization is determined on an individual basis and is based on the goals and timelines identified by the physician, provider, and client, parent, or guardian. PDN services will not be authorized for more than six months at a time.

The TMHP Prior Authorization Department or MCO is required to notify the client, parent, or guardian, physician, and provider of the approval, denial, or other action taken in response to the authorization request by fax or mail.

Home Health Agencies must provide written notice to clients of their intent to voluntarily terminate PDN services at least five calendar days prior to terminating services, except in situations of a potential threat to the provider’s personal safety.

Independently enrolled RNs or LVNs must provide written notice to clients of his or her intent to voluntarily terminate services at least 30 calendar days prior to terminating services, except in situations where there is a potential threat to the nurse’s personal safety.

For clients who are receiving PDN services who also require TPN administration education, intermittent SN visits may be separately authorized when the SN services are for client or client caregiver training in TPN administration, and the PDN provider is not an RN appropriately trained in the administration of TPN, and the PDN provider is not able to perform the function.

Refer to: Subsection 3.3, “Home Health Skilled Nursing and Home Health Aide Services Providers” in this handbook for detailed information about SN benefits.

If the client has no skilled nursing need other than provision of education for self administration of prescribed injections (IM, SQ, or IV), then the client does not qualify for private duty nursing services. Nursing hours for the sole purpose of education to the client and caregiver may be considered through intermittent home health skilled nursing visits.

4.1.4.5 Initial Authorization

An initial PDN prior authorization request requires all of the following documentation:

- A completed THSteps-CCP Prior Authorization Request form signed and dated by the primary physician within 30 calendar days prior to the SOC date.
- A completed POC form, signed and dated by the primary physician within 30 calendar days prior to the SOC date.
- A completed Nursing Addendum to Plan of Care for Private Duty Nursing and/or Prescribed Pediatric Extended Care Centers form signed and dated by the primary physician, RN completing the assessment, and client, parent, guardian, or responsible adult within 30 calendar days prior to the SOC date. The completed Nursing Addendum form must include all the following:
  - An updated problem list
  - An updated rationale or summary page
  - A contingency plan
  - A 24-hour daily care flowsheet
  - A signed Acknowledgement

Note: An advanced practice registered nurse (APRN) or a physician assistant (PA) may sign all documentation related to the provision of private duty nursing services on behalf of the client’s physician when the physician delegates this authority to the APRN or PA.
All documentation must be maintained by the requesting PDN provider. The PDN provider may be asked to submit additional documentation including, but not limited to, nurse’s notes, medication administration records, seizure logs, and ventilator logs to support medical necessity as defined in this handbook.

Initial requests must be submitted within three business days of the SOC date.

Initial requests may be prior authorized for a maximum of 90 days.

Completed initial requests must be received and dated by the TMHP Prior Authorization Department within three business days of the SOC. The request must be received by the TMHP Prior Authorization Department no later than 5 p.m., Central Time, on the third day to be considered received within three business days. If a request is received more than three business days after the SOC, or after 5 p.m., Central Time, on the third day, authorization is given for dates of service beginning three business days before receipt of the completed request.

4.1.4.6 Revisions

The provider may request a revision at any time during the authorization period if medically necessary. The provider must notify the TMHP Prior Authorization Department at any time during an authorization period if the client’s condition changes and the authorized services are not commensurate with the client’s medical needs.

Requests for revisions must be submitted within three business days of the revised SOC date.

Revisions during a current authorization period must fall within that authorization period. If the revision is requested outside of an authorization period, the provider must request a new authorization and submit the following documentation:

- A completed THSteps-CCP Prior Authorization Request form signed and dated by the primary physician within 30 calendar days prior to the SOC date.
- A completed POC form, signed and dated by the primary physician within 30 calendar days prior to the SOC date.
- A completed Nursing Addendum to Plan of Care for Private Duty Nursing and/or Prescribed Pediatric Extended Care Centers form signed and dated by the primary physician, RN completing the assessment, and parent, guardian, client, or responsible adult within 30 calendar days prior to the SOC date.

Note: An advanced practice registered nurse (APRN) or a physician assistant (PA) may sign all documentation related to the provision of private duty nursing services on behalf of the client’s physician when the physician delegates this authority to the APRN or PA.

Note: All documentation must be maintained by the requesting PDN provider. The PDN provider may be asked to submit additional documentation including, but not limited to nurse’s notes, medication administration records, seizure logs, and ventilator logs to support medical necessity.

Revised services may be prior authorized for up to a maximum of six months.

A request for a client that does not satisfy the criteria listed above for a six-month authorization may be authorized for a period up to three months.

The provider is responsible for ensuring that the physician reviews and signs the POC within 30 calendar days of the start date of the revised authorization period, or more often if required by the client’s condition or agency licensure. The provider must maintain the physician-signed POC in the client’s record. PDN providers should not submit a revised POC unless they are requesting a revision.

Completed requests for revision of PDN hours during the current authorization period must be received by TMHP Prior Authorization Department within three business days of the revised SOC. The request must be received by TMHP Prior Authorization Department no later than 5 p.m., Central Time, on the
seventh day to be considered received within three business days. If a request is received more than three
business days after the revised SOC or after 5 p.m., Central Time, on the third day, authorization is given
for dates of service beginning three business days before receipt of the completed request.

Revisions to a current certification must fall within the certification period. If the revision extends
beyond the current certification period, new authorization documentation must be submitted to TMHP
Prior Authorization Department.

4.1.4.7 Required Coordination between PDN and Prescribed Pediatric Extended
Care Centers (PPECCs)

When a client or client’s physician notifies the PDN provider that the client also receives services from
a Medicaid enrolled PPECC provider, the PDN provider must coordinate services with the PPECC
provider. Both PDN and PPECC services are considered ongoing skilled nursing. A client has a choice
of PDN, PPECC, or a combination of both PDN and PPECC for ongoing skilled nursing where PPECC
services are available. Skilled Nursing services are authorized for a set number of hours based on the
client’s medical necessity at the time of the prior authorization request. Skilled nursing hours are not
expected to increase when the client utilizes a combination of both PDN and PPECC services, unless
there is a documented change in medical condition, or the authorized hours are not commensurate to
the client’s medical needs and additional hours are medically necessary.

PDN and PPECC providers must collaborate in developing their respective 24-hour flow charts found
in the Nursing Addendum to Plan of Care for Private Duty Nursing and/or Prescribed Pediatric
Extended Care Centers form each time a client’s authorization for ongoing skilled nursing is initiated,
renewed and revised.

Both providers must maintain documentation that the client or the client’s responsible adult has partic-
ipated in the development of the POC (e.g., the completed Home Health Plan of Care and Nursing
Addendum, with client or client’s responsible adult signatures.

Both providers must discuss with the client or the client’s responsible adult how care will be coordinated
between the two providers.

When a new service is initiated for ongoing skilled nursing services, and the client wants to receive both
PDN and PPECC services, the TMHP Prior Authorization Department will compare the Nursing
Addendum’s 24 hour daily care flow sheets and medical necessity documentation (e.g., authorization
requests). Upon subsequent approval of PDN or PPECC services, the provider who submitted the initial
prior authorization request that established the number of authorized skilled nursing hours will have
their authorized hours reduced to prevent duplication (e.g., if the client currently has PDN, and then
adds PPECC services, the PDN hours will be reduced).

When hours are reduced, the PDN or PPECC provider affected by the reduction will be notified by the
TMHP Prior Authorization Department when the reduction is effective, and the revised amount of
authorized hours. Providers that are affected by the reduction are only required to submit a revision
request and documentation of medical necessity if there is a change in the client’s medical condition or
the client’s medical needs are not commensurate with authorized hours and additional ongoing skilled
nursing hours are medically necessary. No action is required if additional hours are not medically
necessary.

4.1.4.8 Client Receives both PDN and PPECC and Shifts Services from One to the
Other

A client receiving both PDN and PPECC services may choose to shift approved hours from one ongoing
skilled nursing provider to another.

The receiving provider (PDN or PPECC provider who will gain hours in the shift) must submit all
required documentation for a revision.
The sending provider (PDN or PPECC provider who will lose hours in the shift) will receive a notice from TMHP Prior Authorization Department with revised (decreased) hours and the effective date of the reduction. The sending provider does not need to take any action unless there is a change in the client’s medical needs, and additional ongoing skilled nursing hours are medically necessary. If there is a medical need for additional ongoing skilled nursing hours, the sending provider may submit a revision request.

The total combined hours between PDN and PPECC services are not expected to increase without client medical necessity for additional hours (e.g., change in client condition or authorized hours are not commensurate with the client’s medical needs).

4.1.4.9 Recertifications

Recertifications may be prior authorized for up to a maximum of six months.

The following criteria must be met before a client receives a recertification:

- The client must have received PDN services for at least three months
- No significant changes in the client’s condition for at least three months
- No significant changes in the client’s condition are anticipated
- The client’s parent or guardian, physician, and provider agree the recertification is appropriate

A recertification request must be submitted at least 7 calendar days before, but no more than 30 days before, a current authorization period will expire. The PDN provider must submit the following documentation with the recertification request:

- A completed THSteps-CCP Private Duty Nursing six-month authorization signed and dated by the primary physician, nurse provider, and client, parent, or guardian
- A completed THSteps-CCP Prior Authorization Request form signed and dated by the primary physician within 30 calendar days prior to the SOC date
- A completed POC form, signed and dated by the primary physician within 30 calendar days prior to the SOC date
- A completed Nursing Addendum to Plan of Care for Private Duty Nursing and/or Prescribed Pediatric Extended Care Centers form signed and dated by the primary physician, RN completing the assessment, and parent, guardian, client, or responsible adult within 30 calendar days prior to the SOC date

**Note:** An advanced practice registered nurse (APRN) or a physician assistant (PA) may sign all documentation related to the provision of private duty nursing services on behalf of the client’s physician when the physician delegates this authority to the APRN or PA.

**Note:** All documentation must be maintained by the requesting PDN provider. The PDN provider may be asked to submit additional documentation including, but not limited to nurse’s notes, medication administration records, seizure logs, and ventilator logs to support medical necessity.

The provider is responsible for ensuring that the physician reviews and signs the POC within 30 calendar days of the expiration of the authorization period, and this documentation must be maintained in the client’s record. PDN providers should not submit a revised POC unless requesting a revision.

**Note:** An advanced practice registered nurse (APRN) or a physician assistant (PA) may sign all documentation related to the provision of private duty nursing services on behalf of the client’s physician when the physician delegates this authority to the APRN or PA.
The provider may request a revision of a recertification at any time during the recertification period if medically necessary. The provider must notify the claims administrator at any time during a recertification period if the client’s condition changes and the authorized services are not commensurate with the client’s medical needs.

All authorization timelines apply to recertifications.

Completed extension requests must be received and dated by TMHP Prior Authorization Department at least seven calendar days before, but no more than 30 days before, the current authorization expiration date. The request must be received by TMHP Prior Authorization Department no later than 5 p.m., Central Time, on the seventh day, to be considered received within seven calendar days. If a request is received less than seven calendar days before the current authorization expiration date, or after 5 p.m., Central Time, on the seventh day, authorization is given for dates of service beginning no sooner than seven calendar days after the receipt of the completed request by TMHP Prior Authorization Department.

The nursing care provider must notify TMHP Prior Authorization Department at any time during the authorization period if the client’s condition and need for SN care significantly changes.

4.1.4.10 Special Circumstances
PDN services provided in a school or day care facility, at the request of the family, may be authorized provided the client requires the requested amount of PDN services if in the home.

PDN services may be provided in a hospital, SN facility or intermediate care facility for the individuals with intellectual disabilities, or special care facility with documentation from the facility showing it is unable to meet the SN needs of the client, and the services are medically necessary. These facilities are required by licensure to meet all the medical needs of the client.

4.1.4.11 PDN Services Provided in Group Settings
PDN services may be authorized in a provider or client ratio other than 1:1.

An RN or LVN may be authorized to provide PDN services to more than one client over the span of the day as long as:

- Each client’s care is based on an individualized POC
- Each client’s needs and POC do not overlap with another client’s needs and POC

Settings in which a PDN provider may provide services in a provider-client ratio greater than 1:1 include, but are not limited to, homes with more than one client receiving PDN, foster homes, or independent living arrangements.

4.1.4.12 Termination of Authorization
Authorization for PDN services will be terminated when:

- The client is no longer eligible for Medicaid
- The client no longer meets the medical necessity criteria for PDN services
- The place of service does not support the health and safety of the client
- The client, parent, or guardian refuses to comply with the service plan and compliance is necessary to assure the health and safety of the client

4.1.4.13 Appeal of Authorization Decisions
Providers may appeal denials or modifications of requested PDN services with documentation to support the medical necessity of the requested PDN services. Appeals must be submitted to the TMHP Prior Authorization Department with complete documentation and any additional information within two weeks of the date on the decision letter. If changes are made to the authorization based on this
documentation, TMHP Prior Authorization Department will go back no more than three business days for initial, or revision requests and no more than seven calendar days for recertification requests when additional documentation is submitted.

The client, parent, or guardian will be notified of any denial or modification of requested services and will be given information about how to appeal the TMHP Prior Authorization Department decision.

All documentation must be submitted together, and requests are not reviewed until all documentation is received. If complete documentation is received at TMHP Prior Authorization Department by 3 p.m., Central Time, a response is returned to the provider within one business day. Complete documentation for initial, revision, recertification, and extension requests for PDN authorizations include all of the following:

- Home Health Plan of Care (POC) on the TMHP website at www.tmhp.com.
- Nursing Addendum to Plan of Care for Private Duty Nursing and/or Prescribed Pediatric Extended Care Centers on the TMHP website at www.tmhp.com.

4.1.4.14 Start of Care (SOC)

The SOC is the date that care is to begin, as agreed on by the family, the client’s physician, and the provider, and as listed on the POC and the CCP Prior Authorization Request Form. Providers are responsible for determining whether they can accept the client for services.

Once the provider accepts a client for service and accepts responsibility for providing PDN, the provider is required to deliver those services beginning with the SOC date. Providers are responsible for a safe transition of services when the authorization decision is a denial or a reduction of services. Providers are required to notify the physician and the client’s family on receipt of an authorization, a denial, or a change in PDN.

Providers must submit complete documentation no later than three business days from an SOC date to obtain initial coverage for the SOC date.

Note: Texas Medicaid (Title XIX) home health services does not authorize an SOC date earlier than three business days before contact with TMHP.

For PDN recertification, TMHP Prior Authorization Department must receive complete documentation no later than three business days before the SOC date. It is recommended that recertification requests be submitted up to 30 days before the current authorization ends.

During the prior authorization process for initial and recertification requests, providers are required to deliver the requested services from the SOC date.

4.1.4.15 Client and Provider Notification

When PDN is approved as requested, the provider receives written notification. The provider is responsible for notifying the client or family and the physician of the authorized services.

The TMHP Prior Authorization Department or MCO notifies the client and provider in writing when the following instances occur:

- PDN is denied.
- PDN hours authorized are less than the hours requested on the POC.
- PDN hours are modified (e.g., hours are requested by the week but are authorized by the day).
- The TMHP Prior Authorization Department or MCO receives incomplete information from the provider.
- Dates of service authorized are different from those requested.
The provider is responsible for notification and coordination with the physician and family.

4.1.5 Limitations with Other Services

4.1.5.1 PDN Services with PCS, SHARS, or PPECC

When clients are receiving both PDN services and PCS from an individual person over the same span of time, all services will be reimbursed according to the maximum allowable fee schedule.

Texas Medicaid will not reimburse providers for PDN services that duplicate services provided by school districts under the SHARS program. If a provider bills Texas Medicaid for PDN services provided to a client at school, then the school district may be asked to provide documentation that it is not also providing PDN services to the same client at the same time and seeking reimbursement for the service under the SHARS program.

PDN services may be billed on the same day as PPECC services, but may not be billed simultaneously with PPECC services. PDN may be billed when it occurs before or after PPECC services.

4.1.5.2 PDN Provided During a Skilled Nursing Visit for TPN Administration Education

For clients who are receiving PDN services who also require TPN administration education, intermittent SN visits may be separately reimbursed when the SN services are for the client or caregiver training in TPN administration.

Refer to: Subsection 3.3, “Home Health Skilled Nursing and Home Health Aide Services Providers” in this handbook for detailed information about SN benefits.

PDN and SN should not be routinely performed on the same date during the same time period.

PDN and SN will not be considered for reimbursement when the services are performed on the same date during the same time period without prior authorization approval.

If the SN visit for TPN education occurs during a time period when the PDN provider is caring for the client, both the PDN provider and the nurse educator must document in the client’s medical record the skilled services individually provided, including but not limited to:

- The start and stop time of each nursing provider’s specialized task(s)
- The client condition that requires the performance of skilled PDN tasks during the SN visit for TPN education
- The skilled services that each provided during that time period

Both the intermittent SN visit and the PDN services provided during the same time period may be recouped if the documentation does not support the medical necessity of each service provided.

Intermittent SN visits for clients who receive PDN and who require TPN administration education may be considered for separate prior authorization if:

- The PDN provider is not an RN who has been appropriately trained in the administration of TPN, and the PDN provider is not able to perform the function.
- There is documentation that supports the medical need for an additional skilled nurse to perform TPN.

The SN services may be prior authorized only for the client and caregiver who will be trained in TPN administration.

Clients whose only SN need is the provision of education for self-administration of prescribed subcutaneous (SQ), intramuscular (IM), or intravenous (IV) injections will not qualify for PDN services.

Nursing hours for the sole purpose of providing education to the client and caregiver may be considered through intermittent home health SN visits.
5 Documentation Requirements

All documentation, including that which supports medical necessity, and the comprehensive treatment plan related to the therapy services that were prior authorized and provided, must be maintained in the client’s medical record and made available upon request.

Documentation elements that are routinely assessed for compliance in retrospective review of client records include, but are not limited to, the required documentation noted previously, as well as the following:

- All entries are legible to people other than the author, dated (month, day, year, time), and signed by the author.
- Each page of the record documents the client’s name and Medicaid identification number.
- Client assessment time is documented at the beginning of each shift.
- All nurses’ arrival and departure times are documented with signature and time in the narrative section of the nurses’ notes.
- Entries in the nursing flowsheet or narrative notes must be dated and timed every 1 to 2 hours and must include the following:
  - The client’s condition.
  - The name of the medication, dose, route, time given, client response, and other pertinent information is recorded when medication is administered.
  - The name of treatment, time given, route or method used, client response, and other pertinent information is provided when treatments are administered.
  - The amount, type, times given, route or method used, client response, and other pertinent information is provided when feedings are administered.
  - The POC and documentation of services correlate with and reflect medical necessity for the services provided on any given day.
  - A request for prior authorization must include documentation from the provider to support the medical necessity of the service, equipment, or supply.
  - Client’s arrival or departure from the home setting is documented with the time of arrival, departure, mode of transportation, and who accompanied the client.
  - Documentation of teaching the client or the client’s responsible adult includes the length of time, the subject of the teaching, the understanding of the subject matter by the person receiving the teaching, and other pertinent information.
  - Supervisory visits include specifics of the visit.
  - If a client is receiving SN services through another program or service in addition to PDN, such as MDCP, each provider’s shift notes designate specifically which type of service they are providing during that shift.

6 Claims Filing and Reimbursement

6.1 Claims Filing

Providers may purchase CMS-1500 or UB-04 CMS-1450 paper claim forms from the vendor of their choice. TMHP does not supply them.
When completing a CMS-1500 or a UB-04 CMS-1450 paper claim form, all required information must be included on the claim, as TMHP does not key information from attachments.

Refer to: “Section 3: TMHP Electronic Data Interchange (EDI)” (Vol. 1, General Information) for information on electronic claims submissions.

“Section 6: Claims Filing” (Vol. 1, General Information) for general information about claims filing.

Subsection 6.5, “CMS-1500 Paper Claim Filing Instructions” in “Section 6: Claims Filing” (Vol. 1, General Information) for instructions on completing paper claims.

Subsection 6.6, “UB-04 CMS-1450 Paper Claim Filing Instructions” in “Section 6: Claims Filing” (Vol. 1, General Information) for paper claims completion instructions.

### 6.1.1 Home Health Skilled Nursing and Home Health Aide Providers

Providers must use only type of bill (TOB) 321 in Form Locator (FL) 4 of the UB-04 CMS-1450. Other TOBs are invalid and will result in a claim denial. Home health services must be submitted to TMHP in an approved electronic format or on a CMS-1500 or a UB-04 CMS-1450 paper claim form. Submit home health DME and medical supplies to TMHP in an approved electronic format, or on a CMS-1500 or on a UB-04 CMS-1450 paper claim form.

Outpatient claims must have the appropriate revenue code and, if appropriate, the corresponding Healthcare Common Procedure Coding System (HCPCS) code or narrative description. The prior authorization number must appear on the CMS-1500 paper claim form in Block 23 and in Block 63 of the UB-04 CMS-1450 paper claim form. The certification dates or the revised request date on the POC must coincide with the DOS on the claim. Prior authorization does not waive the 95-day filing deadline requirement.

Home health service claims should not be submitted for payment until Medicaid certification is received and a prior authorization number is assigned.

### 6.1.2 PDN Providers

Independently enrolled RNs or LVNs providing PDN services must submit claims on the HCFA-1500 claim form. PDN Providers who are home health agencies must submit claims on the HCFA-1450 (UB-92) claim form.

PDN providers must submit claims for services in an approved electronic claims format or on the appropriate claim form based on their provider type. Home health agencies must submit claims on the UB-04 CMS-1450 paper claim form. Independently enrolled nurses must submit claims on the CMS-1500 paper claim form.

### 6.1.3 Method for Counting Minutes for Timed Procedure Codes

All claims for reimbursement of procedure codes paid in 15-minute increments are based on the actual amount of billable time associated with the service. For those services for which the unit of service is 15 minutes (1 unit = 15 minutes), partial units should be rounded up or down to the nearest quarter hour.

Time intervals for 1 through 12 units are as follows:

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<tr>
<th>Units</th>
<th>Number of Minutes</th>
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<tbody>
<tr>
<td>0 units</td>
<td>0 minutes through 7 minutes</td>
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<tr>
<td>1 unit</td>
<td>8 minutes through 22 minutes</td>
</tr>
<tr>
<td>2 units</td>
<td>23 minutes through 37 minutes</td>
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<tr>
<td>3 units</td>
<td>38 minutes through 52 minutes</td>
</tr>
<tr>
<td>4 units</td>
<td>53 minutes through 67 minutes</td>
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</table>
6.2 Reimbursement

The reimbursement methodology for professional services delivered by home health agencies is a statewide visit rate calculated in accordance with 1 TAC §355.8021. Home health agencies are reimbursed for DME and medical supplies in accordance with 1 TAC §355.8023. PDN services are reimbursed in accordance with 1 TAC §355.8441.

Providers can refer to the OFL or the applicable fee schedule on the TMHP website at www.tmhp.com. Texas Medicaid implemented mandated rate reductions for certain services. The OFL and static fee schedules include a column titled “Adjusted Fee” to display the individual fees with all mandated percentage reductions applied. Additional information about rate changes is available on the TMHP website at www.tmhp.com/resources/rate-and-code-updates/rate-changes.

Refer to: Subsection 2.2, “Fee-for-Service Reimbursement Methodology” in “Section 2: Texas Medicaid Fee-for-Service Reimbursement” (Vol. 1, General Information).

<table>
<thead>
<tr>
<th>Units</th>
<th>Number of Minutes</th>
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<tr>
<td>5 units</td>
<td>68 minutes through 82 minutes</td>
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<tr>
<td>6 units</td>
<td>83 minutes through 97 minutes</td>
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<tr>
<td>7 units</td>
<td>98 minutes through 112 minutes</td>
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<tr>
<td>8 units</td>
<td>113 minutes through 127 minutes</td>
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<td>143 minutes through 157 minutes</td>
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<td>11 units</td>
<td>158 minutes through 172 minutes</td>
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<tr>
<td>12 units</td>
<td>173 minutes through 187 minutes</td>
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# INPATIENT AND OUTPATIENT HOSPITAL SERVICES HANDBOOK

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1 General Information

The information in this handbook is intended for Texas Medicaid hospital (medical and surgical acute care facility) providers and covers services that take place only in an inpatient or outpatient hospital setting. The handbook provides information about Texas Medicaid’s benefits, policies, and procedures applicable to acute care hospitals, including military hospitals.

Important: All providers are required to read and comply with "Section 1: Provider Enrollment and Responsibilities" (Vol. 1, General Information). In addition to compliance with all requirements specific to Texas Medicaid, it is a violation of Texas Medicaid rules when a provider fails to provide health-care services or items to Medicaid clients in accordance with accepted medical community standards and standards that govern occupations, as explained in Title 1 Texas Administrative Code (TAC) §371.1659. Accordingly, in addition to being subject to sanctions for failure to comply with the requirements that are specific to Texas Medicaid, providers can also be subject to Texas Medicaid sanctions for failure, at all times, to deliver health-care items and services to Medicaid clients in full accordance with all applicable licensure and certification requirements including, without limitation, those related to documentation and record maintenance.

Refer to: The Clinics and Other Outpatient Facility Services Handbook (Vol. 2, Provider Handbooks) for information about services offered in settings such as rural health clinics (RHCs), Federally Qualified Health Centers (FQHCs), dialysis centers, and other similar facilities.

1.1 National Drug Codes (NDC)

Refer to: Subsection 6.3.4, “National Drug Code (NDC)” in “Section 6: Claims Filing” (Vol. 1, General Information).

1.2 Medicaid Managed Care Services

This handbook contains information about Texas Medicaid fee-for-service benefits. For information about managed care benefits, refer to the Medicaid Managed Care Handbook (Vol. 2, Provider Handbooks).

Managed care carve-out services are administered as fee-for-service benefits. A list of all carve-out services is available in section 8, “Carve-Out Services” in the Medicaid Managed Care Handbook (Vol. 2, Provider Handbooks).

2 Enrollment

To be eligible to participate in Texas Medicaid, a hospital must be certified by Medicare, have a valid provider agreement with the Health and Human Services Commission (HHSC), and have completed the Texas Medicaid & Healthcare Partnership (TMHP) enrollment process.

2.1 Hospital Eligibility Through Change of Ownership

Under procedures set forth by the Centers for Medicare & Medicaid Services (CMS) and the U.S. Department of Health and Human Services (HHS), a change in ownership of a hospital does not terminate Medicare eligibility; therefore, Medicaid participation may be continued subject to the following requirements:

• The provider must obtain recertification as a Title XVIII (Medicare) hospital.
• The hospital under new ownership must submit a new signed and dated HHSC Medicaid Provider Agreement between the hospital and HHSC.
Providers can download the HHSC Medicaid Provider Agreement from the TMHP website at www.tmhp.com.

Refer to: Subsection 1.4, “Provider Reenrollment” in “Section 1: Provider Enrollment and Responsibilities” (Vol. 1, General Information).

2.1.1 Hospital-based Ambulatory Surgical Center (HASC) Enrollment

All hospitals enrolling in Texas Medicaid (except psychiatric and rehabilitation hospitals) are issued an HASC provider number at the time of enrollment.

2.2 Hospital-based Rural Health Clinic Enrollment

To enroll in Texas Medicaid and qualify for participation as a Title XIX RHC, RHCs must be enrolled in Medicare. A nine-digit provider identifier is issued to the RHC after a certification letter from Medicare is received, stating that the clinic qualifies for Medicaid participation. An RHC can also apply for enrollment as a family planning agency.

All providers of laboratory services must comply with the rules and regulations of the Clinical Laboratory Improvement Amendments (CLIA). Providers who do not comply with CLIA are not reimbursed for laboratory services.


Subsection 1.1, "Provider Enrollment” in “Section 1: Provider Enrollment and Responsibilities” (Vol. 1, General Information) for more information about enrollment procedures, including information on Changes of Ownership.


3 Inpatient Hospital (Medical/Surgical Acute Care Inpatient Facility)

This section contains benefit, limitation, authorization, and claims filing information for inpatient hospital facility accommodation and ancillary services.

Refer to: “Section 6: Claims Filing” (Vol. 1, General Information) for more comprehensive information about claims filing and appeals.

“Section 7: Appeals” (Vol. 1, General Information) for more comprehensive information about claims filing and appeals.

Hospital providers are encouraged to review the other handbooks for applicable information, prior authorization requirements, and for specific requirements for special programs.

3.1 General Information

Inpatient hospital services include medically necessary items and services ordinarily furnished by a Texas Medicaid hospital or by an approved out-of-state hospital under the direction of a physician for the care and treatment of patients. Services must be medically necessary and are subject to Texas Medicaid’s utilization review requirements. Claims submitted to TMHP must comply with the applicable Texas Medicaid policies and procedures.
3.1.1 Reimbursement Limitations

For clients who are 21 years of age and older, Texas Medicaid reimbursement for acute care inpatient hospital services is limited to $200,000 per client, per benefit year (November 1 through October 31). Claims are reviewed retrospectively, and payments that exceed $200,000 are recouped.

This $200,000 limitation does not apply to the following:

- Services related to certain organ transplants.
- Services rendered to Texas Health Steps (THSteps) clients when provided through the Comprehensive Care Program (CCP).

For clients who are 20 years of age and younger, dollar limitations do not apply.

3.1.2 Spell of Illness

Reimbursement to hospitals for inpatient services is limited to the Medicaid spell of illness. The spell of illness is defined as 30 days of inpatient hospital care, which may accrue intermittently or consecutively.

After 30 days of inpatient care is provided, reimbursement for additional inpatient care is not considered until the client has been out of an acute care facility for 60 consecutive days.

Exceptions to the spell of illness are as follows:

- A prior-approved solid organ transplant. The 30-day spell of illness for transplants begins on the date of the transplant, allowing additional time for the inpatient stay.
- THSteps-eligible clients who are 20 years of age and younger when a medically necessary condition exists.

Texas Medicaid will conduct a quarterly utilization review of inpatient claims to determine whether the claims were paid outside of the spell-of-illness limitation.

The first of these utilization reviews were for claims with dates of service from April 27, 2010, through January 6, 2012.

3.1.3 Take-Home Drugs, Self-Administered Drug, or Personal Comfort Items

Take-home drugs and comfort items that are provided by the hospital during an inpatient hospital stay are included in the hospital reimbursement and are not reimbursed separately.

Take-home drugs and supplies may be a benefit through the Vendor Drug Program (VDP) when supplied by prescription.

Self-administered drugs are defined as drugs that the client administers themselves at home and may include, but are not limited to, prescription drugs, vitamins, and supplements. Self-administered drugs provided by the hospital during an inpatient hospital stay are included in the hospital reimbursement and are not reimbursed separately.

The client cannot be billed for take-home drugs, comfort supplies or self-administered drugs that are provided by the hospital during an inpatient hospital stay.

3.1.4 Services Included in the Inpatient Stay

The following services are included in the inpatient stay and are not separately reimbursed:

- Whole blood and packed red blood cells. Inpatient services include whole blood and packed red blood cells that are reasonable and necessary for treatment of illness or injury. Whole blood and packed red blood cells that are available without cost are not reimbursed by Texas Medicaid.
• *Laboratory, radiology, and pathology services.* Inpatient services include all medically necessary services and supplies ordered by a physician to include laboratory, radiology, and pathology services.

**Note:** *Ultrasound interpretations in the inpatient hospital setting will be denied if they are billed by the attending physician. Services that are billed by the attending physician are included in the facility fee and are not reimbursed separately.*

**Note:** *All providers of laboratory services must comply with the rules and regulations of the Clinical Laboratory Improvement Amendments (CLIA). Providers not complying with CLIA will not be reimbursed for laboratory services.*

**Refer to:** Subsection 1.1, “Provider Enrollment” in “Section 1: Provider Enrollment and Responsibilities” (Vol. 1, General Information) for more information about enrollment procedures. Subsection 2.1.1, “Clinical Laboratory Improvement Amendments (CLIA)” in the Radiology and Laboratory Services Handbook (Vol. 2, Provider Handbooks) for more information about CLIA. The Medicaid Managed Care Handbook (Vol. 2, Provider Handbooks).

### 3.1.5 Outpatient Observation Services

Outpatient observation services are not a substitute for a medically appropriate inpatient admission. If a client meets the medical necessity criteria for an inpatient admission and an inpatient admission is ordered by the practitioner, an inpatient admission is a benefit regardless of the length of stay.

**Refer to:** Subsection 4.2.4, “Outpatient Observation Room Services” in this handbook for more information.

### 3.2 Services, Benefits, Limitations, and Prior Authorization - Acute Care

Inpatient hospital services include the following:

• Bed and board
• Whole blood and packed red blood cells
• All medically necessary services ordered by a physician to include laboratory, radiology, and pathology
• All medically necessary supplies ordered by a physician
• Medically necessary emergency and non-emergency ambulance transports during the inpatient stay
• Maternity care
• Newborn care
• Inpatient surgery and rehabilitation
• Organ and tissue transplant services
• Colorectal cancer screening services

#### 3.2.1 Bed and Board

Inpatient bed and board include semiprivate accommodations or accommodations in an intensive care or coronary care unit. The accommodations include:

• Meals
• Special diets
• General nursing services

Private accommodations including meals, special diets, and general nursing services may be reimbursed up to the hospital’s charge for its most prevalent semiprivate accommodations. Bed and board in private accommodations may be reimbursed in full if required for medical reasons as certified by the physician. The hospital must document the medical necessity for a private room (i.e., the existence of a critical or contagious illness, a condition that could result in disturbance to other patients). The medical necessity for the private accommodations must be included in Block 80 of the UB-04 CMS-1450 paper claim form or added as an attachment to the claim submission.

3.2.2 Hysterectomy Services

Hysterectomy services are considered for reimbursement when the claim is filed with a signed Hysterectomy Acknowledgment Form or submitted documentation indicates that the Hysterectomy Acknowledgment Form could not be obtained.

Claims for services related to the hysterectomy cannot be reimbursed unless the signed Hysterectomy Acknowledgement Form is on file; consequently, to avoid claim denials, each individual provider is encouraged to submit a copy of the valid Hysterectomy Acknowledgement Form and not rely on another provider to do so.


Texas Medicaid - Title XIX Acknowledgment of Hysterectomy Information on the TMHP website at www.tmhp.com.

3.2.3 Maternity Care

Inpatient maternity care includes usual and customary care for all female clients.

3.2.3.1 Emergency Coverage

For women with a family income at or below 198 percent of the Federal Poverty Level (FPL), hospital facility charges are paid through Emergency Medicaid. A client must be determined eligible for Emergency Medicaid by HHSC for a claim to be paid to a Medicaid provider. Claims are sent to Texas Medicaid & Healthcare Partnership (TMHP) for processing.

3.2.3.2 Mother and Newborn Hospital Stay

Circumstances that require the mother and newborn to remain in the hospital longer than two days for a routine vaginal delivery or four days for a cesarean section must be documented in the clients’ medical records.

Continuation of hospitalization is a benefit for the infant when the mother is required to remain hospitalized for medical reasons. The reason for the continuation of hospitalization must be documented in the client’s medical record.

3.2.3.3 Children’s Health Insurance Program (CHIP) Perinatal Coverage

For clients who are eligible for CHIP perinatal services as determined by HHSC, CHIP perinatal services include newborn services and inpatient hospital charges related to the delivery of the newborn. Preterm or false labor that does not result in a birth are not CHIP perinatal services.

Inpatient services limited to labor with delivery for women with income at or below 202 percent of FPL will be covered under CHIP perinatal. Newborn services will also be covered under CHIP perinatal.

For CHIP perinatal newborns with a family income at or below 198 percent of the federal poverty level, TMHP will process newborn transfer hospital claims even if the claim from the initial hospital stay has not been received. The hospital transfer must have occurred within 24 hours of the discharge date from the initial delivery hospital stay.
Transfer claims must be filed to TMHP using the admission type 1, 2, 3, or 5 in block 14; source of admission code 4 or 6 in block 15; and the actual date and time the client was admitted in block 12 of the UB-04 CMS-1450 paper claim form.

**Refer to:** Subsection 6.19.1, “CHIP Perinatal Newborn Transfer Hospital Claims” in “Section 6: Claims Filing” (Vol. 1, General Information).

### 3.2.4 Newborn Care

Newborn care includes routine newborn care, routine screenings, and specialized nursery care for newborns with specific problems.

Hospital providers must provide all state-mandated newborn screenings and vaccinations.

**Refer to:** Subsection 5.3.11.2.3, “Hearing Screening” in the *Children’s Services Handbook* (Vol. 2, Provider Handbooks).

Subsection 5.3.9, “Newborn Examination” in the *Children’s Services Handbook* (Vol. 2, Provider Handbooks).

### 3.2.4.1 Newborn Eligibility

A child is deemed eligible for Texas Medicaid through 12 months of age if the mother is receiving Medicaid at the time of the child’s birth, and the mother continues to be eligible for Medicaid or would be eligible for Medicaid if she were pregnant. Therefore, it is not acceptable for a hospital to require a deposit for newborn care from a Medicaid client. The child’s eligibility ends if the mother relinquishes her parental rights or if it is determined that the child is no longer part of the mother’s household.

Hospitals must complete Hospital Report (Newborn Child or Children) (Form 7484) on the TMHP website at [www.tmhp.com](http://www.tmhp.com) to provide information about each child born to a mother who is eligible for Medicaid. If the newborn’s name is known, the name must be on the form.

**Important:** If the newborn’s name is not known, the name may be left blank. The use of “Baby Boy” or “Baby Girl” delays the assignment of a number.

The form must be completed by the hospital no later than five days after the child’s birth and sent to HHSC at the address identified on the form. The form should not be completed for stillbirths. Hospitals should duplicate the form as needed, because they are not supplied by HHSC or TMHP.

Hospitals that submit the birth certificate information using the Department of State Health Services (DSHS), Vital Statistics Unit (VSU) Texas Electronic Registrar for Birth software and the HHSC Form 7484, receive a rapid and efficient assignment of a newborn Medicaid identification number. This process expedites reimbursement to hospitals and other providers involved in newborn care including pharmacies that provide outpatient prescription benefits for medically-needy newborns.

For additional information about obtaining a newborn Medicaid identification number, providers may call 1-888-963-7111, Ext. 7368 or 1-512-458-7368.

After receiving a completed form, HHSC verifies the mother’s eligibility. Within 10 days of receiving the completed form, HHSC sends notices to the hospital, mother, caseworker, and attending physician, if identified. The notice includes the child’s Medicaid client number and the effective date of coverage. After the child has been added to the eligibility file, HHSC issues a Medicaid Identification (Form H3087).

Claims submitted for services provided to a newborn child who is eligible for Medicaid must be filed using the newborn child’s Medicaid client number.
Newborns who are from families with an income at or below 198 percent of the FPL and who receive CHIP perinatal benefits are assigned a client number for Texas Medicaid. This number is only assigned for reimbursement of the newborn’s hospital facility charges (on a UB-04 CMS-1450 paper claim form) for the initial hospital stay after delivery. Claims for the newborn’s hospital facility charges should be sent to TMHP.

3.2.5 Organ and Tissue Transplant Services

3.2.5.1 Transplant Facilities

A facility that renders organ transplants must be a designated children’s hospital or a facility in continuous compliance with the criteria set forth by the following:

- Organ Procurement and Transportation Network (OPTN)
- United Network for Organ Sharing (UNOS)
- National Marrow Donor Program (NMDP)

Facilities whose status of “good standing” has been suspended for any reason by the national credentialing bodies will not be reimbursed by Texas Medicaid for transplant services until the status of “good standing” is restored.

If a Medicaid client receives a transplant in an in-state or out-of-state facility that is not approved by Texas Medicaid, the client must be discharged from the facility to be considered to receive other medical and hospital benefits under Texas Medicaid. Coverage for other services needed as a result of complications of the transplant may be considered when medically necessary, reasonable, and federally allowable. Texas Medicaid will not pay for routine post-transplant services for transplant patients in facilities that are not approved by Texas Medicaid.

3.2.5.1.1 Out-of-state Transplant Facilities

Out-of-state facilities may be reimbursed for transplants rendered to Texas Medicaid clients under certain conditions. In order for Texas Medicaid to reimburse for an out-of-state transplant, the out-of-state facility and professional providers must enroll as Texas Medicaid providers. The out-of-state transplant facilities must submit proof of transplant facility UNOS or NMDP certification as required by the Texas HHSC.

Prior authorization requests may be submitted to the TMHP Prior Authorization Department by mail, fax, or the electronic portal. Prescribing or ordering providers, dispensing providers, clients’ responsible adults, and clients may sign prior authorization forms and supporting documentation using electronic or wet signatures.

Physicians who are licensed by the state of Texas may request prior authorization for transplant services to be performed at out-of-state facilities when all of the following criteria are met:

- The required organ transplant is not available in Texas
- The facility is nationally recognized as a Center of Excellence
- The services are medically necessary, reasonable, and federally allowable
- The client is enrolled in Texas Medicaid
The pretransplant evaluation must be performed by a Texas facility. If it is medically necessary that the
pretransplant evaluation be performed at the out-of-state facility as well, the prior authorization request
for the out-of-state pretransplant evaluation must be submitted with a copy of the evaluation that was
performed by the Texas facility. The documentation must support the need for an out-of-state pretrans-
plant evaluation.

**Important:** Texas Medicaid does not cover transplant services provided out-of-state that are available in
Texas.

### 3.2.5.2 Transplant Benefits and Limitations

If a transplant has been authorized as medically necessary by HHSC or its designee because of an
emergent, life-threatening situation, a maximum of 30 days of inpatient hospital services during Title
XIX spell of illness may be a benefit, beginning with the actual first day of the transplant. This benefit is
in addition to covered inpatient hospital days provided before the actual first day of the transplant. This
30-day period is considered a separate inpatient hospital admission for reimbursement purposes, but is
included under one hospital stay.

*Refer to:* Subsection 3.1.2, “Spell of Illness” in this handbook for additional information about the
30-day spell of illness period.

Reimbursement for transplant is limited to an initial transplant as a lifetime benefit and one subsequent
re-transplant because of rejection. Expenses incurred by a living donor will not be reimbursed.

All transplants require prior authorization. If a solid organ transplant is not prior authorized, services
that are directly related to the transplant within the three-day pre-operative and six-week postoperative
period will be denied, regardless of who provides the services. Services unrelated to the transplant
surgery will be paid separately.

If the organ is rejected, the re-transplant requires its own prior authorization. If the re-transplant is not
prior authorized, services that are directly related to the re-transplant within the three-day pre-operative
and six-week postoperative period will be denied, regardless of who provides the services. Services
unrelated to the re-transplant surgery will be paid separately.

*Note:* The re-transplant is not included in the prior authorization for the initial transplant. The
subsequent re-transplant must be prior authorized separately.

*Refer to:* Subsection 9.2.47, “Organ/Tissue Transplants” in the *Medical and Nursing Specialists,

### 3.2.5.3 Prior Authorization for Organ and Transplant Services

All solid organ transplant services provided by facilities and professionals must be prior authorized. If a
solid organ transplant is not prior authorized, services directly related to the transplant within the three
day pre-operative and six-week postoperative period also will be denied, regardless of who provides the
service, (e.g., laboratory services, status-post visits, and radiology services). Services unrelated to the
transplant surgery will be paid separately.

A transplant request signed by a physician associated with transplant facilities is considered for prior
authorization after the client has been evaluated and meets the guidelines of the institution’s transplant
protocol.

*Refer to:* Subsection 9.2.47, “Organ/Tissue Transplants” in the *Medical and Nursing Specialists,

Prior authorization requests may be submitted to the TMHP Prior Authorization Department by mail,
fax, or the electronic portal. Prescribing or ordering providers, dispensing providers, clients’ responsible
adults, and clients may sign prior authorization forms and supporting documentation using electronic
or wet signatures.
3.2.5.4 Transplants for Medicare-Eligible Clients

Transplants are also a benefit under the Medicare program; therefore, for clients eligible for Medicare and Medicaid, Texas Medicaid will pay only the deductible or coinsurance portion as applicable according to current payment guidelines. Prior authorization must be obtained for Medicaid-only clients; authorization will not be given for Medicare/Medicaid-eligible clients. Texas Medicaid will not pay for a transplant service denied by Medicare for a Medicare-eligible client.

3.2.5.5 Experimental or Investigational Services

Benefits are not available for any experimental or investigational services (including xenotransplantation and artificial/bioartificial liver transplants), supplies, or procedures.

3.2.5.6 Reimbursement for Transplant Services

The hospital diagnosis related group (DRG) payment for the transplant includes procurement of the organ and services associated with the organ procurement. Section 1138 of the Social Security Act defines the conditions of participation for institutions in the organ procurement program. Organ procurement costs are not reimbursed to a hospital that fails to meet the conditions of participation. The specific guidelines may be found in the appropriate areas of the Code of Federal Regulations (CFR) Title 42, Parts 405, 413, 441, 482, and 485. Documentation of organ procurement must be maintained in the hospital’s medical record.

Refer to: Subsection 2.2, “Fee-for-Service Reimbursement Methodology” in “Section 2: Texas Medicaid Fee-for-Service Reimbursement” (Vol. 1, General Information) for more information about reimbursement.

3.2.5.7 Nonsolid Organ Transplants

Under current Texas Medicaid policy, procedures are considered to be medically necessary and reasonable based on safety and efficacy, as demonstrated by scientific evidence and controlled by clinical studies.

Nonsolid organ transplants that are benefits of Texas Medicaid include:

- Allogeneic and autologous stem cell transplantation
- Allogeneic and autologous bone marrow transplantation
- Autologous islet cell transplantation

All nonsolid organ transplants require prior authorization and must be performed in a Texas facility that is a designated Children’s Hospital or a facility in compliance with the criteria set forth by the Organ Procurement and Transportation Network (OPTN), the United Network for Organ Sharing (UNOS), or the National Marrow Donor Program (NMDP).

Experimental or investigational services, supplies, or procedures are not a benefit of Texas Medicaid.

3.2.5.7.1 Inpatient Hospitalization

For a nonsolid organ transplant that has been prior authorized for clients who are 21 years of age and older, a maximum of 30 days of inpatient hospital services during a Title XIX spell of illness is covered beginning with the actual first day of the transplant. This coverage is in addition to covered inpatient hospital days provided before the actual first day of the transplant. This 30-day period is considered a separate inpatient hospital admission for reimbursement purposes but is included under one DRG payment.

Autologous harvesting of stem cells (single or multiple sessions) are reimbursed to the facility when prior authorized by HHSC or its designee and performed in the outpatient facility setting. Harvesting of stem cells performed in a hospital inpatient setting is included in the DRG and is not reimbursed separately.
3.3 Services, Benefits, Limitations, and Prior Authorization - Inpatient Rehabilitation Services

Inpatient rehabilitation services are a benefit of Texas Medicaid when provided as part of a general acute care inpatient admission, or with prior authorization for clients who are 20 years of age and younger in a freestanding rehabilitation facility.

Inpatient rehabilitation services in an acute care setting are included in the hospital DRG payment. All rehabilitation services are subject to Medicaid benefit limitations including the spell of illness. Exceptions to those limitations may be offered under CCP.

Refer to: Subsection 9.2.47.9, “Nonsolid Organ Transplants” in the Medical and Nursing Specialists, Physicians, and Physician Assistants Handbook (Vol. 2, Provider Handbooks).

Refer to: Subsection 1.7.17, “Physical, Occupational, and Speech Therapy Providers” in “Section 1: Provider Enrollment and Responsibilities” (Vol. 1, General Information).

Refer to: Subsection 2.17, “Inpatient Rehabilitation Facility (Freestanding) (CCP)” in the Children’s Services Handbook (Vol. 2, Provider Handbooks).

3.4 Services, Benefits, Limitations, and Prior Authorization - Inpatient Psychiatric Services

3.4.1 Enrollment

Acute care hospitals and state psychiatric facilities must be certified by Medicare, have a valid provider agreement with the HHSC, and have completed the TMHP enrollment process.

Refer to: Subsection 5.1, “Enrollment” in this handbook for more information about acute care hospital enrollment.

Freestanding psychiatric facilities must be licensed by DSHS or by the appropriate state board where services are rendered. The provider must be approved by The Joint Commission (TJC).

Providers cannot be enrolled if their licenses are due to expire within 30 days.

To be eligible to participate in the Comprehensive Care Inpatient Psychiatric (CCIP) Program to render services to Texas Health Steps (THSteps) clients, a freestanding or state psychiatric facility must be accredited by TJC, have a valid provider agreement with HHSC, and have completed the TMHP enrollment process. Facilities certified by Medicare must also meet TJC accreditation requirements.

Note: Acute care hospitals cannot enroll as CCIP facilities.

3.4.2 General Information

Inpatient admissions to acute care hospitals, freestanding psychiatric facilities, and state psychiatric facilities for psychiatric conditions may be a benefit of Texas Medicaid as outlined in the following table:

<table>
<thead>
<tr>
<th>Hospital Type</th>
<th>CCIP Clients 0-20 Years of Age</th>
<th>Medicaid Clients of Any Age</th>
</tr>
</thead>
<tbody>
<tr>
<td>Acute care hospital</td>
<td>No</td>
<td>Yes</td>
</tr>
<tr>
<td>Freestanding psychiatric facility (IMD)</td>
<td>Yes</td>
<td>Yes (clients 65 years of age and older)</td>
</tr>
<tr>
<td></td>
<td></td>
<td>No (clients 21 through 64 years of age)</td>
</tr>
</tbody>
</table>

(IMD) Institution for Mental Diseases.
When a client requires admission, or once the client becomes Medicaid eligible while in the facility, a certification of need must be completed and placed in the client’s record within 14 days of the admission.

Inpatient psychiatric treatment is a benefit of Texas Medicaid if all of the following are met:

- The client has a psychiatric condition that requires inpatient treatment.
- The inpatient treatment is directed by a psychiatrist.
- The inpatient treatment is provided in a nationally accredited facility or hospital.
- The provider is enrolled in Texas Medicaid.

Client services must be provided in the most appropriate setting and in a timely manner to meet the mental health needs of the client.

Inpatient admissions to acute care hospitals, freestanding, and state psychiatric facilities are subject to the Texas Medicaid retrospective utilization review (UR) requirements. The UR requirements are applicable, regardless of the hospital’s designation as a psychiatric unit versus a medical or surgical unit.

### 3.4.2.1 Professional Services Rendered in the Inpatient Setting

Services rendered in the inpatient hospital setting may be reimbursed to the professional that provides the service.

**Refer to:** Subsection 7, “Psychiatric Services for Hospitals” in the Behavioral Health and Case Management Services Handbook (Vol. 2, Provider Handbooks) for benefit and limitation information about services that are rendered by psychiatrists, psychologists, LPAs, APNs, and PAs in the inpatient setting.

### 3.4.2.2 Documentation Requirements

All services require documentation to support the medical necessity of the service rendered, including all hospital services. Hospital services are subject to retrospective review and recoupment if documentation does not support the service that was submitted for reimbursement.

Documentation of medical necessity for inpatient psychiatric care must specifically address the following issues:

- Why the ambulatory care resources in the community cannot meet the treatment needs of the client.
- Why inpatient psychiatric treatment under the care of a psychiatrist is required to treat the acute episode of the client.
- How the services can reasonably be expected to improve the condition or prevent further regression of the client’s condition in a proximate time period.

Supporting documentation (certification of need) must be documented in the individual client’s record. This documentation must be maintained by each facility as applicable to state and federal guidelines and be readily available for review whenever requested by the HHSC or its designee.

Psychological or neuropsychological testing, when performed in an acute care hospital or in a freestanding or state psychiatric facility does not require prior authorization; however, these facilities must maintain documentation that supports medical necessity for the testing and the testing results of any psychological or neuropsychological testing services performed while the client is an inpatient.
3.4.2.3 Noncovered Services

Inpatient admissions including, but not limited to, the following are not benefits of Texas Medicaid without an accompanying medical complication or condition:

- Single diagnosis of chemical dependency or abuse (such as alcohol, opioids, barbiturates, and amphetamines).
- Chronic diagnoses (such as intellectual or developmental disability [IDD], organic brain syndrome, or chemical dependency or abuse).

3.4.2.4 CLIA Certification for Laboratory Services

All providers of laboratory services must comply with the rules and regulations of the Clinical Laboratory Improvement Amendments of 1988 (CLIA). Providers that do not comply with CLIA are not reimbursed for laboratory services.

Texas Medicaid follows the Medicare categorization of tests for CLIA certificate-holders.


For waived tests, providers must use modifier QW as indicated on the CMS website.

3.4.3 Acute Care Hospital Psychiatric Services

Acute care hospital psychiatric services are those services that are rendered to Texas Medicaid clients of any age who are admitted as an inpatient to an acute care hospital for treatment of a psychiatric condition.

Admissions to acute care hospitals must be medically necessary.


3.4.3.1 Prior Authorization Requirements

Prior authorization is not required for fee-for-service clients admitted to psychiatric units in acute care hospitals. Prior authorization cannot be used as a mechanism to deny, reduce, or controvert required court ordered services for Medicaid beneficiaries, as described in Subsection 4.6, “Court-Ordered Services” in the Behavioral Health and Case Management Services Handbook (Vol. 2, Provider Handbooks).

3.4.4 Freestanding and State Psychiatric Facilities

Psychiatric facility services are those services that are rendered in an Institutions for Mental Diseases (IMD). Freestanding and state psychiatric facilities are enrolled in Texas Medicaid as IMDS. According to TAC Rule §419.373(6) and based on 42 CFR §435.1009, an IMD is a hospital, nursing facility, or other institution of more than 16 beds that is primarily engaged in providing diagnosis, treatment, or care of individuals with mental illness, including medical attention, nursing care, and related services.

3.4.4.1 CCIP Services

Inpatient psychiatric treatment in a nationally accredited freestanding psychiatric facility or a nationally-accredited state psychiatric hospital is a benefit of Texas Medicaid for clients who are 20 years of age and younger, and who are eligible for THSteps benefits at the time of the service request and service delivery.

Admissions to freestanding and state psychiatric facilities must be medically necessary, unless they are court-ordered services for mental health commitments or they are a condition of probation.
Revenue code 124 must be used for inpatient psychiatric services that are rendered to children and adolescents in freestanding and state psychiatric facilities.

**Note:** Outpatient services for hospital-based psychiatric day treatment programs or psychiatric facilities are not a benefit of Texas Medicaid.

### 3.4.4.1.1 Prior Authorization Requirements for Children and Adolescents

Prior authorization is not required for initial admission to freestanding psychiatric facilities or state psychiatric hospitals for clients who are 20 years of age and younger for a maximum of five days based on Medicaid eligibility and documentation of medical necessity. Extended stay requests beyond the initial 5 days require prior authorization.

Extended stay prior authorization requests for fee-for-service clients may be submitted to the TMHP Prior Authorization Department via mail, fax, or the electronic portal. The supervising psychiatrist may sign prior authorization forms and supporting documentation using electronic or wet signatures.

**Refer to:** Subsection 5.5.1.2, “Document Requirements and Retention” in “Section 5: Fee-for-Service Prior Authorizations” (Vol. 1, General Information) for additional information about electronic signatures.

To facilitate a determination of medical necessity and avoid unnecessary denials, the physician must provide correct and complete information, including accurate documentation of medical necessity for the services requested.

Authorization procedures and approved providers may be different for managed care clients. Contact the client’s specific health plan for details.

The prior authorization requests will be reviewed as follows:

- Providers must submit a Psychiatric Inpatient Extended Stay Request Form to the TMHP CCIP unit requesting prior authorization for an extended stay. Requests for an extension of stay must be received on or before the fifth day of an initial admission and/or on or before the last day authorized or denied for subsequent stay requests. If an extended stay is requested and the fifth day of the initial admission or the last date authorized or denied of the previous stay falls on a holiday or a weekend, the request for an extended stay is due by 5 p.m. of the next business day. The provider is notified of the decision in writing by the CCIP unit.
- All psychiatric admission extended stay requests for clients who are 11 years of age or younger will be reviewed by a psychiatrist.
- Psychiatric admission extended stay requests for clients who are 12 through 20 years of age will be reviewed by a mental health professional. Any requests for psychiatric extended stays that do not meet the medical necessity criteria for extended stays will be referred to a psychiatrist for final determination.
- The Psychiatric Inpatient Extended Stay Request Form must reflect the need for extended stay in relation to the original need for admission. Any change in the client’s diagnosis must be noted on the request. Additional documentation or information supporting the need for extended stay may be attached to the form. Up to seven days may be authorized for an extension request.

**Medicaid Clinical Criteria for the Initial Inpatient Psychiatric Stay**

The client must have a valid diagnosis as listed in the current version of the DSM as the principal admitting diagnosis and one of the following:

- Outpatient therapy or partial hospitalization has been attempted and failed
- A psychiatrist has documented reasons why an inpatient level of care is required.
- The client must meet at least one of the following criteria:
The client is presently a danger to self, demonstrated by at least one of the following:

- Recent suicide attempt or active suicidal threats with a deadly plan, and there is an absence of appropriate supervision or structure to prevent suicide.

- Recent self-mutilative behavior or active threats of same with likelihood of acting on the threat, and there is an absence of appropriate supervision or structure to prevent self-mutilation (e.g., intentionally cutting/burning self).

- Active hallucinations or delusions directing or likely to lead to serious self-harm or debilitating psychomotor agitation or impairment resulting in a significant inability to care for self.

- Significant inability to comply with prescribed medical health regimens due to concurrent psychiatric illness and such failure to comply is potentially hazardous to the life of the client. The medical diagnosis must be treatable in a psychiatric setting.

The client is a danger to others. This behavior must be attributable to the client’s specific diagnosis as listed in the current version of the DSM, and can be adequately treated only in a hospital setting. This danger is demonstrated by one of the following:

- Recent life-threatening action or active homicidal threats of same with a deadly plan, availability of means to accomplish the plan, and with likelihood of acting on the threat.

- Recent serious assaultive or sadistic behavior or active threats of same with likelihood of acting on the threat, and there is an absence of appropriate supervision or structure to prevent assaultive behavior.

- Active hallucinations or delusions directing or likely to lead to serious harm of others.

- The client exhibits acute onset of psychosis or severe thought disorganization, or there is significant clinical deterioration in the condition of someone with a chronic psychosis, rendering the client unmanageable and unable to cooperate in treatment, and the client is in need of assessment and treatment in a safe and therapeutic setting.

- The client has a severe eating or substance use disorder that requires 24-hour-a-day medical observation, supervision, and intervention.

- The client exhibits severe disorientation to person, place, or time.

- The client’s evaluation and treatment cannot be carried out safely or effectively in other settings due to severely disruptive behaviors and other behaviors, which may also include physical, psychological, or sexual abuse.

- The client requires medication therapy or complex diagnostic evaluation where the client’s level of functioning precludes cooperation with the treatment regimen.

- The client is involved in the legal system, manifests psychiatric symptoms, and is ordered by a court to undergo a comprehensive assessment in a hospital setting to clarify diagnosis and treatment needs.

- The proposed treatment or therapy requires 24-hour-a-day medical observation, supervision, and intervention and must include all of the following:

  - Active supervision by a psychiatrist with the appropriate credentials as determined by the Texas Medical Board (TMB) and with documented specialized training, supervised experience, and demonstrated competence in the care and treatment of children and adolescents. Treatment/therapy plans must be guided by the standards of treatment specified by the Texas Society of Child and Adolescent Psychiatry.

  - Implementation of an individualized treatment plan.
• Provision of services that can reasonably be expected to improve the client’s condition or prevent further regression so that a lesser level of care can be implemented.

• Documentation that proper treatment of the client’s psychiatric condition requires services on an inpatient basis under the direction of a psychiatrist and is being provided in the least restrictive environment available, and ambulatory care resources available in the community do not meet the client’s needs.

Medicaid Clinical Criteria for Extended Stays

Extended stays are considered for THSteps clients in freestanding and state psychiatric hospitals when the client meets at least one of the criteria from above and have a treatment or therapy regimen, which must include all of the following:

• Active supervision by a psychiatrist with the appropriate credentials as determined by the Texas Medical Board (TMB) and with documented specialized training, supervised experience, and demonstrated competence in the care and treatment of children and adolescents. Treatment/therapy plans must be guided by the standards of treatment specified by the Texas Society of Child and Adolescent Psychiatry.

• Treatment/therapy requires an inpatient level of care.

• Initial discharge plans have been formulated and actions have been taken toward implementation, including documented contact with a local mental health provider.

Extended stays are also considered for children and adolescents whose discharge plan does not include returning to their natural home. If the party responsible for placement has provided the inpatient provider with three documented placement options for which the child meets admission criteria, but cannot accept the child, up to five days may be authorized, per request, to allow alternative placement to be located. Up to three 5-day extensions may be authorized.

Court-Ordered Services

A request for prior authorization of court-ordered services must be submitted no later than seven calendar days after the date on which the services began.

Court-ordered services are not subject to the five-day admission limitation or the seven-day extended stay limitation. Court-ordered services include:

• Mental health commitments

• Condition of probation (COP)

For court-ordered admissions, a copy of the doctor’s certificate and all court-ordered commitment papers signed by the judge must be submitted with the Psychiatric Inpatient Extended Stay Request Form.

Prior Authorization Appeals

All prior authorization requests not submitted or received by the TMHP CCIP unit in accordance with established policies are denied through the submission date, and claim payment is not made for the dates of service denied.

All denials may be appealed. The TMHP CCIP unit must receive these appeals within 15 days of the TMHP CCIP unit denial notice.

Appeals of a denial for an extended stay, must be accompanied by the documentation supporting medical necessity that the provider believes warrants reconsideration.

Appeals of a denial for late submission of information, must be accompanied by documentation that the provider believes supports the compliance with HHSC claims submission guidelines.
Appeals are reviewed first by an experienced psychiatric licensed clinical social worker (LCSW) or a registered nurse (RN) to determine if the required criteria is documented and then forwarded to a psychiatrist for final determination. The TMHP CCIP unit will notify the provider of all denial determinations in writing.

3.4.4.2 Psychiatric Services for Clients 65 Years of Age and Older

IMD services for clients who are 65 years of age and older must be medically necessary and do not require prior authorization.

3.4.4.3 Reimbursement for Services Rendered in an IMD

The following services will not be reimbursed during an inpatient stay when they are rendered to clients who are admitted as inpatients to an IMD:

- Ambulance
- Case management
- Acute care hospital
- Mental health rehabilitation
- School Health and Related Services (SHARS)

IMD providers may be reimbursed only for services that are rendered to clients who are 20 years of age and younger or 65 years of age and older. IMD services for clients who are 21 through 64 years of age are not eligible for fee-for-service reimbursement. If delivered through managed care and the managed care organization and client agree to an IMD as the setting for inpatient services, IMD services for clients who are 21 through 64 years of age are eligible for reimbursement for a total of 15 days per calendar month. The 15-day limitation is counted per calendar month, not per stay. As such, a stay of up to 15 days in a single month and a stay of up to 15 days in a following month may both be eligible for reimbursement.

Note: Acute care hospitals are not considered IMDs, even if they have a psychiatric ward.

Services that are rendered in an IMD facility must be identified in the client’s plan of care. Services that are not included in the client’s plan of care are subject to recoupment.

If the client has not been discharged from the IMD, the IMD provider is responsible for acute care services that are rendered to the client in an acute care facility, and claims that are submitted for these services will be denied as a duplicate service that has been paid to another provider.

Services that are rendered on the date of admission to the IMD and the date of discharge from the IMD may be reimbursed.

Important: Claims for professional services rendered during an inpatient stay in an Institution for Mental Disease (IMD) must include the IMD facility’s ten-digit National Provider Identifier (NPI). Claims that do not include the IMD Facility’s NPI will be denied.

3.4.4.3.1 Medicare Coinsurance and Deductible Reimbursement

Freestanding psychiatric hospitals that are enrolled in Medicare may also receive Medicaid payment for the Medicare coinsurance or deductible according to current Medicaid guidelines.

Exception: IMD services for clients who are 21 through 64 years of age are not eligible for fee-for-service reimbursement. If delivered through managed care and the managed care organization and client agree to an IMD as the setting for inpatient services, IMD services for clients who are 21 through 64 years of age are eligible for reimbursement for a total of 15 days per calendar month. The 15-day limitation is counted per calendar month, not per stay. As such, a stay of up to 15 days in a single month and a stay of up to 15 days in a following month may both be
eligible for reimbursement. Medicaid will not reimburse coinsurance and deductible payments for psychiatric services that are rendered to these clients in an IMD beyond 15 days per calendar month.

Refer to: Subsection 2.7, “Medicare Crossover Claim Reimbursement” in “Section 2: Texas Medicaid Fee-for-Service Reimbursement” (Vol. 1, General Information) additional information about Medicaid guidelines for Medicare coinsurance and deductible payments.

3.4.4.4 Providing IMD Client Information to TMHP
IMD providers are requested to inform TMHP of the Medicaid clients who are residing in their facilities before submitting inpatient claims for those clients.

IMD providers can use the TMHP secure web page to enter client information and the admission and discharge dates by going to My Account and choosing the Manage IMD Clients Segment link in the Acute Care Online Portal field.

IMD providers can search for Medicaid client records that are associated with their provider identifiers. Providers will be asked to submit the client’s identification number and admission date. After the client is discharged, providers will be requested to enter the discharge date on the same Manage IMD Clients Segment screen.

Providers will not be able to change previously reported client information except for the To Date of Service information.

Providers that submit inaccurate information can call the Electronic Data Interchange (EDI) Help Desk at 1-888-863-3638 to have that client segment inactivated if the request is made within 24 hours of submission. After the erroneous client segment has been inactivated by TMHP, providers may submit a new client segment to replace it. After 24 hours have elapsed, providers must contact HHSC to request a correction to the information. This change must include the appropriate documentation of the client’s patient control number (PCN) and the admission and discharge dates.

The change request and appropriate documentation must be submitted in writing to the following address:

Texas Health and Human Services Commission
Mail Code 91X PO Box 204077
Austin, Texas 78720-4077

3.4.5 Medicaid Clinical Criteria for Inpatient Psychiatric Care for Clients
The client must have a valid DSM diagnosis as the principle admitting diagnosis and outpatient therapy or partial hospitalization has been attempted and failed, or a psychiatrist has documented reasons why an inpatient level of care is required. The client’s diagnosis must also be included on the request for inpatient psychiatric treatment.

The client must meet at least one of the following criteria:

- The client is presently a danger to self, demonstrated by at least one of the following:
  - Recent suicide attempt or active suicidal threats with a deadly plan and an absence of appropriate supervision or structure to prevent suicide.
  - Recent self-mutilative behavior or active threats of same with likelihood of acting on the threat and an absence of appropriate supervision or structure to prevent self-mutilation (i.e., intentionally cutting or burning self).
  - Active hallucinations or delusions directing or likely to lead to serious self-harm or debilitating psychomotor agitation or intellectual disability resulting in a significant inability to care for self.
• Significant inability to comply with prescribed medical health regimens due to concurrent psychiatric illness and such failure to comply is potentially hazardous to the life of the client. The medical diagnosis must be treatable in a psychiatric setting.

• The client is a danger to others. This behavior should be attributable to the client’s specific DSM diagnosis and can be adequately treated only in a hospital setting. This danger is demonstrated by one of the following:
  • Recent life-threatening action or active homicidal threats of same with a deadly plan and availability of means to accomplish the plan with likelihood of acting on the threat.
  • Recent serious assaultive or sadistic behavior or active threats of same with likelihood of acting on the threat and an absence of appropriate supervision or structure to prevent assaultive behavior.
  • Active hallucinations or delusions directing or likely to lead to serious harm of others.

• The client exhibits acute onset of psychosis or severe thought disorganization, or there is significant clinical deterioration in the condition of someone with a chronic psychosis, rendering the client unmanageable and unable to cooperate in treatment, and the client is in need of assessment and treatment in a safe and therapeutic setting.

• The client has a severe eating or substance use disorder which requires 24-hour-a-day medical observation, supervision, and intervention.

• The client exhibits severe disorientation to person, place, or time.

• The client’s evaluation and treatment cannot be carried out safely or effectively in other settings due to severely disruptive behaviors and other behaviors which may also include physical, psychological, or sexual abuse.

• The client requires medication therapy or complex diagnostic evaluation where the client’s level of functioning precludes cooperation with the treatment regimen.

• The client is involved in the legal system, manifests psychiatric symptoms, and is ordered by court to undergo a comprehensive assessment in a hospital setting to clarify the diagnosis and treatment needs.

The proposed treatment or therapy requires 24-hour-a-day medical observation, supervision, and intervention and must include all of the following:

• Active supervision by a psychiatrist with the appropriate credentials as determined by the Texas Medical Board (TMB) and with documented specialized training, supervised experience, and demonstrated competence in the care and treatment of children and adolescents. Treatment or therapy plans must be guided by the standards of treatment specified by the Texas Society of Child and Adolescent Psychiatry.

• Implementation of an individualized treatment plan.

• Provision of services which can reasonably be expected to improve the client’s condition or prevent further regression so that a lesser level of care can be implemented.

Proper treatment of the client’s psychiatric condition requires services on an inpatient basis under the direction of a psychiatrist and is being provided in the least restrictive environment available, and ambulatory care resources available in the community do not meet the client’s needs.
3.4.6 Extended Stays

Extended stays are considered when the client meets at least one of the criteria from above and has a treatment or therapy regimen that includes all of the following:

- Active supervision by a psychiatrist with the appropriate credentials as determined by the TMB and with documented specialized training, supervised experience, and demonstrated competence in the care and treatment of children and adolescents. Treatment or therapy plans must be guided by the standards of treatment specified by the Texas Society of Child and Adolescent Psychiatry.
- Treatment or therapy requires an inpatient level of care.
- Initial discharge plans have been formulated and actions have been taken toward implementation, including documented contact with a local mental health provider.

Extended stays are considered for children and adolescents whose discharge plan does not include returning to their natural home. If the party responsible for placement has provided the provider with three documented placement options for which the child meets admission criteria, but which cannot accept the child, up to five days may be authorized, per request, to allow alternative placement to be located. Up to three five-day extensions may be authorized.

3.4.7 Court-Ordered Services

Refer to: Subsection 4.6, “Court-Ordered Services” in the Behavioral Health and Case Management Services Handbook (Vol. 2, Provider Handbooks) for additional information.

3.4.8 Denials

All prior authorization requests not submitted or received by the TMHP CCIP Unit in accordance with established policies are denied through the submission date, and claim payment is not made for the dates of service denied.

All denials may be appealed. The TMHP CCIP Unit must receive these appeals within 15 days of the TMHP CCIP Unit denial notice.

- Appeals of a denial for an extended stay must be accompanied by the documentation supporting medical necessity that the provider believes warrants reconsideration.
- Appeals of a denial for late submission of information must be accompanied by documentation which the provider believes supports the compliance with HHSC claims submission guidelines.
- Appeals are reviewed first by an experienced psychiatric LCSW (Licensed Clinical Social Worker) or an RN to determine if the required criteria is documented and then forwarded to a psychiatrist for final determination. The TMHP CCIP Unit will notify the provider of all denial determinations in writing by fax.

3.5 Inpatient Utilization Review

UR activities of all Medicaid services provided by hospitals reimbursed under the DRG prospective payment system or the Tax Equity and Fiscal Responsibility Act (TEFRA) of 1982 are required by Title XIX of the Social Security Act, Sections 1902 and 1903. The review activities are accomplished through a series of monitoring systems developed to ensure services are appropriate to need of optimum quality and quantity, and rendered in the most cost-effective mode. Clients and providers are subject to UR monitoring. The monitoring focuses on the appropriate screening activities, medical necessity of all services, and quality of care as reflected by the choice of services provided, type of provider involved, and settings in which the care was delivered. This monitoring ensures the efficient and cost-effective administration of Texas Medicaid.

The HHSC Office of Inspector General (OIG) UR Unit is responsible for retrospective review of inpatient DRG and TEFRA admissions. These reviews are accomplished through onsite visits, electronic access, or mail-in.
3.6 Utilization Review Process

The inpatient UR process for admissions reimbursed under the DRG prospective payment system consists of sampling medical records of paid Medicaid claims. The review process consists of three major components:

- **Admission review.** Determination of the medical necessity of the admission. For purposes of the Texas Medical Review Program (TMRP), TEFRA, and facility-specific per diem methodology reviews, medical necessity means the client has a condition requiring treatment that can be safely provided only in the inpatient setting.

- **Quality review.** Assessment of the quality of care provided to determine if it meets generally accepted standards of medical and hospital care practices or puts the client at risk of unnecessary injury or death. Quality of care review includes the use of discharge screens and generic quality screens.

- **DRG validation.** Determination that the critical elements necessary to assign a DRG are present in the medical record and procedures are sequenced correctly. The critical elements are age, sex, admission date, discharge date, patient discharge status, principal diagnosis, secondary diagnoses (complications or comorbidities), and principal and secondary procedures.

The HHSC OIG UR Unit staff reviews the complete medical record to make decisions about the medical necessity of the admission, validity of the DRG, including the present on admission (POA) indicator, and quality of care. The medical record must reflect that any services reimbursed by Texas Medicaid were ordered by a physician or non-physician provider.

When an admission denial or a denial of extended stay is issued, or when a technical denial becomes final, all money is recouped from the hospital for the admission or days of stay that are denied. When a DRG is reassigned as a result of UR, the payment to the hospital is adjusted.

**Refer to:** Subsection 3.6.3, “Technical Denials (DRG Prospective Payment)” in this handbook.

If an admission is denied, but a physician’s order is present documenting the client originally was placed in observation, the UR unit may authorize the resubmission of services rendered during the first 48 hours on an outpatient claim.

Compliance with the DRG prospective payment system and aspects of the review as stated above are evaluated quarterly. Identified problems may result in an educational visit or action, such as recoupment or referral to the HHSC OIG Medicaid Program Integrity (MPI) or Sanctions Unit.

3.6.1 Admission Review

All services, supplies, or items submitted as certified on the claim submission must be medically necessary for the client’s diagnosis or treatment. Health and Human Services Commission-Office of Inspector General-Utilization Review (HHSC-OIG-UR) personnel evaluate the medical necessity of an admission by comparing the medical record documentation to Change Healthcare InterQual® evidence-based guidelines. Non-physician reviewers use this criteria to determine medical necessity for initial approval of admission. If the case does not meet the initial approval criteria, the reviewer refers the case to a physician consultant who will determine the medical necessity of the inpatient admission. If the initial approval criteria are met but the medical necessity of the admission is still questionable, the non-physician reviewer refers the case to a physician consultant for a determination. If a physician consultant determines the admission is not medically necessary, HHSC-OIG-UR will issue a denial.

3.6.1.1 Readmission Review

If a hospital admission or readmission occurs within 30 days of a discharge from the same or a different hospital for the same or closely related diagnosis, or for a condition identified during the previous admission, it may be reviewed for medical necessity, quality, and DRG validation including POA indicators.
Transfers from one facility to another and readmissions are also subject to review.

3.6.1.2 Hospital-Based Ambulatory (HASC) Surgical Procedures

Inpatient admissions for surgical procedures listed as ambulatory surgical codes in the current fee schedule are denied if documentation does not support the need for the inpatient admission.

3.6.1.3 Quality Review

Each Medicaid case is evaluated for quality of client care, adequacy of discharge planning, and medical stability of the client at discharge. Quality of care review includes the use of discharge screens and generic quality screens. Potential quality of care issues are identified by the physician. HHSC contracts with physician consultants to review medical records for quality of care. Physician consultants, of the specialty related to the care rendered, may make clinical recommendations or determine corrective actions when deemed appropriate. Child and adolescent psychiatrists may make recommendations based on review of inpatient psychiatric services provided to Medicaid clients younger than 21 years of age. Failure to verify completion of any corrective action recommendation within the specified time frame may result in referral of the case to the HHSC OIG MPI or Sanctions Unit.

3.6.1.4 Diagnosis-Related Group Validation

Each medical record is reviewed to validate the elements critical to the DRG assignment. These elements are the client’s age, sex, admission date, patient discharge date, patient discharge status, principal diagnosis, secondary diagnoses (complications or co-morbidities), POA indicators, and principal and secondary procedures. Documentation of these critical DRG elements in the medical record is evaluated for the correlation to the information provided on the claim form.

The principal diagnosis is the diagnosis (condition) established after study to be chiefly responsible for causing the admission of the client to the hospital for care. The condition must be treated or evaluated during this admission to the hospital.

The secondary diagnoses are conditions that affect client care in terms of requiring clinical evaluation, therapeutic treatment, diagnostic procedures, extended length of hospital stay, increased nursing care and monitoring, or have clinically significant implications for future health-care needs.

The coding of diagnoses that have clinically significant implications for future health-care needs applies only to newborns and must be identified by the physician. Normal newborn conditions or routine procedures are not to be considered as complications or co-morbidities for DRG assignment.

Refer to: Subsection 1.12, “Texas Medicaid Limitations and Exclusions” in “Section 1: Provider Enrollment and Responsibilities” (Vol. 1, General Information).

The POA review will validate the POA indicator assigned to the principal and secondary diagnoses codes reported on claim forms. If it is determined that the principal and/or secondary diagnoses were not present at the time the order for inpatient admission occurs, the Commission will revise the POA indicator for the diagnosis code. Conditions that develop during an outpatient encounter, including emergency department, observation, or outpatient surgery are considered POA.

If the principal diagnosis, secondary diagnoses (complications or co-morbidities), or procedures are not substantiated in the medical record; sequenced correctly; have an incorrect POA indicator; or have been omitted, codes may be deleted, changed, or added. All diagnosis or procedure coding changes potentially resulting in a DRG change are referred to a physician consultant. When it is determined that the diagnoses and procedures are substantiated and sequenced correctly, the information will be entered into the applicable version of the Grouper software for a DRG determination. The CMS-approved DRG software considers each diagnosis and procedure and the combination of all codes and elements to make a determination of the final DRG assignment. When the DRG is reassigned, the payment to the provider is adjusted.
3.6.2 Recommendations to Enhance Compliance with Texas Medicaid Fee-for-Service Hospital Claims Submission

The following information highlights an area for physician and hospital providers where collaboration in client care delivery exists but can improve. Texas Medicaid, through its hospital UR activities, has identified this area for both compliance with provider responsibilities and the reduction of the submission of inappropriate inpatient hospital claims.

To enhance compliance with Texas Medicaid fee-for-service hospital claims submission and decrease the submission of inappropriate inpatient hospital claims, providers should adhere to the following:

- The hospital may admit clients in observation status if the physician has the reasonable expectation that the client will be discharged within 48 hours. If an inpatient claim was denied per retrospective UR, the hospital may resubmit the claim for the first 48 hours as an outpatient claim if the client was initially admitted in observation status (per physician order) and the stay was more than 48 hours.

- When a client is admitted to the hospital as an inpatient and is discharged in less than 48 hours, the hospital may request that the physician change the admission order from inpatient status to outpatient observation status. This practice is acceptable when the physician makes changes to the admitting order before the hospital submits the claim for payment.

- The correction in admission status, when the above criteria are met avoids errors in claims submission and the potential need for a more lengthy appeal process.

- If the physician admitting orders do not accurately reflect the services provided, the hospital inpatient claim may be denied and the inappropriate payment recovered from both the hospital and the admitting physician.

3.6.3 Technical Denials (DRG Prospective Payment)

3.6.3.1 On-Site Reviews

The following information describes on-site reviews:

- If the complete medical record is not made available during the on site review, a preliminary technical denial is issued on site. The hospital is allowed 60 calendar days from the date of the exit conference to provide the complete medical record to HHSC. If the complete medical record is not received by HHSC within this specified time frame, a final technical denial is issued and payment is recouped.

- If a complete medical record is made available on site, but a copy is required for further review, and the copy is not received by HHSC within the specified time frame, a preliminary technical denial is issued by certified mail or fax. The hospital has 60 calendar days from the date of receipt of the notice to submit the complete medical record. If the complete medical record is not received by HHSC within this specified time frame, a final technical denial is issued and payment is recouped.

  Note: A notarized business record affidavit in the format approved by HHSC is required for paper and electronic copies of requested medical records. A provider failing to provide this documentation must resubmit the requested records with the affidavit.


3.6.3.2 Mail-In Reviews

If the complete medical record is not received by HHSC within the specified time frame, a preliminary technical denial is issued by certified mail or fax. The hospital has 60 calendar days from the date of receipt of the notice to submit the complete medical record. If the complete medical record is not received by HHSC within this specified time frame, a final technical denial is issued and payment is recouped.
Hospital inpatient claim payments that have been recouped because of a technical denial may not be resubmitted on an outpatient claim.

**Note:** A notarized business record affidavit in the format approved by HHSC is required for paper and electronic copies of requested medical records. A provider who fails to provide this documentation must resubmit the requested records with the affidavit.

**Refer to:** Subsection 1.7.3, “Retention of Records and Access to Records and Premises” in “Section 1: Provider Enrollment and Responsibilities” (Vol. 1, General Information).

### 3.6.4 Acknowledgment of Penalty Notice

Hospitals must have on file a signed acknowledgment from the physician stating that the physician received the following notice:

**Notice to Physicians:** Medicaid payment to hospitals is based, in part, on each client’s principal and secondary diagnoses and the major procedures performed on the client, as attested to by the client’s attending physician by virtue of his or her signature in the medical record. Anyone who misrepresents, falsifies, or conceals essential information required for payment of federal or state funds, may be subject to fine, imprisonment, or civil penalty under applicable federal and state laws.

The acknowledgment of penalty notice must be specific to Texas Medicaid. Medicare penalty notices are not accepted.

### 3.6.5 Sanctions

Compliance with the DRG prospective payment system and aspects of the review as stated above are evaluated quarterly. Identified problems may result in an educational visit or action such as recoupment or referral to HHSC OIG MPI or Sanctions Unit.

### 3.6.6 Utilization Review Appeals

Hospitals may appeal adverse decisions by HHSC OIG UR Unit to the HHSC UR Medical Appeals Unit. A UR Medical Appeals decision is the final administrative decision of HHSC. Neither HHSC OIG UR Unit nor TMHP are responsible for Medical UR appeals.

**Refer to:** Subsection 7.3.3, “Utilization Review Appeals” in “Section 7: Appeals” (Vol. 1, General Information).

### 3.7 Claims Filing and Reimbursement

#### 3.7.1 Medicaid Relationship to Medicare

Texas Medicaid may make deductible or coinsurance payments according to current Medicaid payment guidelines on valid, assigned Part A (hospital) and Part B (medical) Medicare claims.

**Refer to:** Subsection 2.7, “Medicare Crossover Claim Reimbursement” in “Section 2: Texas Medicaid Fee-for-Service Reimbursement” (Vol. 1, General Information) for information about coinsurance and deductible payment guidelines.

Texas Medicaid provides reimbursement for 30 inpatient benefit days per spell of illness. When the 30 days coincide with the first 30 days of the Medicare benefit period and the client is eligible for both Medicare and Medicaid, Texas Medicaid pays the:

- Inpatient hospital deductible under Medicare Part A.
- Medicare Part A deductible for the first three pints of whole blood or packed red cells.

When the client only has Medicare Part B coverage, the hospital must follow these guidelines:

- Submit to Medicare the charges for certain inpatient ancillary services on a Medicare Claim Form 1483 for payment under the client’s Part B coverage. The ancillary charges include the following:
• Diagnostic X-ray tests, diagnostic laboratory tests, and other diagnostic tests
• X-ray, radium, and radioactive isotope therapy, including materials and services of technicians
• Surgical dressings, splints, casts, and other devices used for reduction of fractures and dislocations
• Prosthetic devices (other than dental) that replace all or part of an internal body organ or member (including contiguous tissue) or all or part of the function of a permanently inoperative or malfunctioning internal body organ or member including replacement or repairs of such devices (e.g., cardiac pacemakers, breast prostheses, maxillofacial devices, colostomy bags, and prosthetic lenses)
• Leg, arm, back, and neck braces, and artificial legs, arms, and eyes, including replacements and adjustments (if required) because of a change in the client’s physical condition
• Physical therapy (PT) services
• Speech pathology services
• Dialysis treatments

• Submit to TMHP the remaining Part A charges on a UB-04 CMS-1450 paper claim form (or its electronic equivalent) indicating in Block 80 that the client is eligible for Medicare Part B benefits only. The client’s Medicare number must appear on the Medicaid claim in Block 80. TMHP must receive these charges within 95 days of the last date of service on the claim. 

Referto: Subsection 2.7, “Medicare Crossover Claim Reimbursement” in “Section 2: Texas Medicaid Fee-for-Service Reimbursement” (Vol. 1, General Information).

3.7.2 Inpatient Claims Information

Medicaid present on admission (POA) reporting is required for all inpatient hospital claims.

All hospital providers are required to submit a POA value for each diagnosis on the claim form, and no hospital is exempt from this POA requirement. Medicare crossover hospital claims must also comply with the Medicaid requirement to include the POA values.

POA is defined as present at the time the order for inpatient admission occurs. Conditions that develop during an outpatient visit, including emergency department, observation, or outpatient surgery, are considered POA.

The following table shows the POA values:

<table>
<thead>
<tr>
<th>POA Value</th>
<th>Description</th>
<th>Payment</th>
</tr>
</thead>
<tbody>
<tr>
<td>Y</td>
<td>Diagnosis was present at the time of admission</td>
<td>Payment will be made by Medicare when a hospital-acquired condition (HAC) is present</td>
</tr>
<tr>
<td>N</td>
<td>Diagnosis was not present at the time of admission</td>
<td>No payment will be made by Medicare when an HAC is present</td>
</tr>
<tr>
<td>U</td>
<td>Documentation was insufficient</td>
<td>No payment will be made by Medicaid when an HAC is present</td>
</tr>
<tr>
<td>W</td>
<td>Clinically undetermined</td>
<td>Payment will be made by Medicaid when an HAC is present:</td>
</tr>
<tr>
<td>(blank)</td>
<td>Exempt from POA reporting</td>
<td>Exempt from POA reporting</td>
</tr>
</tbody>
</table>

Note: Texas Medicaid follows Medicare guidelines for payments referenced in this table.

Note: If a diagnosis code is exempt from POA reporting, providers should leave the POA indicator field blank on the claim.
TMHP will not recalculate the DRG based on POA indicator values for Medicare crossover claims or MCOs.

Depending on the POA indicator value, the DRG may be recalculated, resulting in a lower payment to the hospital facility provider. If the number of days on an authorization is higher than the number of days allowed as a result of a POA DRG recalculation, the lesser of the number of days will be reimbursed.

Refer to: “Section 3: TMHP Electronic Data Interchange (EDI)” (Vol. 1, General Information).

“Section 6: Claims Filing” (Vol. 1, General Information).

A complete list of POA exempt diagnosis codes can be found on the CMS website.

Claims for inpatient hospital services must be submitted to TMHP in an approved electronic format or on the UB-04 CMS-1450 paper claim form. Providers may purchase UB-04 CMS-1450 paper claim forms from the vendor of their choice. TMHP does not supply the forms.

When completing a UB-04 CMS-1450 paper claim form, all required information must be included on the claim, as TMHP does not key any information from attachments. Superbills, or itemized statements, are not accepted as claim supplements.

In Block 44 of the UB-04 CMS-1450, enter the accommodation rate per day. Match the appropriate diagnoses listed in Blocks 67A through 67Q corresponding to each procedure. If a procedure corresponds to more than one diagnosis, enter the primary diagnosis. Each service and supply must be itemized on the claim.

Hospitals may submit information only claims to TMHP when one of the following situations exists. Hospitals must use TOB 110 to file these claims:

- Inpatient 30-day spell of illness benefit is exhausted.
- Payment made by a third party resource or other insurance exceeds the Medicaid allowed amount.

Additional claims information can be found within individual topic areas in this section.

Refer to: “Section 3: TMHP Electronic Data Interchange (EDI)” (Vol. 1, General Information).

“Section 6: Claims Filing” (Vol. 1, General Information).

The Claim Form Examples page on the TMHP website at www.tmhp.com.

3.7.3 Inpatient Claims Submission

Inpatient hospital claims for traditional Medicaid clients must be submitted to TMHP if the client is enrolled in traditional Medicaid at the time of service or during the time of the admission.

3.7.4 Inpatient Reimbursement

3.7.4.1 Prospective Payment Methodology

Inpatient hospital stays except state-owned teaching hospitals, and psychiatric facilities (CCP) are reimbursed according to a prospective payment methodology based on diagnosis-related groups (DRGs). The reimbursement method itself does not affect inpatient benefits and limitations. Inpatient admissions must be medically necessary and are subject to Texas Medicaid’s UR requirements.

The DRG reimbursement includes all facility charges (e.g., laboratory, radiology, and pathology). Hospital-based laboratories and laboratory providers who deliver referred services outside the hospital setting must obtain reimbursement for the technical portion from the hospital. The technical portion includes the handling of specimens and he automated or technician-generated reading and reporting of results. The technical services are not billable to Texas Medicaid clients.
Texas Medicaid does not distinguish types of beds or units within the same acute care facility for the same inpatient stay (e.g., psychiatric or rehabilitation). Because all Medicaid inpatient hospitalizations are included in the DRG database that determines the DRG payment schedule, psychiatric and rehabilitation admissions are not excluded from the DRG payment methodology. To ensure accurate payment, Texas Medicaid requires that only one claim be submitted for each inpatient stay with appropriate diagnosis and procedure code sequencing. The discharge and admission hours (military time) are required on the UB-04 CMS-1450 paper claim form, to be considered for payment.

The number of days of care charged to a beneficiary for inpatient hospital or skilled nursing facility (SNF) care services is always in units of full days. A day begins at midnight and ends 24 hours later. The midnight-to-midnight method is to be used in counting days of care for reporting purposes even if the hospital or SNF uses a different definition of day for statistical or other purposes.

A part of a day, including the day of admission and day on which a patient returns from leave of absence, counts as a full day. However, the day of discharge, death, or a day on which a patient begins a leave of absence is not counted as a day unless discharge or death occur on the day of admission.

If admission and discharge or death occur on the same day, the day is considered a day of admission and counts as one inpatient day.

Reimbursement to acute care hospitals for inpatient services is limited to $200,000 per client, per benefit year (November 1 through October 31). Claims may be subject to retrospective review, which may result in recoupment. This limitation does not apply to services related to certain organ transplants or services to THSteps clients when provided through CCP.

A new provider is given a reimbursement inpatient interim rate of 50 percent until a cost audit has been performed. A default standard dollar amount (SDA) rate is assigned for newly enrolled providers or newly constructed facilities.

Payment is calculated by multiplying the SDA for the hospital’s payment division indicator times the relative weight associated with the DRG assigned by Grouper.

Hospital reimbursement is made in accordance with the following TAC rules:

- 1 TAC §355.761 - Reimbursement Methodology for Institutions of Mental Diseases (IMD)
- 1 TAC §355.8052 - Inpatient Hospital Reimbursement
- 1 TAC §355.8056 - State-Owned Teaching Hospital Reimbursement Methodology
- 1 TAC §355.8058 - Inpatient Direct Graduate Medical Education (GME) Reimbursement
- 1 TAC §355.8060 - Reimbursement Methodology for Freestanding Psychiatric Facilities
- 1 TAC §355.8061 - Outpatient Hospital Reimbursement
- 1 TAC §355.8065 - Disproportionate Share Hospital Reimbursement Methodology
- 1 TAC §355.8066 - Hospital-Specific Limit Methodology

Medicaid providers that are cost-reimbursed are subject to cost reporting, cost reconciliation, and cost settlement processes, as defined in the following TAC rules:

- 1 TAC §355.8061 (a)(2) - Outpatient Hospital Reimbursement
- 1 TAC §355.8052 (i)(i) - Inpatient Hospital Reimbursement
- 1 TAC §355.8056 - State-Owned Teaching Hospital Reimbursement Methodology
3.7.4.2 Client Transfers

3.7.4.2.1 Admission Dates

To ensure correct payor identification, providers that receive transfer patients from another hospital must enter the actual date on which the client was admitted into each facility in Block 12 on the UB-04 CMS-1450.

3.7.4.2.2 Continuous Stays – Client Transfers and Readmissions

Client transfers within the same facility are considered one continuous stay and receive only one DRG payment. Texas Medicaid does not recognize specialty units within the same hospital as separate entities; therefore, these transfers must be submitted as one admission under the provider identifier. Readmissions to the same facility within 24 hours of a previous acute hospital or facility discharge are also considered one continuous stay and receive only one DRG payment.

Readmissions are considered a continuous stay regardless of the original or readmission diagnosis. Admissions submitted inappropriately are identified and denied during the UR process and may result in intensified review.

When more than one hospital provides care for the same client, the hospital providing the most significant amount of care receives consideration for a full DRG payment. The other hospitals are paid a per diem rate based on the lesser of either the mean length of stay for the DRG or the eligible days in the facility. The DRG modifier, PT, on the R&S Report indicates per diem pricing related to a client transfer. Services must be medically necessary and are subject to Texas Medicaid’s UR requirements.

HHSC performs a postpayment review to determine if the hospital providing the most significant amount of care received the full DRG. If the review reveals that the hospital providing the most significant amount of care did not receive the full DRG, an adjustment is initiated.

To ensure correct payor identification, providers that receive transfer patients from another hospital must enter the actual date that the client was admitted into each facility in Block 12 on the UB-04 CMS-1450. Inpatient authorization requirements are based on the requirements that are specified by the program in which the client is enrolled on the date of the original admission. Providers must adhere to the authorization requirements for claims to be considered for reimbursement. Providers are reimbursed at the rate in effect on the date of discharge.

3.7.4.3 Observation Status to Inpatient Admission

The dates of the inpatient admission must be reported as follows:

- **Date of inpatient admission:** The date of admission must reflect the date that the client was admitted to the hospital as an inpatient.

- **Dates of service:** The from date of service (FDOS) must reflect the date that the client first presented at the hospital for services including, but not limited to, emergency room (ER), observation, labor and delivery, or inpatient services.

If services that are rendered before the inpatient admission must be submitted on the inpatient claim, the number of preadmission days that are related to the inpatient admission cannot exceed the days of allowed for the rendered services:

<table>
<thead>
<tr>
<th>Services</th>
<th>Days Allowed</th>
<th>Units</th>
</tr>
</thead>
<tbody>
<tr>
<td>Emergency room (ER) services</td>
<td>One day (24 hours) before the inpatient admission</td>
<td>Submitted per day</td>
</tr>
<tr>
<td>Observation services</td>
<td>Up to two days (48 hours) before the inpatient admission</td>
<td>Submitted in hours</td>
</tr>
<tr>
<td>Labor and Delivery</td>
<td>Up to three days before the inpatient admission</td>
<td>Submitted per day</td>
</tr>
</tbody>
</table>
Diagnosis-Related Group (DRG) hospital claims allow for a total of three days of pre-admit services. Non-DRG hospital claims are allowed one day of pre-admit services, and a second day if additional observation hours occurred.

**Note:** If the client is admitted as an inpatient more than 24 hours after presenting in the ER without being placed in observation status or more than 48 hours after being placed in observation status, the ER and observation services may be reimbursed separately as outpatient services and must not be included on the inpatient claim.

### 3.7.4.4 Outliers

TMHP makes outlier payment adjustments to DRG hospitals for admissions that meet the criteria for exceptionally high costs or exceptionally long lengths of stay for clients who are 20 years of age and younger as of the date of the inpatient admission. If a client’s admission qualifies for both a day and a cost outlier, the outlier resulting in the higher payment to the hospital is paid.

Providers can view their day and cost outlier payment information for inpatient hospital claims on the Electronic Remittance and Status (ER&S) Report. The R&S Report reflects the outlier reimbursement payment and defines the type of outlier paid. To view the day and cost outlier payment information, providers, facilities, and third party vendors may need to update their 835 electronic file format. For information about how to update the 835 electronic file format, refer to the revised electronic data interchange (EDI) companion guide (ANSI ASC X12N 835 Healthcare Claim Payment/Advice-Acute Care Companion Guide) on the TMHP website at [www.tmhp.com](http://www.tmhp.com).

#### 3.7.4.4.1 Day Outliers

The following criteria must be met to qualify for a day outlier payment:

- Inpatient days must exceed the DRG day threshold for the specific DRG.
- Additional payment is based on inpatient days that exceed the DRG day threshold multiplied by 60 percent of the per diem amount of a full DRG payment.
- The per diem amount is established by dividing the full DRG payment amount by the arithmetic mean length of stay for the DRG.

In compliance with 1 TAC §355.8052, all DRG inpatient hospital day outlier payments must not exceed the allowed cost for the service. All hospitals except in-state children’s hospitals, both day and cost outlier payments have been reduced by 10 percent.

TMHP calculates payments as follows:

1) Calculate the day outlier.

   - Calculate the allowed cost for the service (i.e., the cap amount) by taking the difference between the Tax Equity and Fiscal Responsibility Act (TEFRA) and DRG-payable amounts.
   - Take the lesser of the day outlier or the cap amount.
   - Reduce the day outlier by 10 percent for all hospitals except in-state children’s hospitals.

2) Calculate the cost outlier:

   - Reduce the cost outlier by 10 percent for all hospitals except in-state children’s hospitals.

Reimbursement is made for day or cost outliers on claims that qualify. If a client’s admission qualifies for both a day and a cost outlier, the outlier resulting in the higher payment to the hospital is paid.
Example

<table>
<thead>
<tr>
<th>Calculations</th>
<th>Example amount</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>Day Outlier Calculation:</strong></td>
<td>$500.00</td>
</tr>
<tr>
<td>([SDA x DRG relative weight / Mean Length of Stay] x Outlier Days) x 0.6 = day outlier amount</td>
<td></td>
</tr>
<tr>
<td><strong>Apply Cap and Reduce by 10 Percent</strong></td>
<td>$450.00</td>
</tr>
<tr>
<td>Calculate the allowed cost for the service (i.e., the cap amount) by taking the difference between the TEFRA and DRG payable amounts (in this example, $600.00).Take the lesser of the day outlier or the cap amount (in this example it would be the day outlier of $500.00).Reduce by 10 percent.</td>
<td></td>
</tr>
<tr>
<td><strong>Calculation1: Cost Threshold</strong></td>
<td>$550.00</td>
</tr>
<tr>
<td>11.14 x Universal Mean ($6,505.52) = &lt;amount A&gt;</td>
<td></td>
</tr>
<tr>
<td>11.14 x SDA = &lt;amount B&gt;</td>
<td></td>
</tr>
<tr>
<td>1.5 x DRG Relative Weight x SDA = &lt;amount C&gt;</td>
<td></td>
</tr>
<tr>
<td>Cost threshold = The greater of &lt;amount C&gt; compared with (the lesser of &lt;amount A&gt; and &lt;amount B&gt;)</td>
<td></td>
</tr>
<tr>
<td><strong>Calculation2: Cost Outlier</strong></td>
<td>$495.00</td>
</tr>
<tr>
<td>Allowed amount x reimbursement rate = TEFRA amount</td>
<td></td>
</tr>
<tr>
<td>TEFRA amount - cost threshold x 0.6 percent = cost outlier amount</td>
<td></td>
</tr>
<tr>
<td>Reduce by 10 percent.</td>
<td></td>
</tr>
</tbody>
</table>

The calculations in this example would result in one of the following payments:

- If the claim qualifies for the day outlier payment only, payment will be made up to $450.00.
- If the claim qualifies for the cost outlier payment only, payment will be made up to $495.00.
- If the claim qualifies for both the day outlier and cost outlier payment, the payment will be made up to $495.00, which is the greater of the day outlier or the cost outlier payment.

### 3.7.4.5 Neonatal Level of Care Designation for Inpatient Services

Hospitals enrolled in Texas Medicaid may be reimbursed for inpatient neonatal services only if the hospitals have received a neonatal level of care designation from DSHS in accordance with Title 25 Texas Administrative Code §§133.181-133.190.

A neonatal service is any inpatient hospital service rendered to a client who is 28 days of age or younger.

*Refer to:* The [DSHS website](#) for more information on Neonatal Level of Care Designation.

### 3.7.4.6 Hospitals that Do Not Meet Minimum Requirements for Neonatal Level of Care Designation

A hospital that does not meet the minimum requirements for any level of care designation for neonatal services will not be reimbursed for inpatient neonatal services rendered to Texas Medicaid and CSHCN Services Program clients. Hospitals without a neonatal level of care designation may be reimbursed for emergency services to stabilize an infant prior to transport to a facility capable of providing the appropriate level of care.

Claims for inpatient neonatal services submitted by hospitals that do not have a neonatal level of care designation on file will be denied. Providers can appeal claims by providing documentation that emergency services were required.
If neonatal inpatient services are rendered by a facility that has applied for (but not yet received) a neonatal designation, the facility must still adhere to existing claim filing deadlines (95 days from the date of discharge). While awaiting neonatal level of care designation the facility is responsible for maintaining active claims appeals to adhere to the 120-day claim appeal deadline.

Requirements to obtain a neonatal level of care designation only apply to facilities located in Texas. Those entities that are physically located outside of Texas and enrolled in Texas Medicaid (i.e., out-of-state or border state facilities) are exempt from requiring a neonatal level of care designation for inpatient services rendered to neonatal clients.

**Note:** When submitting paper claims for inpatient neonatal services rendered at a facility with an address that is different from the provider’s physical address, providers must enter the address of the facility where services were rendered in the remarks field.

**Refer to:** “Section 7: Appeals” (Vol. 1, General Information) for more information.

### 3.7.4.7 Other Requirements

The submitted facility address on the claim must match the physical address of the location that has been issued a neonatal level of care designation. If the facility address is not included on the claim, the submitted billing address must match the physical address of the location that was issued a neonatal level of care designation.

**Important:** The hospital address on the health facilities license must match the address billed on the claim. Claims will be denied if the address submitted on the claim does not match the address on file. Providers should refer to the DSHS approval letter to verify the correct address.

**Refer to:** The [DSHS website](https://www.dshs.state.tx.us) for more information on address updates.

### 3.7.4.8 Transfers

When Texas Medicaid or CSHCN Services Program clients are 28 days of age and younger on the date of admission and are subsequently transferred to another facility, neonatal level of care designation requirements will apply to all facilities involved in that client’s continuous inpatient stay.

### 3.7.4.9 Texas Provider Identifier Change Due to Split or Merge

Hospital providers with a Texas Provider Identifier (TPI) change that is due to a split or merge are responsible for notifying DSHS. Neonatal level of care designation providers must notify DSHS of any address changes.

### 3.7.4.10 Crossover Claims for Dual Eligible Clients

Neonatal level of care designation requirements will also apply to services rendered to a client who is less than 28 days old and has dual Medicare and Medicaid eligibility on the date of admission.

### 3.7.4.11 Children’s Hospitals

Children’s hospitals are reimbursed using the prospective payment methodology based on APR-DRG methodology.

With the exception of designated children’s hospitals, hospitals that are reimbursed by APR-DRG payment methodology receive one SDA rate.

Designated children’s hospitals receive two SDA rates:

- One rate for obstetric delivery services rendered to clients who are 18 years of age and older.
- One rate for all other services rendered to clients who are 18 years of age and older and all services rendered to clients who are 17 years of age and younger, including obstetric delivery services.
3.7.4.12 Potentially Preventable Complications (PPC) and Potentially Preventable Readmissions (PPR)

Potentially Preventable Complications (PPCs)

By definition, potentially preventable complications (PPCs) are harmful events or negative outcomes that develop after hospital admission and may result from processes of care and treatment rather than from the natural progression of the underlying illness. A PPC is an inpatient hospital complication that was potentially preventable based on criteria such as hospital characteristics, reason for admission, procedures, and the interrelationships between underlying medical conditions.

S.B. 7, Chapter 526, the 82nd Texas Legislature, 2011, establishes the authority of HHSC to identify PPCs in the Medicaid population. HHSC must confidentially report the results to each hospital that serves Texas Medicaid clients, and each of those hospitals must distribute the information to its care providers.

HHSC also produces a public version of the report, which does not specifically identify any of the hospitals. A statewide average PPC rate is calculated for all hospitals within Texas. Each hospital has an individual rate. Hospitals are able to compare their rate of PPC to the statewide average.

The PPC analysis is performed in accordance with TAC, §354.1446 Potentially Preventable Complications.

Potentially Preventable Readmission (PPRs)

By definition, potentially preventable readmissions (PPRs) are return hospitalizations of a person within a period specified by HHSC that results from deficiencies in the care or treatment provided to the person during a previous hospital stay or from deficiencies in post-hospital discharge follow-up.

Texas Medicaid uses a 15 day readmission interval.

Section 531.913, House Bill (H.B.) 1218, 81st Legislature, 2009, requires the HHSC to identify PPRs in the Medicaid population. HHSC must confidentially report the results to each hospital that serves Texas Medicaid clients, and each of those hospitals must distribute the information to its care providers.

HHSC delivers an annual, confidential report of the results to each hospital that is enrolled in Texas Medicaid, and each of those hospitals must distribute the information to their care providers. HHSC also produces a public version of the report, which does not specifically identify any of the hospitals. Patients are never identified in the reports.

PPR Analysis

The PPR analysis is performed in accordance with TAC, §354.1445 Potentially Preventable Readmissions.

Refer to: The HHS website at hhs.texas.gov/about-hhs/process-improvement/improving-services-texans/medicaid-chip-quality-efficiency-improvement/potentially-preventable-events for more information on Potentially Preventable Events (PPE).

3.7.4.13 State-owned Teaching Hospitals

Inpatient hospital stays in designated state-owned teaching hospitals are reimbursed according to the TEFRA payment methodology.

State-owned teaching hospitals are defined specifically in 1 TAC §355.8052 as the following hospitals: University of Texas Medical Branch (UTMB); University of Texas Health Center Tyler; and M.D. Anderson Hospital.
3.7.4.14 Payment Window Reimbursement Guidelines

3.7.4.14.1 Guidelines for Services Preceding an Inpatient Admission

The following payment window reimbursement guidelines apply to services that are rendered by the hospital or an entity that is wholly owned or operated by the hospital. The three-day (or one-day) payment window does not apply if:

- The hospital and the physician office or other entity are both owned by a third party, such as a health system.
- The hospital is not the sole or 100 percent owner of the entity.

Texas Medicaid inpatient hospital providers must submit, as part of the client’s inpatient hospital claim, all related professional and outpatient services that were rendered on the date of the client’s inpatient admission or one of the following dates immediately before admission:

- Within three calendar days before the client’s inpatient admission for hospitals that receive DRG reimbursement, with the exception of children’s hospitals.
- Within one calendar day before the client’s inpatient admission for hospitals that receive reimbursement other than DRG.
- Within one calendar day before the client’s inpatient admission for children’s hospitals.

Professional and outpatient services that must be submitted as part of the inpatient hospital claim include the following services if they are rendered by the hospital or an entity that is wholly owned or operated by the hospital:

- Diagnostic services. Diagnostic services include outpatient laboratory and radiology services that are related to the inpatient admission and submitted by physician and outpatient hospital providers. Affected services will include the total and technical components. The professional interpretation component will not be included in the payment windows identified above.
- Non-diagnostic services. Non-diagnostic services include surgeries and other non-diagnostic procedures and services that are related to the inpatient admission and submitted by physician, outpatient hospital, or other providers.

**Important:** Related professional and outpatient services that were rendered within the specified time frames must be submitted on the inpatient hospital claim and not on an outpatient hospital claim. An outpatient hospital claim for these services will be denied as part of the payment for the inpatient hospital stay.

3.7.4.14.2 Exceptions

The following services are excluded from the payment window and may be submitted and reimbursed separately from the inpatient admission:

- Services rendered by federally qualified health center (FQHC) providers
- Services rendered by rural health clinic (RHC) providers
- Professional services that are rendered in the inpatient hospital setting (place of service 3)
- Non-emergency and emergency ambulance services

The outpatient emergency and maintenance renal dialysis procedure codes in the tables below are also exceptions to the one-day payment window reimbursement guidelines:

<table>
<thead>
<tr>
<th>Procedure Codes</th>
<th>ESRD Physician Services</th>
</tr>
</thead>
<tbody>
<tr>
<td>G0257</td>
<td></td>
</tr>
</tbody>
</table>
3.7.4.14.3 Professional and Outpatient Claims for Services Related to the Inpatient Admission

Professional and outpatient services that are rendered on the date of admission, or within one of the one-day or three-day timeframes indicated above by the hospital or an entity that is wholly owned or operated by the hospital, are considered part of the inpatient stay. Professional and outpatient claims submitted for services that are related to the inpatient admission will be denied or recouped if they are submitted with the specified payment window.

When modifier PD is appended to a professional or outpatient service, the modifier indicates that the service is related to the inpatient admission. The total and technical components for professional and outpatient services that are related to the inpatient admission will be denied when submitted with modifier PD.

*Note:* The professional interpretation component for professional and outpatient services that are related to the inpatient stay may be reimbursed separately even if accompanied by PD modifier.

3.7.4.14.4 Professional and Outpatient Claims for Services Unrelated to the Inpatient Admission

Professional and outpatient services that are rendered within the specified timeframe by the hospital or an entity that is wholly owned or operated by the hospital may be reimbursed if they are identified as unrelated to the inpatient admission as follows:

- Professional and outpatient claims for diagnostic services that are unrelated to the inpatient admission must be submitted with modifier U4, which indicates the service is unrelated to the inpatient admission.
- Professional claims for non-diagnostic services that are unrelated to the inpatient admission will be identified by comparing the referenced diagnosis code that is on the professional claim to the principal inpatient diagnosis. Professional services must be submitted with modifier U4 if the services are unrelated and up to six digits of the referenced professional diagnosis code match the principle inpatient diagnosis code.
• Outpatient claims for non-diagnostic services that are unrelated to the inpatient admission will be identified by comparing the referenced diagnosis code that is on the outpatient claim to the principal inpatient diagnosis. The outpatient services must be submitted with condition code 51 if the services are unrelated and up to six digits of the referenced outpatient diagnosis code match the principle inpatient diagnosis code.

Unrelated services that are denied as part of the inpatient admission can be appealed with modifier U4 or condition code 51, which indicates that the service is unrelated to the inpatient admission.

**Note:** Claims that are submitted with modifier U4 or condition code 51 will be subject to retrospective review and may be recouped if there is not sufficient documentation to indicate the service was unrelated to the inpatient admission.

These benefit changes do not impact services rendered by providers that are not wholly owned or operated by the hospital.

### 3.7.4.15 Potentially Preventable Readmissions (PPR)

H.B. 1218, 81st Legislature, Regular Session 2009, requires that HHSC identify potentially preventable readmissions (PPRs) in the Medicaid population and report results confidentially to each hospital. The law also requires each hospital to distribute the information to its care providers.

### 3.7.5 Provider Cost and Reporting

The method of determining reasonable cost is similar to that used by Title XVIII (Medicare). Hospitals must include inpatient and outpatient costs in the cost reports submitted annually. The provider must prepare one copy of the applicable CMS Cost Report Form along with the required PCCM supplemental worksheets. The PCCM supplemental worksheets include the Inpatient PCCM D-4 worksheet, available from CMS, and the Outpatient PCCM D, Part V worksheet. A sample of the Outpatient PCCM D, Part V is available on the TMHP website at [www.tmhp.com](http://www.tmhp.com).

**Refer:** Subsection 2.2.2, “Cost Reimbursement” in “Section 2: Texas Medicaid Fee-for-Service Reimbursement” (*Vol. 1, General Information*).

If a change of ownership or provider termination occurs, the cost report is due within five months after the date of the change in ownership or termination. Any request for an extension of time to file must be made on or before the cost report due date and sent to TMHP Medicaid Audit at the address indicated under “Written Communication With TMHP” in the “Appendix A: State, Federal, and TMHP Contact Information” (*Vol. 1, General Information*). For questions or assistance, call TMHP Medicaid Audit at 1-512-514-3648.

Annual cost reports must be filed as follows:

- Submit one copy of the cost report to TMHP Medicaid Audit within five months of the end of the hospital’s fiscal year along with any amount due to Texas Medicaid.
- TMHP Medicaid Audit performs a desk review of the cost report and makes a tentative settlement with the hospital. A tentative settlement letter requests payment for any balance due to Texas Medicaid or instructs TMHP to pay the amount due to the provider. Interim payment rates are changed at this time based on the cost report.
- Field audits are conducted when necessary.
- Medicaid final settlement is made after a copy of all the following information is received from the provider or the Medicare intermediary. The provider must send TMHP a copy of one of the following:
  - Audited or settled without audit Medicare Cost Report
  - Medicare Notice of Amount of Program Reimbursement
  - Medicare Audit Adjustment Report, if applicable
Medicaid hospitals may request copies of their claim summaries for their cost reporting fiscal year. The summaries for tentative settlements include three additional months of claim payments for the fiscal year. The summaries for final settlements include ten months of claim payments for the fiscal year. TMHP Medicaid Audit uses this data to determine the tentative and final settlements and interim rates.

The Medicaid claim summary data are only generated once each month, and the logs are received by the 15th of the following month. Requests for tentative settlement logs are submitted within 30 days after the fiscal year-end. Final settlement log requests are submitted within nine months after the fiscal year-end.

The Medicaid logs can be requested through the provider's administrator account on the TMHP website at www.tmhp.com. Medicaid logs can also be requested by calling 1-512-506-6117 or by sending a written request to the following address:

Texas Medicaid & Healthcare Partnership
Medicaid Audit
PO Box 200345
Austin, TX 78720-0345

Allow 45 days for receipt of these logs.

3.7.6 Third Party Liability

Refer to: “Section 8: Third Party Liability (TPL)” (Vol. 1, General Information).

Other Insurance Form on the TMHP website at www.tmhp.com.

Tort Response Form on the TMHP website at www.tmhp.com.

4 Outpatient Hospital (Medical and Surgical Acute Care Outpatient Facility)

This section contains benefit, limitation, authorization, and claims filing information for outpatient hospital facility emergency, observation, and other services.

Refer to: “Section 6: Claims Filing” (Vol. 1, General Information) for more comprehensive information about claims filing and appeals.

“Section 7: Appeals” (Vol. 1, General Information) for more comprehensive information about claims filing and appeals.

Hospital providers are encouraged to review the other handbooks for applicable information, prior authorization requirements, and for specific requirements for special programs.

4.1 General Information

Outpatient diagnostic, therapeutic, and surgical services that are rendered in an acute care hospital setting are services that are provided to clients by or under the direction of a physician.

Outpatient hospital services include those services that are rendered:

- In the emergency room (ER)
- As day surgery
- In the observation room
- By ancillary departments such as the laboratory, radiology, physical or occupational therapy, cardiac rehabilitation, hyperbaric chamber, infusion services, and other areas able to provide services in the outpatient setting.
4.1.1 Drugs and Supplies

4.1.1.1 Self-Administered Drugs
Self-administered drugs are defined as drugs that the client administers themselves at home and may include, but are not limited to, prescription drugs, vitamins, and supplements.

These drugs that are provided by the hospital during an outpatient hospital visit are included in the hospital reimbursement and are not reimbursed separately. The client cannot be billed for self-administered drugs that are provided by the hospital during an outpatient hospital stay.

4.1.1.2 Take-Home Drugs and Supplies
Benefits do not include drugs and biologicals provided by the hospital and taken home by the client. Supplies provided by a hospital for use in physicians’ offices are not reimbursable.

Take-home drugs and supplies are a benefit for services rendered to clients in the outpatient setting when supplied by prescription through the VDP.

4.1.2 Outpatient Services Provided Without Charge
Texas Medicaid pays the clinic registration fee in lieu of other benefits when a hospital provides outpatient services without charge, and if the registration fee is less than the allowed Medicaid payment.

Refer to: TAC Rule §354.1073 for information about authorized outpatient hospital services.
Subsection 1.12, “Texas Medicaid Limitations and Exclusions” in “Section 1: Provider Enrollment and Responsibilities” (Vol. 1, General Information) for more information about noncovered items or services.

4.1.3 Payment Window Reimbursement Guidelines for Services Preceding an Inpatient Admission
According to the three-day and one-day payment window reimbursement guidelines, most professional and outpatient diagnostic and nondiagnostic services that are rendered within the designated timeframe of an inpatient hospital stay and are related to the inpatient hospital admission will not be reimbursed separately from the inpatient hospital stay if the services are rendered by the hospital or an entity that is wholly owned or operated by the hospital.

Refer to: Subsection 3.7.4.14, “Payment Window Reimbursement Guidelines” in this handbook for additional information about the payment window reimbursement guidelines.

4.2 Services, Benefits, Limitations, and Prior Authorization

4.2.1 Prior Authorization Requirements
The hospital is responsible for requesting prior authorization for the non-emergency transport to the client’s home or to a nursing home after a non-scheduled outpatient visit.


4.2.2 Emergency Department Services
An emergency department is defined as an organized hospital-based facility for the provision of unscheduled episodic services to clients who present for immediate medical attention. The facility must be available 24 hours a day, 7 days a week.

Hospital-based emergency departments are reimbursed for services based on a reasonable cost, based on the hospital’s most recent tentative Medicaid cost report settlement. The reasonable cost is reduced by a percentage determined by the state.
All claims that are submitted by outpatient hospital providers must include a procedure code with each revenue code for services that are rendered to Texas Medicaid clients. This procedure code must be listed on the same claim detail line as the emergency department revenue code.

The procedure code billed may include, but is not limited to, E/M, surgical or other procedure, or any other service rendered to the client in the emergency room. The procedure code must accurately reflect the services rendered in the hospital’s emergency department.

Emergency department reimbursement may include room changes and ancillary changes. Emergency department room charges may be submitted using the following revenue codes:

<table>
<thead>
<tr>
<th>Revenue Code</th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td>450</td>
<td>Emergency room</td>
</tr>
<tr>
<td>451</td>
<td>Emergency room-EMTALA emergency medical screening</td>
</tr>
<tr>
<td>456</td>
<td>Emergency room, urgent care</td>
</tr>
<tr>
<td>459</td>
<td>Emergency room, other</td>
</tr>
</tbody>
</table>

Emergency department ancillary services include, but are not limited to, the following:
- Laboratory services
- Radiology services
- Respiratory therapy services
- Diagnostic studies (including, but not limited to, ECGs, computed tomography (CT) scans, and supplies)

The administration of an injection may be reimbursed to the provider who administers the injection. The administration of the injection will not be reimbursed to outpatient hospital providers. An injection or infusion administered by a nurse is included in the emergency room charge and is not reimbursed separately to the outpatient facility.

Ancillary services must be submitted on the UB-04 CMS-1450 paper claim form using the appropriate procedure codes or revenue codes for rendered services.

If a client visits the emergency room more than once in one day, the times must be given for each visit.

If the client ultimately is admitted as an inpatient within 48 hours of treatment in the ER or clinic, the ER or clinic charges must be submitted on the inpatient hospital claim form as an ancillary charge. The date of inpatient admission is the date the client initially was seen in the ER or clinic.

According to the Emergency Medical Treatment and Active Labor Act (EMTALA) of 1986, if any individual presents at the hospital’s emergency department requesting an examination or treatment, the hospital must provide an appropriate medical screening examination and stabilization services within the capability of the hospital’s emergency department, including ancillary services routinely available to the emergency department, to determine whether an emergency medical condition exists.

EMTALA medical screening code (451) may be considered for reimbursement when submitted as a stand-alone service and provided by a qualified medical professional as designated by the facility. Ancillary, professional, or facility services will not be considered for separate reimbursement. Services beyond screening (451) can be submitted with the appropriate corresponding emergency services code (450).

Medicaid claims administrators are prohibited from requiring prior authorization or primary care provider notification for emergency services including those needed to evaluate or stabilize an emergency medical condition or emergency behavioral health condition.
Texas Medicaid provides that certain undocumented aliens and legalized aliens who require treatment of an emergency medical condition or emergency behavioral health condition are eligible to receive that treatment. After the emergency condition requiring care is stabilized and is no longer an emergency, the coverage ends. If the alien continues to receive ongoing treatment after the emergency ceases, the ongoing treatment is not a benefit.

Texas Medicaid provides for medical services for eligible clients while out-of-state. The attending physician or other provider must document that the client was treated for an emergency condition. Out-of-state emergency services are also a benefit when the client’s health would be in danger if he or she were required to travel back to Texas.

Emergency department services are subject to retrospective review.

In instances of sudden illness or injury, the client may receive treatment in the ER and be discharged, placed on observation status, or admitted as an inpatient.

4.2.2.1 Emergency Department Payment Reductions

Nonemergent and nonurgent evaluation and management (E/M) services rendered in the emergency room may be reimbursed at 125 percent of the adult, physician office visit fee for procedure code 99202. Reimbursement is based on the E/M procedure code submitted on the same line item as the emergency room revenue code.

Imaging services rendered by outpatient hospital providers are reimbursed at the flat fee that is based on the procedure code submitted on the same line item as the imaging revenue code.

**Note:** Evaluation and management services that are rendered in the emergency room for critically ill or critically injured Texas Medicaid clients of any age, are not subject to reduction in payment.

**Exception:** Rural hospitals, nonemergent and nonurgent E/M services rendered in the emergency room may be reimbursed at 65 percent of the allowed rate.

4.2.3 Day Surgery

Inpatients may occasionally require a surgery that has been designated as an outpatient procedure. The physician must document the need for this surgery as an inpatient procedure before the procedure is performed. These claims are subject to retrospective review.

These procedures are for clients who are scheduled for a day surgery procedure and are not inpatient at the time the day surgery is performed.

4.2.3.1 Inpatient Admissions for Day Surgeries

If a client is admitted for a day surgery procedure—whether scheduled or emergency—one of the following classifications may be considered an inpatient procedure.

- ASA Classification of Physical Status of III (P3), IV (P4), or V (P5)
- Classification of Heart Disease IV

The day surgery services must be submitted on an inpatient claim (TOB 111) using the hospital’s provider identifier. The reason for the surgery (principal diagnosis), any additional substantiated conditions, and the procedure must be included on one inpatient claim.

**Refer to:** The Anesthesia standards at [www.asahq.org/standards-and-guidelines/asa-physical-status-classification-system](http://www.asahq.org/standards-and-guidelines/asa-physical-status-classification-system) for a description of the ASA classes of physical status.

The descriptions for ASA classes of physical status are as follows:

- **Class I.** A normal healthy patient, without organic, physiological, or psychiatric disturbance.

  **Example:** Healthy patient with good exercise tolerance.
• **Class II.** A patient with mild systemic disease, controlled medical conditions without significant systemic effects.

  **Example:** Controlled hypertension or diabetes mellitus without system effects, cigarette smoking without evidence of chronic obstructive pulmonary disease (COPD), anemia, mild obesity, age less than 1 or greater than 70 years, or pregnancy.

• **Class III.** A patient exhibiting severe systemic disturbance that may or may not be associated with the surgical complaint and that seriously interferes with the patient’s activities.

  **Example:** Severely limiting organic heart disease, severe diabetes with vascular complications; moderate to severe degrees of pulmonary insufficiency; angina pectoris or healed myocardial infarction.

• **Class IV.** A patient exhibiting extreme systemic disturbance that may or may not be associated with the surgical complaint, that interferes with the patient’s regular activities, and that has already become life-threatening.

  **Example:** Organic heart disease with marked signs of cardiac insufficiency present (for example, cardiac decompensation); persistent anginal syndrome, or active myocarditis; advanced degrees of pulmonary, hepatic, renal, or endocrine insufficiency present.

• **Class V.** The rare person who is moribund (in a dying state) before operation, whose pre-operative condition is such that he or she is expected to die within 24 hours even if not subjected to the additional strain of operation.

  **Example:** Burst abdominal aneurysm with profound shock; major cerebral trauma with rapidly increasing intracranial pressure; massive embolus.

The Classification of Heart Disease consists of four classes:

• **Class I.** No limitation of physical activity. Ordinary physical activity does not cause undue fatigue, palpitation, dyspnea, or anginal pain.

• **Class II.** Slight limitation of physical activity. Comfortable at rest, but ordinary physical activity results in fatigue, palpitation, dyspnea, or anginal pain.

• **Class III.** Marked limitation of physical activity. Comfortable at rest, but less than ordinary activity causes fatigue, palpitation, dyspnea, or anginal pain.

• **Class IV.** Unable to carry on any physical activity without discomfort. Symptoms of cardiac insufficiency, or of the anginal syndrome, may be present even at rest. If any physical activity is undertaken, discomfort occurs.

### 4.2.3.2 Complications Following Elective or Scheduled Day Surgeries

If a condition of the scheduled day surgery requires additional care beyond the recovery period, the client may be placed in outpatient observation (stay less than 48 hours). The observation period must be submitted on an outpatient claim (TOB 131) using the hospital’s provider identifier. If the client requires inpatient admission following the observation stay, the admission date for the inpatient claim is the date the client was placed in observation. All charges for services provided from the time of observation placement (excluding the surgical procedure) must be included on the inpatient claim (TOB 111) using the hospital’s provider identifier. The principal diagnosis to be used on the inpatient claim is the complication of the surgery that necessitated the extended stay. The day surgery procedure must still be submitted as an outpatient procedure under the HASC provider identifier.

### 4.2.3.3 Inpatient Admissions After Day Surgery

If a complication occurs for which the client requires inpatient admission immediately following the day surgery (no observation period), the day surgery must be submitted as an outpatient procedure (TOB 131), using the appropriate hospital or HASC provider identifier. The inpatient admission is to be
submitted as an inpatient claim (TOB 111), using the hospital’s provider identifier. The principal
diagnosis to be used on the inpatient claim is the complication of the surgery that necessitated the
extended stay. The day surgery procedure must not be included on the inpatient claim. The inpatient
admission must be medically necessary and is subject to retrospective review.

4.2.3.4 Emergency or Unscheduled Day Surgeries

These procedures are for clients who require an unscheduled (emergency) day surgery procedure and
are not inpatient at the time the day surgery is performed.

If a client is first treated in the ER and then requires emergency surgery as an outpatient, claims for
emergency, unscheduled outpatient surgical procedures must be filed itemizing each service, such as
room charge, laboratory, radiology, anesthesia, and supplies. Providers must submit claims for
unscheduled day surgery procedures and emergency services as outpatient procedures using the hospital
provider identifier. If a condition of the unscheduled day surgery requires additional care beyond the
recovery period, the client may be placed on outpatient observation status. The observation period must
be submitted on the same outpatient claim.

Providers must submit claims for the unscheduled day surgery procedures and emergency services as
outpatient procedures (TOB 131) using the hospital’s provider identifier. If a condition of the
unscheduled day surgery requires additional care beyond the recovery period, the client may be placed
on outpatient observation status (stay less than 48 hours). The observation period must be submitted on
the same outpatient claim (TOB 131) using the hospital’s provider identifier.

4.2.3.5 Complications Following Emergency or Unscheduled Day Surgery

If the client requires inpatient admission following the observation stay, the admission date for the
inpatient claim is the date the client was placed in observation. All charges for services provided from
the time of observation status (excluding surgical procedures and emergency services) must be included
on the inpatient claim (TOB 111) using the hospital’s provider identifier. The principal diagnosis to be
used on the inpatient claim is the complication of the surgery that necessitated the extended stay. The
day surgery and emergency services must not be included on the inpatient claim since they are to be
submitted using TOB 131 as outpatient procedures under the hospital’s provider identifier.

4.2.3.6 Incomplete Day Surgeries

Facilities must use either one of the following diagnosis codes or one of the following modifiers to
indicate that a surgical procedure (type of service F) was not completed:

<table>
<thead>
<tr>
<th>Diagnosis Codes</th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td>Z5309</td>
<td>Procedure and treatment not carried out because of other contraindication</td>
</tr>
<tr>
<td>Z5329</td>
<td>Procedure and treatment not carried out because of patient’s decision for other reasons</td>
</tr>
<tr>
<td>Z538</td>
<td>Procedure and treatment not carried out for other reasons</td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th>Modifier</th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td>73</td>
<td>Discontinued outpatient procedure prior to anesthesia administration</td>
</tr>
<tr>
<td>74</td>
<td>Discontinued outpatient procedure after anesthesia administration</td>
</tr>
</tbody>
</table>

Providers must submit the following documentation with the claim:

- The operative report
- The anesthesia report
- The reason that the operation was not complete
Reimbursement to HASC facilities for canceled or incomplete surgeries because of patient complications, is made according to the following criteria, depending on the extent to which the anesthesia or surgery proceeded:

- Reimburse at 0 percent of HASC group payment schedule for a procedure that is terminated for nonmedical or medical reasons before the facility has expended substantial resources.
- Reimburse at 33 percent of HASC group payment schedule up to the administration of anesthesia.
- Reimburse at 67 percent of HASC group payment schedule after the administration of anesthesia but before incision.
- Reimburse at 100 percent of HASC group payment schedule after incision.

Surgeries canceled because of incomplete pre-operative procedures are not reimbursed.

### 4.2.4 Outpatient Observation Room Services

Observation care is defined by the CMS as “a well-defined set of specific, clinically appropriate services, which include ongoing short term treatment, assessment, and reassessment, that are furnished while a decision is being made regarding whether clients will require further treatment as hospital inpatients or if they are able to be discharged from the hospital.”

Outpatient observation services are usually ordered for clients who present to the emergency department and who then require a significant period of treatment or monitoring in order to make a decision about their admission or discharge. The decision whether to discharge a client from the hospital following resolution of the reason for the observation care or to admit the client as an inpatient can be made in less than 48 hours, usually in less than 24 hours.

Outpatient observation services require the use of a hospital bed and periodic monitoring by the hospital’s nursing or other ancillary staff to evaluate the client’s condition and to determine the need for an inpatient admission. Outpatient observation services can be provided anywhere in the hospital. The level of care, not the physical location of the bed, dictates the observation status.

Outpatient observation services (revenue codes 760, 761, 762, and 769) are a benefit only when medically necessary and when provided under a practitioner’s order or under the order of another person who is authorized by state licensure law and hospital bylaws to admit clients to the hospital and to order outpatient services.

Outpatient observation services are considered medically necessary if the following conditions are met (this list is not all-inclusive):

- The client is clinically unstable for discharge and one of the following additional conditions apply:
  - Laboratory, radiology, or other testing is necessary to assess the client’s need for an inpatient admission.
  - The treatment plan is not established or, based on the client’s condition, is anticipated to be completed within a period not to exceed 48 hours.
  - The client had a significant adverse response to therapeutic services, invasive diagnostic testing, or outpatient surgery and requires short-term monitoring or evaluation.
- The medical necessity for inpatient treatment is unclear, that is:
  - The client’s medical condition requires careful monitoring and evaluation, or treatment to confirm or refute a diagnosis in order to determine whether an inpatient admission is necessary
  - There is a delayed or slow progression of the client’s signs and symptoms that makes diagnosis difficult and the monitoring or treatment does not meet the criteria for an inpatient level of care.
  - The client is undergoing treatment for a diagnosed condition, and continued monitoring of clinical response to therapy may prevent an inpatient admission.
• The admitting practitioner anticipates that the client will require observation care for a minimum of eight hours.

Medically necessary services that do not meet the definition of observation care should be submitted separately or included as part of the emergency department or clinic visit, and are not reimbursed as observation care.

Outpatient observation services are not a substitute for a medically appropriate inpatient admission. If a client meets the medical necessity criteria for an inpatient admission and an inpatient admission is ordered by the practitioner, an inpatient admission is a benefit regardless of the length of stay. Claims for observation services may be denied in their entirety if the services should have initially been inpatient admissions or if a reason for an inpatient admission developed, but the observation stay was not converted to inpatient.

The determination of an inpatient or outpatient status for any given client is specifically reserved to the admitting practitioner. The decision must be based on the practitioner’s expectation of the care that the client will require.

4.2.4.1 Direct Outpatient Observation Admission

A client may be directly admitted to outpatient observation from the evaluating practitioner’s office without being seen in the emergency room by a hospital-based practitioner. The practitioner’s order should clearly specify that the practitioner wants the client to be admitted to outpatient observation status. An order for “direct admission” will be considered an inpatient admission unless otherwise specified by the practitioner’s orders.

Brief observation periods following an office visit or at the direction of an off-site practitioner that involve a simple procedure (e.g., a breathing treatment) would be more appropriately coded as a treatment room visit.

4.2.4.2 Observation Following Emergency Room

A client may be admitted to outpatient observation through the emergency room if the client presents to the facility with an unstable medical condition and the evaluating practitioner determines that outpatient observation is medically necessary to determine a definitive treatment plan. An unstable medical condition is defined as one of the following:

• A variance in laboratory values from what is considered the generally accepted, safe values for the individual client.

• Clinical signs and symptoms that are above or below those of normal range and that require extended monitoring and further evaluation.

• Changes in the client’s medical condition are anticipated, and further evaluation is necessary.

If a client is admitted to observation status from the emergency room, the hospital is reimbursed only for the observation room charges. The emergency room charges are not reimbursed separately, but must be submitted on a separate detail on the same claim as the observation room charges.

Brief observation periods following an emergency room evaluation will not be reimbursed if the service would normally have been provided within the time frames and facilities of an emergency room visit.

4.2.4.3 Observation Following Outpatient Day Surgery

If a medical condition or complication of a scheduled day surgery requires additional care beyond the routine recovery period, the client may be placed in outpatient observation. The observation period should be submitted as an outpatient claim.
Reimbursement for outpatient observation after a scheduled day surgery is limited to situations in which the client exhibits an unusual reaction to the surgical procedure and requires monitoring or treatment beyond what is normally provided in the immediate post-operative period. Examples include, but are not limited to:

- Difficulty in awakening from anesthesia.
- A drug reaction.
- Other post-surgical complications.

### 4.2.4.4 Observation Following Outpatient Diagnostic Testing or Therapeutic Services

A client may be admitted to outpatient observation if the client develops a significant adverse reaction to a scheduled outpatient diagnostic test or to a therapeutic service, such as chemotherapy, that requires further monitoring. Observation services begin when the reaction occurred and end when the practitioner determines that the client is stable for discharge, or that an inpatient admission is appropriate.

### 4.2.4.5 Documentation Requirements for Outpatient Observation

Documentation that supports the medical necessity of the outpatient observation services must be maintained by the facility in the client’s medical record. Documentation must include:

- The order of the ordering practitioner for admission to observation care, which must be dated and timed.
- The practitioner’s admission and progress notes, which must be dated and timed, confirm the need for observation care, and outline the client’s condition, treatment, and response to treatment.
- Nurse’s notes, which must be dated and timed, reflect the time at which the client was admitted to the observation bed, and the reason for the observation stay.
- All supporting diagnostic and ancillary testing reports, including orders for the testing or any preadmission testing.
- Procedure notes and operative notes that address any complication that would support admission to observation status and must be dated and timed.
- Anesthesia and recovery room/post anesthesia care unit notes from the practitioner and the nurse, which must be dated and timed and detail orders and any complications that require admission to observation status.
- Documentation related to an outpatient clinic visit or critical care service that was provided on the same date of service as the observation service. The documentation must address any need for observation services and be dated and timed.
- All of the client education that was provided during the observation stay.
- The order for discharge from observation care, which must be signed, dated, and timed.
- The discharge notes, including nurse’s notes that reflect the date and time at which the client was discharged from observation.

The client must be in the care of a practitioner during the period of observation, as documented in the medical record by admission, discharge, and other appropriate progress notes that are dated, timed, written, and signed by the practitioner.

Claims submitted for outpatient procedures in which the original intention was to keep the client for an extended period of time, such as overnight or for a 48-hour period, will be denied unless significant medical necessity is documented.
Retrospective review may be performed to ensure that the documentation supports the medical necessity of the outpatient observation services. Medical records will be evaluated to determine whether the practitioner's order (practitioner intent) and the services that were actually provided were consistent.

The medical records must clearly support the medical necessity of the outpatient observation services and must include a timed order for observation services that will support the number of hours that the client was under observation care and the hours that were submitted for payment.

### 4.2.4.6 Reporting Hours of Observation

Providers must submit the number of observation hours the client was under observation care.

Observation time begins at the clock time documented in the client’s medical record. This time should coincide with the time that the client is placed in a bed for the purpose of initiating observation care in accordance with the practitioner’s order.

Observation time ends when all medically necessary services related to observation care are completed. The end time of observation services may coincide with the time the client is actually discharged from the hospital or is admitted as an inpatient.

Hospitals should round clock times for the beginning and end of observation to the nearest hour and submit the total number of hours for the observation stay on the claim. For the purposes of submitting claims for observation services, one unit equals one hour. Partial units or hours should be rounded up or down to the nearest hour. Claims submitted with observation room units exceeding 48 hours will be denied.

Any service that was ordered within the observation period may be included on the outpatient claim if a practitioner’s order for the service was made within the observation period time frame but hospital scheduling limitations prevented the service from being performed before the 48 hours expired. Any services ordered after 48 hours must not be included on the outpatient claim nor billed to the client. If a period of observation spans more than one calendar day (i.e., extends past midnight), all of the hours for the entire period of observation must be included on a single line, and the date of service for that line is the date on which the observation care began.

Observation time may include medically necessary services and follow-up care that is provided after the time the practitioner writes the discharge order, but before the client is discharged. Reported observation time does not include the time the client remains in the observation area after treatment is completed for reasons such as waiting for transportation home.

Observation services must not be submitted concurrently with diagnostic or therapeutic services for which active monitoring is part of the procedure. In situations where a diagnostic or therapeutic procedure interrupts the observation stay, hospitals should record for each period of observation services the beginning and ending times of the observation period and add the lengths of time for the periods of observation services together to reach the total number of units reported on the claim.

Recovery room hours that are associated with an outpatient procedure must not be submitted simultaneously with hours of observation time.

Revenue code 761 will be denied if it is submitted for the same date of service by the same provider as revenue code 760, 762, or 769.

### 4.2.4.7 Client Status Change

When a client’s status changes from outpatient observation to inpatient admission within the allowed 48-hour observation period, both the outpatient observation service and the inpatient admission must be submitted as separate details on the same inpatient claim.

The inpatient claim must include:

- The provider must submit the correct FDOS and Date of Admission (DOA) on the claim header.
• Charges for the observation room on the inpatient claim may be coded using the appropriate revenue code (760, 761, 762, or 769).

• The observation services are considered part of the facility’s DRG payment, and are not separately reimbursed.

The practitioner’s order for a change in client status from outpatient observation to inpatient admission must be written, dated, and timed before the outpatient observation claim is submitted for reimbursement.

4.2.4.8 Inpatient Admission to Outpatient Observation

When a client is admitted to the hospital as an inpatient and a subsequent internal utilization review (UR) determines that the services did not meet inpatient criteria, the hospital may change the client’s status from inpatient to outpatient observation. The order to change from an inpatient to outpatient observation admission is effective for the same date and time as the inpatient order. This practice is acceptable under Texas Medicaid if all of the following conditions are met:

• The change in client status is made before the claim is submitted.

• The hospital has not submitted a claim for the inpatient admission.

• The practitioner responsible for the care of the client concurs with the hospital UR determination to change to outpatient status.

• The practitioner’s concurrence with the UR decision is documented in the client’s medical record.

Note: When the hospital has determined that it may submit an outpatient claim according to the conditions described above, the entire episode of care should be billed as an outpatient episode of care.

Reimbursement for emergency room (ER) and observation services are considered part of the inpatient DRG payment and must be submitted as separate details on the inpatient claim when the client is admitted as an inpatient under one or both of the following circumstances:

• The client has spent fewer than 24 hours after presenting in the ER without being placed in observation status.

• The client has spent fewer than 48 hours in observation status after presenting in the ER.

The date of admission on the inpatient claim must reflect the date the client presents at the hospital. If the client is admitted as an inpatient more than 24 hours after presenting in the ER without being placed in observation status or more than 48 hours after being placed in observation status, the ER and observation services may be reimbursed separately as outpatient services.

Examples

The following examples indicate the appropriate dates of admission and claim submissions for different scenarios:

Scenario 1

In scenario 1, the ER and outpatient observation services must be submitted on the inpatient hospital claim, because the ER services are within 24 hours of the observation services, and the observation services are within 48 hours of the inpatient admission, and the client was not discharged and sent home before being admitted as an inpatient.

The inpatient admission date reflects the date the patient presented at the ER.

<table>
<thead>
<tr>
<th>Date and Time</th>
<th>Patient Services</th>
</tr>
</thead>
<tbody>
<tr>
<td>5/1/12 (11:50 p.m.)</td>
<td>Patient presents in the ER</td>
</tr>
</tbody>
</table>
Claims submissions are as follows:

- **ER visit**: Submitted on the inpatient claim as a separate detail (part of the DRG payment)
- **Observation services**: Submitted on the inpatient claim as a separate detail (part of the DRG payment)
- **Date of inpatient admission**: May 1, 2012

### Scenario 2

In scenario 2, the ER service was more than 24 hours before the observation period began and must be submitted on an outpatient hospital claim. The observation service must be billed on the inpatient hospital claim because the service was within 48 hours of the inpatient admission, and the client was not discharged and sent home before being admitted as an inpatient.

The inpatient admission date reflects the date the patient was placed in observation status.

<table>
<thead>
<tr>
<th>Date and Time</th>
<th>Patient Services</th>
</tr>
</thead>
<tbody>
<tr>
<td>5/1/12 (11:50 p.m.)</td>
<td>Patient presents in the ER</td>
</tr>
<tr>
<td>5/2/12 (11:55 p.m.)</td>
<td>24 + hours later, patient is placed in observation status</td>
</tr>
<tr>
<td>5/3/12 (4:00 a.m.)</td>
<td>4 hours later, patient admitted as an inpatient</td>
</tr>
</tbody>
</table>

Claims submissions are as follows:

- **ER visit**: Submitted on an outpatient claim and reimbursed separately from the observation and inpatient services
- **Observation services**: Submitted on the inpatient claim as a separate detail (part of the DRG payment)
- **Date of inpatient admission**: May 2, 2012

### Scenario 3

In scenario 3, the ER service must be submitted on an outpatient claim as part of the observation service because the ER service was within 24 hours of the observation service. The observation service may be reimbursed separately from the inpatient admission because the observation service was more than 48 hours before the inpatient admission, and the client was not discharged and sent home before being admitted as an inpatient.

The inpatient admission date reflects the date the patient was admitted as an inpatient.

<table>
<thead>
<tr>
<th>Date and Time</th>
<th>Patient Services</th>
</tr>
</thead>
<tbody>
<tr>
<td>5/1/12 (11:50 p.m.)</td>
<td>Patient presents in the ER</td>
</tr>
<tr>
<td>5/2/12 (12:30 a.m.)</td>
<td>40 minutes later, patient placed in observation status</td>
</tr>
<tr>
<td>5/4/12 (12:45 a.m.)</td>
<td>48 + hours later patient admitted as an inpatient</td>
</tr>
</tbody>
</table>

Claims submissions are as follows:

- **ER visit**: Submitted on an outpatient claim and reimbursed as part of the outpatient observation services
- **Observation services**: Submitted on the outpatient claim and reimbursed separately from the inpatient services

- **Date of inpatient admission**: May 4, 2012

**Scenario 4**

In scenario 4, the ER service may be reimbursed separately because it was more than 24 hours before the client was placed in observation status. The observation service may be reimbursed separately because it was more than 48 hours before the client was admitted as an inpatient.

The inpatient admission date reflects the date the patient was admitted as an inpatient.

<table>
<thead>
<tr>
<th>Date and Time</th>
<th>Patient Services</th>
</tr>
</thead>
<tbody>
<tr>
<td>5/1/12 (11:50 p.m.)</td>
<td>Patient presents in the ER</td>
</tr>
<tr>
<td>5/2/12 (11:55 p.m.)</td>
<td>24 + hours later, patient is placed in observation status</td>
</tr>
<tr>
<td>5/4/12 (12:00 p.m.)</td>
<td>48 + hours later, patient admitted as an inpatient</td>
</tr>
</tbody>
</table>

Claims submissions are as follows:

- **ER visit**: Submitted on an outpatient claim and reimbursed separately from the observation and inpatient services.

- **Observation services**: Submitted on an outpatient claim and reimbursed separately from the inpatient services.

- **Date of inpatient admission**: May 4, 2012

**Scenario 5**

In scenario 5, the ER service must be submitted on an outpatient claim as part of the observation service because the ER service was within 24 hours of the observation service. The observation service may be reimbursed separately from the inpatient admission because the client was discharged and sent home without being admitted as an inpatient.

The inpatient admission date reflects the date the patient presented at the ER after being discharged and sent home 14 hours earlier.

<table>
<thead>
<tr>
<th>Date and Time</th>
<th>Patient Services</th>
</tr>
</thead>
<tbody>
<tr>
<td>5/1/12 (11:50 p.m.)</td>
<td>Patient presents in the ER</td>
</tr>
<tr>
<td>5/2/12 (12:30 a.m.)</td>
<td>40 minutes later, patient is placed in observation status</td>
</tr>
<tr>
<td>5/2/12 (10:00 a.m.)</td>
<td>9.5 hours later, patient is discharged and sent home</td>
</tr>
<tr>
<td>5/3/12 (12:05 a.m.)</td>
<td>14 hours later, patient presents at the ER again and is admitted as an inpatient</td>
</tr>
</tbody>
</table>

Claims submissions are as follows:

- **ER visit**: Submitted on an outpatient claim and reimbursed as part of the observation services

- **Observation services**: Submitted on the outpatient claim and reimbursed separately from the inpatient services

- **Date of inpatient admission**: May 3, 2012

When the hospital has determined that it may submit an outpatient claim according to the conditions described above, the entire episode of care should be submitted as an outpatient episode of care.
4.2.4.9 Observation Services that are not a benefit
Outpatient observation services that are not medically necessary or appropriate are not benefits of Texas Medicaid, including, but not limited to, services provided under the following circumstances:

- As a substitute for an inpatient admission.
- Without a practitioner’s order, including services ordered as inpatient services by the ordering practitioner, but submitted as outpatient by the billing office.
- For clients awaiting transfer to another facility, such as for nursing home placement.
- For clients with lack of or delay in transportation.
- As a convenience to the client, client’s family, the practitioner, hospital, or hospital staff.
- For routine preparation before, or recovery after, outpatient diagnostic or surgical services.
- When an overnight stay is planned before diagnostic testing.
- To medically stable clients who need diagnostic testing or outpatient procedures that are routinely provided in an outpatient setting.
- Following an uncomplicated treatment or procedure.
- As standing orders for observation following outpatient surgery.
- For postoperative monitoring during a standard recovery period of four to six hours, which is considered part of the recovery room service.
- For outpatient blood or chemotherapy administration and concurrent services.
- For services that would normally require an inpatient admission.
- Beyond 48 hours from the time of the observation admission.
- For a medical examination for clients who do not require skilled support.

4.2.5 Hospital-Based Rural Health Clinic Services
Hospital-based RHCs must use the encounter code T1015. A hospital-based RHC is paid based on an all-inclusive encounter rate. One of the following modifiers must be submitted for general medical services: AH, AJ, AM, SA, TD, TE, or U7.

The services listed below must be submitted using the RHC provider identifier and the appropriate benefit code:

- THSteps medical checkups
- Family planning services (including implantable contraceptive capsules provision, insertion, or removal)
- Immunizations provided in hospital-based RHCs

These services must be submitted with an AM, SA, or U7 modifier if performed in an RHC setting. Claims are paid under the Prospective Payment System (PPS) reimbursement methodology.

When submitting a claim on the CMS-1500 paper claim form, providers must use the appropriate national POS (72) for an RHC setting.

Outpatient hospital services (including emergency room services) and inpatient hospital services provided outside the RHC setting are to be submitted using the individual or group physician provider identifier.
Hospital-based RHCs must submit claims for pneumococcal and influenza vaccines as non-RHC services, under their hospital provider identifier.

**Note:** A visit is a face-to-face encounter between an RHC client and a physician, PA, nurse practitioner (NP), certified nurse-midwife (CNM), visiting nurse, or clinical NP. Encounters with more than one health professional and multiple encounters with the same health professional that take place on the same day and at a single location constitute a single visit, except where one or the other of the following conditions exists:

- After the first encounter, the client suffers illness or injury requiring additional diagnosis or treatment.
- The RHC client has a medical visit and an other health visit.

An other health visit includes, but is not limited to, a face-to-face encounter between an RHC client and a clinical social worker.

### 4.2.6 Cardiac Rehabilitation

Cardiac rehabilitation is a physician-supervised program that furnishes physician-prescribed exercise, cardiac risk factor modification, psychosocial assessment, and outcomes assessment.

Outpatient cardiac rehabilitation is considered reasonable and necessary for clients who have had one of the following within 12 months of beginning the cardiac rehabilitation program:

- Acute myocardial infarction
- Coronary artery bypass surgery (CABG)
- Percutaneous transluminal coronary angioplasty or coronary stenting
- Heart valve repair or replacement
- Major pulmonary surgery
- Sustained ventricular tachycardia or fibrillation
- Class III or class IV congestive heart failure
- Chronic stable angina

**Note:** A cardiac rehabilitation program in which the cardiac monitoring is done using telephonically transmitted electrocardiograms to a remote site is not covered by Texas Medicaid.

Cardiac rehabilitation must be provided in a facility that has the necessary cardiopulmonary, emergency, diagnostic, and therapeutic life-saving equipment (i.e. oxygen, cardiopulmonary resuscitation equipment, or defibrillator) available for immediate use. If no clinically significant arrhythmia is documented during the first three weeks of the program, the provider may have the client complete the remaining portion without telemetry monitoring by the physician’s order.

Although cardiac rehabilitation may be considered a form of physical therapy, it is a specialized program conducted by non-physician personnel who are trained in both basic and advanced cardiac life support techniques and exercise therapy for coronary disease, and provide the services under the direct supervision of a physician.

Direct supervision of a physician means that a physician must be immediately available and accessible for medical consultations and emergencies at all times when items and services are being furnished under cardiac rehabilitation programs. Outpatient cardiac rehabilitation begins after the client has been discharged from the hospital. A physician’s prescription is required after the acute convalescent period.
and after it has been determined that the client’s clinical status and capacity will allow for safe participation in an individualized progressive exercise program. Outpatient cardiac rehabilitation requires close monitoring and direct supervision by a physician and includes:

- Medical evaluation performed by the physician responsible for prescribing the client’s rehabilitation program and includes a clinical examination, a medical history, and an initial plan or goal.
- An education and counseling program to modify risk factors (nutritional counseling, stress reduction, smoking cessation, weight loss, etc.).
- Prescribed exercise concurrent with and without electrocardiogram (ECG) monitoring.
- Services performed in an approved facility by trained professionals.

**Note:** Direct supervision is met when the services are performed on hospital premises or within 250 yards of the hospital.

Cardiac rehabilitation will be limited to a maximum of 2 one-hour sessions per day and 36 sessions over 18 weeks per rolling year.

Providers must obtain prior authorization for additional cardiac rehabilitation sessions, which will be limited to a maximum of 36 sessions in an extended period of time in a 52-week period from the date of authorization of additional sessions.

To confirm that a continuation of cardiac rehabilitation is at the request of, and coordinated with the attending physician, the medical record must include evidence of communication between the cardiac rehabilitation staff and either the medical director or the referring physician. If the physician responsible for such follow-up is the medical director, then his or her notes must be evident in each client’s medical record.

Cardiac rehabilitation may be considered medically necessary beyond 36 sessions if the medical record contains documentation that the client has had another cardiac event, or if the prescribing physician documents that a continuation of cardiac rehabilitation is medically necessary. Medical necessity documentation must include the following:

- Progress made from the beginning of the cardiac rehabilitation period to the current service request date, including progress towards previous goals
- Information that supports the client’s capability of continued measurable progress
- A proposed treatment plan for the requested extension dates with specific goals related to the client’s individual needs

Prior authorization must be obtained through the TMHP Special Medical Prior Authorization (SMPA) Department. Providers must send prior authorization requests, along with documentation to support medical necessity, to the following address:

Texas Medicaid & Healthcare Partnership  
Special Medical Prior Authorization  
12357-B Riata Trace Parkway  
Austin, TX 78727  
Fax: 1-512-514-4213

Requests for prior authorization can also be submitted online through the TMHP website at [www.tmhp.com](http://www.tmhp.com).

The evaluation provided by the cardiac rehabilitation team at the beginning of each cardiac rehabilitation session is not considered a separate service and will be included in the reimbursement for the cardiac rehabilitation session. Evaluation and management (E/M) services unrelated to cardiac rehabilitation may be submitted with modifier 25 appended to the E/M code when supporting documentation in the medical record demonstrates a separately identifiable E/M service was provided on the same day by the same provider who renders the cardiac rehabilitation.
Physical and occupational therapy will not be reimbursed separately when furnished in addition to cardiac rehabilitation exercise program services unless there is also a diagnosis of a non-cardiac condition requiring such therapy.

**Example:** If a client is recuperating from an acute phase of heart disease and has had a stroke that requires physical or occupational therapy, the physical or occupational therapy for the stroke may be reimbursed separately from the cardiac rehabilitation services for the acute phase of heart disease.

When provided as part of the cardiac rehabilitation program, client education services, such as formal lectures and counseling on diet, nutrition, and sexual activity to assist the client in adjusting living habits because of the cardiac condition, will not be separately reimbursed.

Procedure code S9472 (cardiac rehabilitation program nonphysician provider per diem) is used for hospitals submitting claims for cardiac rehabilitation, and it must be submitted with revenue code 943 (other therapeutic services-cardiac rehabilitation) and one of the following diagnosis codes:

<table>
<thead>
<tr>
<th>Diagnosis Codes</th>
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</thead>
<tbody>
<tr>
<td>I110</td>
</tr>
<tr>
<td>I2102</td>
</tr>
<tr>
<td>I219</td>
</tr>
<tr>
<td>I2720</td>
</tr>
<tr>
<td>I5020</td>
</tr>
<tr>
<td>I5040</td>
</tr>
<tr>
<td>I50814</td>
</tr>
<tr>
<td>Z951</td>
</tr>
</tbody>
</table>

Coverage of cardiac rehabilitation programs is considered reasonable and necessary only for clients who have documentation of acute myocardial infarction, coronary artery bypass surgery (CABG), percutaneous transluminal coronary angioplasty or coronary stenting heart valve repair/replacement, major pulmonary surgery, sustained ventricular tachycardia or fibrillation, class III or class IV congestive heart failure, or chronic stable angina within the past twelve (12) months prior to the beginning of the program.

### 4.2.7 Chemotherapy Administration

Hospitals must submit outpatient charges using the appropriate revenue codes for room charges, supplies, IV equipment, and pharmacy.

Revenue code 636 may be reimbursed for the technical component of prolonged infusion of chemotherapeutic agents. The most appropriate chemotherapy procedure code must be billed with revenue code 636.

### 4.2.8 Colorectal Cancer Screening

Fecal occult blood tests, multi-targeted stool DNA (mt-sDNA) tests, screening colonoscopies, and sigmoidoscopies evidence-based methods of colorectal cancer screening.

#### 4.2.8.1 Fecal Occult Blood Tests

Procedure codes G0328 (with modifier QW) and 82270 may be reimbursed one service per rolling year for clients who are 45 years of age and older.

#### 4.2.8.2 MT-sDNA Tests

Procedure code 81528 may be reimbursed once every 3 years for clients who are 45 years of age and older for services rendered in the laboratory setting.
4.2.8.3 Sigmoidoscopies
Procedure code G0104 may be reimbursed once every 5 years and is limited to one of the following diagnosis codes:

<table>
<thead>
<tr>
<th>Diagnosis Codes</th>
</tr>
</thead>
<tbody>
<tr>
<td>Z0000*</td>
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<tr>
<td>Z0001*</td>
</tr>
<tr>
<td>Z1210</td>
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<tr>
<td>Z1211</td>
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<tr>
<td>Z1213</td>
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<tr>
<td>Z859</td>
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<tr>
<td>Z86002</td>
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<tr>
<td>Z86003</td>
</tr>
<tr>
<td>Z86004</td>
</tr>
<tr>
<td>Z86006</td>
</tr>
<tr>
<td>Z86007</td>
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<td>Z86010</td>
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</tbody>
</table>

*Diagnosis code Z0000 or Z0001 may be used for screening if no other diagnosis is appropriate for the service rendered, but no more frequently than recommended by the U.S. Preventive Services Task Force (USPSTF).

4.2.8.4 Colonoscopies
Procedure code G0105 may be reimbursed once every two years for clients who meet the definition of high-risk. Procedure code G0105 must be submitted with one of the following diagnosis codes:

<table>
<thead>
<tr>
<th>Diagnosis Codes</th>
</tr>
</thead>
<tbody>
<tr>
<td>K5010</td>
</tr>
<tr>
<td>K5011</td>
</tr>
<tr>
<td>K50112</td>
</tr>
<tr>
<td>K50113</td>
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<tr>
<td>K50114</td>
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<tr>
<td>K50118</td>
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<tr>
<td>K501211</td>
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<tr>
<td>K501212</td>
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<td>K501213</td>
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<tr>
<td>K51213</td>
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<td>K51214</td>
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<td>K51218</td>
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<td>K51311</td>
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<td>K51313</td>
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<td>K52832</td>
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<td>K52838</td>
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<td>K5289</td>
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<tr>
<td>K529</td>
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<tr>
<td>Z800</td>
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<tr>
<td>Z8371</td>
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<tr>
<td>Z85038</td>
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<tr>
<td>Z85048</td>
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<td>Z8509</td>
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</tbody>
</table>


4.2.8.5 Exclusions
Barium enemas for colorectal cancer screening are not a benefit of Texas Medicaid.

4.2.9 Computed Tomography and Magnetic Resonance Imaging
Prior authorization is required for all outpatient nonemergent (i.e., those that are scheduled) CT, computed tomography angiography (CTA), magnetic resonance imaging (MRI), and magnetic resonance angiography (MRA) studies before services are rendered. Authorization is not required for the emergency department or inpatient hospital radiology services. Retroactive authorization may be required for some outpatient emergent studies.

Reimbursement for procedures with descriptions that specify "with contrast" include payment for contrast materials. Some diagnostic radiopharmaceuticals are benefits of Texas Medicaid. Outpatient hospitals may submit the total component of the following procedure codes:

<table>
<thead>
<tr>
<th>Procedure Codes</th>
</tr>
</thead>
<tbody>
<tr>
<td>77371</td>
</tr>
<tr>
<td>77372</td>
</tr>
<tr>
<td>77373</td>
</tr>
<tr>
<td>77422</td>
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<tr>
<td>77423</td>
</tr>
<tr>
<td>77520</td>
</tr>
<tr>
<td>77525</td>
</tr>
<tr>
<td>G0339</td>
</tr>
</tbody>
</table>

Procedure code 77399 may be submitted as either the total component or the technical component.
Providers can refer to the OFL or the applicable fee schedules on the TMHP website at www.tmhp.com to review the diagnostic radiopharmaceuticals that are reimbursed by Texas Medicaid. OFL and static fee schedules available on the TMHP website display fees after applicable rate reductions have been applied. Previously, the OFL and static fee schedules did not reflect all rate reductions, and providers were required to calculate the 1- and 2-percent reductions implemented.

Refer to: Subsection 4.2.9, “Computed Tomography and Magnetic Resonance Imaging” in this handbook for additional information about prior authorization requirements.

Subsection 3.2.6, “Authorization Requirements for CT, CTA, MRI, fMRI, MRA, PET, and Cardiac Nuclear Imaging Services” in the Radiology and Laboratory Services Handbook (Vol. 2, Provider Handbooks) for additional information about emergency outpatient imaging services.

4.2.10 Electrodiagnostic (EDX) Testing
Electromyography (EMG) and nerve conduction studies (NCS), collectively known as EDX testing, must be medically indicated and may be reimbursed to outpatient hospitals. Testing must be performed using EDX equipment that provides assessment of all parameters of the recorded signals. Studies performed with devices designed only for screening purposes rather than diagnoses are not a benefit of Texas Medicaid.

NCS and EMG studies are diagnosis restricted and may require prior authorization.


4.2.11 Fluocinolone Acetonide
The fluocinolone acetonide intravitreal implant may be reimbursed for services rendered to clients who are 12 years of age and older. Procedure code J7311 is only payable with a posterior uveitis diagnosis of more than six months duration and the condition has been unresponsive to oral or systemic medication treatment.

4.2.11.1 Prior Authorization for Fluocinolone Acetonide
To request prior authorization, providers must submit requests by fax or mail to the SMPA Department at:

Texas Medicaid & Healthcare Partnership
Special Medical Prior Authorization
12357-B Riata Trace Parkway
Austin, TX 78727
Fax: 1-512-514-4213

Requests for prior authorization can be submitted online through the TMHP website at www.tmhp.com.

4.2.12 Fetal Nonstress Testing and Contraction Stress Test
Claims for nonstress and contraction stress testing conducted in the outpatient setting must be submitted with revenue code 729. Services during an inpatient hospital stay are reimbursed under the hospital’s DRG.

Refer to: Section 4.1.1, “Antepartum and Fetal Invasive Procedures” in the Gynecological, Obstetrics, and Family Planning Title XIX Services Handbook (Vol. 2, Provider Handbooks).

4.2.13 Hyperbaric Oxygen Therapy (HBOT)
HBOT is a type of therapy that increases the environmental oxygen pressure to promote the movement of oxygen from the environment into the client’s body tissues. Such treatment may be a benefit of Texas Medicaid when it is performed in specially constructed hyperbaric chambers, pressurized to 1.4
atmosphere absolute (atm abs) or higher, which may hold one or more clients (sea-level pressure is equal to 1 atm.abs). Although oxygen may be administered by mask, cannula, or tube in addition to the hyperbaric treatment, this use of oxygen is not considered hyperbaric oxygen treatment in itself.

HBOT procedure codes 99183 and G0277 require prior authorization before the date that service is initiated.

The number of billable units of procedure code G0277 is based upon the time that the client receives treatment with hyperbaric oxygen.

In calculating how many 30-minute intervals to report, hospitals should take into consideration the time spent under pressure during descent, air breaks, and ascent, (in minutes), as follows:

- The first unit is for the time spent in the chamber receiving hyperbaric oxygen and must be for a minimum of 16 minutes.
- To bill for a second (or subsequent unit), all previous units of time must have been for the full thirty minutes, and the last unit must be for 16-30 minutes.

Procedure code 99183 equates to one total treatment (one professional session).

Procedure code G0277 must be billed with revenue code B-413 on the same claim. If procedure code G0277 is not on the same claim as revenue code B-413, the claim will be denied.


4.2.14 Laboratory Services

Routine laboratory services, directly related to the surgical procedure being performed, are not reimbursed separately. Claims for nonroutine laboratory services provided with emergency conditions may be submitted separately with documentation that the complicating condition arose after the initiation of the surgery. Outpatient claims for laboratory services must reflect only tests actually performed by the hospital laboratory.

Exception: Hospital laboratories may submit claims for all the tests performed on a specimen if some but not all the tests are done by another laboratory on referral from the hospital submitting the claim.

The billing hospital must enter the name and provider identifier of the performing laboratory in Block 80 of the UB-04 CMS-1450 paper claim form and must enter the performing laboratory’s provider identifier next to the service provided by the performing laboratory.

Hospitals may submit claims for a handling fee (procedure code 99001) for collecting and forwarding a specimen to a referral laboratory when the laboratory handling fee is not being billed through other methods. Only one handling fee may be charged per day, per client, unless specimens are sent to two or more different laboratories; this must be documented on the claim.


4.2.14.1 Clinical Laboratory Improvement Amendments (CLIA)

All providers of laboratory services must comply with the rules and regulations of CLIA. Providers not complying with CLIA will not be reimbursed for laboratory services.
Refer to: Subsection 1.1, “Provider Enrollment” in “Section 1: Provider Enrollment and Responsibilities” (Vol. 1, General Information) for more information about enrollment procedures.

Subsection 2.1.1, “Clinical Laboratory Improvement Amendments (CLIA)” in the Radiology and Laboratory Services Handbook (Vol. 2, Provider Handbooks) for more information about CLIA.

The Medicaid Managed Care Handbook (Vol. 2, Provider Handbooks).

### 4.2.15 Lung Volume Reduction Surgery (LVRS)

LVRS surgery must be performed in a facility that meets at least one of the following requirements:

- Certified under the Disease Specific Care Certification Program for LVRS by the Joint Commission on Accreditation of Health Care Organization (JCAHO)
- Approved by Medicare as a lung or heart-lung transplant facility

The surgery must be both preceded and followed by a program of diagnostic and therapeutic services that are consistent with those provided in the National Emphysema Treatment Trial (NETT) and designed to maximize the client’s potential to successfully undergo and recover from surgery. The program must meet all of the following requirements:

- Include a 6-to 10-week series of at least 16, and no more than 20, pre-operative sessions, each lasting a minimum of 2 hours
- Include at least 6, and no more than 10, post-operative sessions, each lasting a minimum of 2 hours, within 8 to 9 weeks after the LVRS
- Be consistent with the care plan that was developed by the treating physician following the performance of a comprehensive evaluation of the client’s medical, psychosocial, and nutritional needs
- Be arranged, monitored, and performed under the coordination of the facility where the surgery takes place. Prior authorization is required for the LVRS procedure. However, prior authorization is not required for the pre-operative and post-discharge pulmonary services

LVRS must be prior authorized and is limited to clients who have severe emphysema, disabling dyspnea, and evidence of severe air trapping. Prior authorization is not required for the associated preoperative pulmonary surgery services for preparation of LVRS (procedure codes G0302, G0303, and G0304) or the associated postdischarge pulmonary surgery services after LVRS (procedure code G0305).

LVRS and post-discharge LVRS are restricted to the following emphysema diagnosis codes:

<table>
<thead>
<tr>
<th>Diagnosis Codes</th>
</tr>
</thead>
<tbody>
<tr>
<td>J430</td>
</tr>
</tbody>
</table>

Procedure codes 32491, G0302, G0303, G0304 and G0305 are limited to one per rolling year per client any provider.

Procedure code G0305 may be reimbursed only if a claim for LVRS (procedure code 32491) has been submitted within the past 12 months.


### 4.2.16 Magnetoencephalography (MEG) Services

Inpatient and outpatient hospitals must use revenue code 860 or 861 for reimbursement of magnetoencephalography (MEG) services. The appropriate MEG procedure code must be listed on the claim.

**Note:** Reimbursement to an outpatient hospital will be based on the submitted procedure code.

4.2.17 Neurostimulators

Neurostimulators may be a benefit in the outpatient hospital setting when medically necessary. All procedures require prior authorizations.


Subsection 4.2.17.1, “Prior Authorization for Neurostimulators” in this handbook.

4.2.17.1 Prior Authorization for Neurostimulators

All devices and related procedures for the initial application or surgical implantation of the stimulator device require prior authorization. Requests for prior authorization must be submitted to the SMPA Department.


4.2.18 Occupational and Physical Therapy Services

Refer to: The Physical Therapy, Occupational Therapy, and Speech Therapy Services Handbook (Vol. 2, Provider Handbooks) for more information about therapy services.

4.2.19 Radiation Therapy Services

Take-home drugs given during the course of therapy can be reimbursed separately through the VDP. Hospitals use revenue code 333, Radiation therapy, on the UB-04 CMS-1450 paper claim form when submitting charges for these services.

The following radiation therapy services provided in an outpatient setting are allowed only once per day unless documentation of medical necessity supports the need for repeated services:

- Therapeutic radiation treatment planning
- Therapeutic radiology simulation-aided field setting
- Teletherapy
- Brachytherapy isodose calculation
- Treatment devices
- Proton beam delivery/treatment
- Intracavity radiation source application
- Interstitial radiation source application
- Remote afterloading high intensity brachytherapy
- Radiation treatment delivery
- Localization, and radioisotope therapy

4.2.19.1 Radiopharmaceuticals

Radiopharmaceuticals may be considered for separate reimbursement when used for therapeutic treatment.
The following procedure codes are payable to outpatient hospitals:

<table>
<thead>
<tr>
<th>Procedure Codes</th>
</tr>
</thead>
<tbody>
<tr>
<td>79403*</td>
</tr>
</tbody>
</table>
| *Total or technical component

Strontium-89 chloride may be billed using procedure code A9600 and will be limited to a total of 10 mci intravenously injected every 90 days, any provider.

Strontium-89 chloride is reimbursed as one service per 90 days for any provider.

Strontium-89 chloride will be considered when submitted with diagnosis code C7951 or C7952.

Sodium phosphate p-32, therapeutic, will be considered when submitted with one of the following diagnoses:

<table>
<thead>
<tr>
<th>Diagnosis Codes</th>
</tr>
</thead>
<tbody>
<tr>
<td>C7951</td>
</tr>
<tr>
<td>C9512</td>
</tr>
</tbody>
</table>

Chromic phosphate p-32 suspension will be considered when submitted with diagnosis code C782 (secondary malignant neoplasm of the pleura) and diagnosis code C786 (secondary malignant neoplasm of the retroperitoneum and peritoneum).


Prior Authorization for Therapeutic Radiopharmaceuticals

Procedure codes A9542 and A9543 require prior authorization. Ibritumomab tiuxetan will be considered when submitted with one of the following diagnosis codes:

<table>
<thead>
<tr>
<th>Diagnosis Codes</th>
</tr>
</thead>
<tbody>
<tr>
<td>C8259</td>
</tr>
<tr>
<td>C8599</td>
</tr>
</tbody>
</table>

Prior authorization is required for procedure codes A9542 and A9543, which will be considered with documentation of all of the following:

- A diagnosis of either a low-grade follicular or transformed B-cell non-Hodgkin’s lymphoma.
- Client has failed, relapsed, or become refractory to conventional chemotherapy.
- Marrow involvement is less than 26 percent.
- Platelet count is 100,000 cell/mm3 or greater.
- Neutrophil count is 1,500 cells/mm3 or greater.
- Client has failed a trial of rituximab.

Lutetium lu 177 dotatate (Lutathera) intravenous injection (procedure code A9513) is indicated for the treatment of clients who are 18 years of age or older with a diagnosis of gastroenteropancreatic-neuroendocrine tumors. For all other indications, Lutetium lu 177 dotatate (Lutathera) injection for intravenous use is not proven to be medically effective and is considered experimental.
Lutetium lu 177 dotatate (Lutathera) procedure code A9513 must be administered under the control of an oncologist or a nuclear medicine specialist who is licensed and authorized to administer radiopharmaceuticals in an outpatient setting. Prior authorization requests must be submitted to the Special Medical Prior Authorization (SMPA) department at TMHP using the Special Medical Prior Authorization (SMPA) Request Form.

An SMPA Request Form must be completed, signed, and dated by the prescribing provider. The SMPA form will not be accepted after 90 days from the date of the prescribing provider’s signature.

The completed SMPA Request Form must be maintained by the prescribing provider in the client’s medical record and is subject to retrospective review.

Section C of the SMPA Request Form under the Statement of Medical Necessity must contain the following:

- Documentation of the client’s dosage
- The administration schedule
- The number of injections to be administered during the prior authorization period
- The requested units and millicuries per injection
- The dosage calculation

Prior authorization must be requested through the SMPA department with appropriate documentation. Requests can be mailed or faxed to:

Texas Medicaid & Healthcare Partnership
Special Medical Prior Authorization
12357-B Riata Trace Parkway
Austin, TX 78727
Fax: 1-512-514-4213

Requests for prior authorization can be submitted online through the TMHP website at www.tmhp.com.

Refer to: Subsection 9.2.73.2, “Other Limitations on Therapeutic Radiopharmaceuticals” in the Medical and Nursing Specialists, Physicians, and Physician Assistants Handbook (Vol. 2, Provider Handbooks) for more information.

4.2.20 Respiratory Services

4.2.20.1 Aerosol Treatment

Nebulized aerosol treatments with short-acting beta-agonists provided in the outpatient setting are considered medically necessary for certain acute medical problems when breathing is compromised. Authorization is not required for aerosol treatments.

Outpatient facilities must submit claims for aerosol treatments using revenue code 412 and the appropriate beta-agonist procedure code.

4.2.20.2 Diagnostic Testing

Nitric oxide expired gas determination (FeNO) measurement provided in the physician’s office or outpatient hospital setting is considered medically necessary as an adjunct to the established clinical and laboratory assessments for diagnosing and assessing asthma, predicting exacerbations, and evaluating the response of a patient with asthma to anti-inflammatory therapy.

Claims for nitric oxide treatments may be submitted using procedure code 95012. Hospital providers must include the following when submitting claims for procedure code 95012:

- Revenue code 419 must appear on the same line as procedure code 95012.
The claim must have a line item for either procedure code 94010 or 94060. This line item must also indicate revenue code 419.

*Note:* Procedure code 94010 or 94060, when submitted in conjunction with procedure code 95012, may only be reimbursed in an office or outpatient hospital setting, and is not reimbursed for critical care, emergency care, or anesthesiology.

### 4.2.20.3 Pulmonary Function Studies

Pulmonary function studies considered for reimbursement to outpatient hospitals include, but are not limited to, the following procedures when submitted with the total component (TOS 5):

<table>
<thead>
<tr>
<th>Procedure Codes</th>
</tr>
</thead>
<tbody>
<tr>
<td>94010</td>
</tr>
<tr>
<td>94618</td>
</tr>
</tbody>
</table>

High Altitude Simulation Test (HAST) procedure codes 94452 and 94453 must be submitted with one of the following diagnosis codes:

<table>
<thead>
<tr>
<th>Diagnosis Codes</th>
</tr>
</thead>
<tbody>
<tr>
<td>E840*</td>
</tr>
<tr>
<td>I2721</td>
</tr>
<tr>
<td>I279*</td>
</tr>
<tr>
<td>J438*</td>
</tr>
<tr>
<td>J700*</td>
</tr>
<tr>
<td>J8401</td>
</tr>
<tr>
<td>J9610</td>
</tr>
<tr>
<td>Q334</td>
</tr>
</tbody>
</table>

*Note:* When billing for HAST (procedure code 94452 or 94453) with one of the diagnosis codes indicated in the table with an asterisk (*), evidence of hypoxemia must be documented in the client’s medical record.

When multiple procedure codes are submitted, the most inclusive code of the related codes will be reimbursed and all other related codes will be denied.

When unrelated pulmonary function studies are submitted together, each will be considered for reimbursement.

### 4.2.21 Screening, Brief Intervention, and Referral to Treatment (SBIRT)


### 4.3 Documentation Requirements

All services require documentation to support the medical necessity of the service rendered, including hospital services. Hospital services are subject to retrospective review and recoupment if documentation does not support the service that was submitted for reimbursement.

### 4.4 Outpatient Utilization Review

UR activities of all Medicaid services provided by hospitals reimbursed under the DRG prospective payment system are required by Title XIX of the Social Security Act, Sections 1902 and 1903. The review activities are accomplished through a series of monitoring systems developed to ensure services are
appropriate to need, of optimum quality and quantity, and rendered in the most cost-effective mode. Clients and providers are subject to UR monitoring. The monitoring focuses on the appropriate screening activities, medical necessity of all services, and quality of care as reflected by the choice of services provided, type of provider involved, and settings in which the care was delivered. This monitoring ensures the efficient and cost-effective administration of Texas Medicaid.

TMHP is responsible for a comprehensive integrated review process to identify misuse and inappropriate claim submission patterns by outpatient hospitals and HASCs. All providers are subject to TMHP’s UR monitoring. Providers are selected for review based on a comparison of their individual resource utilization with a peer group of similar specialty and geographic locality. The main goal of the required utilization control is to identify those providers whose practice patterns are aberrant from their peers and provide the necessary educational actions to help the provider achieve Texas Medicaid compliance. An analysis of UR data is completed by a registered nurse analyst for review by the medical director and staff. If the analyst substantiates that a provider’s practice and claim submission patterns are inconsistent with the federal requirements and Texas Medicaid’s scope of benefits, a TMHP representative contacts the provider. The purpose of the contact is to discuss appropriate claim submission guidelines and to assist the provider in resolving the inappropriate claim submission patterns identified in the review.

TMHP uses the following criteria when reviewing all hospital outpatient medical records. Services must be:

- Medically necessary.
- Ordered by a physician, signed, and dated. Signature stamps are valid if initialed and dated by the physician.
- Submitted in the quantities ordered and documented as provided.
- Program benefits.
- Specifically identified on the charge tickets or itemized statement submitted with the claim or by the HCPCS procedure code on the claim.
- Indicated by the documentation in the medical record.
- Submitted to Texas Medicaid only after other medical insurance resources have been exhausted.

Refer to: Subsection 4.1.1, “Your Texas Benefits Medicaid Card” in “Section 4: Client Eligibility” (Vol. 1, General Information).

The determination of the TMHP UR process may result in the following:

- Educational letters and visits
- Mail-in of medical records for review
- On-site medical record review (outpatient, HASC, or inpatient records not reviewed)
- Referral of questionable claims to HHSC or HHSC OIG
- Recoupment
- Prepayment review

The intent of these actions is to ensure the most effective and appropriate use of available services and facilities and provide appropriate, cost-effective care to clients with Medicaid coverage.
4.5 Claims Filing and Reimbursement

4.5.1 Outpatient Claims Information

Claims for scheduled procedures that are performed in a HASC must be submitted using the HASC provider identifier with type of bill (TOB) 131. Claims for emergency or unscheduled procedures performed in a hospital when the client is an outpatient must be submitted using the hospital provider identifier and appropriate revenue and HCPCS code (if required) with TOB 131.

Claims for outpatient hospital services must be submitted to TMHP in an approved electronic format or on the UB-04 CMS-1450 paper claim form.

Freestanding ambulatory surgical centers must submit claims on the CMS-1500 claim form. The performing surgeon or referring physician name and number must be identified in Block 17. Identification of outpatient charges must be in Block 44 if submitting by HCPCS code. If appropriate, the revenue code must be indicated in Block 42. Texas Medicaid recommends the use of specific procedure codes for claim submission. Do not use the revenue code description in Block 43; the HCPCS narrative description must be identified in this block. For example, when submitting charges for physical therapy, do not use the description associated with revenue code 420. To receive reimbursement for physical therapy services, providers must identify the specific modality used (e.g., gait training).

Examples:

- **Emergency Room.** Submit as “Emergency room” or “Emergency room charge per use.” If the client visits the emergency room more than once in one day, the time must be given for each visit. The time of the first visit must be identified in Block 13, using 00 to 23 hours military time (e.g., 1350 for 1:50 p.m.). Indicate other times on the same line as the procedure code. Claims for emergency CT, CTA, MRI, or MRA studies provided in the emergency department must have the appropriate corresponding emergency services revenue code (450, 451, 456, or 459) to be considered for payment.

- **Observation Room.** Submit as “observation room.” (Revenue code 762).

- **Operating Room.** Submit as “Operating Room.” (Revenue code 360, 361, or 369).

- **Recovery Room.** Submit as “Recovery Room” or “Cast Room” as appropriate. (Revenue code 710 or 719).

- **Injections.** Must have “Inj.-name of drug; route of administration; the dosage and quantity” or the injection code.

- **Drugs and Supplies.** The drug description must include the name, strength, and quantity. Take-home drugs and supplies are not a benefit of Texas Medicaid:
  - Take-home drugs must be submitted with revenue code 253.
  - Take-home supplies must be submitted with revenue code 273.
  - Self-administered drugs must be submitted with revenue code 637.

- **Radiology.** Facilities must submit claims using the most appropriate revenue and HCPCS code. The physician must submit claims for professional services by a physician separately. The license number of the ordering physician must be in Block 83. If the client receives the same radiology procedure more than once in one day, the time must be given for each visit. The time of the first visit must be identified in Block 13, using 00 to 23 hours military time (such as 1350 for 1:50 p.m.). Indicate other times on the same line as the procedure code.

- **Laboratory.** Provide a complete description or use the procedure codes for the laboratory procedures. The physician must submit claims for professional services by a physician separately. Blocks 78–79 must have the license number of the ordering physician. If laboratory work is sent out, enter the name of the test and name and address or Medicaid number of the laboratory where the work
was forwarded. If the client receives the same laboratory procedure more than once in one day, give
the time for each visit. The time of the first visit must be identified in Block 13, using 00 to 23 hours
military time (e.g., 1350 for 1:50 p.m.). Indicate other times on the same line as the procedure code.

- **Nuclear Medicine.** Provide a complete description.
- **Day Surgery.** Day surgery must be submitted as an inclusive charge using TOS F. Providers must not
submit claims for separate services that were provided in conjunction with the surgery (e.g., lab,
radiology, and anesthesia). File claims for unscheduled emergency outpatient surgical procedures
with separate charges (e.g., lab, radiology, anesthesia, and emergency room) for all services using
TOB 131 and the hospital’s provider identifier.

Claims for emergency or unscheduled procedures performed in a hospital when the client is an outpa-
tient must be submitted using the hospital provider identifier and appropriate revenue and HCPCS code
(if required) with TOB 131.

Refer to the ambulatory surgical center/hospital-based ambulatory surgical center (ASC/HASC) section
for information on scheduled procedures. Additional claims information can be found within individual
topic areas within this section.

Charges on claims must be itemized on the face of the UB-04 CMS-1450 paper claim form instead of
submitting attachments or charge details. TMHP uses information attached to the claim for clarification
purposes only.

If a claim contains more than 28 details, continue the claim on additional UB-04 CMS-1450 paper claim
forms or electronic equivalent. Total each claim form as a stand-alone claim. If you do not total each
page, your claim may be denied for being over the limitation, and must be resubmitted with 28 or less
details.

Providers may purchase UB-04 CMS-1450 paper claim forms from the vendor of their choice. TMHP
does not supply the forms.

When completing a UB-04 CMS-1450 paper claim form, all required information must be included on
the claim, as TMHP does not key any information. Superbills, or itemized statements, are not accepted
as claim supplements.

**Refer to:** “Section 3: TMHP Electronic Data Interchange (EDI)” (*Vol. 1, General Information*) for
information on electronic claims submissions.

“Section 6: Claims Filing” (*Vol. 1, General Information*) for general information about
claims filing.

Subsection 6.6, “UB-04 CMS-1450 Paper Claim Filing Instructions” in “Section 6: Claims
Filing” (*Vol. 1, General Information*). Blocks that are not referenced are not required for
processing by TMHP and may be left blank.

Outpatient hospital services must be itemized by date of service. Procedures repeated over a period of
time must be submitted for each separate date of service. Do not combine multiple dates of service on
the same line detail.

**4.5.2 Outpatient Reimbursement**

Outpatient services are reimbursed on a reasonable cost based on a percentage of the hospital’s most
recent tentative Medicaid cost report settlement.

The reimbursement rate for non-high-volume hospitals is as follows with the application of the hospital
specific interim rate:

<table>
<thead>
<tr>
<th>Non-high-volume Provider</th>
<th>Current Allowable Rate</th>
</tr>
</thead>
<tbody>
<tr>
<td>Children’s hospitals</td>
<td>72.27 percent of the allowable charges</td>
</tr>
</tbody>
</table>
The reimbursement rate for high-volume hospitals is as follows with the application of the hospital specific interim rate:

<table>
<thead>
<tr>
<th>High-volume Provider</th>
<th>Current Allowable Rate</th>
</tr>
</thead>
<tbody>
<tr>
<td>Children’s hospitals</td>
<td>76.03 percent of the allowable charges</td>
</tr>
<tr>
<td>Rural hospitals</td>
<td>100 percent of the allowable charges</td>
</tr>
<tr>
<td>State-owned teaching hospitals</td>
<td>76.03 percent of the allowable charges</td>
</tr>
<tr>
<td>Other hospitals</td>
<td>72 percent of the allowable charges</td>
</tr>
<tr>
<td>ASCs/HASCs that qualify as high-volume providers</td>
<td>Additional 5.2 percent increase in payment rates</td>
</tr>
</tbody>
</table>

High-volume providers are eligible for additional payments on Texas Medicaid fee-for-service claims. A high-volume outpatient hospital provider is defined as one that was paid at least $200,000 during calendar year 2004.

All clinical laboratory services are reimbursed at a percentage of the prevailing charge. Hospitals that are identified by Medicare as sole community hospitals are reimbursed at a higher percentage of the prevailing charges for services that are provided to clients in the outpatient setting.

Clinical pathology consultations are also allowed for reimbursement.

Refer to: The HHSC Rate Analysis web page at [https://rad.hhs.texas.gov/hospitals-clinic-services](https://rad.hhs.texas.gov/hospitals-clinic-services) for additional information about hospital reimbursement.

Subsection 3.7.5, “Provider Cost and Reporting” in this handbook for more information about the calculation of the interim rate.


4.5.3 Provider Cost and Reporting

Refer to: Subsection 3.7.5, “Provider Cost and Reporting” in this handbook.

4.5.4 National Correct Coding Initiative (NCCI) and Medically Unlikely Edit (MUE) Guidelines

The HCPCS and CPT codes included in the Texas Medicaid Provider Procedures Manual are subject to National Correct Coding Initiative (NCCI) relationships, which supersede any exceptions to NCCI code relationships that may be noted in the manuals. Providers should refer to the CMS NCCI web page for correct coding guidelines and specific applicable code combinations.

In instances when Texas Medicaid limitations are more restrictive than NCCI MUE guidance, Texas Medicaid limitations prevail.
### 4.5.5 Outpatient Hospital Revenue Codes

UB-04 CMS-1450 revenue codes must be used to submit claims for outpatient hospital facility services. In some instances, a HCPCS procedure code is required in addition to the revenue code for accurate claims processing:

<table>
<thead>
<tr>
<th>Revenue Code</th>
<th>Description</th>
<th>Comments</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>Special Charges – Canceled Surgery</strong></td>
<td></td>
<td></td>
</tr>
<tr>
<td>220</td>
<td>Special Charges</td>
<td>Procedure code required</td>
</tr>
<tr>
<td><strong>Pharmacy</strong></td>
<td></td>
<td></td>
</tr>
<tr>
<td>250</td>
<td>General classification</td>
<td></td>
</tr>
<tr>
<td>251</td>
<td>Generic drugs</td>
<td></td>
</tr>
<tr>
<td>252</td>
<td>Nongeneric drugs</td>
<td></td>
</tr>
<tr>
<td>253</td>
<td>Take-home drugs</td>
<td>Not a benefit</td>
</tr>
<tr>
<td>254</td>
<td>Drugs incident to other diagnostic services</td>
<td></td>
</tr>
<tr>
<td>255</td>
<td>Drugs incident to radiology</td>
<td></td>
</tr>
<tr>
<td>256</td>
<td>Experimental drugs</td>
<td>Not a benefit</td>
</tr>
<tr>
<td>257</td>
<td>Nonprescription drugs</td>
<td></td>
</tr>
<tr>
<td>258</td>
<td>IV solutions</td>
<td></td>
</tr>
<tr>
<td>259</td>
<td>Other pharmacy</td>
<td></td>
</tr>
<tr>
<td>260</td>
<td>General classification</td>
<td></td>
</tr>
<tr>
<td>261</td>
<td>Infusion pump</td>
<td></td>
</tr>
<tr>
<td>262</td>
<td>IV therapy/pharmacy services</td>
<td></td>
</tr>
<tr>
<td>263</td>
<td>IV therapy/drug/supply delivery</td>
<td></td>
</tr>
<tr>
<td>264</td>
<td>IV therapy/supplies</td>
<td></td>
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<td><strong>Labor Room/Delivery</strong></td>
<td></td>
</tr>
<tr>
<td>720</td>
<td>General classification</td>
<td></td>
</tr>
<tr>
<td>721</td>
<td>Labor</td>
<td></td>
</tr>
<tr>
<td>722</td>
<td>Delivery</td>
<td></td>
</tr>
<tr>
<td>723</td>
<td>Circumcision</td>
<td></td>
</tr>
<tr>
<td>724</td>
<td>Birthing center</td>
<td></td>
</tr>
<tr>
<td>729</td>
<td>Other labor room/delivery</td>
<td></td>
</tr>
<tr>
<td></td>
<td><strong>Electrocardiogram (EKG/ECG)</strong></td>
<td></td>
</tr>
<tr>
<td>730</td>
<td>General classification</td>
<td>Procedure code required</td>
</tr>
<tr>
<td>731</td>
<td>Holter monitor</td>
<td>Procedure code required</td>
</tr>
<tr>
<td>732</td>
<td>Telemetry</td>
<td>Procedure code required</td>
</tr>
<tr>
<td>739</td>
<td>Other EKG/ECG</td>
<td>Procedure code required</td>
</tr>
<tr>
<td></td>
<td><strong>Electroencephalogram (EEG)</strong></td>
<td></td>
</tr>
<tr>
<td>740</td>
<td>General classification</td>
<td>Procedure code required</td>
</tr>
<tr>
<td>741</td>
<td>Other EEG</td>
<td>Procedure code required</td>
</tr>
<tr>
<td></td>
<td><strong>Gastrointestinal Services</strong></td>
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</tr>
<tr>
<td>750</td>
<td>General classification</td>
<td></td>
</tr>
<tr>
<td>759</td>
<td>Other gastrointestinal</td>
<td></td>
</tr>
<tr>
<td>Revenue Code</td>
<td>Description</td>
<td>Comments</td>
</tr>
<tr>
<td>--------------</td>
<td>------------------------------</td>
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</tr>
<tr>
<td>760</td>
<td>General classification</td>
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<tr>
<td>761</td>
<td>Treatment room</td>
<td></td>
</tr>
<tr>
<td>762</td>
<td>Observation room</td>
<td></td>
</tr>
<tr>
<td>769</td>
<td>Other treatment/observation room</td>
<td></td>
</tr>
</tbody>
</table>

**Preventive Care Services**

<table>
<thead>
<tr>
<th>Revenue Code</th>
<th>Description</th>
<th>Comments</th>
</tr>
</thead>
<tbody>
<tr>
<td>770</td>
<td>General classification</td>
<td>Procedure code required</td>
</tr>
<tr>
<td>771</td>
<td>Vaccine administration</td>
<td>Procedure code required</td>
</tr>
<tr>
<td>779</td>
<td>Other preventive care services</td>
<td>Procedure code required</td>
</tr>
</tbody>
</table>

**Lithotripsy**

<table>
<thead>
<tr>
<th>Revenue Code</th>
<th>Description</th>
<th>Comments</th>
</tr>
</thead>
<tbody>
<tr>
<td>790</td>
<td>General classification</td>
<td>Not a benefit</td>
</tr>
<tr>
<td>799</td>
<td>Other lithotripsy</td>
<td>Not a benefit</td>
</tr>
</tbody>
</table>

**Miscellaneous Dialysis**

<table>
<thead>
<tr>
<th>Revenue Code</th>
<th>Description</th>
<th>Comments</th>
</tr>
</thead>
<tbody>
<tr>
<td>880</td>
<td>Miscellaneous dialysis</td>
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</tr>
<tr>
<td>881</td>
<td>Ultrafiltration</td>
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</tr>
</tbody>
</table>

**Other Diagnostic Services**

<table>
<thead>
<tr>
<th>Revenue Code</th>
<th>Description</th>
<th>Comments</th>
</tr>
</thead>
<tbody>
<tr>
<td>920</td>
<td>General classification</td>
<td>Procedure code required</td>
</tr>
<tr>
<td>921</td>
<td>Peripheral vascular lab</td>
<td>Procedure code required</td>
</tr>
<tr>
<td>922</td>
<td>Electromyelogram</td>
<td>Procedure code required</td>
</tr>
<tr>
<td>923</td>
<td>Pap smear</td>
<td>Procedure code required</td>
</tr>
<tr>
<td>924</td>
<td>Allergy test</td>
<td>Procedure code required</td>
</tr>
<tr>
<td>925</td>
<td>Pregnancy test</td>
<td>Procedure code required</td>
</tr>
<tr>
<td>929</td>
<td>Other diagnostic service</td>
<td>Procedure code required</td>
</tr>
<tr>
<td>943</td>
<td>Cardiac rehabilitation</td>
<td>Procedure code required</td>
</tr>
</tbody>
</table>

*Refereto:* The National Uniform Billing Committee (NUBC) website at [www.nubc.org](http://www.nubc.org) for additional information.

### 4.5.6 Third Party Liability

*Refereto:* Subsection 3.7.1, “Medicaid Relationship to Medicare” in this handbook.

“Section 8: Third Party Liability (TPL)” (*Vol. 1, General Information*).

Other Insurance Form on the TMHP website at [www.tmhp.com](http://www.tmhp.com).

Tort Response Form on the TMHP website at [www.tmhp.com](http://www.tmhp.com).

### 5 Ambulatory Surgical Center and Hospital Ambulatory Surgical Center

#### 5.1 Enrollment

To enroll in Texas Medicaid, an ASC must:

- Meet and comply with applicable state and federal laws, rules, regulations, and provisions of the state plan under Title XIX of the Social Security Act.
• Be enrolled in Medicare.
• Meet and comply with state licensure requirements for ASCs.

Providers cannot be enrolled if their license is due to expire within 30 days; a current license must be submitted.

All hospitals enrolling in Texas Medicaid (except psychiatric and rehabilitation hospitals) are issued an HASC provider number at the time of enrollment.

An out-of-state provider may enroll in Texas Medicaid if it is the customary or general practice for clients in a particular locality to use medical resources in another state. An out-of-state provider located within 50 miles of the Texas border is automatically considered to meet this criterion.

Refer to: Subsection 1.1, "Provider Enrollment" in "Section 1: Provider Enrollment and Responsibilities" (Vol. 1, General Information) for more information about enrollment procedures.

Subsection 2.1.1, "Clinical Laboratory Improvement Amendments (CLIA)” in the Radiology and Laboratory Services Handbook (Vol. 2, Provider Handbooks).

The Medicaid Managed Care Handbook (Vol. 2, Provider Handbooks).

5.2 Services, Benefits, Limitations, and Prior Authorization

ASCs, both freestanding and hospital-based, provide same day elective surgery for clients who do not require a hospital admission and who are not expected to require extensive postoperative care.

5.2.1 Drugs and Supplies

Outpatient prescribed medications are a benefit to eligible clients when obtained through a pharmacy contracted with the Medicaid Vendor Drug Program. Prescribed take-home supplies are a benefit to eligible clients when obtained through Medicaid durable medical equipment (DME).

Refer to: Section 1.1, “About the Vendor Drug Program” in the Outpatient Drug Services Handbook (Vol. 2, Provider Handbooks)) for information about this program.

Subsection 2.2.4, “Medical Supplies” in the Durable Medical Equipment, Medical Supplies, and Nutritional Products Handbook (Vol. 2, Provider Handbooks).

5.2.2 Incomplete Surgical Procedures

Refer to: Subsection 4.2.3.6, “Incomplete Day Surgeries” in this handbook for information about incomplete surgical procedures.

5.2.3 Complications Following Day Surgery Requiring Outpatient Observation or Inpatient Admission

If the client is placed in outpatient observation or inpatient status following an HASC day surgery, the day surgery procedure must still be submitted as an outpatient procedure under the HASC provider identifier.

Refer to: Subsection 4.2.3.2, “Complications Following Elective or Scheduled Day Surgeries” in this handbook.

Subsection 4.2.3.4, “Emergency or Unscheduled Day Surgeries” in this handbook in this handbook.

5.2.4 Planned Admission for Day Surgery

Inpatients may occasionally require a surgery that has been designated as an outpatient procedure. The physician must document the need for this surgery as an inpatient procedure before the procedure is performed. These claims are subject to retrospective review.
5.2.5 Cochlear Implants
A cochlear implant is a benefit of Texas Medicaid when medically indicated. ASC and HASC providers may be reimbursed for the implantation procedure using procedure code 69930, and for the cochlear implant devices using procedure code L8614.


5.2.6 Colorectal Cancer Screening
Procedure codes G0104, G0105, and G0121 are a benefit of Texas Medicaid in the ASC or HASC setting.

5.2.6.1 Sigmoidoscopies
Procedure code G0104 may be reimbursed once every 5 years and is limited to one of the following diagnosis codes:

<table>
<thead>
<tr>
<th>Diagnosis Codes</th>
</tr>
</thead>
<tbody>
<tr>
<td>Z0000*</td>
</tr>
<tr>
<td>Z0001*</td>
</tr>
<tr>
<td>Z1210</td>
</tr>
<tr>
<td>Z1211</td>
</tr>
<tr>
<td>Z1213</td>
</tr>
<tr>
<td>Z859</td>
</tr>
<tr>
<td>Z86002</td>
</tr>
<tr>
<td>Z86003</td>
</tr>
</tbody>
</table>

*Diagnosis code Z0000 or Z0001 may be used for screening if no other diagnosis is appropriate for the service rendered, but no more frequently than recommended by the USPSTF.

5.2.6.2 Colonoscopies
Procedure code G0121 may be reimbursed once every ten rolling years for clients who are 45 years of age and older. Procedure code G0121 must be submitted with one of the following diagnosis codes:

<table>
<thead>
<tr>
<th>Diagnosis Codes</th>
</tr>
</thead>
<tbody>
<tr>
<td>Z0000*</td>
</tr>
<tr>
<td>Z0001*</td>
</tr>
<tr>
<td>Z1210</td>
</tr>
<tr>
<td>Z1211</td>
</tr>
<tr>
<td>Z1213</td>
</tr>
</tbody>
</table>

*Diagnosis code Z0000 or Z0001 may be used for screening if no other diagnosis is appropriate for the service rendered, but no more frequently than recommended by the USPSTF.

Procedure code G0105 may be reimbursed once every two years for clients who meet the definition of high-risk. Procedure code G0105 must be submitted with one of the following diagnosis codes:

<table>
<thead>
<tr>
<th>Diagnosis Codes</th>
</tr>
</thead>
<tbody>
<tr>
<td>K5000</td>
</tr>
<tr>
<td>K50011</td>
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<tr>
<td>K50012</td>
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<tr>
<td>K50013</td>
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<tr>
<td>K50014</td>
</tr>
<tr>
<td>K50018</td>
</tr>
<tr>
<td>K5010</td>
</tr>
<tr>
<td>K50111</td>
</tr>
<tr>
<td>K50112</td>
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<tr>
<td>K50113</td>
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<tr>
<td>K50114</td>
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<tr>
<td>K50118</td>
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<td>K5018</td>
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<td>K50812</td>
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<td>K50813</td>
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<td>K50818</td>
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<tr>
<td>K5090</td>
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<tr>
<td>K50911</td>
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<td>K50912</td>
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<td>K50913</td>
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<tr>
<td>K50914</td>
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<td>K50918</td>
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<td>K5289</td>
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<td>K529</td>
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<tr>
<td>Z800</td>
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<td>Z8371</td>
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<td>Z85038</td>
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<td>Z85048</td>
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<td>Z859</td>
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<tr>
<td>Z86002</td>
</tr>
<tr>
<td>Z86003</td>
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<tr>
<td>Z86004</td>
</tr>
<tr>
<td>Z86006</td>
</tr>
<tr>
<td>Z86007</td>
</tr>
<tr>
<td>Z86010</td>
</tr>
</tbody>
</table>

5.2.7 **Dental Therapy Under General Anesthesia**

Facilities must use procedure code 41899 with modifier EP to submit claims for dental therapy under general anesthesia. Prior authorization is not required for ASCs and HASCs unless the client is enrolled in a Medicaid managed care organization.

**Refer to:** Subsection 4.2.29.2.1, “Dental Therapy Under General Anesthesia” in the *Children’s Services Handbook* (Vol. 2, Provider Handbooks).


5.2.8 **Fluocinolone Acetonide**

Procedure code 67027 for implantation may be reimbursed to HASCs. This benefit is limited to clients who are 12 years of age and older and requires prior authorization.

**Refer to:** [Non-emergency Ambulance Prior Authorization Request](https://www.tmhp.com) on the TMHP website.

Subsection 4.2.11, “Fluocinolone Acetonide” in this handbook.

5.2.9 **Implantable Infusion Pumps**

Implantable infusion pumps are a benefit of Texas Medicaid. Implantable infusion pumps may be medically necessary in the following circumstances:

- Administration of intrathecal or epidural antispasmodic drugs to treat refractory intractable spasticity
- Administration of intrathecal, epidural, or central venous analgesic (opioid or non-opioid) drugs for treatment of severe chronic intractable pain
- Administration of intrahepatic chemotherapy for primary liver cancer or metastatic cancer with metastases limited to the liver

An IIP is not a benefit for the following uses:

- Continuous insulin infusion for diabetes
- Continuous heparin infusion for recurrent thromboembolic disease
- Continuous intralesional infusion for severe chronic intractable pain
- Continuous intra-arterial infusion
- Continuous intra-articular infusion for severe chronic intractable pain
- Administration of antibiotics for osteomyelitis

All supplies associated with an IIP are included with the reimbursement for the surgery to implant the infusion pump and are not reimbursed separately.

Providers may be reimbursed for implantable infusion pumps using procedure codes E0782, E0783, and E0786.

If procedure codes E0782 and E0783 are billed with the same date of service, only one may be reimbursed.

**Refer to:** Subsection 9.2.38, “Implantable Infusion Pumps” in the *Medical and Nursing Specialists, Physicians, and Physician Assistants Handbook* (Vol. 2, Provider Handbooks).

5.2.9.1 **Prior Authorization for Implantable Infusion Pump**

Implantable infusion pumps (procedure codes E0782, E0783, and E0786) require prior authorization.
Prior authorization is not required for the physician services associated with the insertion, revision, removal, refilling, or maintenance of the IIP.

Providers must request prior authorization for the implantable infusion pump through the SMPA department with the supporting documentation for medical necessity. Send authorization requests to:

Texas Medicaid & Healthcare Partnership  
Special Medical Prior Authorization  
12357-B Riata Trace Parkway  
Austin, TX 78727  
Fax: 1-512-514-4213

Requests for prior authorization can be submitted online through the TMHP website at [www.tmhp.com](http://www.tmhp.com).


5.2.10 Brachytherapy

The following procedure codes are payable to ASC and HASC facilities:

| Procedure Codes |
|-----------------|-----------------|-----------------|-----------------|
| 19296 | 19297 | 19298 | 31643 | 55860 | 55874 | 55875 | 57155 | 58346 |

Prior authorization is not required for brachytherapy services.


5.2.11 Neurostimulators

Neurostimulators are a benefit of Texas Medicaid when medically necessary. All procedures require prior authorization.


Neurostimulator devices may be reimbursed separately from the global fee.

Refer to: The Texas Medicaid fee schedules on the TMHP website at [www.tmhp.com](http://www.tmhp.com) for procedure codes that may be reimbursed to ASC providers.

5.2.12 Prior Authorization

Some procedures require the performing provider to obtain prior authorization. When prior authorization is required, providers can mail or fax the request to:

Texas Medicaid & Healthcare Partnership  
Special Medical Prior Authorization  
12357-B Riata Trace Parkway  
Austin, TX 78727  
Fax: 1-512-514-4213

Requests for prior authorization can be submitted online through the TMHP website at [www.tmhp.com](http://www.tmhp.com).
5.2.13  Gynecological and Reproductive Health and Family Planning Services

The following gynecological and reproductive health services and family planning services procedure codes may be reimbursed to ASC and HASC providers:

<table>
<thead>
<tr>
<th>Procedure Codes</th>
</tr>
</thead>
<tbody>
<tr>
<td>11976 11981 11982 11983 55250 56620 56805 57335 57410 58260</td>
</tr>
<tr>
<td>58262 58353 58356 58541 58545 58546 58550 58552 58555 58558</td>
</tr>
<tr>
<td>58559 58560 58561 58562 58563 58565 58600 58615 58670 58671</td>
</tr>
<tr>
<td>58673 58674 59150 59151 59840 59841</td>
</tr>
</tbody>
</table>

*Refer to:* The Gynecological, Obstetrics, and Family Planning Title XIX Services Handbook (Vol. 2, Provider Handbooks) for additional information about these procedure codes.

The Texas Medicaid fee schedules on the TMHP website at [www.tmhp.com](http://www.tmhp.com) for additional information about procedure codes that may be reimbursed to ASC providers.

The online fee lookup (OFL) or the static fee schedules at [www.tmhp.com](http://www.tmhp.com) for additional information about rate information for individual procedure codes.

5.3 Documentation Requirements

All services require documentation to support the medical necessity of the service rendered, including ASC and HASC services. ASC and HASC services are subject to retrospective review and recoupment if documentation does not support the service submitted for reimbursement.

5.4 Claims Filing and Reimbursement

5.4.1 Claims Information

Freestanding ASC claims must be submitted to TMHP in an approved electronic claims format or on a CMS-1500 paper claim form. Hospital-based ASCs must submit claims to TMHP in an approved electronic claims format or on a UB-04 CMS-1450 paper claim form.

Claims must contain the billing provider’s complete name, address, and a provider identifier. When completing a UB-04 CMS-1450 or a CMS-1500 paper claim form, providers must include all required information on the claim; TMHP does not key any information from claim attachments. Providers must purchase UB-04 CMS-1450 and CMS-1500 paper claim forms from the vendor of their choice. TMHP does not supply them.

Scheduled procedures performed in a HASC must be submitted for reimbursement using the HASC provider identifier with TOB 131. Emergency or unscheduled procedures performed in a hospital when the client is an outpatient must be submitted for reimbursement using the hospital provider identifier with TOB 131.

To submit claims for services performed by certified registered nurse anesthetists (CRNAs), an ASC must enroll as a CRNA group provider and indicate the CRNA performing provider identifier on claims for those services.

*Refer to:* Section 4, “Certified Registered Nurse Anesthetist (CRNA)” in the Medical and Nursing Specialists, Physicians, and Physician Assistants Handbook (Vol. 2, Provider Handbooks) for specific billing instructions for CRNA services.

“Section 6: Claims Filing” (Vol. 1, General Information).

Section 10, “Claim Form Examples” in this handbook.
5.4.2 Reimbursement

Reimbursement of ASC and HASC procedures is based on the CMS-approved Ambulatory Surgical Code Groupings (1 through 9 per CMS and Group 10 per HHSC) payment schedule. Reimbursement is limited to the lesser of the amount reimbursed to an ASC for similar services, the hospital’s actual charge, or the allowable cost determined by HHSC. When multiple surgical procedures are performed on the same day, only the procedure with the highest surgical code grouping is reimbursed. A complete list of approved ASC and HASC procedure codes with the assigned payment group can be found on the TMHP website at [www.tmhp.com](http://www.tmhp.com). Click on Resources and then Online Fee Lookup. This list can also be obtained by calling the TMHP Contact Center at 1-800-925-9126.

Claims for physician and CRNA services performed in an ASC or HASC must be submitted under the physician or CRNA provider identifier and are reimbursed separately.

5.4.2.1 ASC and HASC Global Services

The ASC or HASC payment represents a global payment and includes room charges and supplies. Covered services provided are submitted as one inclusive charge. All facility services provided in conjunction with the surgery (e.g., laboratory, radiology, anesthesia supplies, medical supplies) are considered part of the global payment and cannot be itemized or submitted separately.

Routine X-ray and laboratory services directly related to the surgical procedure being performed are not reimbursed separately. All nonroutine laboratory and X-ray services provided with emergency conditions may be submitted separately with documentation that the complicating condition arose after the initiation of the surgery.

Medical and prosthetic devices such as intraocular lenses may be supplied by the ASC or HASC and implanted, inserted, or otherwise applied during a covered surgical procedure and is considered part of the global surgical fee.

**Exception:** Certain pieces of equipment, (e.g., cochlear implants, implantable infusion pumps, and neurostimulator devices) may be reimbursed separately from the ASC or HASC global rate.

**Refer:** Subsection 2.2, “Fee-for-Service Reimbursement Methodology” in “Section 2: Texas Medicaid Fee-for-Service Reimbursement” (Vol. 1, General Information) for more information about reimbursement.

Subsection 4.2.3, “Day Surgery” in this handbook for information about HASCs.

5.4.2.2 NCCI and MUE Guidelines

The HCPCS and CPT codes included in the Texas Medicaid Provider Procedures Manual are subject to NCCI relationships, which supersede any exceptions to NCCI code relationships that may be noted in the manuals. Providers should refer to the [CMS NCCI web page](http://www.cms.gov) for correct coding guidelines and specific applicable code combinations.

In instances when Texas Medicaid limitations are more restrictive than NCCI MUE guidance, Texas Medicaid limitations prevail.

6 Military Hospitals

6.1 Military Hospital Enrollment

To enroll in Texas Medicaid, a military hospital must be certified by Medicare, have a valid provider agreement with HHSC, and have completed the TMHP enrollment process. Veterans Administration (VA) hospitals are eligible to receive Texas Medicaid payment only on claims that have crossed over from Medicare.
Military hospital providers must comply with CLIA rules and regulations. Providers who do not comply with CLIA will not be reimbursed for laboratory services.

6.2 Services, Benefits, Limitations and Prior Authorization

6.2.1 Military Hospital Inpatient Services

Inpatient hospital services include medically necessary items and services ordinarily furnished by a Medicaid hospital or by an approved out-of-state hospital under the direction of a physician for the care and treatment of inpatient clients. Reimbursement to hospitals for inpatient services is limited to the Medicaid “spell of illness.” The spell of illness is defined as “30 days of inpatient hospital care, which may accrue intermittently or consecutively.”

After 30 days of inpatient care have been provided, reimbursement for additional inpatient care is not considered until the client has been out of an acute care facility for 60 consecutive days. Exceptions are made in the following instances:

- THSteps-eligible clients do not have a 30-day spell of illness limitation, if medically necessary conditions exist (covered under THSteps-CCP).

Refer to: The Medicaid Managed Care Handbook (Vol. 2, Provider Handbooks).


Hospitals may submit information only claims to TMHP when one of the following situations exists:

- The inpatient 30-day spell of illness benefit is exhausted.
- Payment that was made by a third party resource or other insurance exceeds the Medicaid allowed amount.

For clients who are 21 years of age and older, there is an inpatient expenditure cap of $200,000 per benefit year (November 1 through October 31). Claims are reviewed retrospectively, and payments exceeding $200,000 will be recouped.

It is appropriate to submit information only claims using TOB 110.

The following hospital services must be medically necessary and are subject to the utilization review requirements of Texas Medicaid. Medicaid reimbursement for services cannot exceed the limitations of Texas Medicaid.

Inpatient hospital services include the following items and services:

- Bed and board in semiprivate accommodations or in an intensive care or coronary care unit, including meals, special diets, and general nursing services; or an allowance for bed and board in private accommodations, including meals, special diets, and general nursing services up to the hospital’s charge for its most prevalent semiprivate accommodations. Bed and board in private accommodations are provided in full if required for medical reasons, as certified by the physician. Additionally, the hospital must document the medical necessity for a private room, such as the existence of a critical or contagious illness or a condition that could result in disturbance to other patients. This type of information is included in Block 80 or attached to the claim.

- Whole blood and packed red cells that are reasonable and necessary for treatment of illness or injury, provided they are not available without cost.

- All medically necessary services or supplies ordered by a physician.

Medicaid benefits are not available for take-home or self-administered drugs or personal comfort items except when received by prescription through the VDP.

Only inpatient claims that have an emergency diagnosis on the claim are considered for reimbursement.
6.2.2 Military Hospital Outpatient and Physician Services
Although Medicare reimburses for emergency outpatient and inpatient services, Medicaid does not reimburse for either outpatient or physician services. Military hospitals are not reimbursed for outpatient day surgery.

6.2.3 Prior Authorization
Prior authorization is not required for services rendered in military hospitals.

6.3 Documentation Requirements
All services require documentation to support the medical necessity of the service rendered, including military hospital services. Military hospital services are subject to retrospective review and recoupment if documentation does not support the service submitted for reimbursement.

6.3.1 Documentation for Nursing Facility Admissions
The admission Minimum Data Set (MDS) must be used for admissions to a nursing facility. There are instances in which hospital social workers and discharge nurses might also complete the admission MDS, such as:

- If the client is in a long-term care acute center.
- If the potential receiving nursing facility wants a better clinical picture of the client, a paper copy of the admission MDS is completed by the hospital staff before the client is accepted for admission into the nursing facility.

Refer to: The Long Term Care Program page on the TMHP website at www.tmhp.com for additional information, including instructions for all forms and assessments.

6.4 Claims Filing and Reimbursement
6.4.1 Military Hospital Claims Information
If TOB 110 is used to submit a claim, all charges must be noncovered and the claim will finalize with EOB 217, “Payment reduced through hospital action.”

It is appropriate to submit information only claims using TOB 110.

Military hospitals may submit total charges in one line with appropriate accommodation revenue codes. Emergency hospital services must be submitted to TMHP in an approved electronic format or on the UB-04 CMS-1450 paper claim form. Providers may purchase claim forms from the vendor of their choice. TMHP does not supply the forms.

Refer to: “Section 3: TMHP Electronic Data Interchange (EDI)” (Vol. 1, General Information) for information on electronic claims submissions.

“Section 6: Claims Filing” (Vol. 1, General Information) for general information about claims filing.

When completing a UB-04 CMS-1450 paper claim form, all required information must be included on the claim, as TMHP does not key any information from claim attachments. Superbills, or itemized statements, are not accepted as claims supplements.

Refer to: Subsection 6.6, “UB-04 CMS-1450 Paper Claim Filing Instructions” in “Section 6: Claims Filing” (Vol. 1, General Information) for paper claims completion instructions. Blocks that are not referenced are not required for processing by TMHP and may be left blank.

Military Hospital (Emergency Inpatient) on the TMHP website at www.tmhp.com for a claim form example.
6.4.2 Military Hospital Reimbursement

Reimbursement is limited to claims submitted for emergency inpatient care only.

Allowed inpatient hospital stays are reimbursed according to a prospective payment methodology based on DRGs. The reimbursement method itself does not affect inpatient benefits and limitations. Texas Medicaid requires that one claim be submitted for each inpatient stay with appropriate diagnosis and procedure code sequencing. Providers must submit only one claim per inpatient stay to Medicaid, regardless of the diagnosis, to ensure accurate payment. The DRG reimbursement includes all facility services provided to the client while registered as an inpatient.

Reimbursement to hospitals for inpatient services is limited to $200,000 per client, per benefit year (November 1 through October 31). This limitation does not apply to services related to certain organ transplants or services to clients who are 20 years of age and younger and covered by the CCP.

Military hospitals should keep a Medicaid client as an inpatient for only the length of time necessary to stabilize the client. The Medicaid client, once stabilized, should be transferred to the nearest Medicaid acute care hospital facility for further treatment.

When more than one hospital provides care for the same client, the hospital that furnishes the most significant amount of care receives consideration for a full DRG payment.

The other hospital is paid a per diem rate based on the lesser of the mean length of stay for the DRG or eligible days in the facility.

Client transfers within the same facility or readmissions to the same facility within 24 hours of a previous acute hospital or facility discharge are considered one continuous stay. These readmissions are considered a continuous stay regardless of the original or readmission diagnosis. Texas Medicaid does not recognize specialty units within the same hospital as separate entities; therefore, these transfers must be included in one submission under the provider identifier. Admissions that were submitted inappropriately are identified and denied during the utilization review process and may result in an intensified review.

After all hospital claims have been submitted, TMHP performs a post-payment review to determine if the hospital furnishing the most significant amount of care received the full DRG. If the review reveals that the hospital furnishing the most significant amount of care did not receive the full DRG, an adjustment is initiated.

The inpatient DRG reimbursement includes payment for all radiology and laboratory services, including those sent to referral laboratories.

Refer to: Subsection 2.2, “Fee-for-Service Reimbursement Methodology” in “Section 2: Texas Medicaid Fee-for-Service Reimbursement” (Vol. 1, General Information) for more information about reimbursement.

7 Claims Resources

Refer to the following sections and forms when filing claims:

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<td>Subsection A.10, “TMHP Telephone and Fax Communication” in “Appendix A: State, Federal, and TMHP Contact Information” (Vol. 1, General Information)</td>
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CMS-1500 Paper Claim Filing Instructions | Subsection 6.5, “CMS-1500 Paper Claim Filing Instructions” in “Section 6: Claims Filing” (Vol. 1, General Information)
State, federal, and TMHP contact information | “Appendix A: State, Federal, and TMHP Contact Information” (Vol. 1, General Information)
TMHP electronic claims submission information | Subsection 6.2, “TMHP Electronic Claims Submission” in “Section 6: Claims Filing” (Vol. 1, General Information)
TMHP Electronic Data Interchange (EDI) information | “Section 3: TMHP Electronic Data Interchange (EDI)” (Vol. 1, General Information)

### 8 Contact TMHP

*Note:* The TMHP Contact Center at 1-800-925-9126 is available Monday through Friday from 7 a.m. to 7 p.m., Central Time.

### 9 Forms

The following linked forms can also be found on the [Forms](#) page of the Provider section of the TMHP website at [www.tmhp.com](http://www.tmhp.com):

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<td>Sterilization Consent Form (English)</td>
</tr>
<tr>
<td>Sterilization Consent Form (Spanish)</td>
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### 10 Claim Form Examples

The following linked claim form examples can also be found on the [Claim Form Examples](#) page of the Provider section of the TMHP website at [www.tmhp.com](http://www.tmhp.com):

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1 General Information

The information in this handbook is intended for Texas Medicaid managed care providers, including providers who are enrolled in a managed care organization (MCO) that is contracted by Texas Medicaid to provide managed care coverage for Texas Medicaid clients.

This handbook provides information about the following managed care programs and services:

- STAR
- STAR+PLUS
- STAR Kids
- STAR Health
- Children’s Medicaid Dental Services

Refer to: The Medicaid and CHIP Programs page of the HHS website at hhs.texas.gov.

2 Overview of Medicaid Managed Care

Texas Medicaid, which is administered by the Texas Health and Human Services Commission (HHSC), operates Medicaid managed care under the authority of federal waivers and state plan amendments that were approved by the Centers for Medicare & Medicaid Services (CMS).

Medicaid managed care is administered by MCOs and dental maintenance organizations (DMOs) that are contracted by HHSC to provide services for Medicaid managed care clients. The Medicaid managed care MCOs and DMOs provide most of the same benefits that Texas Medicaid provides for Medicaid fee-for-service clients. Some plans may also offer value-added services.

Refer to: Subsection 8, “Carve-Out Services” in this handbook.

The principal objectives of Medicaid managed care are early intervention and improved access to quality care, which significantly improve health outcomes for clients, with a special focus on prenatal and well-child care.

In Medicaid managed care, clients assume responsibility for their personal health care by choosing a health plan and primary care provider and by making use of preventive primary care services. Eligible clients may also choose a DMO and a main dentist. This collaborative approach to health-care delivery helps to reduce costs by eliminating duplicate services and unnecessary emergency and inpatient care.

Medicaid managed care uses MCOs to cover services in 13 service areas (SAs).

Medicaid managed care consists of the following programs:

- The STAR program uses MCOs to cover acute care services for children, newborns, pregnant women, and some families. STAR is available statewide, and clients have the choice of at least two STAR MCOs in each SA. The STAR program operates under a federal 1115 waiver.
- The STAR Kids program uses MCOs to cover integrated acute care and long-term services and supports (LTSS) for children and adults who are 20 years of age or younger and have a disability. STAR Kids is available statewide, and clients have the choice of at least two STAR Kids MCOs in each SA. The STAR Kids program operates under a federal 1115 waiver.
- The STAR+PLUS program uses MCOs to cover integrated acute care and LTSS for adults who are 65 years of age or older (including those who are dually eligible for Medicare and Medicaid) and adults who are 21 years of age or older and have a disability. STAR+PLUS also serves those who are...
eligible for Medicaid for Breast and Cervical Cancer (MBCC). The STAR+PLUS program is available statewide, and clients have the choice of at least two STAR+PLUS MCOs in each SA. The STAR+PLUS program operates under a federal 1115 waiver.

- The STAR Health program uses one MCO to deliver health-care and dental services to children who are in foster care throughout the state. STAR Health is administered by Superior HealthPlan and operates under federal 1915(a) authority. The STAR Health program manages the health and dental care of most children in foster care.
- Children’s Medicaid dental services are administered by DMOs that process dental authorization requests and claims for most Medicaid fee-for-service and Medicaid managed care clients who are 20 years of age or younger regardless of their medical benefit plan.

Refer to: Section 7, “Children’s Medicaid Dental Services” in this handbook for exceptions and additional information.

Refer to: The Medicaid Managed Care page of the Provider section of the TMHP website at www.tmhp.com for a current list of managed care service areas.

2.1 Managed Care Services

MCOs and DMOs administer most of the services that are rendered to Medicaid managed care clients, including, but not limited to:

- Professional, inpatient facility, and outpatient facility medical services.
- Prescription drug/pharmacy services.
- Children’s Medicaid dental services for most clients who are 20 years of age or younger.
- Orthodontia services.
- Services rendered to Medicaid managed care SSI clients.
- Value-added services that an individual MCO or DMO elects to provide.

Administrative procedures, such as prior authorization, precertification, referrals, and claims/encounter data filing may differ from traditional Medicaid (fee-for-service) and from MCO to MCO. Providers should contact the client’s specific MCO for details.

2.1.1 Medical Services

Most medical service benefits including professional, inpatient, and outpatient services rendered to Medicaid managed care clients are administered by individual MCOs. Medical services include all those administered by TMHP for fee-for-service clients as well as any value-added services covered by the individual MCOs.

Some services rendered to Medicaid managed care clients are considered “carve-out” services. Carve-out services are administered and paid by TMHP and not by the client’s MCO.

Refer to: Section 8, “Carve-Out Services” in this handbook.

2.1.2 Prescription Drug/Pharmacy Services

The client’s MCO administers and pays for pharmacy services.

Pharmacy providers must first be enrolled by the Medicaid or Children’s Health Insurance Program (CHIP) Vendor Drug Program before they can enroll with an MCO.

Refer to: Subsection 2.2, “Provider Enrollment and Responsibilities” in this handbook.

Generally, there is no monthly prescription limit for managed care clients.
Refer to: The MCO that administers the client’s Medicaid managed care benefits for information about prescription drug and pharmacy benefits.

Each MCO contracts one Pharmacy Benefit Manager (PBM). The MCOs and PBMs must adhere to Medicaid preferred drug list (PDL) and HHSC Medicaid and CHIP formularies. HHSC manages the Texas Medicaid and CHIP formularies.

Important: MCOs and PBMs cannot require clients to use a mail-order pharmacy.

The MCOs must:

- Perform drug utilization review for managed care clients.
- Monitor pharmacy providers for compliance.
- Establish help lines for providers and clients.
- Ensure that all clients have access to a minimum of one network pharmacy:
  - Within 15 miles of the client’s residence.
  - With 24-hour coverage within 75 miles of the client’s residence.
- Provide e-prescribing abilities to:
  - Verify client eligibility.
  - Review medication history.
  - Review formulary and PDL information.
- Process correct pharmacy claims that are submitted electronically within 18 days of submission.

2.1.2.1 Prescription Drug Prior Authorizations

Prescribers may be required to request prior authorization for a prescription drug. The prescriber must contact the client’s MCO or PBM and follow MCO or PBM guidelines and procedures for prior authorization requests.

Important: TMHP does not have access to the MCOs’ or PBMs’ guidelines and procedures for prior authorizations. The provider must contact the MCOs of PBMs for information. Individual PBMs will have their own PA processes and telephone lines.

The MCO must notify the prescriber’s office of a prior authorization approval or denial:

- Within 24 hours of a request submitted by fax or web.
- Immediately for telephone requests.

Prior authorization is required for non-preferred drugs or any drug requiring a clinical prior authorization.

If the pharmacy cannot dispense the client’s prescription because prior authorization is required but it has not been requested, the pharmacy should contact the MCO or PBM to request prior authorization. The prescribing provider is required to submit certain prior authorization requests including, but not limited to, non-preferred drug prior authorizations.

2.1.2.1.1 Emergency 72-Hour Prescriptions

If the prescribing provider cannot be reached or is unable to request a prior authorization, the pharmacy should submit an emergency 72-hour prescription. The request for an emergency 72-hour prescription claim should not be used for routine and continuous overrides.
A 72-hour emergency prescription will be paid in full to pharmacy providers and does not count toward the three-prescription limit for adults who have not already received their maximum prescriptions for the month.

**Reminder:** There is no prescription limit for clients who are 20 years of age or younger.

Federal and Texas law allow a 72-hour emergency supply of prescribed medication to be dispensed any time a prior authorization is not available and the prescription must be filled without delay for a medical condition. This rule applies to non-preferred drugs on the Preferred Drug list and any drug for which prior authorization must be requested by the prescribing physician.

2.1.2.1.2 Formulary

The MCOs or PBMs are responsible for informing network providers about how to access the formulary and PDL.

**Refer to:** The Medicaid and CHIP formularies on the VDP website at [www.txvendordrug.com](http://www.txvendordrug.com) and [www.epocrates.com](http://www.epocrates.com) for more information.

MCOs may also selectively contract with pharmacies for specialty drugs.

2.1.3 Service and Care Coordination

Texas Medicaid has different types of service and care coordination. Clients who are enrolled in Medicaid may receive service or care coordination from MCOs or providers.

2.1.3.1 Service Coordination in STAR Kids and STAR+PLUS

In STAR Kids and STAR+PLUS, service coordination is specialized care management. Service coordination is provided by STAR Kids and STAR+PLUS MCOs and is available to all clients.

MCO service coordinators:

- Identify physical health, mental health, and LTSS needs and develop a service plan.
- Help clients get timely access to providers and covered services.
- Coordinate covered services with non-managed care programs, such as FFS waivers.

2.1.3.2 Service Management in STAR and CHIP

In STAR, service management is an administrative service that is provided by STAR and CHIP MCOs. Service management is available to STAR and CHIP clients who have a serious, ongoing illness, a chronic or complex condition, a disability, or who require regular ongoing therapeutic intervention and evaluation by appropriately trained personnel.

MCO service managers:

- Work with clients to develop a service plan and coordinate services with the client’s primary care provider, specialty providers, and non-medical providers.
- Help clients get access to and use appropriately medically necessary covered services, services with non-managed care programs, such as FFS waivers, and other services and supports.

2.1.3.3 STAR Health Service Coordination and Service Management

In STAR Health, service coordination and service management have distinct functions that are different from the other managed care programs.

STAR Health defines service coordination and service management as follows:

- Service coordination is an administrative service that helps caregivers manage information, such as medical information for court hearings, and coordinate services with non-managed care programs, such as FFS waivers. Service coordination is available to STAR Health clients who have a medical or behavioral need or as requested.
• Service management is a clinical service that is provided to clients who have a complex medical or behavioral need. Service management helps the client implement a service plan, and it coordinates services between the client’s primary care provider and specialty care providers to ensure that the client has access to and appropriately uses medically necessary covered services. Service management may also be requested.

2.1.3.4 Provider Administered Care Coordination

Several types of care coordination are administered at the provider level. These services do not duplicate the service coordination or service management provided by MCOs.

Care coordination services include targeted case management for:
• Early Childhood Intervention (ECI).
• Individuals with intellectual or developmental disabilities.
• Clients who have a mental illness.
• Children and pregnant women.
• Community Living Assistance and Support Services (CLASS) and Deaf Blind with Multiple Disabilities (DBMD).
• Home- and Community-Based Services (HCBS)-Adult Mental Health (AMH) recovery management.

2.2 Provider Enrollment and Responsibilities

Important: All providers are required to read and comply with Section 1: Provider Enrollment and Responsibilities. In addition to required compliance with all requirements specific to Texas Medicaid, it is a violation of Texas Medicaid rules when a provider fails to provide health-care services or items to Medicaid clients in accordance with accepted medical community standards and standards that govern occupations, as explained in Title 1 Texas Administrative Code (TAC) §371.1659. Accordingly, in addition to being subject to sanctions for failure to comply with the requirements that are specific to Texas Medicaid, providers can also be subject to Texas Medicaid sanctions for failure, at all times, to deliver health-care items and services to Medicaid clients in full accordance with all applicable licensure and certification requirements including, without limitation, those related to documentation and record maintenance.

Refer to: “Section 1: Provider Enrollment and Responsibilities” (Vol. 1, General Information) for more information about enrollment procedures.

Subsection 1.1, “About the Vendor Drug Program” in the Outpatient Drug Services Handbook (Vol. 2, Provider Handbooks) for information about this program.

2.2.1 Enrollment, Contracting, and Credentialing

Providers must be enrolled in Texas Medicaid before they can be enrolled by an MCO or DMO. Individual MCOs and DMOs have their own guidelines for contracting providers.

All Medicaid MCOs must use the Texas Association of Health Plans’ (TAHP’s) contracted credentialing verification organization (CVO) as part of its credentialing and recredentialing process regardless of membership in the TAHP.

Important: Enrollment in Texas Medicaid does not guarantee that an MCO or DMO will contract enroll or credential a particular provider.
Providers must refer all questions about enrollment to the MCO or DMO that administers the clients’ managed care benefits. TMHP does not have access to the enrollment requirements for the individual MCOs and DMOs.

All questions about Texas Medicaid enrollment can be referred to the TMHP Contact Center.

**Note:** Providers who render only carve-out services are not required to enroll in Medicaid MCOs and DMOs.

**Refer to:** Section 8, “Carve-Out Services” in this handbook for a list of services that are carved out of the Medicaid Managed Care program.

Subsection 1.1, “About the Vendor Drug Program” in the Outpatient Drug Services Handbook (Vol. 2, Provider Handbooks) for information about this program.

### 2.2.2 Online Provider Lookup (OPL)

Clients can search for providers using a particular county, SA, or name to find providers who participate in a managed care area. Providers that participate in MCOs and DMOs are responsible for declaring themselves as managed care providers on the OPL. Clients can search for providers that are enrolled in Medicaid managed care on the OPL. The OPL has links to the websites of the MCOs and DMOs, which allows clients to search each MCO’s and DMO’s network of participating providers.

### 2.2.3 Terminated Enrollment

Texas Medicaid monitors provider claim activity. Providers that have not submitted a claim to Texas Medicaid or a Medicaid MCO or DMO within an 18-month period are notified that their Texas Medicaid enrollment will be terminated at 24 months if they have not submitted any claims.

If a provider’s Texas Medicaid enrollment is terminated, the provider’s Medicaid managed care contracts with individual MCOs or DMOs will also be terminated.

To reactivate a TPI that has been terminated, the provider must complete the Texas Medicaid Provider Enrollment Application.

### 2.2.4 Excluded Entities and Providers

Title 42 Code of Federal Regulations (CFR) §1003.400(a)(2) states that civil monetary penalties may be imposed against MCOs that employ or enter into contracts with excluded individuals or entities to provide items or services to Medicaid clients. No Medicaid payments can be made to an MCO for any items or services directed or prescribed by an excluded physician or other authorized person if the MCO either knew or should have known of the exclusion. This prohibition applies even when the Medicaid payment itself is made to another provider, practitioner, or supplier that is not excluded.

### 2.2.5 Accounts Receivable

Providers that have outstanding accounts receivables on their weekly Remittance & Status (R&S) reports must settle them with TMHP even if they no longer submit claims to TMHP.

Payments from the MCOs and DMOs may be held until the debt with TMHP is resolved. Providers can refund payments to TMHP as follows:

- If the provider no longer receives claim payments from TMHP, the provider must issue a check for the refund amount to TMHP. Payment options may be available. If a refund check is mailed to TMHP, the provider must also submit Form 7.2, “Texas Medicaid Refund Information Form.”

- If the provider continues to receive claim payments from TMHP, a recoupment of the funds may be requested through the paper appeal process. If the provider requests a recoupment through the paper appeal process, the provider must not issue a check to TMHP. The refund amounts will be deducted from future payments, and the deductions will appear on the provider’s R&S Reports.
2.2.6  Educating Clients about Managed Care

Providers cannot enroll Medicaid clients; however, providers are encouraged to educate clients about Medicaid managed care.

Providers that participate in one or more Texas Medicaid managed care plans should follow these rules when educating clients:

- Providers may not influence clients to choose one MCO or DMO over another.
- Providers must inform clients of all Medicaid managed care health plans and DMOs in which the providers participate.
- Providers and subcontractors may only directly contact potential clients with whom they have an established relationship.
- Providers may inform clients of special services offered by all Medicaid managed care health and DMOs in which the providers participate.
- Providers may inform clients of particular hospital services, specialists, or specialty care available in all plans in which the providers participate.
- Providers may assist a client by contacting a plan (or plans) to determine if a particular specialist or service is available, if the client requests this information.
- Providers may inform clients of special services offered by all Medicaid managed care health and DMOs in which the providers participate.
- Providers may assist a client by contacting a plan (or plans) to determine if a particular specialist or service is available, if the client requests this information.
- Providers may not influence clients based on reimbursement rates or methodology used by a particular plan.
- At the client’s request, providers can provide the necessary information for the client to contact a particular plan but cannot promote any plan over another.
- In no instances can providers stock, reproduce, assist in filling out, or otherwise handle the enrollment form. Information can be provided as outlined on the previous page, and clients can be reminded that they can easily enroll over the telephone with the enrollment broker. However, the call must be made by the client, not by the provider or the provider’s agent.
- Providers may assist clients with completing the Medicaid application.
- Providers may display stickers that indicate that they participate in a particular Medicaid managed care health or DMO as long as they do not indicate anything more than “(health plan or DMO) is accepted or welcomed here” (provided the sticker meets Medicaid/CHIP Marketing Guidelines regarding size limitations).
- Providers may display state-approved, health-related marketing materials in their offices, provided it is done equally for all MCOs and DMOs in which they participate. MCO and DMO providers cannot give out or display plan-specific marketing items or giveaways to clients.

Important:  Providers must comply with their applicable licensing agency’s laws and regulations, including any related to marketing and advertising, and any applicable state and federal laws and regulations, contractual requirements, and other guidance documents. Providers are encouraged to review the Provider Marketing Guidelines page of the HHS website.

2.3  General Information About Client Enrollment in Managed Care

Most of the clients who have been determined to be eligible for Texas Medicaid are first enrolled in fee-for-service. Specific client groups within the Texas Medicaid population are eligible for managed care based on certain established criteria. If the client is eligible for Medicaid managed care, the client will choose an MCO and primary care provider or a DMO and main dentist or both. The managed care enrollment date is generally separate from the Medicaid eligibility date. In most cases, Medicaid managed care enrollment is not retroactive.
Claim and authorization transactions for services rendered during the client’s fee-for-service eligibility must be submitted to TMHP, and claim and authorization transactions for services rendered during the client’s Medicaid managed care enrollment must be submitted to the appropriate entity (i.e., TMHP for carve-out services and the MCO or DMO for managed care services).

If a client loses Medicaid eligibility and then regains eligibility within a certain amount of time, the client is automatically reassigned to the same health plan and primary care provider or DMO that the client had before the client lost Medicaid eligibility.

**Method of Enrollment**

<table>
<thead>
<tr>
<th>Description</th>
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<tbody>
<tr>
<td>Telephone</td>
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<td>Mail</td>
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<td>Onsite</td>
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</tbody>
</table>
2.3.1.1  Managed Care Enrollment Timeline

Example 1
Benefits under Medicaid managed care usually begin on the first day of the next month following the client’s selection of a managed care plan and primary care provider. The following example shows the managed care enrollment date for a client who selects a health plan and primary care provider before the designated cutoff date (approximately the 15th of the month):

<table>
<thead>
<tr>
<th>Task</th>
<th>Date</th>
</tr>
</thead>
<tbody>
<tr>
<td>Client certified for Texas Medicaid</td>
<td>January 1</td>
</tr>
<tr>
<td>Medicaid benefits begin</td>
<td>January 1</td>
</tr>
<tr>
<td>Client selects health plan and primary care provider (before the 15th of the month)</td>
<td>January 10</td>
</tr>
<tr>
<td>Managed care benefits begin</td>
<td>February 1</td>
</tr>
</tbody>
</table>

Example 2
The following example shows the managed care enrollment date for a client who selects a health plan and primary care provider after the designated cutoff date (approximately the 15th of the month):

<table>
<thead>
<tr>
<th>Task</th>
<th>Date</th>
</tr>
</thead>
<tbody>
<tr>
<td>Client certified for Texas Medicaid</td>
<td>January 10</td>
</tr>
<tr>
<td>Medicaid benefits begin</td>
<td>January 10</td>
</tr>
<tr>
<td>Client selects health plan and primary care provider (after the 15th of the month)</td>
<td>January 20</td>
</tr>
<tr>
<td>Managed care benefits begin</td>
<td>March 1</td>
</tr>
</tbody>
</table>

2.3.1.2  Eligibility Verification Resources

The provider must verify the client’s eligibility before providing services except in cases of emergency. If emergency care is provided, the provider should determine the client’s MCO and primary care provider as soon as possible. The provider must also verify and abide by the prior authorization and administrative requirements established by the MCO or DMO.

STAR MCOs provide their clients with an MCO identification card. Providers should check the MCO identification card and the Your Texas Benefits Medicaid card to determine whether the client is a managed care client.

*Refer to:* “Section 4: Client Eligibility” (Vol. 1, General Information) for more information.
The client’s MCO and DMO enrollment information can be verified by:

- Checking the client’s MCO or DMO ID card (if applicable).
- Visiting the client’s Medicaid MCO or DMO’s website.
- Calling the client’s MCO or DMO.
- Visiting TexMedConnect, accessing the Medicaid Client Portal for Providers, or calling the TMHP Contact Center at 1-800-925-9126.

The client’s managed care eligibility can also be verified using:

- The TMHP Automated Inquiry System (AIS) at 1-800-925-9126.
- Third-party software that uses the TMHP EDI Gateway.
- National Council for Prescription Drug Programs (NCPDP) Eligibility Verification (E1) transaction. The E1 transaction is submitted through a pharmacy’s point-of-sale system.
- The Vendor Drug Eligibility Verification Portal (EVP). EVP is a browser-based application that is free for all contracted pharmacy providers.

Refer to: Subsection 4.4.3, “Client Eligibility Verification” in “Section 4: Client Eligibility” (Vol. 1, General Information) for additional information about verifying client eligibility.

The Eligibility Verification page of the Texas Medicaid Vendor Drug Program website at www.txvendordrug.com for more information.

2.3.2 Enrollment in Managed Care for Specific Groups

2.3.2.1 Expedited Enrollment of Pregnant Women (Program Type 40)

If a pregnant woman applies for program type 40 and is eligible for Medicaid managed care, she has 15 days from the date on which the Enrollment Broker receives the eligibility determination to choose an MCO that participates in her managed care program. If she does not choose an MCO, one will be chosen for her.

The Enrollment Broker contacts the client to begin the enrollment process and helps the client select an MCO. The client can also contact the Enrollment Broker directly. To protect continuity of care and client choice, the Enrollment Broker will work with each pregnant woman to select an MCO that includes her current prenatal care provider or to choose an obstetrical care provider that meets her needs.

Clients will be covered under Texas Medicaid fee-for-service until their Medicaid MCO coverage begins. To ensure proper billing, providers should call the Enrollment Broker to obtain the name of the client’s health plan. However, client eligibility should always be verified at the time the service is rendered.

Women who are certified as Medicaid program type 40 may be retroactively enrolled in managed care. Women who are certified as Medicaid program type 40 on or before the 10th of the month will be enrolled in managed care beginning on the first day of the month of certification. Those who are certified after the 10th of the month will be enrolled in Texas Medicaid fee-for-service for the month of certification and will be enrolled in managed care beginning on the first day of the month following the month of certification.

There are exceptions to this rule:

- Women who are certified at any time in their estimated month of delivery will be enrolled in managed care on the first day of the following month (prospective enrollment).
- Women who are certified at any time in their actual month of delivery (if known by HHSC before certification) will be enrolled in managed care on the first day of the following month (prospective enrollment).
• Women who are not identified as a candidate for program type 40 within three days of certification after her enrollment has been processed will be enrolled in managed care prospectively.

• In cases where the Medicaid effective date is a future date, prospective enrollment rules will be used.

**Important:** Providers must verify the client’s plan and primary care provider information.

The following examples show when benefits begin in relation to certification:

<table>
<thead>
<tr>
<th>Example 1: Woman Certified in Her 6th Month</th>
</tr>
</thead>
<tbody>
<tr>
<td>Client certified for Texas Medicaid</td>
</tr>
<tr>
<td>Medicaid benefits begin</td>
</tr>
<tr>
<td>Managed care start date</td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th>Example 2: Woman Certified in Her 6th Month</th>
</tr>
</thead>
<tbody>
<tr>
<td>Client certified for Texas Medicaid</td>
</tr>
<tr>
<td>Medicaid benefits begin as fee-for-service</td>
</tr>
<tr>
<td>Managed care start date</td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th>Example 3: Woman Certified in Her 9th Month</th>
</tr>
</thead>
<tbody>
<tr>
<td>Client certified for Texas Medicaid</td>
</tr>
<tr>
<td>Medicaid benefits begin as fee-for-service</td>
</tr>
<tr>
<td>Managed care start date</td>
</tr>
</tbody>
</table>

**Note:** Expedited enrollments of pregnant women (program type 40) into Medicaid managed care may be retroactive.

### 2.3.2.2 Newborn Enrollment

The effective date of the newborn’s enrollment is the same as the newborn’s date of birth. Claims for services provided to newborns should be filed with the mother’s MCO. Health-care providers should file newborn claims using the newborn’s Medicaid identification number as soon as the number is made available. Providers that file claims for services provided to newborns are still responsible for meeting the Medicaid filing deadlines, which, in most cases, is within 95 days of the date of service.

MCOs must pay providers for labor- and delivery-related inpatient and professional services that are rendered to mothers for:

• Up to 48 hours following an uncomplicated vaginal delivery.

• Up to 96 hours following an uncomplicated Caesarian delivery.

MCOs must provide neonatal care for newborn clients until the time of discharge. Prior authorizations and primary care provider assignments cannot be a reason to deny claims.

MCOs must notify providers involved in the care of mothers and newborn clients, including out-of-network providers and hospitals, of the MCO’s prior authorization requirements. MCOs cannot require prior authorization for services provided to a mother or newborn client for a medical condition that requires emergency services, regardless of when the emergency condition arises.

Authorization requests, utilization review questions, and claim status inquiries and appeals should be directed to the MCO in which the client is enrolled.

**Note:** Telephone numbers and addresses for MCO claims submission and appeals can be found in the appropriate MCO provider policies and procedures manual for the appropriate service.
In the STAR Program, newborns are automatically assigned to the STAR MCO the mother is enrolled with at the time of the newborn’s birth for at least 90 days following the date of birth unless the mother requests a plan change as a special condition. The effective date of the newborn’s enrollment is the same as the newborn’s DOB.

STAR MCOs are responsible for all covered services provided to newborn clients.

**STAR Example**

Enrollments of newborns born to mothers who are enrolled in STAR are retroactive to the newborn’s date of birth. The following example shows the managed care enrollment date for a newborn:

<table>
<thead>
<tr>
<th>Task</th>
<th>Date</th>
</tr>
</thead>
<tbody>
<tr>
<td>Client’s date of birth (mother enrolled in STAR)</td>
<td>January 3</td>
</tr>
<tr>
<td>Client certified for Texas Medicaid</td>
<td>February 1</td>
</tr>
<tr>
<td>Medicaid benefits begin</td>
<td>January 3 (retroactive to DOB)</td>
</tr>
<tr>
<td>STAR enrollment begins (mother’s STAR plan at time of birth)</td>
<td>January 3 (retroactive to DOB)</td>
</tr>
</tbody>
</table>

Children born to STAR+PLUS or STAR Kids clients will be automatically enrolled with the STAR MCO in the SA operated by the same STAR+PLUS or STAR Kids MCO, if available. If the STAR+PLUS or STAR Kids MCO does not also operate a STAR MCO in the SA, the newborn will be enrolled in Texas Medicaid fee-for-service, and the mother will be given the opportunity to choose a STAR MCO for the newborn. Children born to STAR Health clients will be automatically enrolled in the STAR Health MCO. The effective date of the newborn’s enrollment is the same as the newborn’s date of birth. The newborn will remain enrolled for at least 90 days following the date of birth unless the mother requests a plan change as a special condition.

**2.3.2.2.1 Timely Notification and Assignment of Medicaid ID for Newborns**

There may be a delay of up to several months from the DOB for a newborn to receive a Medicaid ID number. Providers should check with each MCO to determine the claim filing requirements for newborns who do not yet have a Medicaid client number. Medicaid MCOs must adjudicate claims for services provided to newborns in accordance with HHSC’s claims processing requirements using the proxy ID number or state-issued Medicaid ID number. The MCO cannot deny claims based on a provider’s non-use of state-issued Medicaid ID number for a newborn. The MCO must accept provider claims for newborn services based on the mother’s name or Medicaid ID number with accommodations for multiple births, as specified by the MCO.

Hospitals that submit their birth certificate information using the DSHS Vital Statistics Unit (VSU) electronic Certificate Manager software and the Hospital Report (Newborn Child or Children) (Form 7484), receive a rapid and efficient assignment of a newborn Medicaid ID number. This process expedites reimbursement to hospitals and other providers involved in newborn care, including pharmacies that provide outpatient prescription benefits for medically needy newborns.

**Note:** The enrollment of newborns that are born to mothers who are enrolled in an MCO on the date of birth are retroactive to the newborn’s date of birth (DOB).

**2.3.2.3 Breast Pump Claims Filing for MCO Services**

Texas Medicaid and CHIP cover breast pumps and equipment after a baby is born when they are medically necessary.
<table>
<thead>
<tr>
<th>Coverage in prenatal period</th>
<th>Coverage at delivery</th>
<th>Coverage for newborn</th>
<th>Breast pump coverage &amp; billing</th>
</tr>
</thead>
<tbody>
<tr>
<td>STAR</td>
<td>STAR</td>
<td>STAR</td>
<td>STAR covers breast pumps when medically necessary for mothers or newborns. Breast pump equipment may be billed under the mother’s Medicaid ID or the newborn’s Medicaid ID.</td>
</tr>
<tr>
<td>CHIP Perinatal, with income at or below 198% of federal poverty level (FPL)*</td>
<td>Emergency Medicaid fee-for-service (FFS) or STAR**</td>
<td>Medicaid fee-for-service (FFS) or STAR**</td>
<td>Medicaid FFS and STAR cover breast pumps when medically necessary for newborns when the mother does not have coverage under CHIP. The breast pump must be billed under the newborn’s Medicaid ID.</td>
</tr>
<tr>
<td>CHIP Perinatal, with income above 198% FPL</td>
<td>CHIP Perinatal</td>
<td>CHIP Perinatal</td>
<td>CHIP covers breast pumps when medically necessary for CHIP Perinatal newborns. Breast pump equipment must be billed under the newborn’s CHIP Perinatal ID.</td>
</tr>
<tr>
<td>STAR Kids</td>
<td>STAR Kids</td>
<td>Medicaid FFS or STAR**</td>
<td>Medicaid FFS, STAR, STAR Kids, and STAR+PLUS cover breast pumps when medically necessary for mothers or newborns. Breast pump equipment may be billed under the mother’s Medicaid ID or the newborn’s Medicaid ID.</td>
</tr>
<tr>
<td>STAR+PLUS</td>
<td>STAR+PLUS</td>
<td>Medicaid FFS or STAR**</td>
<td>Medicaid FFS, STAR, STAR Kids, and STAR+PLUS cover breast pumps when medically necessary for mothers or newborns. Breast pump equipment may be billed under the mother’s Medicaid ID or the newborn’s Medicaid ID.</td>
</tr>
</tbody>
</table>

* CHIP Perinatal clients who have household incomes at or below 198% FPL must apply for Emergency Medicaid coverage for labor and delivery services. HHSC mails the pregnant woman an Emergency Medicaid application 30 days before her reported due date. When Emergency Medicaid covers a birth, the newborn is certified for 12 months of Medicaid coverage, beginning on the date of birth.

** These newborns will be in FFS Medicaid until they are enrolled with a STAR MCO. Claims should be filed with TMHP using the newborn’s Medicaid ID if the mother does not have coverage.
2.3.2.4 Medicaid for Transitioning Foster Care Youth (MTFCY) and Former Foster Care Children (FFCC)

Medicaid for Former Foster Care Children (FFCC) covers Medicaid clients who are 18 years of age or older and were receiving Medicaid when they aged out of foster care, but would otherwise not be eligible to continue Medicaid coverage. FFCC clients are automatically enrolled in STAR Health through the last day of the month in which they turn 21 years of age. FFCC clients may opt out of STAR Health for STAR, which allows clients to choose their MCO.

Medicaid clients who were receiving Medicaid and SSI when they aged out of foster care are automatically enrolled in STAR Kids.

Eligibility for STAR Health and STAR Kids ends on the last day of the month in which they turn 21 years of age and then coverage transfers to STAR unless the client is eligible for STAR+PLUS.

Medicaid for Transitioning Foster Care Youth (MTFCY) covers former foster care youth who were not receiving Medicaid when they aged out of foster care at 18 years of age. MTFCY clients are eligible through the last day of the month in which they turn 21 years of age to receive services through the fee-for-service or managed care models.

2.3.3 Client Rights

In Texas, Medicaid managed care clients have defined rights and responsibilities. Each health plan and primary care provider share the responsibility to ensure and protect client rights and to assist clients in understanding and fulfilling their responsibilities as plan clients.
Medicaid managed care clients have the right to:

- Be treated fairly and with dignity and respect.
- Know that their medical records and discussions with their providers will be kept private and confidential.
- Request changes to their medical records (if incorrect).
- A reasonable opportunity to choose a health-care plan and primary care provider (the doctor or health-care provider they will see most of the time and who will coordinate their care) and to change to another plan or provider in a reasonably easy manner. These opportunities include the right to:
  - Be informed of available health plans and primary care providers in their areas.
  - Be informed of how to choose and change health plans and primary care providers.
  - Choose any health plan that is available in their area and choose a primary care provider.
  - Change their primary care provider at any time for any reason.
  - Change health plans without penalty.
  - Be educated about how to change health plans or primary care providers.
- Know that doctors, hospitals, and others who provide care can advise clients about their health status, medical care, and treatment. The health plan cannot prevent them from giving clients this information, even if the care or treatment is not a covered service.
- Know that clients are not responsible for paying for covered services. Doctors, hospitals, and others cannot require clients to pay copayments or any other amounts for covered services.
- Ask questions and get answers about anything the client doesn’t understand, and that includes the right to:
  - Have their provider explain their health-care needs to them and talk to them about the different ways their health-care problems can be treated.
  - Be told why care or services were denied and not given.
- Consent to or refuse treatment and actively participate in treatment decisions, and that includes the right to:
  - Work as part of a team with their provider in deciding what health care is best for them.
  - Say yes or no to the care recommended by their provider.
- Utilize each available complaint and appeal process through the MCO and through Medicaid, receive a timely response to complaints, appeals, and fair hearings. These processes include the right to:
  - Make a complaint to their health plan or to the state Medicaid program about their health-care, provider, or health plan.
  - Receive a timely answer to their complaint.
  - Access the health plan appeal process and the procedures for doing so.
  - Request a fair hearing from the state Medicaid program and request information about the process for doing so.
- Timely access to care that does not have any communication or physical access barriers. They have the right to:
  - Have telephone access to a medical professional 24 hours a day, 7 days a week in order to obtain any needed emergency or urgent care.
• Receive medical care in a timely manner.

• Be able to get in and out of a health-care provider’s office, including barrier free access for persons with disabilities or other conditions limiting mobility, in accordance with the Americans with Disabilities Act.

• Have interpreters, if needed, during appointments with their providers and when talking to their health plan. Interpreters include people who can speak in their native language, assist with a disability, or help them understand the information.

• Be given an explanation they can understand about their health plan rules, including the health-care services they can get and how to get them.

• Not be restrained or secluded when doing so is for someone else’s convenience, or is meant to force them to do something they are unwilling to do, or to punish them.

### 2.3.3.1 Advance Directives

Federal and state law require providers and MCOs to maintain written policies and procedures for informing and providing written information to all adult clients who are 18 years of age and older about their rights under state and federal law, in advance of their receiving care ([Social Security Act §§1902(a)[57] and 1903(m)[1][A]]). The written policies and procedures must contain procedures for providing written information regarding the client’s right to refuse, withhold, or withdraw medical treatment advance directives.

These policies and procedures must comply with provisions contained in [42 Code of Federal Regulations (CFR) §§434.28 and 489, SubPart I, relating to the following state laws and rules:](4)

- A client’s right to self-determination in making health-care decisions.

- The Advance Directives Act, Chapter 166, Texas Health and Safety Code, which includes:
  - A client’s right to execute an advance written directive to physicians and family or surrogates, or to make a nonwritten directive to administer, withhold or withdraw life-sustaining treatment in the event of a terminal or irreversible condition.
  - A client’s right to make written and nonwritten Out-of-Hospital Do-Not-Resuscitate Orders.
  - A client’s right to execute a Medical Power of Attorney to appoint an agent to make health-care decisions on the client’s behalf if the client becomes incompetent.
  - The Declaration for Mental Health Treatment, Chapter 137, Texas Civil Practice and Remedies Code, which includes a client’s right to execute a Declaration for Mental Health Treatment in a document making a declaration of preferences or instructions regarding mental health treatment.

These policies can include a clear and precise statement of limitation if a participating provider cannot or will not implement a client’s advance directive. A statement of limitation on implementing a client’s advance directive should include at least the following information:

- A clarification of the provider’s conscience objections.

- Identification of the state legal authority permitting a provider’s conscience objections to carrying out an advance directive.

- A description of the range of medical conditions or procedures affected by the conscience objection.

A provider cannot require a client to execute or issue an advance directive as a condition for receiving health-care services. A provider cannot discriminate against a client based on whether or not the client has executed or issued an advance directive.

A provider’s policies and procedures must require the provider to comply with the requirements of state and federal law relating to advance directives.
2.3.4 Client Responsibilities

Medicaid managed care health plans and primary care providers should help clients understand their responsibilities. These include the responsibility to:

- Learn and understand each right they have under Medicaid. That includes the responsibility to:
  - Learn and understand their rights under the Medicaid program.
  - Ask questions if they do not understand their rights.
  - Learn what choice of health plan is available in their area.
- Abide by the health plan and Medicaid managed care policies and procedures. That includes the responsibility to:
  - Learn and follow their health plan rules and Medicaid rules.
  - Choose their health plan and a primary care provider.
  - Make any changes in their health plan and primary care provider in the ways established by Medicaid managed care and by the health plan.
  - Keep their scheduled appointments.
  - Cancel appointments in advance when they cannot keep them.
  - Always contact their primary care provider first for nonemergency medical needs.
  - Be sure they have approval from their primary care provider before going to a specialist (except for self-referred services).
  - Understand when they should and should not go to the ER.
- Share information relating to their health status with their primary care provider and become fully informed about service and treatment options. That includes the responsibility to:
  - Tell their primary care provider about their health.
  - Talk to their providers about their health-care needs and ask questions about the different ways their health-care problems can be treated.
  - Help their providers get their medical records.
- Actively participate in decisions relating to service and treatment options, make personal choices, and take action to maintain their health. That includes the responsibility to:
  - Work as a team with their providers in deciding what health care is best for them.
  - Understand how the things they do can affect their health.
  - Do the best they can to stay healthy.
  - Treat providers and staff with respect.
  - Talk to their provider about all of their medications.

2.4 Primary Care Provider/Main Dentist Guidelines for Medicaid Managed Care Clients

In Medicaid managed care, eligible Medicaid clients choose a primary care provider or a main dentist who will work with the client to coordinate the client’s health care or dental services.

The managed care client’s primary care provider/main dentist is responsible for the following:

- Furnishes primary-care related services
• Arranges for and coordinates referrals for all medically necessary specialty services
• Is available directly or through on-call arrangements 24 hours a day, 7 days a week for urgent or emergency care

Refer to: Subsection 2.4.4, “Continuous Access” in this handbook.

Primary care includes ongoing responsibility for preventive health or dental care, health or dental maintenance, treatment of illness and injuries, and the coordination of access to specialist providers and other services. The primary care provider or main dentist either furnishes or arranges for most of the client’s health-care or dental-care needs, including well-checkups, office visits, referrals, outpatient surgeries, hospitalizations, and health- and dental-related services. Primary care providers/main dentists can choose to contract with various MCOs or DMOs.

Although primary care providers are encouraged to help clients access these services, Medicaid managed care enrollees may self-refer for the following services:

• Emergency services
• Family planning services
• THSteps medical services
• Immunizations
• Early Childhood Intervention (ECI)
• Case Management for Children and Pregnant Women
• Obstetric or gynecological services
• School Health and Related Services (SHARS)
• DSHS case management
• HHSC case management
• Behavioral health services (contact client’s health plan for specific requirements)
• Vision care (including ophthalmologic or therapeutic optometry)

2.4.1 Enrolling as a Primary Care Provider or Main Dentist

Various providers may be eligible to enroll in Medicaid managed care as primary care providers or main dentist. Providers must contact the individual Medicaid managed care health plans or DMOs for enrollment information.

The following provider types are eligible to serve as primary care providers:

• Pediatricians
• Family/general practitioners
• Internists
• Obstetrician/gynecologists
• Advanced Practice Registered Nurses (APRNs) under the supervision of a physician who qualifies as a primary care provider
• Certified nurse-midwives (CNM) practicing under the supervision of a physician
• Physician assistants (PAs) practicing under the supervision of a physician who qualifies as a primary care provider
• Rural health clinics (RHCs)
• Federally Qualified Health Centers (FQHCs)
• Community Clinics
• Specialists willing to provide medical homes to clients who have special needs
• The following provider types are eligible to serve as main dentists:
  • General dentist
  • Pediatric dentist
• Rural health clinics (RHCs)
• Federally Qualified Health Centers (FQHCs)

2.4.2 Primary Care Provider Requirements for THSteps Medical Services
THSteps providers must be enrolled with Medicaid to be reimbursed for services provided to clients. THSteps medical services are self-referred. Medicaid MCOs determine how their clients will access THSteps services. The MCO may require the client to go to an in-network THSteps provider or may allow the client to go to any Medicaid THSteps provider, whether or not they are in the MCO’s network. Providers that render THSteps services must work in collaboration with the client’s primary care provider to ensure continuity of care.

THSteps providers are required to bill claims as an exception to periodicity when the clients visit is outside of the periodicity schedule because of extenuating circumstances.

Refer to: Section 5, “THSteps Medical” in the Children’s Services Handbook (Vol. 2, Provider Handbooks) for more information about billing an exception-to-periodicity checkup
Subsection 5.3.7, “Exception-to-Periodicity Checkups” in the Children’s Services Handbook (Vol. 2, Provider Handbooks) for more information about billing an exception-to-periodicity checkup
Subsection 4.2.12.1, “Exceptions to Periodicity” in the Children’s Services Handbook (Vol. 2, Provider Handbooks) for more information about billing an exception-to-periodicity checkup.

2.4.3 Primary Care Provider and Main Dentist Changes
Primary care provider and main dentist changes may be requested or initiated by any of the following:

• A client who is enrolled in a Medicaid MCO or DMO may request a primary care provider or main dentist change at any time and for any reason.

• The MCO or DMO may reassign the client to another primary care provider or main dentist for any of the following reasons:
  • The primary care provider or main dentist is sanctioned by HHSC.
  • The primary care provider or main dentist exhibits a documented pattern of unacceptable quality of care.
  • The primary care provider or main dentist inappropriately reduces the client’s right to access specialty services covered under Medicaid managed care.
  • The provider leaves Medicaid, retires, or dies.

• A provider may request a client be reassigned to another primary care provider or main dentist for any of the following reasons:
  • The client is not included in the primary care provider’s or main dentist’s scope of practice.
  • The client is noncompliant with medical or dental advice.
• The client consistently displays unacceptable office decorum.
• The client’s relationship with the primary care provider or main dentist is not mutually agreeable.

Any request by a provider to reassign a client to another primary care provider or main dentist must be processed through the applicable Medicaid MCO or DMO. Before a request for reassignment can be initiated, reasonable measures must be taken to correct the client’s behavior. Reasonable measures may include education or counseling by the MCO or DMO staff. The MCO or DMO will notify the client of the reassignment if all attempts to remedy the situation have failed. Providers should also notify the client about the reassignment in writing and send a copy of the notification to the MCO or DMO.

The MCOs and DMOs can affect a primary care provider or main dentist change immediately if necessary; however, the Medicaid client eligibility verification systems may not immediately reflect the change.

2.4.4 Continuous Access
Continuous access is an important feature of Medicaid managed care. Twenty-four-hour primary care provider and main dentist availability enables clients to access and use services appropriately, instead of relying on ERs for after-hours care.

Continuous access can be provided through direct access to a primary care provider’s or main dentist’s office or through on-call arrangements with another office or service. Clients should be informed of the primary care provider’s or main dentist’s normal office hours and should be instructed how to access urgent medical care after normal office hours.

2.4.4.1 After-Hours Guidelines
Primary care providers and main dentists are required to have at least one of the following arrangements in place to provide 24-hour, 7-day a week access for managed care clients:

• The office telephone is answered after-hours by an answering service, which meets language requirements of the major population groups and which can contact the primary care provider, main dentist, or another designated provider. All calls answered by an answering service must be returned within 30 minutes.

• The office telephone is answered after normal business hours by a recording in the language of each of the major population groups served, directing the patient to call another number to reach the primary care provider, main dentist, or another provider designated by the primary care provider or main dentist. Someone must be available to answer the designated provider’s telephone. Another recording is not acceptable.

• The office telephone is transferred after office hours to another location where someone will answer the telephone and be able to contact the primary care provider, main dentist, or another designated medical practitioner, who can return the call within 30 minutes.

2.4.4.2 Unacceptable Telephone Arrangements
The telephone answering procedures listed below are not acceptable:

• The office telephone is only answered during office hours.
• The office telephone is answered after-hours by a recording that tells clients to leave a message.
• The office telephone is answered after-hours by a recording that directs clients to go to an Emergency Room for any services needed.
• Returning after-hours calls outside of 30 minutes.
2.5  Cultural Competency and Sensitivity

HHSC values the diversity of the Texas Medicaid population and requires Medicaid managed care to provide programs to support clients from diverse cultural backgrounds. Helplines are staffed by both Spanish- and English-speaking customer service representatives who, at any time, may access a multi-language translation service for assistance.

Providers must comply with the laws concerning discrimination on the basis of race, color, national origin, or sex.

2.5.1  Limited English Proficiency

Medicaid providers are required to provide services in the languages of the major Medicaid population groups they serve and to ensure quality appropriate translations. Title VI, section 601, of the Civil Rights Act of 1964 states that “no person in the United States shall on the basis of race, color, or national origin, be excluded from participating in, be denied the benefits of, or be subjected to discrimination under any program or activity receiving Federal financial assistance.”

HHSC requires Medicaid providers to ensure persons with limited English proficiency have equal access to the Medicaid services to which they are legally entitled.

Meeting the requirements of Title VI may require the primary care provider or main dentist to take all or some of the following steps at no cost or additional burden to the beneficiary with limited English proficiency:

- Have a procedure for identifying the language needs of patients/clients.
- Have access to proficient interpreters during hours of operation (MCOs or DMOs arrange interpreters).
- Develop written policies and procedures regarding interpreter services (MCOs or DMOs arrange interpreters).
- Disseminate interpreter policies and procedures to staff and ensure staff awareness of these policies and procedures and of their Title VI obligations to persons with limited English proficiency.

To meet interpretation requirements, providers may choose to incorporate into their business practice any of the following (or equally effective) procedures:

- Hire bilingual staff. (Does not apply to MCOs.)
- Hire staff interpreters. (Does not apply to MCOs.)
- Use qualified volunteer staff interpreters. (Does not apply to MCOs.)
- Arrange for the services of volunteer community interpreters—excluding the client’s family or friends. (Does not apply to MCOs.)
- Contract with an outside interpreter service. (MCO or DMO must provide.)
- Use a telephone interpreter service.
- Develop a notification and outreach plan for beneficiaries with limited English proficiency.

It is the provider’s responsibility to ensure that interpretive services are available to his practice to meet requirements on limited English proficiency and communication disabilities. Interpretive services include language and American Sign Language (ASL) interpreters.

Language Line Services operate 24 hours a day, 7 days a week. Language Line Services provides over-the-telephone interpretation, video interpreting, document translation, interpreter testing and training, and other language products as well. Language Line Services charges a fee for the service. For complete details about their billing practices and services, providers should visit the Language Line Services website at www.languageline.com or call 1-800-752-6096.
Complaints and reports of non-compliance with Title VI regulations are handled by the Office for Civil Rights (OCR). Additional information, including the complete guidance memorandum on prohibition of discrimination against persons with limited English proficiency issued by the OCR, can be found on the Internet at www.hhs.gov/ocr/civilrights/resources/specialtopics/lep/index.html.

**Note:** MCOs are responsible for providing interpreter services.

### 2.6 Reimbursement

Providers must read and comply with “Section 2: Texas Medicaid Fee-for-Service Reimbursement” *(Vol. 1, General Information)*.

Reimbursement for benefits that are administered by a Texas Medicaid MCO or DMO is determined by the MCO or DMO. Providers should contact the MCO or DMO for additional information.

**Note:** The MCOs and DMOs are not limited to following the Texas Medicaid fee schedules. There may be some differences in reimbursement based on decisions made by the individual health and DMOs.

Texas Medicaid reimburses carve-out services according to the appropriate reimbursement methodology defined in the applicable Texas Medicaid Provider Procedures Manual handbook and the applicable Texas Medicaid fee schedules, which are available on the TMHP website at www.tmhp.com.

#### 2.6.1 Coinsurance and Deductible Payments for Dual-Eligible Clients

Crossover claims for payment for deductibles or coinsurance according to current payment guidelines are processed by TMHP and not the client’s MCO.

For clients who are enrolled in a Medicare Advantage Plan (MAP) and/or Special Needs Plan (SNP), crossover claims for coinsurance and deductible payments are processed by the MAP and/or SNP. These claims are not processed by TMHP.

#### 2.6.2 Third Party Liability (TPL)

Refer to: “Section 8: Third Party Liability (TPL)” *(Vol. 1, General Information)* for additional information.

#### 2.6.3 Out-of-Network Reimbursement

MCOs must ensure their clients have access to covered services on a timely basis. They are required to have a defined network of providers to meet client needs, and to provide support to clients who need help finding a doctor or setting up appointments. MCOs must maintain access to network providers based on federal and state requirements. If an in-network provider is not available, the MCO is still required to locate a willing provider to ensure clients have access to medically necessary and appropriate services.

If medically necessary covered services are not available through in-network providers, MCOs must allow a referral to an out-of-network Medicaid provider. The referral must be requested by an in-network provider and within the time appropriate to the circumstances relating to the delivery of the services and the condition of the client but no longer than five business days after the request.

The MCO must fully reimburse the non-network provider in accordance with the out-of-network methodology for Medicaid as defined by HHSC in 1 TAC § 353.4.

MCOs must allow pregnant clients, past the 24th week of pregnancy to remain under their current OB/GYN’s care through the client’s postpartum checkup, even if the OB/GYN provider is, or becomes, out-of-network.

For newly enrolled clients, MCOs must pay the existing out-of-network Medicaid providers for medically necessary covered services for up to 90 days, until the client’s records and care can be transferred to an in-network provider, or until the client is no longer enrolled with the MCO.
MCOs are not responsible for payment for unauthorized nonemergency services by out-of-network providers, except when that provider is an Indian Health Care Provider (IHCP) enrolled as a FQHC.

2.7 Managed Care Plan Changes
The MCO or DMO changes can be initiated by clients, MCOs, or DMOs.

2.7.1 Client-Initiated Plan Changes
Clients have the right to change plans. Clients must call the Enrollment Broker at 1-800-964-2777 to initiate a plan change. If a plan change request is received before the middle of the month, the plan change is effective on the first day of the following month. If the request is received after the middle of the month, the plan change will be effective on the first day of the second month following the request, as shown below.

<table>
<thead>
<tr>
<th>Request received on or before</th>
<th>Change effective</th>
</tr>
</thead>
<tbody>
<tr>
<td>Mid-May</td>
<td>June 1</td>
</tr>
</tbody>
</table>

Note: All plan change requests must be processed by the Enrollment Broker.

The STAR Health Program only has one plan choice available. As a result, clients cannot change plans, but may change primary care providers within their assigned STAR Health MCO.

2.7.2 Plan Administrator-Initiated Changes
Each health plan and DMO has a limited right to request that a client be disenrolled without the client’s consent. HHSC must approve any request for such disenrollment.

Health plans and DMOs may request that a client be disenrolled for the following reasons:
- The client loans his or her Your Texas Benefits Medicaid card to another person to obtain services.
- The client continually disregards the advice of his primary care provider or main dentist.
- The client repeatedly uses the ER inappropriately.
- Client is disruptive, unruly, threatening, or uncooperative to the extent that client’s membership seriously impairs MCO’s, DMO’s, or provider’s ability to provide services to the client or to obtain new patients, and the client’s behavior is not caused by a physical or behavioral health condition.
- Client refuses to comply with managed care restrictions (e.g., repeatedly using emergency room in combination with refusing to allow MCO to treat the underlying medical condition).
- For STAR+PLUS MCOs, under limited conditions, the MCO may request disenrollment of clients who are totally dependent on a ventilator or who have been diagnosed with End Stage Renal Disease.

Before a request for disenrollment can be initiated, reasonable measures must be taken to correct the client’s behavior. Reasonable measures may include education or counseling conducted by health plan or DMO staff. HHSC will notify the client in writing of the disenrollment if all attempts to remedy the situation have failed. HHSC will also notify the client in writing of the availability of appeal procedures and the HHSC fair hearing process.

Health plans, DMOs, and providers can not request a client’s disenrollment because of an adverse change in the client’s health or the utilization of services that are medically necessary for the treatment of a client’s condition.
2.7.3 Managed Care Organization (MCO) Clients Who Transition to Medicaid Fee-For Service (FFS)

When clients transition from an MCO to FFS, providers can request that previously approved authorizations for Comprehensive Care Program (CCP) services, occupational therapy (OT), physical therapy (PT), private duty nursing (PDN), and speech therapy (ST) be transferred from the MCO to FFS.

2.7.3.1 Submission Guidelines

TMHP will consider the reimbursement of claims for services that were rendered on or after the MCO’s disenrollment date only when the provider submits a request to TMHP to transfer the previously approved authorization for CCP services. The request to TMHP must be received on or before the end date of the previously approved MCO authorization. Any requests submitted after the MCO’s authorization end date will have to meet the regular submission guidelines for the specific service type.

2.7.3.2 Documentation Requirements

All of the requests to transfer the authorizations from the MCO to FFS must include:

- A copy of the previously approved authorization letter.
- All of the documentation that was sent in the original authorization request, including any physician orders that were used to determine the start of care. TMHP will accept the physician orders as the required documentation for the requested services.
- The completed CCP prior authorization form, Special Medical Prior Authorization (SMPA) form, Home Health Plan of Care, or Texas Medicaid Physical, Occupational, or Speech Therapy (PT, OT, ST) Prior Authorization Form, whichever is applicable for the requested service. The form must include the dates of service and quantities that are being requested from TMHP, which must match the dates of service and quantities that were approved in the original authorization.

Note: It is not necessary to obtain signatures or dates on the forms listed above when submitted to TMHP for the purpose of transferring an authorization from an MCO to FFS Medicaid.

Authorizations for services transferred from an MCO to FFS Medicaid are subject to retrospective review.

TMHP will verify the client’s eligibility, the dates of service, and the quantities requested.

TMHP will process reimbursement claims as follows:

- Claims for services that were rendered before the date on which the transfer request was received will be denied as a late submission, and the provider will be notified of their administrative appeal rights through the Health and Human Services Commission (HHSC).
- Claims for services that were rendered on or after the date of receipt use the required information from the transferred authorization and will be processed as if the request was received in a timely manner.
- Claims for services that were paid by an MCO and then recouped must contain the recoupment EOB from the MCO for consideration of payment. The claims must meet the 95-day deadline from the recoupment disposition date.

Note: Letter requests for refunds will not be accepted. A recoupment EOB with a disposition date is required.

If a request to transfer an MCO authorization is submitted after the end date of the MCO authorization, or the provider does not have an authorization letter from the MCO, TMHP will process the request to transfer the authorization based on established TMHP authorization submission guidelines for CCP services, PDN, OT, PT, and ST.

All new requests for rendered services must meet the documentation requirements.
2.7.3.3 **New Services and Extension of Services**

For new services that occur after the client’s MCO disenrollment change date, the provider is responsible for submitting all TMHP required paperwork and meeting all established submission guidelines for prior authorization.

Requests for the extension of services that occur after the MCO disenrollment change date must include all of the paperwork that is required by TMHP and meet all established submission guidelines for prior authorization.

2.7.3.4 **Loss of Eligibility**

If an MCO disenrolled a client and the client also loses Medicaid eligibility, providers must anticipate, if and when Medicaid eligibility is restored, that the client will initially be considered a Medicaid FFS client and will have a retroactive eligibility period.

All requests for services that require prior authorization and that occur during the client’s retroactive eligibility period, must be submitted to TMHP following the process that is outlined in subsection 5.1.1, “Prior Authorization Requests for Clients with Retroactive Eligibility” in “Section 5: Fee-for-Service Prior Authorizations” (Vol. 1, General Information).

If a client is retroactively disenrolled by an MCO, all of the services that are rendered by the provider during this retroactive disenrollment period (specifically from the date on which the client was eligible for FFS to the date of the client’s MCO eligibility change) will be denied by TMHP, and the provider will be notified of their administrative appeal rights.

**Refer to**: Subsection 6.4.2.9, “Attachments to Claims” in “Section 6: Claims Filing” (Vol. 1, General Information).

TMHP may consider services for the MCO transition beginning on the date of the client’s MCO eligibility change date and going forward. TMHP uses the MCO transition process for the submission of paperwork and the processing of provider requests.

2.8 **Authorizations for Managed Care Services**

MCOs must have a process in place that allows prior authorizations eligible for recertification to be submitted by providers and reviewed by the MCO within the timeframes as outlined in Section 5: Fee-for-Service Prior Authorizations.

Authorization requests for services administered by the client’s MCO or DMO must be submitted to the client’s MCO or DMO according to the guidelines specific to the plan under which the client is covered. Health plan prior authorizations do not automatically transfer with a client who moves to another plan.

When a client or provider notifies the new MCO that there is an existing prior authorization, the new MCO must ensure that the client gets a continued authorization of those services for the same amount, duration, and scope. The new MCO must get the continued authorization within the shortest of the following time periods:

- Within 90 Days of the transition to a new MCO
- Until the end of the current authorization period
- Until the MCO has evaluated and assessed the client and issued or denied a new authorization

If a client who is transitioning from FFS to managed care was receiving a service that did not require a prior authorization in FFS, but does require one in the new MCO, the MCO must ensure that the client receives services for the same amount, duration, and scope. The continued authorization must last for the shortest period of the following:

- Within 90 days of the transition to the new MCO
• Until the MCO has evaluated and assessed the client and issued or denied a new authorization

Dental prior authorizations may transfer from one DMO to another.

### 2.9 Claims Filing for Managed Care Services

Claims for services administered by an MCO or DMO must be submitted to the client’s MCO or DMO. Providers may submit directly to the appropriate MCO or DMO using the methods established by the MCO or DMO.

Providers must contact the appropriate MCO or DMO for information about filing electronic or paper claims directly to the MCO or DMO.

MCOs may also subcontract with behavioral health organizations (BHOs) and pharmacy benefit managers (PBMs). Providers must contact the client’s MCO to verify the claims process.

Refer to:

- The TMHP website at [www.tmhp.com/topics/managed-care](http://www.tmhp.com/topics/managed-care) for additional information, including MCO and DMO contact information.

Important:

Providers must call the client’s MCO or DMO who processed the claim for information about the MCO’s or DMO’s explanation of benefits (EOB), claims payment, claim rejection, how to correct a rejected claim, or any other questions about the MCO or DMO claim guidelines and processes. TMHP does not have any information about the MCO’s or DMO’s claims, benefits, or processes.

Refer to:

- “Section 3: TMHP Electronic Data Interchange (EDI)” (Vol. 1, General Information) for information on electronic claims submissions to TMHP.

Reminder:

Claims for Medicaid managed care clients must be submitted to the MCO or DMO in which the client is enrolled at the time of service (or date of admission for inpatient hospital claims). The MCO or DMO, as a payor of last resort, does not determine payment based on the primary payer’s (i.e., TPR or other primary source of insurance) authorization of services or approval of hospital stays.

Refer to:

- “Section 8: Third Party Liability (TPL)” (Vol. 1, General Information) for additional information.

The TMHP Medicaid Managed Care web page at [www.tmhp.com/topics/managed-care](http://www.tmhp.com/topics/managed-care) for additional information.

### 2.9.1 Filing Deadlines

The following table summarizes the filing deadlines that apply for MCO and DMO claim submissions:

<table>
<thead>
<tr>
<th>Submission</th>
<th>Filing Deadline</th>
</tr>
</thead>
<tbody>
<tr>
<td>Initial submission submitted to the correct plan</td>
<td>95 days from the DOS</td>
</tr>
<tr>
<td>Initial submission submitted to the wrong plan</td>
<td>95 days of the date on the Remittance and Status (R&amp;S) Report from the other (wrong) carrier (documentation of timely filing is required)</td>
</tr>
<tr>
<td>Initial submission to TPR (not the Medicaid MCO or DMO)</td>
<td>95 days from the date of disposition by the other insurance resource</td>
</tr>
<tr>
<td>Initial submission for newborns</td>
<td>Submit to the client’s or mother’s MCO within 95 days of the DOS</td>
</tr>
</tbody>
</table>

Claims must be submitted to the appropriate entity whether TMHP or the MCO or DMO within 95 days of the date of service. If the claim is not received by the MCO or DMO within 95 days, the claim will be denied.
If the provider files with the wrong plan within the 95 day submission requirement (e.g., State Claims Administrator but not with the MCO or DMO), the provider must resubmit the claim with documentation that shows the claim was submitted within the appropriate time frame but to the wrong plan. The MCO or DMO must honor the initial filing date and process the claim without denying the resubmission for the sole reason of passing the filing timeframe. The provider must file the claim with the correct MCO within 95 days of the date on the Remittance and Status (R&S) Report from the other (wrong) carrier.

When a service is billed to other insurance, the claim must be refiled and received by the Medicaid MCO or DMO within 95 days from the date of disposition by the other insurance resource. The MCO or DMO will determine, as a part of its provider claims filing requirements, the documentation required when a provider refiles these types of claims with the MCO or DMO.

MCOs and DMOs are subject to the requirements related to coordination of benefits for secondary payers in the Texas Insurance Code section 843.349 (e) and (f).

Refer to:

Subsection 2.9, “Claims Filing for Managed Care Services” in this handbook for details about managed care claims.

Subsection 6.1, “Claims Information” in “Section 6: Claims Filing” (Vol. 1, General Information) for information about MCO claims processed by TMHP and not the client’s MCO.

2.10 MCO/DMO Appeals, Complaints, and Fair Hearings

Providers can submit their appeals directly to the MCO or DMO that administers the clients’ managed care benefits.

Claims that were originally submitted to TMHP for routing to the appropriate MCO or DMO can be appealed to TMHP using TexMedConnect or EDI. The appeals will be routed to the appropriate entity for processing.

2.10.1 Medicaid Managed Care Complaints and Fair Hearings

Medicaid managed care providers may file complaints with HHSC if they find they did not receive full due process from the respective managed care health plan.

Appeals, grievances, or dispute resolution is the responsibility of each MCO or DMO. Providers are encouraged to exhaust the complaints or grievance process with their MCO or DMO before filing a complaint with HHSC.

Refer to: The respective MCO or DMO for information about specific complaint policies and procedures.

The respective health plan’s policies and procedures for information about the MCO appeals and fair hearing process.

Subsection 7.1.5, “Paper Appeals” in “Section 7: Appeals” (Vol. 1, General Information) for information about paper appeals.

Once the MCO’s or DMO’s complaints or grievance process has been exhausted, complaint requests may be sent to HHSC.

Complaints about STAR, STAR+PLUS, STAR Kids, STAR Health, and DMO can be:

- Emailed to HHSC at HPM_Complaints@hhsc.state.tx.us.
3   **STAR Program**

The principal objectives of the STAR Program are to emphasize early intervention and to promote improved access to quality care thereby significantly improving health outcomes for the target populations. The special focus of the STAR Program is on prenatal and well-child care.

### 3.1 STAR Program Clients

Most clients in Texas Medicaid get their coverage through the STAR Program. STAR provides primary care, acute care, behavioral health care, and pharmacy services for pregnant women, newborns, and children and parents with limited income. Some former foster care children and youth are eligible for the STAR Program as well.

STAR is available statewide in 13 service areas (SAs). STAR Medicaid clients can select from at least two MCOs in each service area.

**Refer to:** Subsection 2.3.2.4, “Medicaid for Transitioning Foster Care Youth (MTFCY) and Former Foster Care Children (FFCC)” in this handbook.

### 3.2 STAR Program Benefits

STAR Program clients receive all the benefits of Texas Medicaid fee-for-service and the following additional benefits:

- Removal of the inpatient spell of illness limitation for adults
- Unlimited medically necessary prescription drugs for adults
- Waiver of the $200,000 individual annual limit on inpatient services

### 3.2.1 Spell of Illness

STAR clients are not limited to the 30-day spell of illness. The spell of illness limitation is defined as 30 days of inpatient hospital care, which may accrue intermittently or consecutively. After 30 days of inpatient care is provided, reimbursement for additional inpatient care is not considered until the client has been out of an acute care facility for 60 consecutive days. All Medicaid clients who are 20 years of age and younger are already not limited to the 30-day spell of illness.

### 3.2.2 Prescriptions

STAR clients who are 21 years of age or older receive unlimited medically necessary prescription drugs. The elimination of the three prescription limit per month for adult clients enrolled in STAR allows the provider greater flexibility in treating and managing a client’s health-care needs. All Medicaid clients who are 20 years of age or younger already receive unlimited medically necessary prescription drugs.
4  STAR Kids Program

The STAR Kids Program is designed to improve access to care, coordinate care across service areas, improve ease of program participation for clients, managed care organizations, and providers, and achieve cost efficiency and cost containment. The STAR Kids program integrates acute care and long-term services and supports (LTSS) into a Medicaid managed care delivery system for children and adults who are 20 years of age or younger and have a disability.

STAR Kids also provides Medically Dependent Children’s Program (MDCP) services for eligible clients.


4.1  STAR Kids Clients

STAR Kids is available statewide in 13 service areas (SAs). STAR Kids Medicaid clients can select from at least two MCOs in each service area.

Participation in the STAR Kids program is required for Medicaid clients who are 20 years of age or younger and meet at least one of the following:

- Receive Supplemental Security Income (SSI).
- Receive SSI and Medicare.
- Receive services through the Medically Dependent Children Program (MDCP) waiver.
- Receive services through the Youth Empowerment Services (YES) waiver.
- Live in a community-based intermediate care facility for individuals with an intellectual disability or related condition (ICF/IID) or nursing facility.
- Receive services through a Medicaid Buy-In program.
- Receive services through any of the following HHSC intellectual and developmental disability (IDD) waiver programs.
  - Community Living Assistance and Support Services (CLASS)
  - Deaf Blind with Multiple Disabilities (DBMD)
  - Home and Community-based Services (HCS)
  - Texas Home Living (TxHmL)

Children and youth who receive SSI or SSI-related Medicaid or are enrolled in MDCP receive all of their Medicaid services through the STAR Kids program.

Children and youth who receive services through other 1915(c) waiver programs receive their basic Medicaid health services through STAR Kids, while receiving their LTSS through their waiver program.

Children who are dually eligible receive most of their acute care services through Medicare but receive LTSS and service coordination through STAR Kids.

4.2  STAR Kids Benefits

STAR Kids Program clients receive all the benefits of Texas Medicaid fee-for-service, including Texas Health Steps and Comprehensive Care Program services. STAR Kids clients who have an assessed need for LTSS, identified by the STAR Kids Screening and Assessment Instrument (SK-SAI), may receive services through their STAR Kids MCO.
4.2.1 Spell-Of-Illness
No spell-of-illness limitation exists for THSteps-eligible clients who are 20 years of age or younger when a medically necessary condition exists.

4.2.2 Prescriptions
All Medicaid clients who are 20 years of age or younger receive unlimited medically necessary prescription drugs.

4.2.3 Service Coordination
*Refer to:* Subsection 2.1.3, “Service and Care Coordination” in this handbook.

The [STAR Kids Handbook](https://hhs.texas.gov) page of the HHS website at hhs.texas.gov.

5 STAR+PLUS Program

The STAR+PLUS Program is designed to improve access to care, provide care in the least restrictive setting, and provide more accountability and control on costs. The STAR+PLUS program integrates acute care and long-term services and supports (LTSS) into a Medicaid managed care delivery system for SSI-eligible Medicaid clients.

The STAR+PLUS Program serves adults with a disability, clients age 65 and older (including those dually eligible for Medicare and Medicaid), as well as women in the Medicaid Breast and Cervical Cancer Program.

5.1 STAR+PLUS Program Clients

Medicaid clients who are 21 years of age or older and enrolled in any of the following HHSC programs must enroll in STAR+PLUS for acute care benefits, unless the member is dual-eligible for Medicaid and Medicare:

- Community-based Intermediate Care Facility of Individuals with Intellectual Disabilities (ICF-IID)
- Home and Community-based Services (HCS)
- Community Living Assistance and Support Services (CLASS)
- Texas Home Living (TxHmL)
- Deaf Blind with Multiple Disabilities (DBMD)

Enrollment in the STAR+PLUS program is required for clients of Medicaid who meet one or more of the following criteria:

- Receive Supplemental Security Income (SSI) benefits or SSI-related Medicaid
- Qualify for STAR+PLUS Home and Community-Based Waiver Services
- Are 21 years of age or older and receive Medicaid because they are in a Social Security Exclusion program and meet financial criteria for STAR+PLUS Home and Community-Based Services Program
- Are 21 years of age or older and reside in a nursing facility
- Qualify for Medicaid for Breast and Cervical Cancer (MBCC)
- Former Foster Care Children (FFCC) clients who are 21 years of age through the end of the month of their 26th birthday, who are dual eligible, and who reside in a community-based ICF-IID or receive services under the following Medicaid 1915(c) waivers:
  - HCS
• CLASS
• TxHmL
• DBMD

**Exception:** Clients who receive Medicare Part B in addition to Medicaid will remain in FFS for all Medicaid services.

**Refer to:** Subsection 4.8, “Medicaid for Breast and Cervical Cancer (MBCC)” in “Section 4: Client Eligibility” (Vol. 1, General Information) for more information about the MBCC Program.

After selecting an MCO, STAR+PLUS Program clients who are not dual-eligible are required to select a primary care provider from the MCO provider directory.

### 5.1.1 STAR+PLUS Program Dual-Eligible Clients

Many STAR+PLUS clients are eligible for Medicaid and Medicare. Dual eligible clients who participate in the STAR+PLUS program receive most acute care services through their Medicare provider and LTSS through the STAR+PLUS MCO. STAR+PLUS program dual eligible clients must select a STAR+PLUS MCO to receive LTSS through STAR+PLUS. Dual eligible clients may receive some additional services through their STAR+PLUS MCO.

Most STAR+PLUS clients with Medicare and Medicaid are Medicaid Qualified Medicare Beneficiaries (MQMBs). MQMBs receive Medicare benefits through a Medicare risk product (MCO) or Medicare fee-for-service insurance program. To reduce confusion, HHSC has mandated that STAR+PLUS MQMBs continue to receive all their acute care services as they do today, with Medicare being the primary payor and Texas Medicaid fee-for-service, through TMHP, the secondary payor.

MQMB clients qualify for Medicaid benefits that are not covered by Medicare.

Providers are to continue billing for Medicare acute care services through the client’s Medicare MCO or fee-for-service insurer following the rules of the Medicare insurer. If the client is in both a Medicare MCO and a Medicaid MCO, the client uses the Medicare primary care provider, and providers follow the Medicare MCO’s medical management rules for authorization, concurrent review, etc. MQMBs choose a Medicaid MCO but do not choose a Medicaid primary care provider.

**Refer to:** Subsection 4.9, “Medicare and Medicaid Dual Eligibility” in “Section 4: Client Eligibility” (Vol. 1, General Information) for more information and further MQMB instructions.

#### 5.1.1.1 Dual Demonstration

The Dual Eligible Integrated Care Demonstration Project, or Dual Demonstration, is a fully integrated managed care model for individuals age 21 and older who are dually eligible for Medicare and Medicaid and required to be enrolled in the STAR+PLUS program. This model involves a three-party contract between an MCO with an existing STAR+PLUS contract, HHSC, and the Centers for Medicare & Medicaid Services (CMS) for the provision of the full array of Medicaid and Medicare services.

Under this initiative, the MCO is responsible for the full array of Medicare and Medicaid covered services, including acute care and LTSS. The demonstration does not include clients who reside in an ICF/IID and individuals with IDD who receive services through CLASS, DBMD, HCS, or TxHmL waivers. The demonstration operates in Bexar, Dallas, El Paso, Harris, Hidalgo, and Tarrant counties.

#### 5.1.1.2 Medicare Advantage Dual Eligible Special Needs Plan

A Dual Eligible Special Needs Plan (D-SNP) is a managed care delivery model specifically designed to coordinate care between Medicare and Medicaid covered services for individuals that are dually eligible for both programs.
Under this managed care delivery option, D-SNPs are responsible for the coordination of care between Medicare and Medicaid covered services. D-SNPs that also operate in STAR+PLUS deliver Medicaid services through the STAR+PLUS program. D-SNPs that do not also operate in STAR+PLUS are only responsible for paying beneficiary cost-sharing.

5.1.2 Clients who are Ineligible for The STAR+PLUS Program

Clients who meet the following criteria are not eligible to enroll in STAR+PLUS and will remain in Texas Medicaid fee-for-service:

- Residents in a State Supported Living Center
- Residents in an ICF-IID who are dual eligible in Medicare and Medicaid
- Residents of state hospitals or institutions for mental diseases
- Program of All-Inclusive Care for the Elderly (PACE)
- In-Home and Family Support Program Services clients
- Qualified Medicare Beneficiaries (QMBs) that do not receive Medicaid benefits other than Medicare deductible or coinsurance liabilities according to current payment guidelines
- Clients who receive limited Medicaid benefits and do not qualify for participation in the VDP

5.2 STAR+PLUS Program Benefits

STAR+PLUS Program clients receive all the benefits of Texas Medicaid fee-for-service and the following additional benefits:

- Unlimited medically necessary prescription drugs for adults who are not dual-eligible
- Waiver of the $200,000 individual annual limit on inpatient services

Refer to: Subsection 3.2.2, “Prescriptions” in this handbook for more information about prescription benefits.

Note: Dual eligible adults continue to be limited to three prescriptions unless they have joined the Medicare MCO also offered by their STAR+PLUS MCO.

5.2.1 Prescriptions

STAR+PLUS clients who are 21 years of age and older and do not receive Medicare receive unlimited medically necessary prescription drugs. The elimination of the three prescription limit per month for adult clients enrolled in STAR+PLUS allows the provider greater flexibility in treating and managing a client’s health care needs. All Medicaid clients who are 20 years of age and younger already receive unlimited medically necessary prescription drugs.

5.2.2 Spell of Illness

The spell-of-illness limitation applies to clients in the STAR+PLUS Program.

A spell-of-illness is defined as 30 days of inpatient hospital care, which may accrue intermittently or consecutively. After 30 days of inpatient care is provided, reimbursement for additional inpatient care is not considered until the client has been out of an acute care facility for 60 consecutive days.

An individual may be discharged from and readmitted to a hospital several times, regardless of the admittance reasons, and still be considered to be in the same spell of illness if 60 days have not elapsed between discharge and readmission.

The spell-of-illness limitation does not apply in the following situations:

- A prior-approved solid organ transplant has an additional 30-day spell of illness, which begins on the date of the transplant.
• No spell-of-illness limitation exists for THSteps-eligible clients who are 20 years of age and younger when a medically necessary condition exists.

• The client is enrolled in the Medicaid managed care STAR program.

• The client is not eligible for Medicare and is admitted to an inpatient facility with a diagnosis of bipolar disorder, major depressive disorder, recurrent depressive disorder, schizoaffective disorder, or schizophrenia.

For confirmation of spell-of-illness limitation contact the client’s MCO.

5.2.3 Service Coordination

The MCO must furnish a service coordinator to all STAR+PLUS clients who request one, or when the MCO determines the need for a service coordinator through an assessment. A service coordinator is the person with primary responsibility for providing service coordination and care management to STAR+PLUS clients.

*Refer to:* Subsection 2.1.3, “Service and Care Coordination” in this handbook.

6 STAR Health Program

The STAR Health program ensures that children and youth in state conservatorship are able to receive all services they need immediately upon entry into conservatorship. Benefits under STAR Health begin the date the client is placed in conservatorship.

HHSC has selected Superior HealthPlan as the MCO for this program. Superior HealthPlan is responsible for assigning a to clients when they are enrolled in the STAR Health Program. Foster care families are given the opportunity to change their primary care provider after this initial assignment.

6.1 STAR Health Program Clients

All Medicaid clients in foster care are placed in this program with the following exceptions:

• Children adjudicated and placed with the Texas Juvenile Justice Department

• Children from other states who are placed in Texas Children in Medicaid-paid facilities such as children in nursing homes, ICF-IIDs, or State-Supported Living Centers

• Children who are active SSI-related Medicaid clients

• Children who are in state conservatorship who are placed outside of Texas

• Children who are in adoption assistance

Clients who participate in the Medicaid for Transitioning Foster Care Youth (MTFCY) program and the Former Foster Care Children (FFCC) program are eligible for the STAR Health program.

The following table shows the age ranges for clients who may be eligible for the STAR Health program:

<table>
<thead>
<tr>
<th>Group Clients Belong to</th>
<th>Age Ranges</th>
</tr>
</thead>
<tbody>
<tr>
<td>Client under DFPS conservatorship</td>
<td>DFPS can retain conservatorship through the month of the client’s 18th birthday. (Eligibility ends the month conservatorship ends.)</td>
</tr>
<tr>
<td>Clients who voluntarily continue in a foster care placement after DFPS conservatorship ends</td>
<td>18 through 21 years of age (Eligibility ends the month of their 22nd birthday.)</td>
</tr>
<tr>
<td>Clients who are participating in the MTFCY program</td>
<td>18 through 20 years of age (Eligibility ends the month of their 21st birthday.)</td>
</tr>
</tbody>
</table>
STAR Health clients can gain eligibility on any day of the month. To ensure the accurate confirmation of STAR Health eligibility, it is essential that all health-care providers verify eligibility by contacting the STAR Health MCO. The STAR Health MCO receives updated eligibility information on a daily basis, so it will have the most current eligibility information.

The Department of Family and Protective Services (DFPS) Form 2085 as well as the Your Texas Benefits Medicaid card may also be used to verify eligibility in the STAR Health Program.

Newborns born to a mother who is enrolled in the STAR Health program are automatically enrolled in STAR Health.

Newborns that are taken into State conservatorship while still in the hospital will be enrolled in STAR Health on the date the State takes conservatorship.

Refer to:
- Subsection 2.3.2.4, “Medicaid for Transitioning Foster Care Youth (MTFCY) and Former Foster Care Children (FFCC)” in this handbook.
- Subsection 2.3.2.2, “Newborn Enrollment” in this handbook.

### 6.2 STAR Health Program Benefits

STAR Health Program clients receive all the benefits of traditional Texas Medicaid as well as service coordination to assist in making appointments and accessing services; and service management to assist with managing the health care of those with ongoing and serious medical needs.

Refer to:
- Subsection 2.1.3, “Service and Care Coordination” in this handbook.

Most Medicaid foster care claims are capitated services and must be submitted to Superior HealthPlan.

Refer to:
- Section 8, “Carve-Out Services” in this handbook for the list of non-capitated services that may be reimbursed by TMHP.

All THSteps dental, medical, vision, and mental health providers should submit claims for services rendered to foster care clients to Superior HealthPlan’s dental, vision, and mental health contractors.

For general provider information or authorizations, from STAR Health subcontractors, contact Superior Health Plan.

#### 6.2.1 Spell-Of-Illness

No spell-of-illness limitation exists for THSteps-eligible clients who are 20 years of age and younger when a medically necessary condition exists.

#### 6.2.2 Prescriptions

All Medicaid clients who are 20 years of age and younger receive unlimited medically necessary prescription drugs.

**Note:** HIPPP program clients who are enrolled in STAR Health should be removed from the HIPPP program and continue to receive their benefits under the STAR Health program.
7 Children’s Medicaid Dental Services

7.1 Overview
The principal objectives of children’s Medicaid managed care dental services are to provide quality, comprehensive dental services in a manner that improves oral health of clients through preventative care, health education, and early intervention and to promote improved access to quality care, thereby significantly improving health outcomes for the target populations.

7.2 Children’s Medicaid Dental Services Model
Clients’ primary and preventive Medicaid dental services are provided statewide through Medicaid managed care DMOs. Each Medicaid managed care DMO is responsible for contracting with general dentists, pediatric dentists, and dental specialists to create a delivery network. Clients who receive their dental services through a Medicaid managed care DMO are required to select a DMO and a main dentist (or main dental home provider or dental home). The client selects the main dentist from a provider directory.

A main dentist means a provider who has agreed with a DMO to provide a dental home to clients and who is responsible for providing routine preventive, diagnostic, urgent, therapeutic, initial, and primary care to patients, maintaining the continuity of patient care, and initiating referral for care. Provider types that can serve as main dental home providers are general dentist and pediatric dentist. RHCs and FQHCs are also eligible to serve as a Main Dentist.

The First Dental Home Initiative is included in this model.

Refer to: Subsection 4.2.9, “First Dental Home” in the Children’s Services Handbook (Vol. 2, Provider Handbooks).

7.3 Client Eligibility
Most children who are 20 years of age and younger will receive their dental services through Medicaid managed care DMO.

Populations that will not receive services through the children’s Medicaid managed care DMOs are:
- Medicaid recipients who are 21 years of age or older.
- Recipients who reside in an institution (i.e. nursing homes, state supported living centers, or ICF-IID).
- Recipients in the STAR Health Program.

7.4 Client Enrollment
Clients choose a DMO and Main Dentist. To maximize enrollment, the children’s Medicaid dental services offer four alternative ways that clients can enroll:
- Telephone Enrollment. A client can enroll in a DMO by calling 1-800-964-2777 (telecommunications device for the deaf (TDD): 1-800-267-5008) A customer care representative will provide essential education about the program and details needed for enrollment.
• Mail-in Enrollment. If calling is not convenient, a client may enroll by completing the an enrollment form and dropping it in the mail using the postage-paid, self-addressed envelope. Enrollment forms are mailed to all eligible mandatory clients along with information explaining the services and how to choose a Main Dentist.

• Onsite Enrollment. In addition to telephone and mail-in enrollment, clients can enroll by talking with customer care representative at a local HHSC office, at Women, Infants, and Children (WIC) classes, community facilities, or during enrollment events.

• Default Enrollment. The final method of enrollment is through an assignment process. If a client does not exercise the right to choose a dental and Main Dentist, the client will be assigned to a DMO. After the default assignment is made, the DMO will assign the client a Main Dentist.

7.5 Authorizations for Children’s Medicaid Managed Care Dental Services (Non-orthodontia Services)

Authorization requests for services administered by the client’s DMO must be submitted to the client’s DMO according to the guidelines specific to the plan under which the client is covered.

If a client is new to a DMO and has an open authorization for covered dental services from TMHP or another HHSC-contracted Medicaid managed care DMO, the DMO must accept that authorization and cannot require additional authorization or review.

TMHP authorizes and processes dental and emergent orthodontic services for clients who are 20 years of age and younger but have not yet enrolled in a DMO.

TMHP also authorizes services for the following clients:

• Dental services for Medicaid clients who are 21 years of age and older
• Dental and orthodontia services for all Medicaid clients, regardless of age, who reside in Medicaid-paid facilities such as nursing homes, state-supported living centers, or ICF-IIDs

Exception:STAR Health Foster Care Program clients receive dental and orthodontic services through DentaQuest.

7.6 Children’s Medicaid Dental Orthodontia Services

The Medicaid managed care DMOs will be responsible for authorizing, processing, and reimbursing any orthodontic services rendered to Texas Medicaid managed care clients. Claims for orthodontic services that were initially authorized by TMHP but later transitioned to a managed care DMO will be processed and reimbursed by the DMO. Providers should check client eligibility to identify the managed care DMO in which the client is enrolled.

TMHP will continue to process claims and claims adjustments for orthodontia services claims for clients who are ICF-IID residents.

If a Medicaid client is enrolled in a DMO for at least one month, is receiving orthodontic treatment, and either ages out of the program or loses eligibility, the DMO is responsible for completion of the course of treatment. The only exception is if the client is disenrolled with cause but is still Medicaid-eligible. For example, if a client goes into a State Supported Living Center, the DMO will no longer be responsible for services rendered.

8 Carve-Out Services

Some services are “carved out” of one or more of the managed care programs. Carved out services are those that are rendered to Medicaid managed care clients but are processed for payment consideration by TMHP rather than an MCO or DMO.
The following tables show the services that are partially or completely carved out of the MCO and DMO managed care program:

<table>
<thead>
<tr>
<th>Carve Out</th>
<th>STAR</th>
<th>STAR Kids</th>
<th>STAR+PLUS</th>
<th>STAR Health</th>
</tr>
</thead>
<tbody>
<tr>
<td>Nutritional products through WIC</td>
<td>TMHP</td>
<td>TMHP</td>
<td>TMHP</td>
<td>TMHP</td>
</tr>
<tr>
<td>DSHS MH rehabilitation (w/ Modifier HZ)</td>
<td>TMHP</td>
<td>TMHP</td>
<td>TMHP</td>
<td>MCO</td>
</tr>
<tr>
<td>County Indigent Health Care Program (CIHCP)</td>
<td>TMHP</td>
<td>TMHP</td>
<td>TMHP</td>
<td>TMHP</td>
</tr>
<tr>
<td>Early Childhood Intervention (ECI) case management</td>
<td>TMHP</td>
<td>TMHP</td>
<td>TMHP</td>
<td>TMHP</td>
</tr>
<tr>
<td>ECI specialized skills training</td>
<td>TMHP</td>
<td>TMHP</td>
<td>TMHP</td>
<td>TMHP</td>
</tr>
<tr>
<td>Family planning services for Dell Children's managed care health plan</td>
<td>TMHP</td>
<td>N/A</td>
<td>N/A</td>
<td>N/A</td>
</tr>
<tr>
<td>Texas School Health and Related Services (SHARS)</td>
<td>TMHP</td>
<td>TMHP</td>
<td>TMHP</td>
<td>TMHP</td>
</tr>
<tr>
<td>Elevated lead investigation services</td>
<td>TMHP</td>
<td>TMHP</td>
<td>TMHP</td>
<td>TMHP</td>
</tr>
<tr>
<td>DSHS TB providers</td>
<td>TMHP</td>
<td>TMHP</td>
<td>TMHP</td>
<td>TMHP</td>
</tr>
<tr>
<td>DSHS targeted case management (w/ Modifier HZ)</td>
<td>TMHP</td>
<td>TMHP</td>
<td>TMHP</td>
<td>TMHP</td>
</tr>
<tr>
<td>HHSC Blind Children’s Vocational Discovery and Development Program</td>
<td>TMHP</td>
<td>TMHP</td>
<td>TMHP</td>
<td>TMHP</td>
</tr>
<tr>
<td>Case Management for Children and Pregnant Women</td>
<td>TMHP</td>
<td>TMHP</td>
<td>TMHP</td>
<td>TMHP</td>
</tr>
<tr>
<td>Community First Choice (CFC) services*</td>
<td>TMHP</td>
<td>MCO</td>
<td>MCO</td>
<td>MCO</td>
</tr>
<tr>
<td>Personal Care Services (PCS)</td>
<td>TMHP</td>
<td>MCO</td>
<td>MCO</td>
<td>MCO</td>
</tr>
<tr>
<td>Youth Empowerment Services (YES) Waiver</td>
<td>TMHP</td>
<td>TMHP</td>
<td>TMHP</td>
<td>TMHP</td>
</tr>
<tr>
<td>Home and Community Based Services Adult Mental Health (HCBS-AMH)</td>
<td>TMHP</td>
<td>TMHP</td>
<td>TMHP</td>
<td>TMHP</td>
</tr>
</tbody>
</table>

* CFC services are carved out for STAR Kids clients who receive services through CLASS, DBMD, HCS, or TxHiMl.

<table>
<thead>
<tr>
<th>Dental Carve Out</th>
<th>Children's Medicaid Dental</th>
<th>STAR Health</th>
</tr>
</thead>
<tbody>
<tr>
<td>THSteps Dental</td>
<td>DMO</td>
<td>MCO</td>
</tr>
<tr>
<td>Orthodontia for most clients who are 20 years of age and younger</td>
<td>DMO</td>
<td>MCO</td>
</tr>
<tr>
<td>ICF-IID Dental Services</td>
<td>TMHP</td>
<td>TMHP</td>
</tr>
</tbody>
</table>

Note: Authorizations and claims for SSI clients who are enrolled in the STAR Program are submitted to the client's MCO or DMO.

Providers must submit authorization requests and claims for services that are carved out of the managed care program to TMHP according to the fee-for-service guidelines that are established for the same service.
Refer to: “Section 6: Claims Filing” (Vol. 1, General Information) for more information about applicable authorization request and claims filing guidelines.

8.1 Family Planning Carve-Out Services

Some family planning services are carved-out services for Texas Medicaid clients whose managed care benefits are administered through Dell Children’s Health Plan. These carved out services may be considered for payment by Texas Medicaid through TMHP if the service has been denied by the health plan as a family planning service.

All Dell Children’s Medicaid providers should submit family planning claims using the CMS-1500 paper claim form, or electronic equivalent, to the client’s managed care health plan to receive the health plan’s denial.

Important: Services that are denied by the health plan for any other reason will not be considered for reimbursement by Texas Medicaid.

8.1.1 Professional and Outpatient Claims

For affected claims to be eligible for reimbursement through TMHP, providers must do the following:

1) Submit the claim to the client’s managed care health plan to receive the health plan’s denial. Claims that are submitted electronically using TexMedConnect will automatically be forwarded to the client’s Medicaid managed care plan.

2) Submit a paper claim to TMHP upon receipt of the health plan’s denial. All applicable documentation must be included with the paper claim, including, but not limited to:

   • The health plan’s EOB document that indicates the denial code with its description and the date the EOB was issued. The denial must indicate that the service was denied because it was a family planning service. The EOB date will be used to calculate the filing deadline for the claim submission.

   • All documentation for family planning services including the Sterilization Consent Form and any other documentation that is required by Texas Medicaid.

Note: A paper claim is required because TMHP automatically forwards electronic claims to the client’s health plan without processing. Providers must comply with all filing deadlines unless otherwise specified below in this article.

Refer to: The Gynecological, Obstetrics, and Family Planning Title XIX Services Handbook (Vol. 2, Provider Handbooks) for Texas Medicaid guidelines for family planning services.

8.1.1.1 Claim Forms for Submission to TMHP

After receiving the health plan’s denial, Medicaid family planning services providers should submit paper claim forms to TMHP as follows:

• Providers that contract with the HHSC Family Planning Program should submit claims on a 2017 paper claim form along with the health plan’s denial.

• Providers that do not contract with the HHSC Family Planning Program should submit claims on a CMS-1500 paper claim form along with the health plan’s denial.

Providers should submit the health plan’s EOB document that indicates the denial code with its description and the date that the EOB was issued. The denial must indicate that the service was denied because it was a family planning service. The EOB date will be used to calculate the filing deadline for the claim submission.

Providers must comply with all filing deadlines.
The initial paper claim will be denied by TMHP. TMHP will automatically reprocess for payment consideration any claim that has been denied only with EOB 00081, “Services billed to TMHP in error. Bill HMO.” TMHP will reprocess only those claims that were denied with EOB 00081 as the only EOB message on the claim. If a claim has been denied with other EOB messages in addition to EOB 00081, the provider must resolve the other reasons for denial through the standard appeals process before TMHP can reprocess the claim for payment of the carved-out services.

8.1.2 Inpatient Claims

For affected claims to be eligible for reimbursement through TMHP, providers must do the following:

1) Submit the claim to the client’s managed care health plan to receive the health plan’s denial. Claims that are submitted electronically using TexMedConnect will automatically be forwarded to the client’s Medicaid managed care plan.

2) Submit a paper claim to HHSC Administrative Appeals upon receipt of the health plan’s denial. All applicable documentation must be included with the paper claim, including, but not limited to:
   - The health plan’s EOB document that indicates the denial code with its description and the date the EOB was issued. The denial must indicate that the service was denied because it was a family planning service. The EOB date will be used to calculate the filing deadline for the claim submission.
   - All documentation for family planning services including Sterilization Consent Forms and Hysterectomy Acknowledgements Forms, and any other documentation that is required by Texas Medicaid.

HHSC Administrative Appeals will send the family planning services inpatient claims to TMHP for reprocessing. Medical portions of the claims will be denied by Texas Medicaid because they are covered under the client’s health plan and will not be considered for reimbursement through TMHP. The services that were denied by the health plan as family planning services will be considered for payment according to Medicaid guidelines.

Refer to: The Gynecological, Obstetrics, and Family Planning Title XIX Services Handbook (Vol. 2, Provider Handbooks) for Texas Medicaid guidelines for family planning services.

“Section 7: Appeals” (Vol. 1, General Information) for additional information about administrative appeals.

9 Other State Health-Care Programs

The services available under the following programs are administered by TMHP or other state programs and not by the client’s MCO or DMO:

- Healthy Texas Women (HTW) program - HHSC/TMHP
- HHSC Family Planning Program contracted services - HHSC/TMHP
- Medicaid Medical Transportation Program (MTP)
- CHIP Perinatal—CHIP Perinatal provides prenatal care to the unborn children of pregnant women who have an income of up to 202 percent of the federal poverty level and who are not eligible for other Medicaid programs or traditional CHIP. CHIP Perinatal covered services include prenatal care, labor with delivery, and two postpartum visits.
• For CHIP perinates in families that have an income at or below the Medicaid eligibility threshold, facility charges associated with labor and delivery are covered by Emergency Medicaid, and professional charges associated with labor and delivery are covered by the CHIP Perinatal MCO.

• For CHIP perinates in families that have an income above the Medicaid eligibility threshold, the facility and professional charges associated with labor with delivery are covered by the CHIP Perinatal MCO.

• Health Insurance Premium Payment (HIPP) Program—The HIPP Program reimburses clients for the cost of medical insurance premiums when Medicaid finds it more cost-effective to reimburse a Medicaid client’s group health insurance premiums than to reimburse the client’s medical bills directly through Medicaid.

Refer to:
- “Section 4: Client Eligibility” (Vol. 1, General Information).

Claims and authorization requests for the services listed above must be submitted according to the established guidelines.

10 Contact Information

The following information can be used to communicate with TMHP:

<table>
<thead>
<tr>
<th>Correspondence</th>
<th>Contact Information</th>
</tr>
</thead>
<tbody>
<tr>
<td>All correspondence for services rendered to clients who are enrolled with a Texas Medicaid health/DMO</td>
<td>Contact the client’s health/DMO.</td>
</tr>
<tr>
<td>Claims, authorizations, and other TMHP correspondence for transactions that are processed by TMHP.</td>
<td>Volume 1, “Written Communication With TMHP,” for the list of post office box addresses that must be used for specific items.</td>
</tr>
<tr>
<td>HHSC contact information for clients of STAR+PLUS, STAR, STAR Kids, STAR Health, and Children’s Medicaid dental services</td>
<td>1-800-252-8263</td>
</tr>
<tr>
<td>TMHP Contact Center</td>
<td>1-800-925-9126</td>
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MEDICAL AND NURSING SPECIALISTS, PHYSICIANS, AND PHYSICIAN ASSISTANTS

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1 General Information

The information in this handbook is intended for Texas chiropractors, nurse practitioners (NP), clinical nurse specialists (CNS), certified nurse midwives (CNM), certified registered nurse anesthetists (CRNA), podiatrists, geneticists, maternity service clinics, physicians, and physician assistants. The handbook provides information about Texas Medicaid’s benefits, policies, and procedures.

Important: All providers are required to read and comply with "Section 1: Provider Enrollment and Responsibilities" (Vol. 1, General Information). In addition to required compliance with all requirements specific to Texas Medicaid, it is a violation of Texas Medicaid rules when a provider fails to provide healthcare services or items to Medicaid clients in accordance with accepted medical community standards and standards that govern occupations, as explained in Title 1 Texas Administrative Code (TAC) §371.1659. Accordingly, in addition to being subject to sanctions for failure to comply with the requirements that are specific to Texas Medicaid, providers may also be subject to Texas Medicaid sanctions for failure, at all times, to deliver healthcare items and services to Medicaid clients in full accordance with all applicable licensure and certification requirements including, without limitation, those related to documentation and record maintenance.

Refer to: “Section 1: Provider Enrollment and Responsibilities” (Vol. 1, General Information).
Subsection 2.2, “Provider Enrollment and Responsibilities” in the Medicaid Managed Care Handbook (Vol. 2, Provider Handbooks).
Section 5, “THSteps Medical” in the Children’s Services Handbook (Vol. 2, Provider Handbooks).

For information on Advanced Practice Registered Nurses (APRNs), refer to:
Section 3, “Certified Nurse Midwife (CNM)” in this handbook.
Subsection 4.1, “Enrollment” in this handbook for information about CRNAs.
Subsection 8.1, “Enrollment” in this handbook for information about NPs and CNSs
Section 9, “Physician” in this handbook.

1.1 Payment Window Reimbursement Guidelines for Services Preceding an Inpatient Admission

According to the three-day and one-day payment window reimbursement guidelines, most professional and outpatient diagnostic and nondiagnostic services that are rendered within the designated timeframe of an inpatient hospital stay and are related to the inpatient hospital admission will not be reimbursed separately from the inpatient hospital stay if the services are rendered by the hospital or an entity that is wholly owned or operated by the hospital.

These reimbursement guidelines do not apply in the following circumstances:

- The professional services are rendered in the inpatient hospital setting.
- The hospital and the physician office or other entity are both owned by a third party, such as a health system.
- The hospital is not the sole or 100-percent owner of the entity.
Refer to: Subsection 3.7.4.14, “Payment Window Reimbursement Guidelines” in the Inpatient and Outpatient Hospital Services Handbook (Vol. 2, Provider Handbooks) for additional information about the payment window reimbursement guidelines.

2 Chiropractic Manipulative Treatment (CMT)

2.1 Enrollment
To enroll in Texas Medicaid, a doctor of chiropractic medicine (DC) must be licensed by the Texas Board of Chiropractic Examiners and enrolled as a Medicare provider.

Providers cannot be enrolled if their license is due to expire within 30 days; a current license must be submitted.

2.2 Services, Benefits, Limitations, and Prior Authorization
CMT performed by a chiropractor licensed by the Texas State Board of Chiropractic Examiners is a benefit of Texas Medicaid.

CMT is limited to an acute condition or an acute exacerbation of a chronic condition for a maximum of 12 visits in a consecutive 12-month period, and a maximum of one visit per day. The 12-month period consists of 12 consecutive months, beginning with the date the client receives the first treatment.

If the condition persists more than 180 days from the start of therapy, the condition is considered chronic, and treatment is no longer considered acute.

CMT is not a benefit of Texas Medicaid for maintenance therapy when:

- Further clinical improvement cannot reasonably be expected from continuous ongoing care.
- The chiropractic treatment becomes supportive rather than corrective in nature.

CMT may be reimbursed when billed using procedure codes 98940, 98941, or 98942.

Procedure codes 98940, 98941, and 98942 must be submitted with the AT modifier. The AT modifier is used to identify treatment provided for an acute condition or an exacerbation of a chronic condition that persists for 180 days or less from the start date of treatment. Providers may file an appeal for a claim denied beyond the 180 days of treatment with documentation supporting that further clinical improvement can be reasonably expected, maximal improvement has not been reached, and further improvement has not ceased.

Procedure code 98940 will be denied as part of another service when billed for the same date of service as 98941 or 98942 by any provider.

Procedure code 98941 will be denied as part of another service when billed for the same date of service as 98942 by any provider.

Texas Medicaid does not reimburse chiropractors for X-ray services, office visits, injections, supplies, appliances, spinalator treatments, laboratory services, physical therapy, or other adjunctive services furnished by themselves or by others under their orders or directions. Additionally, braces or supports, even though ordered by a physician (doctor of medicine [MD] or doctor of osteopathy [DO]) and supplied by a chiropractor are not reimbursable items.

CMT is reimbursed only for a diagnosis of subluxation of the spine. The level of subluxation must be indicated by the appropriate diagnosis codes listed below:

<table>
<thead>
<tr>
<th>Diagnosis Codes</th>
</tr>
</thead>
<tbody>
<tr>
<td>M9900</td>
</tr>
</tbody>
</table>
2.2.1 Prior Authorization

Prior authorization is not required for CMT services.

2.3 Documentation Requirements

Manipulations must be provided in accordance with an ongoing, written treatment plan that supports medical necessity of an acute condition or an acute exacerbation of a chronic condition.

Documentation that supports medical necessity for the treatment plan includes all of the following:

- Diagnosis
- Region(s) treated
- Degree of severity
- Impairment characteristics
- Physical examination findings, X-ray, or other pertinent findings
- Specific statements of short- and long-term goals
- A reasonable estimate of when the goals will be reached (estimated duration of treatment)
- Frequency of treatment (number of times per week)
- Equipment and/or the techniques utilized

The treatment plan must be updated as the client’s condition changes. Treatment plans must be maintained in the medical records and are subject to retrospective review.

2.4 Claims Filing and Reimbursement

2.4.1 Claims Information

Chiropractic services must be submitted to TMHP in an approved electronic claims format or on a CMS-1500 paper claim form. Providers may purchase CMS-1500 claim forms from the vendor of their choice. TMHP does not supply them.

When completing a CMS-1500 claim form, all required information must be included on the claim, as TMHP does not key any information from claim attachments. Superbills, or itemized statements, are not accepted as claim supplements.

Refer to: “Section 3: TMHP Electronic Data Interchange (EDI)” (Vol. 1, General Information) for information on electronic claims submissions.

Subsection, “Section 6: Claims Filing” in “Section 6: Claims Filing” (Vol. 1, General Information) for general information about claims filing.

Subsection 6.5, “CMS-1500 Paper Claim Filing Instructions” in “Section 6: Claims Filing” (Vol. 1, General Information) for instructions on completing paper claims. Blocks that are not referenced are not required for processing by TMHP and may be left blank.

2.4.2 Reimbursement

The Medicaid rates for chiropractic manipulative treatment (CMT) are reimbursed in accordance with 1 TAC §355.8085. See the online fee lookup (OFL) or the applicable fee schedule on the TMHP website at www.tmhp.com.
Texas Medicaid implemented mandated rate reductions for certain services. The Online Fee Lookup (OFL) and static fee schedules include a column titled “Adjusted Fee” to display the individual fees with all mandated percentage reductions applied. Additional information about rate changes is available on the TMHP website at [www.tmhp.com/resources/rate-and-code-updates](http://www.tmhp.com/resources/rate-and-code-updates).

**Note:** Certain rate reductions including, but not limited to, reductions by place of service, client type program, or provider specialty may not be reflected in the Adjusted Fee column.

**Refer to:** Subsection 2.2, “Fee-for-Service Reimbursement Methodology” in “Section 2: Texas Medicaid Fee-for-Service Reimbursement” (*Vol. 1, General Information*) for more information about reimbursement.

### 3 Certified Nurse Midwife (CNM)

#### 3.1 Provider Enrollment

To enroll in Texas Medicaid, a CNM must be licensed as a registered nurse and as an advanced practice registered nurse (APRN) by the Texas Board of Nursing (BON), and be authorized to practice as a nurse-midwife. A registered nurse under the multistate licensure compact may be licensed in another state but certified as an APRN for the state of Texas by the Texas BON. Texas Medicaid accepts a signed letter of certification from the Texas BON as documentation of appropriate licensure and certification for enrollment.

The American Midwifery Certification Board (AMCB) is responsible for the certification requirements of CNMs.

**Refer to:** The HHSC website at [www.healthytexaswomen.org](http://www.healthytexaswomen.org) for information about family planning and the locations of family planning clinics that receive funding from the HHSC Family Planning Program.

Providers cannot be enrolled if their license is due to expire within 30 days; a current license must be submitted.

All providers of laboratory services must comply with the rules and regulations of the Clinical Laboratory Improvement Amendments (CLIA). Providers not complying with CLIA are not reimbursed for laboratory services.

All APRNs (including CNMs, CRNAs, CNSs, and NPs) are enrolled within the categories of practice as determined by the Texas BON. CNSs and NPs must enroll as an APRN; CNMs and CRNAs may enroll using their specific titles.

A CNM must identify the licensed physician or group of physicians with whom there is an arrangement for referral and consultation if medical complications arise. Upon initial enrollment and upon re-enrollment, the CNM must complete and submit to TMHP, along with the Texas Medicaid Provider Enrollment Application, the Physician’s Letter of Agreement form that affirms the CNM’s referring or consulting physician arrangement. A separate letter of agreement must be submitted for each physician or group of physicians with whom an arrangement is made. This agreement must be signed by the CNM and the physician. The collaborating physician does not have to be a participating provider in Texas Medicaid. According to TAC, §354.1252 (3), if the collaborating physician or group is not a participating provider in Texas Medicaid, the CNM must inform clients of their potential financial responsibility. If the arrangement is changed or canceled, the CNM must notify TMHP Provider Enrollment in writing and a new letter of agreement must be completed and submitted to TMHP within 10 business days of the change or cancellation.

CNMs are encouraged to participate in or make referrals to family planning agencies.
Refer to: “Section 1: Provider Enrollment and Responsibilities” (Vol. 1, General Information) for more information about enrollment in Texas Medicaid.

Subsection 5.2, “Enrollment” in the Children’s Services Handbook (Vol. 2, Provider Handbooks) for more information about enrollment in the THSteps Program.


### 3.1.1 Enrollment in Texas Health Steps (THSteps)

CNMs may enroll as providers of THSteps medical checkups for newborns and adolescent females.

### 3.2 Services, Benefits, Limitations, and Prior Authorization

CNM providers may be reimbursed for family planning, obstetrical, neonatal, and primary care services.

#### 3.2.1 Deliveries

CNM providers may be reimbursed for procedure code 59409, 59410, 59612, or 59614 for delivery services.

Refer to: Section 4, “Obstetric Services” in the Gynecological, Obstetrics, and Family Planning Title XIX Services Handbook (Vol. 2, Provider Handbooks) for billing requirements.

#### 3.2.2 Newborn Services

Routine newborn care may be reimbursed to CNM providers.

Refer to: Subsection 5.3.9, “Newborn Examination” in the Children’s Services Handbook (Vol. 2, Provider Handbooks).

Subsection 9.2.44, “Newborn Services” in this handbook for additional guidelines and limitations.

#### 3.2.3 Prenatal and Postpartum Services

CNM and physician providers are limited to a combined total of 20 outpatient prenatal care visits and 1 postpartum care visit per pregnancy. Normal pregnancies are anticipated to require around 11 visits per pregnancy and high-risk pregnancies are anticipated to require around 20 visits per pregnancy. If more than 20 visits are medically necessary, the provider can appeal with documentation supporting pregnancy complications. The high-risk client’s medical record documentation should reflect the need for increased visits and is subject to retrospective review.

When billing for prenatal services, use modifier TH with the appropriate evaluation and management procedure code to the highest level of specificity.

Postpartum care provided after discharge must be billed using procedure code 59430. Only one postpartum visit is allowed per pregnancy.

Refer to: Section 4, “Obstetric Services” in the Gynecological, Obstetrics, and Family Planning Title XIX Services Handbook (Vol. 2, Provider Handbooks) for billing requirements.

#### 3.2.4 Laboratory and Radiology Services

Laboratory (including pregnancy tests) and radiology services that are rendered during pregnancy must be billed separately from prenatal care visits.

#### 3.2.5 Prior Authorization

Prior authorization is not required for any of these services except delivery in the home. For prior authorization of a home delivery and the related supplies (procedure code S8415), the CNM must submit a written request for prior authorization during the client’s third trimester of pregnancy. The CNM must include a statement signed by a licensed physician who has examined the client during the third
trimester and determined at that time that she is not at high risk and is suitable for a home delivery. Documentation must also include a plan for access to emergency transport for mother and neonate, if needed. Requests for home delivery prior authorizations must be submitted to the TMHP Medical Director at the following address:

Texas Medicaid & Healthcare Partnership
Special Medical Prior Authorization
12357-B Riata Trace Parkway, Suite 100
Austin, TX 78727
Fax: 1-512-514-4213

Claims submitted for home deliveries performed by a CNM without prior authorization will be denied.

3.2.6 Documentation Requirements

All services require documentation to support the medical necessity of the service rendered, including CNM services.

CNM services are subject to retrospective review and recoupment if documentation does not support the service billed.

3.2.7 Claims Filing and Reimbursement

CNMs must bill maternity services in one of two ways: itemizing each service individually on one claim form and filing at the time of delivery (the filing deadline is applied to the date of delivery) or itemizing each service individually and submitting claims as the services are rendered (the filing deadline is applied to each individual date of service).

CNM services must be submitted to TMHP in an approved electronic format or on the CMS-1500 claim form. Providers may purchase CMS-1500 claim forms from the vendor of their choice. TMHP does not supply the forms.

When completing a CMS-1500 claim form all required information must be included on the claim, as TMHP does not key any information from claim attachments. Superbills, or itemized statements, are not accepted as claim supplements.

According to 1 TAC §355.8161(a), the Medicaid rate for CNMs is 92 percent of the rate paid to a physician (doctor of medicine [MD] or doctor of osteopathy [DO]) for the same service and 100 percent of the rate paid to physicians for laboratory services, X-ray services, and injections.

Note: CNM providers who are enrolled in Texas Medicaid as THSteps providers also receive 92 percent of the rate paid to a physician for THSteps services when a claim is submitted with their THSteps provider identifier as the billing provider.

Physicians who submit a claim using the physician’s own provider identifier for services provided by a CNM must submit modifier SB on each claim detail if the physician does not make a decision regarding the client’s care or treatment on the same date of service as the billable medical visit.

Physicians may be reimbursed 92 percent of the established reimbursement rate for services provided by a CNM if the physician does not make a decision regarding the client’s care or treatment on the same date of service as the billable medical visit. This 92 percent reimbursement rate does not apply to laboratory services, X-ray services, and injections provided by a CNM.

Providers can refer to the Online Fee Lookup (OFL) or the applicable fee schedule on the TMHP website at www.tmhp.com.

Texas Medicaid implemented mandated rate reductions for certain services. The OFL and static fee schedules include a column titled “Adjusted Fee” to display the individual fees with all mandated percentage reductions applied. Additional information about rate changes is available on the TMHP website at www.tmhp.com/resources/rate-and-code-updates.
Refer to: Subsection 4.1, “General Medicaid Eligibility” in “Section 4: Client Eligibility” (Vol. 1, General Information) for information about crossover payments.

“Section 3: TMHP Electronic Data Interchange (EDI)” (Vol. 1, General Information) for information on electronic claims submissions.

Subsection 6.1, “Claims Information” in “Section 6: Claims Filing” (Vol. 1, General Information) for general information about claims filing.


Subsection 2.2, “Fee-for-Service Reimbursement Methodology” in “Section 2: Texas Medicaid Fee-for-Service Reimbursement” (Vol. 1, General Information) for more information about reimbursement.

4 Certified Registered Nurse Anesthetist (CRNA)

4.1 Enrollment

To enroll in Texas Medicaid, a CRNA must be licensed as a registered nurse (RN) and as an APRN by the Texas BON and must be currently certified by the Council on Certification of Nurse Anesthetists or the Council on Recertification of Nurse Anesthetists. An RN under the multistate licensure compact may be licensed in another state but certified as an APRN for the state of Texas by the Texas BON. Texas Medicaid accepts a signed letter of certification from the Texas BON as acceptable documentation of appropriate licensure and certification for enrollment.

Medicare enrollment is a prerequisite for enrollment as a Medicaid provider. A current copy of the provider’s Council on Certification of Nurse Anesthetists or Recertification of Nurse Anesthetists Certificate must be submitted with the Medicaid provider enrollment application.

Providers cannot be enrolled if their license is due to expire within 30 days; a current license must be submitted.

4.2 Services, Benefits, Limitations, and Prior Authorization

Medically necessary services that are performed by a CRNA are benefits if the services are within the scope of the CRNA’s practice as defined by state law; are prescribed, supervised by, and provided under the direction of a supervising physician (MD or DO), dentist, or podiatrist licensed in the state in which they practice and to the extent allowed by state law; and are provided under one of the following conditions:

- There is no physician anesthesiologist on the medical staff of the facility where the services are provided (e.g., rural settings).
- There is no physician anesthesiologist available to provide the services, as determined by the policies of the facility in which the services are provided.
- The physician, dentist, or podiatrist who performs the procedure that requires the services specifically requests the services of a CRNA.
- The eligible client who requires the services specifically requests the services of a CRNA.
- The CRNA is scheduled or assigned to provide the services according to the policies of the facility in which the services are provided.
- The services are provided by the CRNA in connection with a medical emergency.

Texas Medicaid does not reimburse the CRNA for equipment, drugs, or supplies.

### 4.2.1 Prior Authorization

Services performed by a CRNA are subject to the same prior authorization guidelines as services performed by other provider types.

### 4.3 Documentation Requirements

All services require documentation to support the medical necessity of the services rendered, including CRNA services. CRNA services are subject to retrospective review and recoupment if documentation does not support the service billed.

### 4.4 Claims Filing and Reimbursement

#### 4.4.1 Claims Information

All CRNA services must be billed with a CRNA individual provider identifier or a CRNA group provider identifier. No payment for CRNA services will be made under a hospital or physician provider identifier.

CRNA services must be submitted to TMHP in an approved electronic format or on the CMS-1500 claim form. Providers may purchase CMS-1500 claim forms from the vendor of their choice. TMHP does not supply the forms.

When completing a CMS-1500 claim form, all required information must be included on the claim, as TMHP does not key any information from claim attachments. Superbills, or itemized statements, are not accepted as claim supplements.

Refer to: “Section 3: TMHP Electronic Data Interchange (EDI)” *(Vol. 1, General Information)* for information on electronic claims submissions.

“Section 6: Claims Filing” *(Vol. 1, General Information)* for general information about claims filing.

Subsection 6.5, “CMS-1500 Paper Claim Filing Instructions” in “Section 6: Claims Filing” *(Vol. 1, General Information)* for instructions on completing paper claims. Blocks that are not referenced are not required for processing by TMHP and may be left blank.

Subsection 9.2.7.9.3, “CRNA, AA, and Other Qualified Professional Services” in this handbook for more information on billing for CRNA services.

#### 4.4.1.1 Interpreting the R&S Report

The Billed Qty field on the Remittance and Status (R&S) Report reflects only the number of time units TMHP processes. The Relative Value Units (RVUs) assigned for the procedure code are not shown in the Billed Qty field.

#### 4.4.2 Reimbursement

A CRNA is reimbursed the lesser of either the CRNA’s billed charges or 92 percent of the reimbursement for the same service paid to a physician (M.D. or D.O.) other than an anesthesiologist in accordance with 1 TAC §355.8221. A CRNA under the supervision of an anesthesiologist is reimbursed the lesser of the billed charges or 50 percent of the calculated payment for a supervised anesthesia service.
Texas Medicaid implemented mandated rate reductions for certain services. The OFL and static fee schedules include a column titled “Adjusted Fee” to display the individual fees with all mandated percentage reductions applied. Additional information about rate changes is available on the TMHP website at www.tmhp.com/resources/rate-and-code-updates.

**Note:** Certain rate reductions including, but not limited to, reductions by place of service, client type program, or provider specialty may not be reflected in the Adjusted Fee column.

**Refer to:** Subsection 9.2.7.8, “Reimbursement Methodology” in this handbook for more information about flat fees and time based fees.

## 5 Geneticists

### 5.1 Enrollment

#### 5.1.1 Geneticists

Geneticists may enroll in Texas Medicaid as both a physician or physician group and as a geneticist. Enrollment as a geneticist allows enhanced reimbursement for specific procedure codes when a claim is submitted using the geneticist provider identifier.

A provider of genetic services that wishes to enroll in Texas Medicaid as a geneticist must complete the required Medicaid provider enrollment application forms and enter into a written agreement with HHSC. Texas Medicaid provider enrollment forms are available from TMHP, and may be downloaded on the TMHP website at www.tmhp.com. Completed applications are submitted to:

Texas Medicaid & Healthcare Partnership  
Provider Enrollment  
PO Box 200795  
Austin, TX 78720

Prior to enrollment, applicant qualifications for the provision of genetic services are verified and approved by DSHS. Verification and approval are administered through the Newborn Screening Unit. Basic contract requirements are as follows:

- The provider must be a clinical geneticist (MD or DO) who is board eligible or board certified by the American Board of Medical Geneticists (ABMG).
  
  **Note:** Board eligible providers are required to provide documentation reflecting completion of education requirements in a residency program in genetics.

- The provider must use a team of professionals to provide genetic evaluative, diagnostic, and counseling services. The team rendering the services must consist of professional staff including the clinical geneticist and at least one of the following: nurse, social worker, medical geneticist, or genetic counselor.

- Upon DSHS approval, TMHP issues a provider identifier and a performing provider identifier for the provision of genetic services.

- Providers cannot be enrolled if their license is due to expire within 30 days; a current license must be submitted.
5.2 Services, Benefits, Limitations, and Prior Authorization

Genetic services may be used to diagnose a condition, optimize disease treatment, predict future disease risk, and prevent adverse drug response. Genetic services may be provided by a physician, physician assistant, nurse practitioner, or clinical nurse specialist and typically include one or more of the following:

- Comprehensive physical exams
- Diagnosis, management, and treatment for clients with genetically-related health problems
- Evaluation of family histories for the client and the client’s family members
- Genetic risk assessment
- Genetic laboratory tests
- Interpretation and evaluation of laboratory test results
- Education and counseling of clients, their families, and other medical professionals on the causes of genetic disorders
- Consultation with other medical professionals to provide treatment

Pharmacogenetics encompasses the use of information encoded in DNA to help predict responses to medicines and thereby enhance the effectiveness and safety of medicines for individual clients.

Refer to: Subsection 9.2.40, “Pharmacogenetics” in this handbook for additional information about pharmacogenetics services.

5.2.1 Family History

It is important for primary care providers to recognize potential genetic risk factors in a client so that they can make appropriate referrals to a genetic specialist.

Obtaining an accurate family history is an important part of clinical evaluations, even when genetic abnormalities are not suspected. Knowing the family history may help health-care providers identify single-gene disorders or chromosomal abnormalities that occur in multiple family members or through multiple generations. Some genetic disorders that can be traced through an accurate family history include diabetes, hypertension, certain forms of cancer, and cystic fibrosis. Early identification of the client’s risk for one of these diseases can lead to early intervention and preventive measures that can delay onset or improve health conditions.

Using a genetics-specific questionnaire helps to obtain the information needed to identify possible genetic patterns or disorders. The most commonly used questionnaires are provided by the American Medical Association and include the Prenatal Screening Questionnaire, the Pediatric Clinical Genetics Questionnaire, and the Adult History Form.

5.2.2 Genetic Tests

Diagnostic tests to check for genetic abnormalities must be performed only if the test results will affect treatment decisions or provide prognostic information. Tests for conditions that are treated symptomatically are not appropriate since the treatment would not change. Providers who are uncertain whether a test is appropriate are encouraged to contact a geneticist or other specialist to discuss the client’s needs.

Any genetic testing and screening procedure must be accompanied by appropriate non-directive counseling, both before and after the procedure. Information must be provided to the client and family (if appropriate) about the possible risks and purpose and nature of the tests being performed.

The interpretation of certain tests, such as nuchal translucency, requires additional education and experience. Texas Medicaid supports national certification standards when available.
5.2.3 Laboratory Practices

For many heritable diseases and conditions, test performance and interpretation of test results require information about client race/ethnicity, family history, and other pertinent clinical and laboratory information. To facilitate test requests and ensure prompt initiation of appropriate testing procedures and accurate interpretation of test results, the requesting provider must be aware of the specific client information needed by the laboratory before tests are ordered.

To help providers make appropriate test selections and requests, handle and submit specimens, and provide clinical care, laboratories that perform molecular genetic testing for heritable diseases and conditions must educate providers that request services about the molecular genetic tests the laboratory performs. For each molecular genetic test, the laboratory must provide the following information:

- Indications for testing
- Relevant clinical and laboratory information
- Client race and ethnicity
- Family history
- Pedigree

Testing performed on a client to provide genetic information for a family member, and testing performed on a non-Medicaid client to provide genetic information for a Medicaid client are not benefits of Texas Medicaid.

5.2.4 Genetic Counselors

Genetic counselor services may be billed by a physician when the genetic counselor is under physician supervision and is an employee of the physician. Services provided by independent genetic counselors are not a benefit of Texas Medicaid.

5.2.5 Genetic Evaluation and Counseling by a Geneticist

A provider enrolled in Texas Medicaid as a geneticist may bill the following evaluation and management codes and receive an enhanced reimbursement. All other procedure codes must be billed under the geneticist’s individual, group, or laboratory provider identifier.

<table>
<thead>
<tr>
<th>Procedure Code</th>
<th>Limitations</th>
</tr>
</thead>
<tbody>
<tr>
<td>96040</td>
<td>None</td>
</tr>
<tr>
<td>99213</td>
<td>None</td>
</tr>
<tr>
<td>99214</td>
<td>None</td>
</tr>
<tr>
<td>99215</td>
<td>One per year, any provider</td>
</tr>
<tr>
<td>99244</td>
<td>One every three years, per provider</td>
</tr>
<tr>
<td>99245</td>
<td>One every three years, per provider</td>
</tr>
<tr>
<td>99254</td>
<td>One every three years, per provider</td>
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<tr>
<td>99255</td>
<td>One every three years, per provider</td>
</tr>
<tr>
<td>99402</td>
<td>One per pregnancy, per provider*</td>
</tr>
<tr>
<td>99404</td>
<td>One every three years, per provider</td>
</tr>
</tbody>
</table>

* Exception: Additional services are allowed when documentation of medical necessity to repeat a procedure accompanies a claim.

One office consultation, performed by a geneticist, (procedure code 99244 or 99245) may be considered for reimbursement if procedure code 99244, 99245, 99254, or 99255 has not been submitted by and reimbursed to that geneticist in the previous three years.
Inpatient consultations, performed by a geneticist, (procedure code 99254 or 99255) may be reimbursed once every three years regardless of whether an office consultation has been reimbursed in the previous three years.

5.2.6 Prior Authorization
Prior authorization is not required for services billed by a geneticist.

5.3 Documentation Requirements
All services require documentation to support the medical necessity of the service rendered, including genetic services. Genetic services are subject to retrospective review and recoupment if documentation does not support the service billed.

5.4 Claims Filing and Reimbursement

5.4.1 Claims Information
Genetic services must be submitted to TMHP in an approved electronic format or on a CMS-1500 claim form. Providers may purchase CMS-1500 claim forms from the vendor of their choice. TMHP does not supply the forms.

When completing a CMS-1500 claim form, all required information must be included on the claim, as information is not keyed from attachments. Superbills, or itemized statements, are not accepted as claim supplements.

TMHP representatives are available for provider questions about genetic services, such as reimbursement rates and procedures. For more information, call the TMHP Contact Center at 1-800-925-9126.

Refer to: “Section 3: TMHP Electronic Data Interchange (EDI)” (Vol. 1, General Information) for information on electronic claims submissions.

"Section 6: Claims Filing” (Vol. 1, General Information) for general information about claims filing.

Subsection 6.5, “CMS-1500 Paper Claim Filing Instructions” in “Section 6: Claims Filing” (Vol. 1, General Information) for instructions on completing paper claims. Blocks that are not referenced are not required for processing by TMHP and may be left blank.

5.4.2 Reimbursement
Genetic services providers are reimbursed according to the established allowable maximum fee schedule. Providers can refer to the OFL or the applicable fee schedule on the TMHP website at www.tmhp.com.

Texas Medicaid implemented mandated rate reductions for certain services. The OFL and static fee schedules include a column titled “Adjusted Fee” to display the individual fees with all mandated percentage reductions applied. Additional information about rate changes is available on the TMHP website at www.tmhp.com/resources/rate-and-code-updates.

Refer to: Subsection 2.2, “Fee-for-Service Reimbursement Methodology” in “Section 2: Texas Medicaid Fee-for-Service Reimbursement” (Vol. 1, General Information) for more information about reimbursement.
6   Licensed Midwife (LM)

6.1   Provider Enrollment
To enroll in Texas Medicaid, an LM must be licensed as a midwife by the Texas Department of Licensing and Regulation (TDLR).

Providers cannot be enrolled if their license is due to expire within 30 days; a current license must be submitted.

An LM must identify the licensed physician or group of physicians with whom there is an arrangement for referral and consultation if medical complications arise. Upon initial enrollment and upon re-enrollment, the LM must complete and submit to TMHP, along with the Texas Medicaid Provider Enrollment Application, the Physician’s Letter of Agreement form that affirms the LM’s referring or consulting physician arrangement. A separate letter of agreement must be submitted for each physician or group of physicians with whom an arrangement is made. This agreement must be signed by the LM and the physician.

If the arrangement is changed or canceled, the LM must notify TMHP Provider Enrollment in writing and a new letter of agreement must be completed and submitted to TMHP within 10 business days after the change or cancellation.

The referral physician or group does not have to be a participating provider in Texas Medicaid. According to TAC, §354.1253(c), if the referral physician or group is not a participating provider in Texas Medicaid, the LM must inform clients of their potential financial responsibility.

Refer to: “Section 1: Provider Enrollment and Responsibilities” (Vol. 1, General Information) for more information about enrollment in Texas Medicaid.

6.2   Services, Benefits, Limitations, and Prior Authorization
LM providers may be reimbursed for obstetrical and newborn care services provided in a freestanding birthing center that is also enrolled as a Texas Medicaid provider.

6.2.1   Deliveries
LM providers may be reimbursed for procedure code 59409 for delivery services.

Refer to: Section 4, “Obstetric Services” in the Gynecological, Obstetrics, and Family Planning Title XIX Services Handbook (Vol. 2, Provider Handbooks) for billing requirements.

6.2.2   Newborn Services
Newborn care procedure codes 99460 and 99463 may be reimbursed to LM providers.

Refer to: Subsection 9.2.44, “Newborn Services” in this handbook for additional guidelines and limitations.

6.2.3   Prenatal Services
LM providers must include modifier TH with the appropriate evaluation and management procedure code (99202, 99211, or 99212) for prenatal services.

LM providers are limited to a total of 20 outpatient prenatal care visits, performed in a birthing center, per pregnancy. Normal pregnancies are anticipated to require around 11 visits per pregnancy and high-risk pregnancies are anticipated to require around 20 visits per pregnancy. If more than 20 visits are medically necessary, the provider can appeal with documentation supporting pregnancy complications. The high-risk client’s medical record documentation should reflect the need for increased visits and is subject to retrospective review.
If a client is discharged before delivery, LM providers may submit procedure code 99218, 99219, or 99220 for labor services only. Clinical documentation that clearly demonstrates the level of medical decision-making (i.e., moderate or complex) must be included in the client’s medical record. All medical documentation is subject to retrospective review. Services that are not supported by the medical documentation are subject to recoupment.

Refer to: Section 4, “Obstetric Services” in the Gynecological, Obstetrics, and Family Planning Title XIX Services Handbook (Vol. 2, Provider Handbooks) for billing requirements.

6.2.4 Prior Authorization
Prior authorization is not required for services billed by an LM.

6.2.5 Documentation Requirements
All services require documentation to support the medical necessity of the service rendered, including LM services.

LM services are subject to retrospective review and recoupment if documentation does not support the service billed.

6.2.6 Claims Filing and Reimbursement
LM services must be submitted to TMHP in an approved electronic format or on the CMS-1500 claim form. Providers may purchase CMS-1500 claim forms from the vendor of their choice. TMHP does not supply the forms.

When completing a CMS-1500 claim form all required information must be included on the claim, as TMHP does not key any information from claim attachments. Superbills, or itemized statements, are not accepted as claim supplements.

According to 1 TAC §355.8161 (b), the Medicaid rate for LMs is 70 percent of the rate paid to a physician (doctor of medicine [MD] or doctor of osteopathy [DO]) for the same service.

Providers can refer to the Online Fee Lookup (OFL) or the applicable fee schedule on the TMHP website at www.tmhp.com.

Texas Medicaid implemented mandated rate reductions for certain services. The OFL and static fee schedules include a column titled “Adjusted Fee” to display the individual fees with all mandated percentage reductions applied. Additional information about rate changes is available on the TMHP website at www.tmhp.com/resources/rate-and-code-updates.

7 Maternity Service Clinics (MSC)

7.1 Provider Enrollment
To enroll in Texas Medicaid, MSCs must submit a complete application and meet the following requirements:

- Must be a facility that is not an administrative, organizational, or financial part of a hospital.
- Must be organized and operated to provide maternity clinic services to outpatients.
- Must comply with all applicable federal, state, and local laws and regulations.
- Must employ or have a contractual agreement or formal arrangement with a licensed MD or DO who assumes professional responsibility for the services provided to the clinic’s patients.
- Must adhere to the Bureau of Maternal and Child Health Maternity Guidelines, dated June 20, 1988, and subsequent revisions issued by the Texas Department of State Health Services, unless otherwise specified by the department or its designee.
• Must ensure that services provided to each patient are commensurate with the patient’s risk assessment and are documented in the patient’s medical record.

The supervising physician’s license information must be provided. Providers cannot be enrolled in Texas Medicaid if their licenses are due to expire within 30 days.

Medicare certification is not a prerequisite for MSC enrollment.

7.1.1 Physician Responsibility

To meet the requirement to assume professional responsibility for the services provided to the clinic’s clients, the supervising physician must do the following:

• See the client at least once
• Prescribe the type of care to be provided or approve the client’s plan of care (POC)
• Periodically review the need for continued care (if the services are not limited by the prescription)

The physician must base the POC on a risk assessment completed by the physician or by licensed, professional clinic staff. The assessment must be based on findings obtained through a health history, laboratory or screening services, and a physical examination.

7.1.2 Case Management Services to High-Risk Individuals

An MSC that wants to bill and receive reimbursement for case management services to high-risk individuals including infants, pregnant adolescents, and women must meet the eligibility criteria for case management services. To be considered for reimbursement for case management for these clients, the MSC must enroll as a group in Case Management for Children and Pregnant Women, and each eligible case manager must enroll as a performing provider.


7.2 Services, Benefits, Limitations, and Prior Authorization

Services billed by an MSC are those provided by a physician or by licensed, professional clinic staff and are determined to be reasonable and medically necessary for the care of a pregnant adolescent or woman during the prenatal period and subsequent 60-day postpartum period. MSC benefits do not include deliveries.

MSCs are limited to 20 prenatal care visits and 1 postpartum care visit per pregnancy. Normal pregnancies are anticipated to require around 11 visits per pregnancy and high-risk pregnancies are anticipated to require around 20 visits per pregnancy. If more than 20 visits are medically necessary, the provider can appeal with documentation supporting pregnancy complications. The high-risk client’s medical record documentation must reflect the need for increased visits and is subject to retrospective review.

Procedure codes in the following table are for prenatal and postpartum care visits:

<table>
<thead>
<tr>
<th>Procedure Codes</th>
</tr>
</thead>
<tbody>
<tr>
<td>59430*</td>
</tr>
<tr>
<td>99202-TH</td>
</tr>
<tr>
<td>99203-TH</td>
</tr>
<tr>
<td>99204-TH</td>
</tr>
<tr>
<td>99205-TH</td>
</tr>
<tr>
<td>99211-TH</td>
</tr>
<tr>
<td>99212-TH</td>
</tr>
</tbody>
</table>

* Procedure code 59430 is not submitted with modifier TH

Note: The prenatal visits must be billed with modifier TH
Providers must bill the most appropriate new or established prenatal visit code or postpartum visit code. New patient codes may be used when the client has not received any professional services from the provider, or another provider of the same specialty who belongs to the same group practice, within the past three years (36 months).

An MSC may be reimbursed for prenatal and postpartum care visits only. Hemoglobin, hematocrit, and urinalysis procedures are included in the charge for prenatal care and not separately reimbursed. Services other than prenatal and postpartum care visits will be denied. MSCs that are enrolled in Case Management for Children and Pregnant Women as a group may be reimbursed for these services under the group provider identifier assigned to their facility.

Medical services must be furnished on an outpatient basis by the physician or by licensed, professional clinic staff under the direction of the physician and must be within the staff’s scope of practice or licensure as defined by state law. Although the physician does not necessarily have to be present at the clinic when services are provided, the physician must assume professional responsibility for the medical services provided at the clinic and ensure through approval of the POC that the services are medically appropriate. The physician must spend as much time in the clinic as is necessary to ensure that clients are receiving medical services in a safe and efficient manner in accordance with accepted standards of medical practice.

MSCs must follow the procedures outlined throughout this manual. All service, frequency, and documentation requirements are applicable.

Providers submitting charges for high-risk prenatal care must document the high-risk diagnosis on the claim form and document the condition in the client’s medical record.

### 7.2.1 Initial Prenatal Care Visit Components

The following initial prenatal care visit components should be completed as early as possible in the client’s pregnancy.

#### 7.2.1.1 History

History includes OB-GYN, present pregnancy, medical and surgical, substance use, environmental, nutritional, psychosocial (including violence), and family support system.

#### 7.2.1.2 Physical Examination

Physical examination includes height, weight, blood pressure; head, neck, lymph, breasts, heart, lungs, back, abdomen, pelvis, rectum, extremities, and skin; and uterine size, fetal heart rate, and location.

#### 7.2.1.3 Laboratory Tests

The initial hematocrit or hemoglobin and each subsequent hematocrit or hemoglobin is included in the visit fee and is not separately reimbursable to MSCs.

The laboratory services listed may not be billed using the MSC provider identifier. These services may be ordered by MSC personnel and provided by a reference laboratory.

MSCs must supply the client’s Medicaid number and the MSC provider identifier to the reference laboratory when laboratory services are requested.

The laboratory services requested by an MSC may include, but are not limited to, the following:

- Hemoglobin, hematocrit, or complete blood count (CBC)
- Urinalysis
- Blood type and Rh
- Antibody screen
- Rubella antibody titer
• Serology for syphilis
• Hepatitis B surface antigen
• Cervical cytology
• Other laboratory tests

The following tests may be performed at the initial prenatal care visit, as indicated:
• Pregnancy test
• Gonorrhea test
• Urine culture
• Sickle cell test
• Tuberculosis (TB) test
• Chlamydia test

As stated in the Health and Safety Code §81.090, screening for Hepatitis B virus infection, HIV, and Syphilis must be performed at the initial prenatal care visit. In addition, HIV testing must be performed in the third trimester. HBV and Syphilis must be performed at labor and delivery.

Multiple marker screens for neural tube defects must be offered if the client initiates care between 16 and 20 weeks.

7.2.1.4 Assessment
Assessment includes pregnancy, general health, medical, and psychosocial.

7.2.1.5 Plan
Plan includes pregnancy, preventive health, medical, and referral as indicated.

7.2.1.6 Education and Counseling
Education and counseling includes pregnancy, delivery, nutrition, breast-feeding, family planning, and preventive health. The education and counseling should also include the need for a medical home and information about THSteps medical and dental checkups for the client.

The complete physical examination may be completed at the second visit if the MSC’s routine involves a two-stage initial evaluation.

7.2.2 Subsequent Prenatal Care Visits
The following is a recommended guide for the frequency of subsequent prenatal visits for a regular pregnancy:
• One visit every 4 weeks for the first 28 weeks of pregnancy.
• One visit every 2 to 3 weeks from 28 to 36 weeks of pregnancy.
• One visit per week from 36 weeks to delivery.

More frequent visits may be medically necessary. Physicians, CNMs, and MSCs are limited to 20 prenatal care visits per pregnancy and 1 postpartum care visit per pregnancy after discharge from the hospital, without documentation of a complication of pregnancy.

Each subsequent visit must include the following:
• Interim History
• Problems
• Maternal status
• Fetal status

7.2.2.1 Physical Examination
The physical examination must include the following:
• Weight and blood pressure
• Fundal height, fetal position and size, and fetal heart rate
• Extremities

7.2.2.2 Laboratory Tests
Required laboratory tests include the following:
• Urinalysis for protein and glucose every visit
  Note: The urinalysis for protein and glucose, hemoglobin, and hematocrit is included in the visit fee and is not separately reimbursable to MSCs.
• Hematocrit or hemoglobin repeated once a trimester and at 32 to 36 weeks of pregnancy
• Multiple marker screen for fetal abnormalities offered at 16 to 20 weeks of pregnancy
• Repeated antibody screen for Rh negative women at 28 weeks (followed by Rho immune globulin administration if indicated)
• Gestational diabetes screen at 24 to 28 weeks of pregnancy, one hour post 50 gram glucose load
• Blood sample for HBsAg screening at the first examination and visit followed by a second blood sample for HBsAg screening on admission for delivery
• Other laboratory tests as indicated by the medical condition of the client

7.2.3 Postpartum Care Visit
Postpartum care provided by MSCs must be billed using procedure code 59430. A maximum of 1 postpartum visit is allowed per pregnancy.

Refer to: Section 4, “Obstetric Services” in the Gynecological, Obstetrics, and Family Planning Title XIX Services Handbook (Vol. 2, Provider Handbooks) for billing requirements.

7.2.4 Prior Authorization
Prior authorization is not required for services rendered in MSCs.

7.3 Documentation Requirements
Each client must have a complete and accepted standard medical record with documentation for the initial visit with procedures, as well as each subsequent visit with procedures. Such records must be made available when requested by HHSC or TMHP for utilization and quality assurance reviews as required by federal regulations. The documentation record or a true copy or narrative abstract must be sent to the hospital of delivery by the client’s 35th week of pregnancy. The record must be made available to the client if the client transfers care to another institution. Records completed by licensed professional clinic staff under the direction of a physician must be signed by the supervising physician.

7.4 Claims Filing and Reimbursement
MSC services must be submitted to TMHP in an approved electronic format or on the CMS-1500 claim form. Providers may purchase CMS-1500 claim forms from the vendor of their choice. TMHP does not supply the forms. When completing a CMS-1500 claim form, all required information must be included on the claim, as TMHP does not key any information from claim attachments. Superbills, or itemized statements, are not accepted as claim supplements.
Refer to: “Section 3: TMHP Electronic Data Interchange (EDI)” (Vol. 1, General Information) for information on electronic claims submissions.

“Section 6: Claims Filing” (Vol. 1, General Information) for general information about claims filing.

Subsection 6.5, “CMS-1500 Paper Claim Filing Instructions” in “Section 6: Claims Filing” (Vol. 1, General Information). Blocks that are not referenced are not required for processing by TMHP and may be left blank.

MSCs are reimbursed in accordance with 1 TAC §355.8085. Providers can refer to the OFL or the applicable fee schedule on the TMHP website at www.tmhp.com.

Texas Medicaid implemented mandated rate reductions for certain services. The OFL and static fee schedules include a column titled “Adjusted Fee” to display the individual fees with all mandated percentage reductions applied. Additional information about rate changes is available on the TMHP website at www.tmhp.com/resources/rate-and-code-updates.

Note: Certain rate reductions including, but not limited to, reductions by place of service, client type program, or provider specialty may not be reflected in the Adjusted Fee column.

8 Nurse Practitioner (NP) and Clinical Nurse Specialist (CNS)

For other APRNs, see Section 4, “Certified Registered Nurse Anesthetist (CRNA)” in this handbook for information regarding CRNAs, and Section 3, “Certified Nurse Midwife (CNM)” in this handbook for information about certified nurse midwives (CNMs).

8.1 Enrollment

To enroll in Texas Medicaid, an NP or CNS must be licensed as a registered nurse and as an APRN by the Texas BON. A registered nurse under the multistate licensure compact may be licensed in another state but certified as an APRN for the state of Texas by the Texas BON. Texas Medicaid accepts a signed letter of certification from the Texas BON as documentation of appropriate licensure and certification for enrollment.

Providers cannot be enrolled if their license is due to expire within 30 days.

All providers of laboratory services must comply with the rules and regulations of the Clinical Laboratory Improvement Amendments (CLIA). Providers not complying with CLIA are not reimbursed for laboratory services.

All APRNs (including CNMs, CRNAs, CNSs, and NPs) are enrolled within the categories of practice as determined by the Texas BON. CNSs and NPs must enroll as an APRN; CNMs and CRNAs may enroll using their specific titles.

Refer to: Subsection 2.1.1, “Clinical Laboratory Improvement Amendments (CLIA)” in the Radiology and Laboratory Services Handbook (Vol. 2, Provider Handbooks).

Section 3, “Certified Nurse Midwife (CNM)” in this handbook for more information on CNM enrollment.

Section 4, “Certified Registered Nurse Anesthetist (CRNA)” in this handbook for more information on CRNA enrollment.

APRNs may be included as primary care providers in the provider network for Medicaid and CHIP programs (both fee-for-service and managed care), regardless of whether the physician supervising the APRN is enrolled in Medicaid or in the provider network.
8.1.1 Enrollment in Texas Health Steps (THSteps)

APRNs, including NPs, and CNSs, who are recognized by the Texas BON can enroll as THSteps providers and provide checkup services within their scope of practice. Specific information is found in the Children’s Services Handbook.

Refer to: subsection 5.2, “Enrollment” in the Children’s Services Handbook (Vol. 2, Provider Handbooks) for more information on enrollment procedures.

8.2 Services, Benefits, Limitations, and Prior Authorization

Services performed by NPs and CNSs are benefits if the services meet the following criteria:

- Are within the scope of practice for NPs and CNSs, as defined by Texas state law.
- Are consistent with rules and regulations promulgated by the Texas BON or other appropriate state licensing authority.
- Are covered by Texas Medicaid when provided by a licensed physician (MD or DO).
- Are reasonable and medically necessary as determined by HHSC or its designee.

NPs and CNSs who are employed or remunerated by a physician, hospital, facility, or other provider must not bill Texas Medicaid for their services if the billing results in duplicate payment for the same services.

Physicians who submit a claim using the physician’s own provider identifier for services provided by an NP or CNS must submit modifier SA on each claim detail if the physician does not make a decision regarding the client’s care or treatment on the same date of service as the billable medical visit.

Benefit limitation information for services can be found in Section 9, “Physician” in this handbook, the Children’s Services Handbook (Vol. 2, Provider Handbooks), and the Gynecological, Obstetrics, and Family Planning Title XIX Services Handbook (Vol. 2, Provider Handbooks).

Payment for supplies is not a benefit of Texas Medicaid. Costs of supplies are included in the reimbursement for office visits.

Refer to: Section 2, “Medicaid Title XIX Family Planning Services” in the Gynecological, Obstetrics, and Family Planning Title XIX Services Handbook (Vol. 2, Provider Handbooks).

Section 9, “Physician” in this handbook.

Section 5, “THSteps Medical” in the Children’s Services Handbook (Vol. 2, Provider Handbooks) for more information on THSteps services.

8.2.1 Prior Authorization

Services performed by an NP or CNS are subject to the same prior authorization guidelines as services performed by other provider types.

8.3 Documentation Requirements

All services require documentation to support the medical necessity of the service rendered, including NP and CNS services. NP and CNS services are subject to retrospective review and recoupment if documentation does not support the service billed.
8.4 Claims Filing and Reimbursement

8.4.1 Claims Information

APRN services must be submitted to TMHP in an approved electronic format or on the CMS-1500 claim form. Providers may purchase CMS-1500 claim forms from the vendor of their choice. TMHP does not supply the forms.

When completing a CMS-1500 claim form, all required information must be included on the claim, as TMHP does not key any information from claim attachments. Superbills, or itemized statements, are not accepted as claim supplements.

Refer to: “Section 3: TMHP Electronic Data Interchange (EDI)” (Vol. 1, General Information) for information on electronic claims submissions.

“Section 6: Claims Filing” (Vol. 1, General Information) for general information about claims filing.

Subsection 6.5, “CMS-1500 Paper Claim Filing Instructions” in “Section 6: Claims Filing” (Vol. 1, General Information) for instructions on completing paper claims. Blocks that are not referenced are not required for processing by TMHP and may be left blank.

8.4.2 Reimbursement

According to 1 TAC §355.8281, the Medicaid rate for NPs and CNSs is 92 percent of the rate paid to a physician (MD or DO) for the same professional service and 100 percent of the rate paid to physicians for laboratory services, X-ray services, and injections. When NPs or CNSs bill Medicaid directly for services they performed, they must use their individual provider identifier. If the services are performed by the NP or CNS but billed by a physician or physician group, the billing provider is the physician or physician group. Physicians may be reimbursed 92 percent of the established reimbursement rate for services provided by an NP or CNS but not make a decision regarding the client’s care or treatment on the same date of service as the billable medical visit. This 92 percent reimbursement rate does not apply to laboratory services, X-ray services, and injections provided by an NP or CNS.

Note: NP and CNS providers who are enrolled in Texas Medicaid as THSteps providers also receive 92 percent of the rate paid to a physician for THSteps services when a claim is submitted with their THSteps provider identifier as the billing provider.

Providers can refer to the OFL or the applicable fee schedule on the TMHP website at www.tmhp.com.

Texas Medicaid implemented mandated rate reductions for certain services. The OFL and static fee schedules include a column titled “Adjusted Fee” to display the individual fees with all mandated percentage reductions applied. Additional information about rate changes is available on the TMHP website at www.tmhp.com/resources/rate-and-code-updates.

Refer to: Subsection 1.1, “Provider Enrollment” in “Section 1: Provider Enrollment and Responsibilities” (Vol. 1, General Information).

Subsection 2.2, “Fee-for-Service Reimbursement Methodology” in “Section 2: Texas Medicaid Fee-for-Service Reimbursement” (Vol. 1, General Information) for more information about reimbursement.
9 Physician

9.1 Enrollment

9.1.1 Physicians and Doctors
To enroll in Texas Medicaid to provide medical services, physicians (MD or DO), doctors of dental surgery [DDS], and doctors of podiatric medicine (DPM) must be authorized by the licensing authority of their profession to practice in the state where the services are performed at the time they are provided.

Providers cannot be enrolled in Texas Medicaid if their licenses are due to expire within 30 days. A current Texas license must be submitted.

Important: The Centers for Medicare & Medicaid Services (CMS) guidelines mandate that physicians who provide durable medical equipment (DME) products such as spacers or nebulizers are required to enroll as Texas Medicaid DME providers.

All physicians except gynecologists, pediatricians, pediatric subspecialists, pediatric psychiatrists, and providers performing only Texas Health Steps (THSteps) medical or dental checkups must be enrolled in Medicare before enrolling in Medicaid. TMHP may waive the Medicare enrollment prerequisite for pediatricians or physicians whose type of practice and service may never be billed to Medicare.

9.2 Services, Benefits, Limitations, and Prior Authorization

The Administrative Simplification provisions of the Health Insurance Portability and Accountability Act (HIPAA) of 1996 mandates the use of national coding and transaction standards. HIPAA requires that the American Medical Association’s (AMA) Current Procedural Terminology (CPT) system be used to report professional services, including physician services. Correct use of CPT coding requires using the most specific code that matches the services provided, based on the code’s description. Providers must pay special attention to the standard CPT descriptions for the evaluation and management (E/M) services. The medical record must document the specific elements necessary to satisfy the criteria for the level of services as described in CPT. Reimbursement may be recouped when the medical record documents a different level of service from what is submitted on the claim. The level of service provided and documented must be medically necessary, based on the clinical situation and needs of the client.

To receive reimbursement, providers must document the following information in the client’s medical record:

- The service
- The date rendered
- Pertinent information about the client’s condition supporting the need for the service
- The care given

Physician services include those reasonable and medically necessary services ordered and performed by physicians or under physician supervision that are within the scope of practice of their profession as defined by state law.

9.2.1 Electronic Signatures in Prior Authorizations

Prior authorization requests may be submitted to the TMHP Prior Authorization Department via mail, fax, or the electronic portal. Prescribing or ordering providers, dispensing providers, clients’ responsible adults, and clients may sign prior authorization forms and supporting documentation using electronic or wet signatures.

Refer to: Subsection 5.5.1.2, “Document Requirements and Retention” in “Section 5: Fee-for-Service Prior Authorizations” (Vol. 1, General Information) for additional information about electronic signatures.
9.2.2 Teaching Physician and Resident Physician

The roles of the teaching physician and resident physician occur in the context of an accredited graduate medical education (GME) training program.

The teaching physician is the Medicaid-enrolled physician who is professionally responsible for the particular services that were provided and are being submitted for reimbursement; the physician must be affiliated and in good standing with an accredited GME program and must possess all appropriate licensure.

Physician services must be performed personally by the teaching physician or by the person to whom the physician has delegated the responsibility. The level of supervision required may be direct or personal.

In all cases, the client's medical record must clearly document that the teaching physician provided identifiable supervision of the resident. As defined below, the supervision must be direct or personal depending on the setting and the clinical circumstances:

- **Direct supervision** means that the teaching physician must be in the same office, building, or facility when and where the service is provided and must be immediately available to furnish assistance and direction.

- **Personal supervision** means that the teaching physician must be physically present in the room when and where the service is being provided.

Personal supervision by the teaching physician is required during the key portions of all major surgeries and the key portions of all other physician services billed to Texas Medicaid if the immediate supervision, participation, or intervention of the supervising physician is medically prudent in order to assure the health and safety of the client. Physician services that require personal supervision may include invasive procedures and evaluation and management services that require complex medical decision making. Situations that require personal supervision include those in which:

- The clinical condition of the client is unstable or will likely become unstable during, or as a result of, the planned medical intervention.

- The planned medical intervention, even under optimal conditions will result in a medically reasonable risk for significant morbidity or death following the procedure.

- Deviation from the expected technique at the time the procedure or service is performed presents a medically reasonable, causally-related, foreseeable risk to the patient's life or health.

This criterion applies regardless of the place of service.

The teaching physician must provide medically appropriate, identifiable direct supervision for all other services that do not require personal supervision.

The following prerequisites apply when the teaching physician submits claims for services performed, in whole or in part, by the resident physician in the inpatient hospital setting, the outpatient hospital setting, and surgical services and procedures.

**Note:** When requesting services for prior authorization at patient discharge, the signature of the resident on the actual prescription is permitted as long as the Medicaid enrolled attending/supervising physician's signature appears on the Home Health Services (Title XIX) DME/Medical Supplies Physician Order Form and on any letters or documentation provided to support medical necessity. The resident's order and the Title XIX Form signed by the attending/supervising physician must be for the same service.
9.2.2.1 Teaching Physician Prerequisites

Services provided in an outpatient setting.

All requirements for personal or direct supervision in the outpatient setting must be met for the services to qualify for reimbursement. The following tasks must be performed by the teaching physician and their completion must be documented in the patient’s medical record before the claims are submitted for consideration of reimbursement:

- Review the patient’s history and physical examination.
- Confirm or revise the patient’s diagnosis.
- Determine the course of treatment to be followed.
- Assure that any necessary supervision of interns or residents was provided.
- Confirm that documentation in the medical record supports the level of service provided.

**Exception:** Exception for E/M services furnished in certain primary care centers. Teaching physicians that meet the primary care exception under Medicare are allowed to bill for low-level and mid-level E/M services furnished by residents in the absence of a teaching physician. Facilities that meet the primary care exception under Medicare may bill Texas Medicaid, Family Planning, or the Children with Special Health Care Needs (CSHCN) Services Program for new patient services (procedure codes 99202 and 99203) and established patient services (procedure codes 99211, 99212, and 99213).

**Note:** All services provided in an outpatient setting that do not qualify for the exception above require that the teaching physician examine the patient.

Services provided in an inpatient setting.

For services provided in an inpatient setting, the teaching physician must demonstrate that medically appropriate supervision was provided. The following tasks must be performed and their completion must be documented in the patient’s medical record before the claims are submitted for consideration of reimbursement. The documentation must be made in the same manner as required by federal regulations under Medicare:

- Review the patient’s history, review the resident’s physical examination, and examine the patient no later than 36 hours after the patient’s admission and before the patient’s discharge.
- Confirm or revise the patient’s diagnosis.
- Determine the course of treatment to be followed.
- Document the teaching physician’s presence and participation in the major surgical or other complex and dangerous procedure or situation.
- Confirm that documentation in the medical record supports the level of service provided.
- A face-to-face encounter with the client on the same day as any services provided by the resident physician.

Surgical services and procedures.

The teaching surgeon is responsible for the patient’s preoperative, operative, and postoperative care. The teaching physician must demonstrate that medically appropriate supervision was provided. The following tasks must be performed and their completion must be documented in the patient’s medical record before the claims are submitted for consideration of reimbursement. The documentation must be made in the same manner as required by federal regulations under Medicare:

- Review the patient’s history, review the resident’s physical examination, and examine the patient within a reasonable period of time after the patient’s admission and before the patient’s discharge.
• Confirm or revise the client’s diagnosis.
• Determine the course of treatment to be followed.
• Document the teaching physician’s presence and participation in the major surgical or other complex and dangerous procedure or situation.

**Important:** Reimbursement may be reduced, denied, or recouped if the prerequisites are not documented in the medical record. The documentation must be made in the same manner as required by federal regulations under Medicare.

### 9.2.3 Substitute Physician

Physicians may bill for the service of a substitute physician who sees clients in the billing physician’s practice under either a reciprocal or locum tenens arrangement.

A reciprocal arrangement is one in which a substitute physician covers for the billing physician on an occasional basis when the billing physician is unavailable to provide services. Reciprocal arrangements are limited to a continuous period no longer than 14 days and do not have to be in writing.

A locum tenens arrangement is one in which a substitute physician assumes the practice of a billing physician for a temporary period no longer than 90 days when the billing physician is absent for reasons such as illness, pregnancy, vacation, continuing medical education, or active duty in the armed forces. The locum tenens arrangement may be extended for a continuous period of longer than 90 days if the billing physician’s absence is due to being called or ordered to active duty as a member of a reserve component of the armed forces. Locum tenens arrangements must be in writing.

The substitute physician must be enrolled in Texas Medicaid and must not be on the Texas Medicaid or HHSC Family Planning Program provider exclusion list. The billing provider’s name, address, and national provider identifier must appear in Block 33 of the claim form. The name and office or mailing address of the substitute physician must be documented on the claim in Block 19, not Block 33.

When a physician bills for a substitute physician, modifier Q5 or Q6 must follow the procedure code in Block 24D for services provided by the substitute physician. The Q5 modifier is used to indicate a reciprocal arrangement and the Q6 modifier is used to indicate a locum tenens arrangement.

When physicians in a group practice bill substitute physician services, the performing provider identifier of the physician for whom the substitute provided services must be in Block 24J.

Physicians must familiarize themselves with these requirements and document accordingly. Those services not supported by the required documentation as detailed above will be subject to recoupment.

### 9.2.4 Aerosol Treatment

Nebulized aerosol treatments (procedure codes 94640, 94644, and 94645) with short-acting beta-agonists are a benefit of Texas Medicaid and considered medically necessary when breathing is compromised by certain acute medical conditions. Documentation to support an aerosol treatment for the worsening of an acute or chronic condition must be maintained in the client’s medical record and is subject to retrospective review.

Procedure code 94645 is only a benefit in the outpatient setting, specifically in a hospital emergency department or an urgent care clinic.

Pulse oximetry and evaluation of the client’s use of an aerosol generator, nebulizer, or metered-dose inhaler are considered part of an evaluation and management (E/M) visit and will not be reimbursed separately.

Hypertonic saline used in aerosol therapy will be denied if billed separately.

**Refer to:** Subsection 4.2.20.1, “Aerosol Treatment” in the *Inpatient and Outpatient Hospital Services Handbook* (Vol. 2, Provider Handbooks).
9.2.4.1 Diagnostic Testing
Nitric oxide expired gas determination (FeNO) measurement (procedure code 95012) is a benefit for Texas Medicaid.

FeNO measurement provided in the physician’s office is considered medically necessary as an adjunct to the established clinical and laboratory assessments for diagnosing and assessing asthma, predicting exacerbations, and evaluating the response of a client who has asthma to anti-inflammatory therapy. FeNO measurement may be reimbursed by Texas Medicaid when the test is used as follows:

- To assist in assessing the etiology of respiratory symptoms.
- To help identify the eosinophilic asthma phenotype.
- To assess potential response or failure to respond to anti-inflammatory agents, particularly inhaled corticosteroids (ICS).
- To establish a baseline FeNO during non-exacerbations for subsequent monitoring of chronic persistent asthma.
- To guide changes in dosing of anti-inflammatory medications, i.e., step-down dosing, step-up dosing, or discontinuation of anti-inflammatory medications.
- To assist in the evaluation of adherence to anti-inflammatory medications.
- To assess whether airway inflammation is contributing to respiratory symptoms.

The technical and interpretation components of procedure code 95012 will not be reimbursed separately, as the instrument produces an exhaled nitric oxide (NO) measurement that requires little interpretation. Procedure code 95012 will be limited to once per day and must be submitted with procedure code 94010 or 94060.

If FeNO is measured during an office visit where additional E/M components are fulfilled, a separate E/M procedure code may be reimbursed if it is submitted with modifier 25.

9.2.5 Allergy Services
Texas Medicaid uses the following guidelines for reimbursement of allergy services.

9.2.5.1 Allergy Immunotherapy
Allergen immunotherapy consists of the parenteral administration of allergenic extracts as antigens at periodic intervals, usually on an increasing dosage scale to a dosage which is maintained as maintenance therapy.

Preparation of the allergy vial or extracts is a benefit of Texas Medicaid when preparations are made in accordance with the American Academy of Allergy, Asthma, and Immunology. Claims for preparations should be submitted using the following procedure codes:

<table>
<thead>
<tr>
<th>Procedure Codes for Preparation of Allergy Vial or Extract</th>
</tr>
</thead>
<tbody>
<tr>
<td>95145 95146 95147 95148 95149 95165 95170</td>
</tr>
</tbody>
</table>

Administration of the allergy extract may be reimbursed using procedure codes 95115 and 95117.

Rapid desensitization may be reimbursed using procedure code 95180 when submitted with diagnosis code ZS16.

Allergen immunotherapy is a benefit for clients who have allergy conditions when the following criteria are met:

- A diagnosed hypersensitivity to an allergen can be indicated by one of the valid diagnosis codes listed below.
• Hypersensitivity cannot be managed by avoidance or pharmacologic therapy to control allergic symptoms, or the client has unacceptable side effects with pharmacologic therapy.
• The pharmacologic treatment is refused by the client or leads to significant side effects.
• The allergen content is based on appropriate skin testing, and the allergens are prepared for the client individually.

The preparation of the allergy vial or extract and the administration of an injection may be reimbursed for the following diagnosis codes:

<table>
<thead>
<tr>
<th>Diagnosis Codes</th>
</tr>
</thead>
<tbody>
<tr>
<td>H1045</td>
</tr>
<tr>
<td>H65112</td>
</tr>
<tr>
<td>H65194</td>
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<tr>
<td>H65493</td>
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<tr>
<td>J449</td>
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<tr>
<td>J4541</td>
</tr>
<tr>
<td>J45998</td>
</tr>
<tr>
<td>T531X4S</td>
</tr>
<tr>
<td>T534X4D</td>
</tr>
<tr>
<td>T63001A</td>
</tr>
<tr>
<td>T63003S</td>
</tr>
<tr>
<td>T63012D</td>
</tr>
<tr>
<td>T63021A</td>
</tr>
<tr>
<td>T63023S</td>
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<tr>
<td>T63032D</td>
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<tr>
<td>T63041A</td>
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<tr>
<td>T63043S</td>
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<tr>
<td>T63062D</td>
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<tr>
<td>T63071A</td>
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<tr>
<td>T63073S</td>
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<tr>
<td>T63082D</td>
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<tr>
<td>T63092A</td>
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<tr>
<td>T63094S</td>
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<tr>
<td>T63113D</td>
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<tr>
<td>T63122A</td>
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<tr>
<td>T63124S</td>
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<tr>
<td>T63193D</td>
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<tr>
<td>T632X2A</td>
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<tr>
<td>T632X4S</td>
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<tr>
<td>T63303D</td>
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<tr>
<td>T63312A</td>
</tr>
<tr>
<td>T63314S</td>
</tr>
<tr>
<td>T63323D</td>
</tr>
</tbody>
</table>
9.2.5.1.1 Prior Authorization for Allergy Immunotherapy

Authorization is not required for immunotherapy services; however, requests for services beyond the established limits of 160 doses per one-year period for procedure code 95165 may be considered for prior authorization with documentation of medical necessity. Documentation must be submitted to the Special Medical Prior Authorization Department and include the following information:

- Copy of the allergen testing results
- Severity and periodicity of symptoms
• Physical limitations created by the symptoms
• Concurrent drug treatment
•Explanation of how efficacy has not been achieved with prior treatment and the objectives of the new anticipated treatment program

9.2.5.1.2 Limitations of Allergy Immunotherapy

The quantity billed for the allergy extract preparation procedure must represent the total number of doses to be administered from the vial. If the number of doses is not stated on the claim, a quantity of one is allowed.

**Note:** A “dose” is defined as the amount of antigen(s) administered in a single injection from a multidose vial.

Procedure code 95165 is limited to a total of 160 doses per one-year period, which begins the date the immunotherapy is initiated. Additional doses may be considered for reimbursement through prior authorization with documentation of medical necessity. Procedure code 95165 is limited to no more than ten doses per vial.

When an injection is given from a vial, providers should use an administration-only procedure code (95115 or 95117). Reimbursement for the administration is limited to one per day.

An office visit, clinic visit, or observation room visit is not considered for reimbursement in addition to the fee for the preparation or the administration of the allergy vial or extract unless the additional visit results in a non-allergy-related diagnosis or a re-evaluation of the client’s condition. The following E/M procedure codes may be submitted with modifier 25:

<table>
<thead>
<tr>
<th>Procedure Codes</th>
</tr>
</thead>
<tbody>
<tr>
<td>99202</td>
</tr>
</tbody>
</table>

Allergen immunotherapy that is considered experimental, investigational, or unproven is not a benefit of Texas Medicaid.

Single dose vials (procedure code 95144) are not a benefit of Texas Medicaid.

9.2.5.2 Allergy Testing

Texas Medicaid benefits include allergy testing for clients with clinically significant allergic symptoms. Allergy testing is focused on determining the allergens that cause a particular reaction and the degree of the reaction. Allergy testing also provides justification for recommendations of particular medicines, of immunotherapy, or of specific avoidance measures in the environment.

Evaluation and management E/M services will not be reimbursed on the same date of service as allergy testing. Allergy testing will be paid and the E/M service will be denied as part of another procedure on the same date of service.

The following allergy tests are benefits of Texas Medicaid:

• Percutaneous and intracutaneous skin test. The skin test for IgE-mediated disease with allergenic extracts is used in the assessment of allergy-prone clients. The test involves the introduction of small quantities of test allergens below the epidermis. Procedure codes 95004, 95017, 95018, 95024, 95027, and/or 95028 should be used to submit skin tests for consideration of reimbursement.

• **Patch or application tests.** Patch testing (procedure code 95044) is used for diagnosing contact allergic dermatitis.

• **Photo or photo patch skin test.** Procedure codes 95052 and 95056 may be used for diagnosing contact allergic dermatitis.
• **Ophthalmic mucous membrane or direct nasal mucous membrane tests.** Nasal or ophthalmic mucous membrane tests (procedure codes 95060 and 95065) are used for the diagnosis of either food or inhalant allergies and involve the direct administration of the allergen to the mucosa.

• **Inhalation bronchial challenge testing (not including necessary pulmonary function tests).** Bronchial challenge testing with methacholine, histamine, or allergens (procedure code 95070) is used for defining asthma or airway hyperactivity when skin testing results are not consistent with the client’s medical history. Results of these tests are evaluated by objective measures of pulmonary function.

Procedure code 95199 may be used for an unlisted allergy or clinical immunologic service or procedure if there is not a specific procedure code that describes the service performed. Prior authorization is required for unlisted procedure codes. Every effort must be used to bill with the appropriate CPT code that describes the procedure being performed. If a code does not exist to describe the service performed, prior authorization may be requested using unlisted procedure code 95199 and must be submitted with documentation to assist in determining coverage. The documentation submitted must include all of the following:

- The client’s diagnosis
- Medical records indicating prior treatment for this diagnosis and the medical necessity of the requested procedure
- A clear, concise description of the procedure to be performed
- Reason for recommending this particular procedure
- A CPT or HCPCS procedure code that is comparable to the procedure being requested
- Documentation that this procedure is not investigational or experimental
- Place of service (POS) the procedure is to be performed
- The physician’s intended fee for this procedure

Prior authorization requests for Texas Medicaid fee-for-service clients must be submitted by the physician to the Special Medical Prior Authorization (SMPA) department.

The number of allergy tests performed must be indicated on the claim. When the number of tests is not specified, a quantity of one is allowed.

**9.2.5.2.1 Allergy Blood Tests**

Allergy blood testing procedure codes 86001, 86003, 86005, and 86008 are a benefit when the test is performed for a reason that includes, but is not limited to, the following:

- The client is unable to discontinue medications
- An allergy skin test is inappropriate for the client for the following reasons:
  - The client is pediatric
  - The client is disabled
  - The client suffers from a skin condition such as dermatitis

Radioallergosorbent tests (RAST) and multiple antigen simultaneous tests (MAST) are benefits of Texas Medicaid. RAST testing is used to detect specific allergens. RAST testing is usually performed by an independent lab; however, there are physicians who have the capability of performing these tests in their offices. Physicians who submit RAST/MAST tests performed in the office setting must use modifier SU to be considered for reimbursement. Without the use of the SU modifier, RAST/MAST testing submitted with POS 1 (office) is denied with the message, “Lab performed outside of office must be billed by the performing facility.”

RAST/MAST tests must be submitted using procedure codes 86003, 86005, and 86008.
Procedure code 86001 is limited to 20 allergens per rolling year, any provider.
Procedure code 86003 and 86008 are limited to 30 allergens per rolling year, any provider.
Procedure code 86005 is limited to 4 multiallergen tests per rolling year, same provider.

9.2.5.2.2 Collagen Skin Test
Collagen skin tests are a benefit of Texas Medicaid using procedure code Q3031. Collagen skin tests are administered to detect a hypersensitivity to bovine collagen. This skin test is given four weeks prior to any type of surgical procedure that utilizes collagen.

Collagen injections that are used for cosmetic surgery are not considered medically necessary and are not a benefit of Texas Medicaid.

9.2.5.2.3 Prior Authorization
Prior authorization is required for collagen skin test procedure code Q3031.

Prior authorization requests for Texas Medicaid fee-for-service clients must be submitted by the physician to the Special Medical Prior Authorization (SMPA) department.

Prior authorization is not required for other allergy testing procedure codes unless the limits are exceeded. The following medical documentation must be submitted to the SMPA Department with the prior authorization request for additional procedures:

- Results of any previous treatment
- Documentation that explains why the client’s treatment could not be completed within the policy limits for the requested procedures
- Client diagnosis and conditions that support the medical necessity for the additional procedures requested
- Client outcomes that the requested procedures will achieve

9.2.5.2.4 Ingestion Challenge Test
Ingestion challenge tests are a benefit of Texas Medicaid using procedure code 95076. Ingestion challenge tests are used to confirm an allergy to a food or food additive.

Procedure code 95076 is limited to one service per day, any provider.

9.2.6 Ambulance Transport Services - Nonemergency
Nonemergency ambulance services require prior authorization in circumstances not involving an emergency. Facilities and other providers must request and obtain prior authorization before contacting the ambulance provider for nonemergency ambulance services.


Subsection 5.1.8, “Prior Authorization for Nonemergency Ambulance Transport” in “Section 5: Fee-for-Service Prior Authorizations” (Vol. 1, General Information) for more information about nonemergency ambulance transport prior authorization.

9.2.7 Anesthesia
Anesthesia services are a benefit of Texas Medicaid with specific benefits and limitations to reimbursement.
Medicaid may reimburse anesthesiologists, certified registered nurse anesthetists (CRNAs), and anesthesiologist assistants (AAs) for administering anesthesia as defined within their individual scope of practice.

9.2.7.1 Medical Direction by an Anesthesiologist

Medical direction by an anesthesiologist of an anesthesia practitioner (CRNA, AA, or other qualified professional) is a benefit of Texas Medicaid if the following criteria are met:

• No more than four anesthesia procedures are being performed concurrently.
• The anesthesiologist is physically present in the operating suite.

Exception: Anesthesiologists may be considered for reimbursement when they medically direct more than four anesthesia services or simultaneously supervise a combination of more than four CRNAs, AAs, or other qualified professionals under emergency circumstances only.

Medical direction provided by an anesthesiologist is a benefit of Texas Medicaid if the following criteria are met:

• The anesthesiologist performs a preanesthetic examination and evaluation.
• The anesthesiologist prescribes the anesthesia plan.
• The anesthesiologist personally participates in the critical portions of the anesthesia plan, including induction and emergence.
• The anesthesiologist ensures that a qualified professional can perform the procedures in the anesthesia plan that the anesthesiologist does not perform personally.
• The anesthesiologist monitors the course of anesthesia administration at intervals.
• The anesthesiologist provides direct supervision when medically directing an anesthesia procedure. Direct supervision means the anesthesiologist must be immediately available to furnish assistance and direction.
• The anesthesiologist provides postanesthesia care.

The anesthesiologist does not perform any other services (except as noted below) during the same time period. The anesthesiologist who directs the administration of no more than four anesthesia procedures may provide the following without affecting the eligibility of the medical direction services:

• Address an emergency of short duration in the immediate area
• Administer an epidural or caudal anesthetic to ease labor pain
• Provide periodic, rather than continuous, monitoring of an obstetrical patient
• Receive clients entering the operating suite for the next surgery
• Check or discharge clients in the recovery room
• Handle scheduling matters

As noted above, an anesthesiologist may concurrently medically direct up to four anesthesia procedures. Concurrency is defined as the maximum number of procedures that the anesthesiologist is medically directing within the context of a single procedure and whether those other procedures overlap each other. Concurrency is not dependent on each of the cases involving a Medicaid client. For example, if three procedures are medically directed but only two involve Medicaid clients, the Medicaid claims must be billed as concurrent medical direction of three procedures.

For medical direction, the anesthesiologist must document in the client’s medical record that he or she did the following:

• Performed the pre-anesthetic exam and evaluation.
• Provided indicated post-anesthesia care.
• Was present during the critical and key portions of the anesthesia procedure, including, if applicable, induction and emergence.
• Was present during the anesthesia procedure to monitor the client’s status.

The following information must be available to state agencies upon request and is subject to retrospective review:
• The name of each CRNA, AA, or other qualified professional that was concurrently medically directed or supervised and a description of the procedure that was performed must be documented and maintained.
• Signatures of the anesthesiologist, CRNA, AA, or other qualified professional involved in administering anesthesia services must be documented in the client’s medical record.

### 9.2.7.2 Anesthesia for Sterilization

**Refer to:** Subsection 2.2, “Services, Benefits, Limitations, and Prior Authorization” in the *Gynecological, Obstetrics, and Family Planning Title XIX Services Handbook* (Vol. 2, Provider Handbooks) for the complete list of family planning diagnosis codes.


Section 4, “Federally Qualified Health Center (FQHC)” in the *Clinics and Other Outpatient Facility Services Handbook* (Vol. 2, Provider Handbooks) for more information about FQHCs and billing the annual family planning examination for Title XIX clients.

### 9.2.7.3 Anesthesia for Labor and Delivery

Providers must bill the most appropriate procedure code for the service provided. Other time-based procedure codes cannot be submitted if either 01960 or 01967 is the most appropriate procedure code.

The following procedure codes must be used for obstetrical anesthesia:

<table>
<thead>
<tr>
<th>Procedure Codes</th>
</tr>
</thead>
<tbody>
<tr>
<td>01960</td>
</tr>
</tbody>
</table>

Procedure codes 01960 and 01967 are limited to once every 210 days when billed by any provider and are reimbursed a flat fee. The time reported must be in minutes. Providers should refer to the definition of time in the CPT manual in the “Anesthesia Guidelines—Time Reporting” section.

Procedure code 01968 or 01969 may be considered for reimbursement when submitted with procedure code 01967. For a Cesarean delivery following a planned vaginal delivery, the anesthesia administered during labor must be billed with procedure code 01967 and must indicate the time in minutes that represents the time between the start and stop times for the procedure. The additional anesthesia services administered during the operative session for a Cesarean delivery must be submitted using procedure code 01968 or 01969 and must indicate the time spent administering the epidural and the actual face-to-face time spent with the client. The insertion and injection of the epidural are not considered separately for reimbursement.

All time must be documented in block 24D of the claim form or the appropriate field of the chosen electronic format.

For continuous epidural analgesia procedure codes (other than procedure codes 01960 and 01967), Texas Medicaid reimburses providers for the time when the physician is physically present and monitors the continuous epidural. Reimbursable time refers to the period between the catheter insertion and when the delivery commences.
Texas Medicaid reimburses the epidural anesthesia services and the delivery at full allowance when they are provided by the delivering obstetrician.

9.2.7.4 Anesthesia Provided by the Surgeon (Other Than Labor and Delivery)

Local, regional, or general anesthesia provided by the operating surgeon is not reimbursed separately from the surgery. A surgeon billing for a surgery will not be reimbursed for the anesthesia when billing for the surgery, even when using the CPT modifier 47. The anesthesia service is included in the global surgical fee.

9.2.7.5 Complicated Anesthesia

The following procedure codes may be reimbursed in addition to an anesthesia procedure or service: 99100, 99116, 99135, and 99140. Documentation supporting the medical necessity for use of the procedure codes may be subject to retrospective review.

Procedure code 99140 is not reimbursed for diagnosis code O80 or O82 when one of these diagnoses is documented as the referenced diagnosis on the claim. The referenced diagnosis must indicate the complicating condition. An emergency is defined as existing when delay in treatment of the client would lead to a significant increase in the threat to life or body part.

9.2.7.6 Multiple Procedures

When billing for anesthesia and other services on the same claim, the anesthesia charge must appear in the first detail line for correct reimbursement. Any other services billed on the same day must be billed as subsequent line items.

When billing for multiple anesthesia services performed on the same day or during the same operative session, use the procedure code with the higher RVU. For accurate reimbursement, apply the total minutes and dollars for all anesthesia services rendered on the higher RVU code. Multiple services reimbursement guidelines apply.

9.2.7.7 Monitored Anesthesia Care

Monitored anesthesia care may include any of the following:

- Intraoperative monitoring by an anesthesiologist or qualified professional under the medical direction of an anesthesiologist
- Monitoring of the client’s vital physiological signs in anticipation of the need for general anesthesia
- Monitoring of the client’s development of an adverse physiological reaction to a surgical procedure

Anesthesiologists, CRNAs, AAs, or other qualified professionals may use modifier QS to report monitored anesthesia care.

The QS modifier is an informational modifier.

9.2.7.8 Reimbursement Methodology

There are two types of reimbursement for anesthesia procedure codes.

- Flat fee
- Time-based fees, which require documentation of the exact amount of face-to-face time with the client

Anesthesiologists directing one or multiple CRNAs and/or AAs during medical procedures will be reimbursed at 50 percent of the established reimbursement rate.

An AA under the supervision of an anesthesiologist is reimbursed the lesser of the billed charges or 50 percent of the calculated payment for a supervised anesthesia service.
If multiple CRNAs, anesthesiologists, or anesthesiologist assistants under anesthesiologist supervision are providing anesthesia services for a client, only one CRNA or AA and one anesthesiologist may be reimbursed.

Both the flat-fee and time-based-fee procedure codes must be submitted with modifiers and are subject to medical direction/supervision reimbursement adjustments.

**Flat Fees**

Both OB related anesthesia procedure codes 01960 and 01967 are considered for reimbursement with a flat-fee rate.

- Flat fees are subject to medically-directed modifier combination adjustments based on the modifier submitted with the anesthesia procedure code.

- The time-based add-on procedure code 01968 must be billed in addition to the flat fee when anesthesia for Cesarean delivery following neuraxial labor analgesia/anesthesia has occurred.

For flat-fee anesthesiology codes, anesthesia time begins when the anesthesia practitioner begins to prepare the client for the induction of anesthesia in the operating room or the equivalent area and ends when the anesthesia practitioner is no longer in personal attendance, that is, when the client may be safely placed under postoperative supervision.

**Time-Based Fees**

For time-based anesthesiology procedure codes, anesthesia time is the time during which an anesthesia practitioner is present with the client. Anesthesia time begins when the anesthesia practitioner begins to prepare the client for the induction of anesthesia in the operating room or the equivalent area and ends when the anesthesia practitioner is no longer in personal attendance (e.g., when the client may be safely placed under postoperative supervision).

For time-based anesthesiology codes, anesthesia practitioners must document interruptions in anesthesia time in the client’s medical record.

The documented time must be the same in the records or claims of the anesthesiologist and other anesthesia practitioners who were medically directed by the anesthesiologist.

One time unit is equal to 15 minutes of anesthesia. Providers must submit the total anesthesia time in minutes on the claim. The claims administrator will convert total minutes to time units.

Reimbursement of time-based anesthesia services is derived by adding the RVUs (e.g., base units) for the procedures performed (when multiple procedures are performed use the procedure with the highest RVUs) to the total face-to-face anesthesia time in minutes divided by 15 minutes, multiplied by the appropriate conversion factor:

\[
[RVUs + (Minutes / 15)] \times \text{Conversion Factor} = \text{Anesthesia Reimbursement}
\]

<table>
<thead>
<tr>
<th>Provider Type Description - Physician Pricing Example</th>
</tr>
</thead>
<tbody>
<tr>
<td>Time: 120 minutes = 120/15 = 8 (quantity billed)</td>
</tr>
<tr>
<td>Procedure code: 00851 = (6 RVUs) 6.00 + 8 = 14.00</td>
</tr>
<tr>
<td>Conversion factor: $19.58 = 14.00 \times 19.58 = $274.12 (physician reimbursement)</td>
</tr>
</tbody>
</table>

**Conversion Factor**

A conversion factor is the multiplier that transforms relative values into payment amounts. There is a standard conversion factor for anesthesia services.
9.2.7.9 Anesthesia Modifiers
Each anesthesia procedure code must be submitted with the appropriate anesthesia modifier combination whether billing as the sole provider or for the medical direction of CRNAs, AAs, or other qualified professionals.

When an anesthesia procedure is billed without the appropriate reimbursement modifiers or is billed with modifier combinations other than those listed below in the Modifier Combinations section, the claim will be denied.

A procedure billed with a modifier indicating that the anesthesia was personally performed by an anesthesiologist (modifier AA) will be denied if another claim has been paid indicating the service was personally performed by, and reimbursed to, a CRNA (modifier QZ) for the same client, date of service, and procedure code. The opposite is also true—a CRNA-administered procedure will be denied if a previous claim was paid to an anesthesiologist for the same client, date of service, and procedure code. Denied claims may be appealed with supporting documentation of any unusual circumstances.

9.2.7.9.1 State-Defined Modifiers
Modifiers U1 (indicating one Medicaid claim billed by an anesthesia practitioner) and U2 (indicating two Medicaid claims) are state-defined modifiers that must be billed by an anesthesiologist, CRNA, AA, or other qualified professional.

Modifier U1, indicating that only one Medicaid claim will be submitted, cannot be billed by two providers for the same procedure, client, and date of service. Modifier U2, indicating that two Medicaid claims will be submitted, can only be billed by two providers for the same procedure, client, and date of service if one of the providers was medically directed by the other. Denied claims may be appealed with supporting documentation of any unusual circumstances.

Anesthesia providers must submit modifier U1 or U2 in combination with an appropriate pricing modifier (AA, GC, QY, QK, AD, QZ, QX) when billing for any payable anesthesia procedure codes.

9.2.7.9.2 Modifier Combinations
When a single claim per client is billed by the anesthesiologist for personally performing the anesthesia service, the AA and U1 modifier combination must be billed together.

Anesthesiologists may be reimbursed for medical direction of CRNAs, AAs, or other qualified professional by using one of the following modifier combinations:

<table>
<thead>
<tr>
<th>Modifier Combination</th>
<th>Who will submit claims?</th>
</tr>
</thead>
<tbody>
<tr>
<td>Anesthesiologist Directing Other Qualified Professionals</td>
<td>Only the anesthesiologist</td>
</tr>
<tr>
<td>QY and U1</td>
<td>When a single claim per client is billed by the anesthesiologist for medically directing anesthesia services of an anesthesia procedure provided by one CRNA, AA, or other qualified professional, the QY + U1 modifier combination must be billed together when the CRNA, AA, or qualified professional are a part of a clinic/group.</td>
</tr>
</tbody>
</table>

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<thead>
<tr>
<th>Modifier Combination</th>
<th>When is it used?</th>
</tr>
</thead>
<tbody>
<tr>
<td>Anesthesiologist Directing Other Qualified Professionals</td>
<td>When a single claim per client is billed by the anesthesiologist for medically directing anesthesia services of an anesthesia procedure provided by one CRNA, AA, or other qualified professional, the QY + U1 modifier combination must be billed together when the CRNA, AA, or qualified professional are a part of a clinic/group.</td>
</tr>
<tr>
<td>Modifier Combination Submitted by Anesthesiologist</td>
<td>When is it used?</td>
</tr>
<tr>
<td>--------------------------------------------------</td>
<td>-----------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------</td>
</tr>
</tbody>
</table>
| AA, U1, and GC                                    | When a single claim per client is billed by the anesthesiologist for medically directing anesthesia services of an anesthesia procedure provided by one resident physician.  
  
  **Note:** For procedure code 01967 medical supervision of resident physicians rather than medical direction is required, however, modifiers AA−U1-GC must still be noted on the claim. | Only the anesthesiologist |
| QK and U1                                         | When a single claim per client is billed by the anesthesiologist for medically directing anesthesia services of two, three, or four concurrent anesthesia procedures provided by CRNAs, AAs, or other qualified professionals. | Only the anesthesiologist |
| AD and U1 (Emergency circumstances only)          | When a single claim per client is billed by the anesthesiologist for medical supervision of anesthesia services for more than four concurrent anesthesia procedures provided by CRNAs, AAs, or other qualified professionals. Used in emergency circumstances only and limited to 6 units (90 minutes) per case for each occurrence requiring five or more concurrent procedures. | Only the anesthesiologist |
| Anesthesiologist Directing CRNAs or AAs          |                                                                                                                                                                                                                |                        |
| QY and U2                                         | When two claims per client are billed, one by the medically directing anesthesiologist and one by the CRNA, AA, or other qualified professional.                                                                    | Both the anesthesiologist and CRNA, AA, or other qualified professional |
| QK and U2                                         | When two claims per client are billed for medically directed anesthesia services of two, three, or four concurrent anesthesia procedures provided by CRNAs, AAs, or other qualified professionals.                                                    | Both the anesthesiologist and CRNA, AA, or other qualified professional |
| AD and U2 (Emergency circumstances only)          | When two claims per client are billed for the medical supervision of more than four concurrent anesthesia procedures provided by CRNAs, AAs, or other qualified professionals. Used in emergency circumstances only and limited to 6 units (90 minutes) per case for each occurrence requiring five or more concurrent procedures. | Both the anesthesiologist and CRNA, AA, or other qualified professional |

### 9.2.7.9.3 CRNA, AA, and Other Qualified Professional Services

Modifiers QZ and U1 must be submitted when a CRNA has personally performed the anesthesia services, is not medically directed by the anesthesiologist, and is directed by the physician.

Modifiers QX and U2 must be submitted by a CRNA, AA, or other qualified professional who provided services under the medical direction of an anesthesiologist.
9.2.7.10  Prior Authorization for Anesthesia

9.2.7.10.1  Anesthesia for Medical Services

Anesthesia services provided in combination with most medical surgical procedures do not require prior authorization. However, some medical surgical procedures may require prior authorization. Anesthesia may be reimbursed if prior authorization for the surgical procedure was not obtained, but services provided by the facility, surgeon, and assistant surgeon will be denied.

9.2.7.11  Claims Filing

Texas Medicaid reimburses anesthesiologists based on the Tax Equity and Fiscal Responsibility Act (TEFRA) of 1982. Anesthesiologists must identify the following information on their claims:

- Procedure performed (CPT anesthesia code in Block 24 of the CMS-1500 paper claim form).
- Person (physician, CRNA, or AA) administering anesthesia (modifiers must be used to designate this provider type).
- Time in minutes.
- Any other appropriate modifier (refer to subsection 6.3.5, “Modifiers” in “Section 6: Claims Filing” (Vol. 1, General Information) for a list of the most common modifiers).

9.2.7.12  Anesthesia (General) for THSteps Dental

Refer to: Section 4 *, “Texas Health Steps (THSteps) Dental” in the Children’s Services Handbook (Vol. 2, Provider Handbooks) for additional information.

9.2.8  Bariatric Surgery

Bariatric surgery is considered medically necessary when used as a means to treat covered medical conditions that are caused or significantly worsened by the client’s obesity in cases where those comorbid conditions cannot be adequately treated by standard measures unless significant weight reduction takes place. The pathophysiology of the covered comorbid conditions must be sufficiently severe that the expected benefits of weight loss subsequent to this surgery significantly outweigh the risks associated with bariatric surgery.

The following procedure codes may be reimbursed for medically necessary bariatric surgery services with prior authorization: 43644, 43645, 43659, 43770, 43771, 43772, 43773, 43774, 43775, 43842, 43843, 43845, 43846, 43847, 43848, 43886, 43887, and 43888.

Bariatric surgery is not a benefit when the primary purpose of the surgery is any of the following:

- For weight loss for its own sake
- For cosmetic purposes
- For reasons of psychological dissatisfaction with personal body image
- For the client’s or provider’s convenience or preference

9.2.8.1  Prior Authorization for Bariatric Surgery

All clients must meet the criteria outlined below.

The same contraindications exist for bariatric surgery as for any other elective abdominal surgery. Documentation provided for prior authorization must attest that none of the following additional contraindications exist:

- Endocrine cause for obesity, inflammatory bowel disease, chronic pancreatitis, cirrhosis, portal hypertension, or abnormalities of the gastrointestinal tract
- Chronic, long-term steroid treatment
• Pregnant, or plans to become pregnant within 18 months
• Noncompliance with medical treatment
• Significant psychological disorders that would be exacerbated or interfere with the long-term management of the client after the operation
• Active malignancy

All clients must undergo preoperative psychological evaluation by a behavioral health provider and have clearance for surgery if any of the following conditions exist:
• They have a history of psychiatric or psychological disorders.
• They are currently under the care of a psychologist or psychiatrist.
• They are on psychotropic medications.

The client’s medical record must include documentation of the evaluation.

Clients without a history of psychiatric or psychological disorder must also undergo a preoperative psychological evaluation by a behavioral health provider and have clearance for surgery. The client’s medical record must include documentation that the client is psychologically mature and able to cope with the postsurgical changes of the surgery.

Documentation must be submitted with the prior authorization request that is signed by the surgeon and attests that the services are provided by a facility in Texas that is one of the following:
• Accredited by the Metabolic and Bariatric Surgery Accreditation and Quality Improvement Program (MBSAQIP).
• A children’s hospital that has a bariatric surgery program and provides access to an experienced surgeon who employs a team that is capable of long-term follow-up of the metabolic and psychosocial needs of the client and family.

Bariatric surgery for clients who are 20 years of age and younger may be prior authorized when the client meets all of the following criteria:
• The client has reached a Tanner Scale stage IV or V plus 95 percent of adult height based on bone age.
• The client has a body mass index (BMI) of greater than or equal to 40 kg/m2.
• The client has one or more comorbid conditions that are exacerbated by or attributable to obesity.
• Female clients must be at least 13 years of age and menstruating.
• Male clients must be at least 15 years of age.

Bariatric surgery for clients who are 21 years of age and older may be prior authorized when the client meets all of the following criteria:
• The client has a BMI of greater than or equal to 35 kg/m2.
• The client has one or more of the following comorbid conditions that are exacerbated by or attributable to obesity:
  • Obesity-associated hypoventilation
  • Moderate to severe sleep apnea (defined as apnea/hypoapnea index of 16 or more events per hour)
  • Congestive heart failure
  • Obesity-induced cardiomyopathy
• Refractory hypertension resistant to pharmacotherapy (defined as blood pressure greater than 140mmHg systolic or greater than 90mmHg diastolic, despite maximally tolerated doses of at least three different classes of antihypertensive medications)

• Pseudotumor cerebri (documented idiopathic intracerebral hypertension)

• Adult onset (Type II) diabetes (with or without complications) with Hgb A1c greater than 9 percent, regardless of therapy, or 7 to 9 percent on maximal medical therapy (defined as taking insulin or maximally tolerated doses of at least two different classes of oral hypoglycemic medications)

• Cardiovascular or peripheral vascular disease

• Refractory hyperlipidemia (defined as triglycerides greater than 250 mg/dl, cholesterol greater than 220/mg/dl, HDL less than 35 mg/dl, or LDL greater than 200 mg/dl, despite maximally tolerated doses of at least two different classes of lipid-lowering medications)

• Recurrent or chronic skin ulcerations with infection

• Pulmonary hypertension

• Chronic joint disease, deterioration of the joint cartilage, and the formation of new bone (bone spurs) at the margins of the joints, with symptoms that severely affect work or leisure activities, on maximal medical therapy (defined as maximally tolerated dose of a non-steroidal anti-inflammatory drug (NSAID) or COX-II inhibitor or acetaminophen and the completion of at least one physical-therapist-supervised exercise program)

• Hepatic steatosis without evidence of active inflammation

Documentation must include a summary of the treatment provided for the client’s comorbid conditions, including descriptions of how the client’s response to standard treatment measures are unsatisfactory and why the bariatric surgery is medically necessary in the context of current treatment and medically-reasonable alternatives that are available.

Referral for bariatric surgery to the bariatric surgeon is required from the practitioner who is treating the comorbid condition(s). The bariatric surgeon will determine the client’s eligibility for bariatric surgery. Documentation of the referral must be submitted with the prior authorization request.

The client must have had previous unsuccessful medical treatment for obesity, as documented in the medical record. All of the following minimal requirements must be met:

• The client has made a diligent effort to achieve healthy body weight with such efforts described in the medical record and certified by the operating surgeon.

• The client has failed to maintain a healthy weight despite a minimum of 6 months documented regular participation in a structured dietary program overseen by a physician (M.D. or D.O.) within 12 months of the request date.

Documentation that is submitted for prior authorization must also include all of the following:

• The process by which the client will receive postoperative surgical, nutritional, and psychological services.

• Affirmation that the client and the parent/guardian (if applicable) understand and will support the changes in eating habits that must accompany the surgery and the extensive postoperative follow-up.

Repeat bariatric surgery may be considered medically necessary in either of the following circumstances:

• To correct complications from bariatric surgery such as band malfunction, obstruction, or stricture
• To convert to a Roux-en-Y gastroenterostomy or to correct pouch failure in an otherwise compliant client when the initial bariatric surgery met medical necessity criteria

**Note:** Conversion to a Roux-en-Y gastroenterostomy may be considered medically necessary for clients who have not had adequate success (defined as a loss of more than 50 percent of excess body weight) two years following the primary bariatric surgery procedure, and the client has been compliant with a prescribed nutrition and exercise program following the procedure.

All documentation required for prior authorization is to be maintained in the client’s medical record and is subject to retrospective review. This includes medical records from both the practitioner treating the comorbid condition(s) and the bariatric surgeon.

Providers may fax or mail prior authorization requests for bariatric surgery services for clients who are 20 years of age and younger to the TMHP Comprehensive Care Program (CCP) Prior Authorization Department. Prior authorization requests for clients who are 21 years of age and older may be faxed or mailed to the TMHP Special Medical Prior Authorization Department.

Clients may be eligible under Texas Medicaid or CCP for separate reimbursement for nutritional and psychological assessment and counseling associated with bariatric surgery.

Behavioral health services provided as part of the preoperative or postoperative phase of bariatric surgery are subject to behavioral health guidelines, and are not considered part of the bariatric surgery.

**Refer to:** Subsection 7, “Psychiatric Services for Hospitals” in the *Behavioral Health and Case Management Services Handbook* (Vol. 2, Provider Handbooks) for information about behavioral health services.

### 9.2.9 Bacillus Calmette-Guérin (BCG) Intravesical for Treatment of Bladder Cancer

Live BCG for intravesical (procedure code 90586) or transvesical (procedure code J9030) are benefits of Texas Medicaid for the following diagnosis codes:

<table>
<thead>
<tr>
<th>Diagnosis Codes</th>
<th>C670</th>
<th>C671</th>
<th>C672</th>
<th>C673</th>
<th>C674</th>
<th>C675</th>
<th>C676</th>
<th>C677</th>
</tr>
</thead>
<tbody>
<tr>
<td>C678</td>
<td>C679</td>
<td>C7911</td>
<td>D090</td>
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</tbody>
</table>

Procedure code 90585 is a benefit of Texas Medicaid for diagnosis code Z23. Authorization is not required for the BCG vaccine.

Bladder instillation of anticarcinogenic agent (procedure code 51720) may be reimbursed separately when billed with BCG instillation (procedure code 90586 or J9030).

### 9.2.10 Behavioral Health Services

**Refer to:** The *Behavioral Health and Case Management Services Handbook* (Vol. 2, Provider Handbooks).

### 9.2.11 Biopsy

A biopsy refers to the surgical excision of tissue for pathological examination.

If a surgeon bills separate charges for a surgical procedure and a biopsy on the same organ or structure on the same day, the charges are reviewed and reimbursed only for the service with the higher of the allowed amounts.
9.2.12 Biofeedback Services

Biofeedback services are a benefit of Texas Medicaid for clients who are 4 years of age and older with the following conditions:

- Urinary incontinence
- Fecal incontinence
- Migraine and tension headache

Biofeedback services may be reimbursed using procedure codes 90901, 90912, and 90913.

Biofeedback services are limited to a maximum of 18 sessions rendered by any provider for the lifetime of each client for each condition.

Biofeedback services that are not a benefit of Texas Medicaid are the following:

- Biofeedback performed in the home setting
- Neurofeedback (such as, but not limited to, electroencephalography [EEG])
- Treatment for muscle tension, except tension headache
- Psychological, psychophysiological, and behavioral health therapy and psychosomatic conditions
- Investigational or experimental biofeedback services and procedures

Procedure codes 90901, 90912, and 90913 are limited to one service per day. The reimbursement for procedure codes 90901, 90912, and 90913 include all modalities of the biofeedback training performed on the same day, regardless of the time increments or the number of modalities performed.

Any device used during a biofeedback session is considered part of the procedure and will not be reimbursed separately.

9.2.12.1 Biofeedback Certification

A staff member who is certified by Biofeedback Certification International Alliance (BCIA) must perform biofeedback services.

The certification types accepted by Texas Medicaid are the following:

- General biofeedback certification (BCB)
- Pelvic muscle dysfunction biofeedback certification (BCB-PMD)

Providers must maintain documentation in the client’s medical record to support the medical necessity of the biofeedback service provided. Documentation must include the name of the staff person who provided the biofeedback and the prescribing physician must maintain in the office a record of the current certification of the staff member(s) who perform biofeedback. Documentation is subject to retrospective review.

9.2.12.2 Prior Authorization for Biofeedback Services

Prior authorization is required for biofeedback services.

- Any combination of procedure codes 90901, 90912, and 90913 are a benefit for biofeedback sessions for urinary or fecal incontinence conditions in clients who are 4 years of age and older.
- Procedure code 90901 is a benefit for biofeedback sessions for migraine or tension headache conditions.

The initial request may include up to 12 visits and not exceed a total duration of 12 weeks. Documentation of the following must be submitted for consideration of prior authorization:

- Conventional treatments that were given but were not successful, including, but not limited to, pharmacotherapy, exercise, rest, and heating and cooling modalities.
• Statements from the prescribing physician that the client is capable of understanding the requirements and agrees actively to participate in the biofeedback sessions.

• Name and certification information for the person performing the training.

In addition, documentation must be submitted to support the specific type of biofeedback requested.

**Urinary and Fecal Incontinence**

• Diagnosis of fecal or urinary stress, urge, overflow, or a mix of stress and urge incontinence in a client who is 4 years of age or older.

• Exclusion by the physician of any underlying medical conditions that could be causing the problem.

• Failed pelvic floor muscle exercise (PME) training for clients who are 21 years of age and older.

  **Note:** Failed trial of PME training is defined as no clinically significant improvement in urinary incontinence after completing four weeks of an ordered plan of PME exercises.

**Migraine and tension headache**

• A diagnosis of migraine, tension headache, or mixed migraine and tension headache.

• Symptoms that occur with a duration of at least 4 hours for at least 15 days a month over at least 3 months.

• Failure of first-line approaches, including avoidance of precipitating stimuli and pharmacological prophylaxis.

Prior authorization requests must be submitted by the physician to the Special Medical Prior Authorization (SMPA) Department. The request must be submitted with documentation that supports medical necessity. Providers may submit prior authorization requests online through the TMHP website at [www.tmhp.com](http://www.tmhp.com), by fax to 1-512-514-4213, or by mail to the following address:

Texas Medicaid & Healthcare Partnership  
Special Medical Prior Authorization  
12357-B Riata Trace Parkway  
Austin, TX 78727

After the client completes the initial biofeedback treatment course, prior authorization may be considered for a total of six follow-up sessions not to exceed three sessions per week and total duration not to exceed eight weeks. Providers must submit prior authorization documentation for the same condition as the original request, and must include each original symptom and how it has objectively improved. Documentation may include, but is not limited to, the following:

• For treatment of urinary incontinence, improvement in continence scores, vitality, health, a decrease in high-grade stress incontinence, nocturnal enuresis, and urine loss with activity. In clients who are 21 years of age and older, evidence of increased pelvic floor contraction strength and the ability to hold the contractions longer and to perform more repetitions.

• For treatment of fecal incontinence, improvement in continence scores, squeeze and anal pressures, squeeze duration, vitality, and health. In clients who are 21 years of age and older, evidence of increased pelvic floor contraction strength and the ability to hold the contractions longer and to perform more repetitions.

• For migraine and tension headaches, diminished intensity, frequency, and duration of the headache activity.

**9.2.13 Blepharoplasty Procedures**

Procedure codes 15820, 15821, 67911, 67961, 67966, 67971, 67973, 67974, and 67975 are not diagnosis-restricted.
Procedure codes 67901, 67902, 67903, 67904, 67906, 67908, and 67909 may be reimbursed for clients who are 20 years of age and younger without prior authorization when performed for one of the following diagnosis codes:

<table>
<thead>
<tr>
<th>Diagnosis Codes</th>
</tr>
</thead>
<tbody>
<tr>
<td>Q100</td>
</tr>
</tbody>
</table>

Procedure codes 67901, 67902, 67903, 67904, 67906, and 67908 do not require prior authorization for clients who are 21 years of age and older when billed for the following diagnosis codes:

<table>
<thead>
<tr>
<th>Diagnosis Codes</th>
</tr>
</thead>
<tbody>
<tr>
<td>H0231</td>
</tr>
<tr>
<td>H02422</td>
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</table>

Blepharoplasty for clients who are 21 years of age and older requires mandatory prior authorization. The following information from the physician is required at the time of the request for blepharoplasty for procedure codes 15820, 15821, 67901, 67902, 67903, 67904, 67906, 67908, 67909, 67911:

- A brief history and physical evaluation
- Photographs of the eyelid problem
- Visual field measurements
- Diagnosis code

The following blepharoplasty and eyelid repair procedures do not require prior authorization:

<table>
<thead>
<tr>
<th>Procedure Codes</th>
</tr>
</thead>
<tbody>
<tr>
<td>67916</td>
</tr>
</tbody>
</table>

All supporting documentation must be included with the request for authorization. Send requests and documentation to the following address:

Texas Medicaid & Healthcare Partnership  
Special Medical Prior Authorization  
12357-B Riata Trace Parkway, Suite 100  
Austin, TX 78727  
Fax: 1-512-514-4213

Retroactive authorization may be granted on an appeal basis when submitted with the appropriate documentation.

### 9.2.14 Bone Growth Stimulation

Professional services for bone growth stimulation (procedure codes 20974, 20975, and 20979) are a benefit of Texas Medicaid.

Prior authorization is required for a bone growth stimulator device (procedure codes E0747, E0748, E0749, and E0760).

**Refer to:** Subsection 2.2.8, “Bone Growth Stimulators” in the *Durable Medical Equipment, Medical Supplies, and Nutritional Products Handbook (Vol. 2, Provider Handbooks)* for prior authorization criteria.
9.2.14.1 Invasive Bone Growth Stimulation

Invasive bone growth stimulation (procedure code 20975) is indicated for the following conditions:

- Nonunion of long bone fractures (i.e., clavicle, humerus, radius, ulna, femur, tibia, fibula, and metacarpal, metatarsal, carpal, and tarsal bones). Nonunion of long bone fractures is considered to exist only when serial radiographs have confirmed that fracture healing has ceased for three or more months prior to starting treatment with the bone growth stimulator. Serial radiographs must include a minimum of 2 sets of radiographs separated by a minimum of 90 days. Each set of radiographs must include multiple views of the fracture site.
- Failed fusion of a joint other than the spine when a minimum of three months has elapsed since the joint fusion was performed.
- Congenital pseudoarthrosis.
- An adjunct to spinal fusion surgery for patients at high risk for pseudoarthrosis due to previously failed spinal fusion at the same site.
- An adjunct to multiple-level fusion, which involves three or more vertebrae (e.g., L3-L5, L4-S1, etc.).

9.2.14.2 Non-invasive Bone Growth Stimulation

Non-invasive bone growth stimulation (procedure code 20974) is indicated for the following conditions:

- Nonunions, failed fusions, and congenital pseudoarthrosis where there is no evidence of progression of healing for three or more months despite appropriate fracture care.
- Delayed unions of fractures of failed arthrodesis at high risk sites (e.g., open or segmental tibial fractures, carpal navicular fractures).

Documentation must also indicate all of the following:

- Serial radiographs have confirmed that no progressive signs of healing have occurred.
- The fractured gap is 1 cm or less.
- The individual can be adequately immobilized and is likely to comply with non-weight-bearing restrictions.

Non-invasive bone growth stimulation for spinal application is indicated for the following conditions:

- One or more failed fusions.
- Grade II or worse spondylolisthesis.
- A multiple-level fusion with extensive bone grafting is required.
- Other risk factors for fusion failure are present, including gross obesity, degenerative osteoarthritis, severe spondylolisthesis, current smoking, previous fusion surgery, previous disc surgery, or gross instability.

9.2.14.3 Ultrasound Bone Growth Stimulation

Ultrasound bone growth stimulation (procedure code 20979) is indicated for nonunion of a fracture, other than the skull or vertebrae, in a skeletally mature person, which is documented by a minimum of two sets of radiographs that were:

- Obtained prior to starting treatment with the osteogenesis stimulator.
- Separated by a minimum of 90 days.
- Taken with multiple views of the fracture site.
- Accompanied by a written interpretation by a physician who states that there has been no clinically significant evidence of fracture healing between the two set of radiographs.
Documentation must also indicate evidence of all of the following:

- The fracture is not tumor-related.
- The fracture is not fresh (less than 7 days), closed or grade I open, tibial diaphyseal fractures, or closed fractures of the distal radius (Colles fracture).

### 9.2.14.4 Reimbursement

Professional claims that are submitted for bone growth stimulation (procedure codes 20974, 20975, and 20979) may be reimbursed if the claim includes documentation of one of the following:

- Documentation of medical necessity as outlined for each type of bone growth stimulation.
- The corresponding bone growth stimulator device was submitted within 95 days of the date the bone growth stimulation procedure was performed.

The appropriate evaluation and management (E/M) procedure code must be billed for monitoring the effectiveness of bone growth stimulation treatment.

Procedure codes 20974, 20975, and 20979 are limited to one per six months. During the six-month limitation period, a subsequent fracture that meets the criteria for a bone growth stimulator may be reimbursed after the submission of an appeal with documentation of medical necessity that demonstrates the criteria have been met.

### 9.2.15 Cancer Screening and Testing

#### 9.2.15.1 BRCA Testing


#### 9.2.15.2 Colorectal Cancer Screening

Colorectal cancer screening is a benefit of Texas Medicaid. Fecal occult blood tests, multi-targeted stool DNA (mt-sDNA) tests, screening colonoscopies, and sigmoidoscopies are evidenced based methods of colorectal cancer screening. Screening refers to the testing of asymptomatic persons to assess their risk for the development of colorectal cancer. Screening has been shown to decrease mortality due to this cancer by detecting cancers at earlier stages and allowing the removal of adenomas, thus preventing the subsequent development of cancer.

The American Cancer Society (ACS) recommends screening people at average risk for colorectal cancer beginning at 45 years of age by any of the following methods:

- A fecal occult blood test (FOBT)* or fecal immunochemical test (FIT) every year, or
- A multi-targeted stool DNA test (mt-sDNA) every three years, or
- Flexible sigmoidoscopy every five years, or
- A Flexible sigmoidoscopy every ten years, in addition to annual FIT screening, or
- Colonoscopy every ten years

**Note:** For FOBT, the take-home multiple sample method with three samples should be used.

The U.S. Preventative Services Task Force (USPSTF) guidelines indicate that the net benefit of colorectal cancer screening in adults who are 76 years of age and older who have been previously screened is small. The risks should be considered on an individual basis, as screening in this age group is most appropriate for those healthy enough to undergo treatment.

The ACS and USPSTF recommends screening for people at high-risk for colorectal cancer once every two years.

Indications/characteristics of a high-risk individual may include one or more of the following:
- A close relative (sibling, parent or child) has had colorectal cancer or an adenomatous polyp.
- There is a family history of familial adenomatous polyposis.
- There is a family history of hereditary nonpolyposis colorectal cancer.
- There is a personal history of adenomatous polyps.
- There is a personal history of colorectal cancer.
- There is a personal history of colonic polyps.
- There is a personal history of inflammatory bowel disease, including Crohn’s disease and ulcerative colitis.

**Note:** “Relative” means close blood relatives including first degree male or female relatives (parents, siblings, or children), second-degree relatives (aunts, uncles, grandparents, nieces, nephews), and third-degree relatives (first cousins, great-grandparents) who are on the same side of the family as the clients.

Colorectal screening services are considered for reimbursement when submitted using procedure codes G0328 (with modifier QW), G0104, G0105, and G0121, by associated risk category based on the ACS and USPSTF frequency recommendations. Reimbursement for these procedure codes is considered when medical necessity is documented in the client’s record.

**Fecal Occult Blood Tests**
Procedure code G0328 (with modifier QW) and 82270 may be reimbursed once per rolling year for clients who are 45 years of age and older.

**MT-sDNA Test**
Procedure code 81528 is considered for reimbursement once every three years for clients who are 45 years of age and older.

**Sigmoidoscopies**
Procedure code G0104 is considered for reimbursement once every five years for clients who are 45 years of age and older when submitted with diagnosis code Z0000, Z0001, Z1210, Z1211, Z1213, Z859, Z86002, Z86009, Z86006, Z86007, or Z86010, as recommended by the ACS and USPSTF. Diagnosis code Z0000 or Z0001 may be used for screening if no other diagnosis is appropriate for the service rendered, but not more frequently than recommended by the USPSTF.

If a lesion or growth is detected that results in a biopsy or removal of the growth during a screening flexible sigmoidoscopy, the appropriate diagnostic procedure classified as a flexible sigmoidoscopy with biopsy or removal must be reported.

**Colonoscopies: Average Risk**
Procedure code G0121 is considered for reimbursement once every ten rolling years for clients who are 45 years of age and older when submitted with diagnosis code Z0000, Z0001, Z1210, Z1211, or Z1213. Diagnosis code Z0000 or Z0001 may be used for screening if no other diagnosis is appropriate for the service rendered, but not more frequently than recommended by the USPSTF.

**Colonoscopies: High-Risk**
Procedure code G0105 is considered for reimbursement once every two years for clients who meet the definition of high-risk. Procedure code G0105 must be submitted with one of the following diagnosis codes:

<table>
<thead>
<tr>
<th>Diagnosis Codes</th>
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</thead>
<tbody>
<tr>
<td>K5000</td>
</tr>
</tbody>
</table>
9.2.15.2.1 Prior Authorization for Colorectal Cancer Screening
Prior authorization is not required for colorectal screening.

9.2.15.2.2 Exclusions
Barium enemas for colorectal cancer screening are not a benefit of Texas Medicaid.

9.2.15.3 Genetic Testing for Colorectal Cancer
Genetic testing for colorectal cancer may be considered for reimbursement to independent laboratories with prior authorization.

Genetic testing may be provided to clients who have a known predisposition (i.e., having a first- or second-degree relative) for colorectal cancer. Results of the testing may indicate whether the client has an increased risk of developing colorectal cancer. A first-degree relative is defined as a sibling, parent, or offspring. A second-degree relative is defined as an uncle, aunt, grandparent, nephew, niece, or half-sibling.

Genetic test results, when informative, may influence clinical management decisions. Documentation in the medical record must reflect that the client or family members have been given information on the nature, inheritance, and implications of genetic disorders to help them make informed medical and personal decisions before the genetic testing.

Genetic testing for colorectal cancer may be considered for reimbursement with the following procedure codes:

<table>
<thead>
<tr>
<th>Procedure Codes</th>
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<tbody>
<tr>
<td>81201</td>
</tr>
<tr>
<td>81294</td>
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<tr>
<td>81319</td>
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</tbody>
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Diagnosis code Z800 is acceptable as a diagnosis for the procedure codes in the table above. Prior authorization is still required and must be obtained for these services. Interpretation of gene mutation analysis results is not reimbursed separately. Interpretation is part of the physician E/M service.

The genetic testing for colorectal testing procedure codes in the table above are limited to once per lifetime for any procedure code by any provider. Testing is limited to once per lifetime for any procedure code by any provider, regardless of whether additional services are authorized.

Providers must maintain the following documentation in the client’s medical record for genetic testing for colorectal cancer:

- Documentation of formal pre-test counseling, including assessment of the client’s ability to understand the risks and limitations of the test.
• The client’s informed choice to proceed with the genetic testing for colorectal cancer.

The provider must order the most appropriate test based on familial medical history and the availability of previous family testing results.

The medical record is subject to retrospective review.

9.2.15.3.1 Testing for Familial Adenomatous Polyposis

Testing for familial adenomatous polyposis (procedure codes 81201, 81202, and 81203) may be offered to clients who have well-defined hereditary cancer syndromes and for whom a positive or negative result will change medical care. Testing for familial adenomatous polyposis may be considered for reimbursement with documentation of at least one of the following:

• The client has more than 20 polyps.
• The client has a first-degree relative with familial adenomatous polyposis and a documented mutation.
• For clients who are 7 years of age and younger, testing must be medically necessary and supported by documentation with a clear rationale for testing, which must be retained in the client’s medical record.

9.2.15.3.2 Hereditary Nonpolyposis Colorectal Cancer (HNPCC)

Testing for HNPCC (procedure codes 81288, 81292, 81293, 81294, 81295, 81296, 81297, 81298, 81299, 81300, 81301, 81317, 81318, and 81319) is used to determine whether a client has an increased risk of colorectal cancer or other HNPCC-associated cancers, including Lynch Syndrome. Results of the test may influence clinical management decisions. Testing for HNPCC may be considered for reimbursement with documentation of at least one of the following:

• The client has three or more family members, one of whom is a first-degree relative, with colorectal cancer; two successive generations are affected; one or more of the colorectal cancers was diagnosed before the family member was 50 years of age; and familial adenomatous polyposis has been ruled out for the client.
• The client has had two previous HNPCCs.
• The client has colorectal cancer and a first-degree relative who has one of the following:
  • Colorectal cancer or HNPCC extracolonic cancer at 50 years of age and younger
  • Colorectal adenoma at 40 years of age and younger
• The client has had colorectal cancer or endometrial cancer at 50 years of age and younger.
• The client has had right-sided colorectal cancer with an undifferentiated pattern of histology at 50 years of age and younger.
• The client has had signet-cell type colorectal cancer at 50 years of age and younger.
• The client has had a colorectal adenoma at 40 years of age and younger.
• The client is asymptomatic and has a first- or second-degree relative who has a documented HNPCC mutation.
• The client has a family history of malignant neoplasm in the gastrointestinal tract.
• For clients who are 20 years of age and younger, testing must be medically necessary and supported by documentation with a clear rationale for testing, which must be retained in the client’s medical record.
9.2.15.3.3 Prior Authorization for Genetic Testing for Colorectal Cancer

Prior authorization is required for genetic testing for colorectal cancer. A completed Special Medical Authorization Request Form must be signed, dated, and submitted by the ordering provider rendering direct care. Requests from laboratories will not be processed. The provider should then share the authorization number with the laboratory submitting the claim.

A provider’s signature, including the prescribing provider’s, on a submitted document indicates that the provider certifies, to the best of the provider’s knowledge, the information in the document is true, accurate, and complete.

Medical documentation that is submitted by the physician must verify the client’s diagnosis or family history. Requisition forms from the laboratory are not sufficient for verification of the personal and family history.

To complete the prior authorization process, the provider must mail or fax the request to the TMHP Special Medical Prior Authorization Unit and include documentation of medical necessity. The form may be faxed to 1-512-514-4213 or mailed to the following address:

Texas Medicaid & Healthcare Partnership
Special Medical Prior Authorization Department
12357-B Riata Trace Parkway, Suite 100
Austin, TX 78727

A request for retroactive authorization must be submitted no later than 7 calendar days after the lab draw is performed. To facilitate a determination of medical necessity and avoid unnecessary denials, the ordering physician rendering care must provide correct and complete information, including the accurate medical necessity of the services requested.

9.2.15.4 Mammography (Screening and Diagnostic Studies of the Breast)

The following breast imaging studies are benefits of Texas Medicaid:

- Screening mammogram
- Diagnostic mammogram
- Diagnostic breast ultrasound

The American Cancer Society recommends that women discuss when to start breast cancer screening mammography with their provider beginning at 40 years of age.

By the age of 45 all women should begin annual breast cancer mammography screening.

By the age of 55 women may transition to screening with mammography every other year, or in some cases may continue annual screenings in consultation with their healthcare provider.

Digital breast tomosynthesis (DBT), also known as three-dimensional (3D) mammography, provides 3D images and is a modification of conventional mammography. Screening DBT is used, along with conventional screening mammography, to detect breast changes in women who have no signs or symptoms of breast cancer.

Diagnostic DBT is used, along with conventional diagnostic mammography, to diagnose breast disease in women or men who have breast symptoms or findings on physical examination or screening mammogram.

A screening mammogram may be billed using procedure code 77067.

Procedure code 77063 must be billed with primary procedure code 77067. Reimbursement may be considered for procedure code 77063 when performed on the same date of service, by any provider, as procedure code 77067.

Procedure codes 77063 and 77067 are limited to one per rolling year, any provider.
A diagnostic mammogram may be billed using procedure code 77065 or 77066. Procedure code 77065 will be denied if it is submitted for the same date of service as procedure code 77066 by any provider.

Procedure code G0279 must be billed with primary procedure code 77065 or 77066. Reimbursement may be considered for procedure code G0279 when performed on the same date of service, by any provider, as procedure code 77065 or 77066.

Reimbursement may be considered for a screening mammogram (procedure code 77063 or 77067) performed on the same patient on the same date of service as a diagnostic mammogram (procedure code 77065, 77066, or G0279), by submitting the diagnostic mammography with the modifier GG.

A mammogram may be indicated for a male client based on medical necessity due to existing signs and symptoms. In such rare circumstances, procedure codes 77065, 77066, and G0279 may be considered for reimbursement.

Other breast diagnostic radiology procedures may be medically necessary based on existing signs and symptoms. When indicated, such procedures may be considered for reimbursement using procedure code 76098, 77053, or 77054. Procedure code 77053 will be denied if it is submitted for the same date of service as procedure code 77054 by any provider. Procedure code 76098 may be reimbursed for both male and female clients.

Breast ultrasound may be considered for reimbursement using procedure code 76641 or 76642. Authorization is not required for these services.

The prescribing physician must maintain documentation of medical necessity in the client’s medical record.

The radiologist or interpreting physician at the testing facility may determine and document that, because of the abnormal result of the diagnostic test performed, additional studies are medically necessary. The radiologist or interpreting physician ordering the additional studies must provide documentation to the prescribing physician.

9.2.15.5 Prognostic Breast and Gynecological Cancer Studies

Prognostic breast and gynecological cancer studies are benefits of Texas Medicaid when ordered by a physician for the purpose of determining the best course of treatment for a patient with breast/gynecological cancers.

Prognostic breast and gynecological cancer studies are divided into three categories: Receptor assays, Her-2/neu, and gene expression profiling.

- **Receptor Assays** (procedure codes 84233 and 84234) - The estrogen receptor assay (ERA) and the progesterone receptor assay (PRA) are tests in which a tissue sample is exposed to radioactively tagged estrogen or progesterone. The presence of these receptors can have prognostic significance in breast and endometrial cancer.

- **Her-2/neu** (procedure codes 83950, 88237, 88239, 88271, 88274, 88291, 88341, 88342, 88344, 88345, 88346, 88347, 88348, 88360, 88361, 88364, 88365, 88366, 88367, 88368, 88369, 88373, 88374, and 88377) - Human epidermal growth factor receptor 2 (Her-2/neu) is responsible for the production of a protein that signals cell growth. The overexpression of Her-2/neu in breast cancer is associated with decreased overall survival and response to some therapies. Each procedure used in the analysis should be coded separately.

- **Gene expression profiling** (procedure code 81519) - Gene expression profiling using the Oncotype DX® Breast Cancer Assay analyzes the expression of a panel of 21 genes to predict the likelihood of breast cancer recurrence in clients with newly diagnosed early stage invasive breast cancer.
Reimbursement for procedure codes 88360 and 88361 is limited to claims with a diagnosis of breast or uterine cancer as listed in the following table:

<table>
<thead>
<tr>
<th>Diagnosis Codes</th>
</tr>
</thead>
<tbody>
<tr>
<td>C50011</td>
</tr>
<tr>
<td>C50211</td>
</tr>
<tr>
<td>C50411</td>
</tr>
<tr>
<td>C50611</td>
</tr>
<tr>
<td>C50911</td>
</tr>
<tr>
<td>C7981</td>
</tr>
</tbody>
</table>

Testing for other diagnoses will be denied.

Interpretation of receptor assays, and Her-2/neu results is not considered separately for reimbursement. Interpretation is part of the physician’s E/M service.

Gene expression profiling (procedure code 81519) is a benefit when all of the following criteria are met:

- The test is ordered by an oncologist.
- The client has newly diagnosed breast cancer. ("Newly diagnosed" means that not more than six months have elapsed since the initial diagnosis.)
- The clinical stage of the breast cancer is I or II.
- Axillary node biopsy is negative for tumor, and there is no evidence of metastatic breast cancer.
- The primary tumor is estrogen receptor-positive, and Her-2/neu receptor negative, or the primary tumor is Her-2/neu receptor positive and less than 1 cm in diameter.
- The client is a candidate for adjuvant chemotherapy.
- The outcome of the test will guide decision-making regarding adjuvant chemotherapy.
- The client has one of the following diagnosis codes:

<table>
<thead>
<tr>
<th>Diagnosis Codes</th>
</tr>
</thead>
<tbody>
<tr>
<td>C50011</td>
</tr>
<tr>
<td>C50211</td>
</tr>
<tr>
<td>C50411</td>
</tr>
<tr>
<td>C50611</td>
</tr>
<tr>
<td>C50911</td>
</tr>
<tr>
<td>D0581</td>
</tr>
</tbody>
</table>

Gene expression profiling is limited to once per lifetime, but may be considered for reimbursement more than once per lifetime for the same client on appeal. The provider must submit documentation that demonstrates that the client has a new, second, primary breast cancer diagnosis that meets the criteria described above.

The provider must maintain documentation of medical necessity in the client’s medical record. Retrospective review may be performed to ensure that the documentation supports the medical necessity of the service.

Gene expression profiling is not covered for repeat testing or testing of multiple tumor sites in the same client.
Tests for gene expression profiling other than Oncotype DX® are considered experimental and investigational, and are not benefits of Texas Medicaid.

**9.2.16 Capsulotomy**

A capsulotomy is a benefit when not performed with a joint surgery.

**9.2.17 Cardiac Rehabilitation**

Cardiac rehabilitation is a physician-supervised program that furnishes physician-prescribed exercise, cardiac risk factor modification, psychosocial assessment, and outcomes assessment. Cardiac rehabilitation programs must include all of the following:

- Physician-prescribed exercise for each day on which cardiac rehabilitation items and services are furnished
- Cardiac risk factor modification, including education, counseling, and behavioral intervention, tailored to a client’s individual needs
- Psychosocial assessment
- Outcomes assessment
- An individual treatment plan that specifies how components are used for a client and that is reviewed and signed by the prescribing physician every 30 days

Cardiac rehabilitation procedure codes 93797 and 93798 are benefits of Texas Medicaid.

The appropriate procedure code must be billed with one of the following diagnosis codes:

<table>
<thead>
<tr>
<th>Diagnosis Codes</th>
</tr>
</thead>
<tbody>
<tr>
<td>I110</td>
</tr>
<tr>
<td>I2102</td>
</tr>
<tr>
<td>I219</td>
</tr>
<tr>
<td>I2720</td>
</tr>
<tr>
<td>I5020</td>
</tr>
<tr>
<td>I5040</td>
</tr>
<tr>
<td>I50814</td>
</tr>
<tr>
<td>Z951</td>
</tr>
</tbody>
</table>

Coverage of cardiac rehabilitation programs is considered reasonable and necessary only for clients for whom there is documentation of any of the following conditions within the 12 months immediately preceding the beginning of the program:

- Acute myocardial infarction
- Coronary artery bypass surgery (CABG)
- Percutaneous transluminal coronary angioplasty or coronary stenting
- Heart valve repair or replacement
- Major pulmonary surgery
- Sustained ventricular tachycardia or fibrillation
- Class III or class IV congestive heart failure
• Chronic stable angina

**Note:** A cardiac rehabilitation program in which the cardiac monitoring is done using telephonically transmitted electrocardiograms (ECGs) to a remote site is not a benefit of Texas Medicaid.

Cardiac rehabilitation must be provided in a facility that has the necessary cardiopulmonary, emergency, diagnostic, and therapeutic life-saving equipment (e.g., oxygen, cardiopulmonary resuscitation equipment, or defibrillator) available for immediate use.

Cardiac rehabilitation is limited to 2 one-hour sessions per day for 18 weeks per rolling year and can not exceed 36 sessions.

Cardiac rehabilitation may be considered medically necessary beyond 36 sessions if the client has another documented cardiac event or if the prescribing physician documents that a continuation of cardiac rehabilitation is medically necessary. To confirm that a continuation of cardiac rehabilitation is at the request of or is coordinated with the prescribing physician, the medical record must include evidence of communication between the cardiac rehabilitation staff and the prescribing physician. If the physician responsible for such follow-up is the medical director, then the physician’s notes must be evident in each client’s chart.

Additional cardiac rehabilitation sessions must be prior authorized and must not exceed a total of 36 sessions for 52 weeks from the date of authorization of additional sessions.

If no clinically-significant arrhythmia is documented during the first three weeks of the program, the physician may give the order for the client to complete the remaining portion of the cardiac rehabilitation without telemetry monitoring.

Although cardiac rehabilitation may be considered a form of physical therapy, it is a specialized program that is conducted by personnel who are not physicians but are trained in both basic and advanced cardiac life support techniques and exercise therapy for coronary disease and who provide the services under the direct supervision of a physician.

Direct supervision of a physician means that a physician must be immediately available and accessible for medical consultations and emergencies at all times when items and services are being furnished under cardiac rehabilitation programs.

9.2.17.1 Prior Authorization for Cardiac Rehabilitation

Prior authorization is not required for the initial 36 sessions of cardiac rehabilitation.

Cardiac rehabilitation may be considered medically necessary beyond 36 sessions in the following circumstances:

- The medical record must support the client has had another cardiac event; or
- The prescribing physician documents that a continuation of cardiac rehabilitation is medically necessary. Documentation must include the following:
  - Progress made from the beginning of cardiac rehabilitation period to the current service request date, including progress towards previous goals.
  - Information that supports the client’s capability of continued measurable progress.
  - A proposed treatment plan for the requested extension dates with specific goals related to the client’s individual needs.

Requests for prior authorization for additional sessions that exceed a total of 36 sessions in 52 weeks will not be granted. Prior authorization must be obtained through the TMHP Special Medical Prior Authorization (SMPA) Department.
9.2.17.2 Reimbursement

The evaluation provided by the cardiac rehabilitation team at the beginning of each cardiac rehabilitation session is not considered a separate service and will be included in the reimbursement for the cardiac rehabilitation session. Evaluation and management (E/M) services unrelated to cardiac rehabilitation may be billed with modifier 25 appended to the E/M code when a separately identifiable E/M service was provided on the same day by the provider that rendered cardiac rehabilitation. Documentation that supports the provision of a significant, separately identifiable E/M service must be maintained in the client’s medical record and made available to Texas Medicaid upon request.

Physical and occupational therapy will not be reimbursed when furnished in addition to cardiac rehabilitation exercise program services unless there is also a diagnosis of a non-cardiac condition that requires such therapy, e.g., a client who is recuperating from an acute phase of heart disease and may have had a stroke that requires physical and/or occupational therapy.

Client education services, such as formal lectures and counseling on diet, nutrition, and sexual activity, that help a client adjust living habits because of the cardiac condition; will not be separately reimbursed when the services are provided as part of the cardiac rehabilitation program.

9.2.18 Casting, Splinting, and Strapping

Casting, splinting, and strapping supplies are considered part of the procedure and are not reimbursed separately. The following procedure codes for casting, splinting, and strapping are a benefit of Texas Medicaid:

<table>
<thead>
<tr>
<th>Procedure Codes</th>
</tr>
</thead>
<tbody>
<tr>
<td>29000 29010 29015 29035 29040 29044 29046 29049 29055 29058</td>
</tr>
<tr>
<td>29065 29075 29085 29086 29105 29125 29126 29130 29131 29200</td>
</tr>
<tr>
<td>29220 29240 29260 29280 29305 29325 29345 29355 29358 29365</td>
</tr>
<tr>
<td>29405 29425 29435 29440 29445 29450 29505 29515 29520 29530</td>
</tr>
<tr>
<td>29540 29550 29580</td>
</tr>
</tbody>
</table>

When a claim for casting, splinting, or strapping is submitted with the same date of service as a surgery, the surgery may be reimbursed and the procedure codes listed in the table above will be denied as part of another procedure.

The replacement of a cast, splint, or strapping is not included in the original surgical fee and may be reimbursed separately. Reimbursement for cast removal, windowing, wedging, or repair will be denied if submitted for reimbursement within six weeks of the initial cast application, splinting, or strapping by the same provider.

<table>
<thead>
<tr>
<th>Procedure Codes</th>
</tr>
</thead>
<tbody>
<tr>
<td>29700 29705 29710 29720 29730 29740</td>
</tr>
</tbody>
</table>

The following procedure codes for cast removal, windowing, wedging, or repair may be reimbursed to a provider other than the provider who applied the initial cast, splint, or strap:

<table>
<thead>
<tr>
<th>Procedure Codes</th>
</tr>
</thead>
<tbody>
<tr>
<td>29700 29705 29710 29720 29730 29740 29750 29799</td>
</tr>
</tbody>
</table>

Authorization is not required for casting, splinting, or strapping services.
The following table includes the procedure codes that will be denied when submitted for reimbursement with other casting, splinting, and strapping procedure codes:

<table>
<thead>
<tr>
<th>Procedure Codes That Will Be Denied</th>
<th>When Submitted With Any of These Procedure Codes</th>
</tr>
</thead>
<tbody>
<tr>
<td>36000, 36410, 37202, 51701, 51702, 51703, 64415, 64416, 64417, 64450, 96360, 96365, 96372, 96374, or 96375</td>
<td>29000, 29010, 29015, 29035, 29040, 29044, 29046, 29049, 29055, 29058, 29065, 29075, 29085, 29086, 29105, 29125, 29126, 29130, 29131, 29200, 29220, 29240, 29260, 29280, 29305, 29325, 29345, 29355, 29358, 29365, 29405, 29425, 29435, 29440, 29445, 29450, 29505, 29515, 29520, 29530, 29540, 29550, 29580, 29700, 29705, 29710, 29720, 29730, 29740, 29750, or 29799</td>
</tr>
<tr>
<td>29035</td>
<td>29040, 29044, or 29046</td>
</tr>
<tr>
<td>29044</td>
<td>29046</td>
</tr>
<tr>
<td>29075</td>
<td>29065, 29105, or 29425</td>
</tr>
<tr>
<td>29085, 29125, 29126, or 29705</td>
<td>29065 or 29075</td>
</tr>
<tr>
<td>29105</td>
<td>29065</td>
</tr>
<tr>
<td>11055, 11056, 11057, or 29125</td>
<td>29425</td>
</tr>
<tr>
<td>12001, 12002, 12035, 29125, or 29705</td>
<td>29105</td>
</tr>
<tr>
<td>12001, 28190, 28192, 28193, 29130, 29131, 29260, or 29700</td>
<td>29075</td>
</tr>
<tr>
<td>29705</td>
<td>29435</td>
</tr>
<tr>
<td>12002</td>
<td>29125, 29530, or 29580</td>
</tr>
<tr>
<td>12001, 12032, 12042, 12044, 13121, 13132, 29130, or 29260</td>
<td>29125</td>
</tr>
<tr>
<td>29305</td>
<td>29325</td>
</tr>
<tr>
<td>29365 or 29425</td>
<td>29345</td>
</tr>
<tr>
<td>29405</td>
<td>29345, 29425, or 29740</td>
</tr>
<tr>
<td>29345, 29365, 29405, or 29425</td>
<td>29355</td>
</tr>
<tr>
<td>29440, 29580, 29700, or 29705</td>
<td>29405 or 29425</td>
</tr>
<tr>
<td>29580</td>
<td>29515 or 29705</td>
</tr>
<tr>
<td>29730</td>
<td>29405</td>
</tr>
<tr>
<td>29540</td>
<td>29425, 29505, 29515, or 29580</td>
</tr>
<tr>
<td>29730 or 29740</td>
<td>29445</td>
</tr>
<tr>
<td>29515</td>
<td>29505</td>
</tr>
<tr>
<td>11055, 11056, or 29550</td>
<td>29515</td>
</tr>
<tr>
<td>11900, 12004, or 29550</td>
<td>29540</td>
</tr>
<tr>
<td>12004, 15852, 29550, or 29700</td>
<td>29580</td>
</tr>
<tr>
<td>G0127, 11719, or 11900</td>
<td>29550</td>
</tr>
<tr>
<td>15852</td>
<td>29705</td>
</tr>
</tbody>
</table>

### 9.2.19 Cardiopulmonary Resuscitation (CPR)

CPR (procedure code 92950) is a benefit of Texas Medicaid and may be reimbursed when medical necessity is documented in the client's medical record. Only the primary provider performing CPR may be reimbursed for procedure code 92950. CPR billed as an ambulance service by an ambulance provider will be denied.
CPR may be billed with the same date of service as critical care when reported as a separately identifiable procedure. The time spent performing CPR must not be included in the time reported as critical care.

### 9.2.20 Circumcisions

Texas Medicaid may provide reimbursement for circumcisions billed with procedure code 54150 or procedure code 54161. Circumcisions performed on clients who are 1 year of age and older must be documented with medical necessity.

*Refer to:* Subsection 9.2.44.1, “Circumcisions for Newborns” in this handbook for additional benefit information.

### 9.2.21 Closure of Wounds

The repair of wounds is defined as simple, intermediate, or complex. Simple repair involves the dermis and subcutaneous tissue and requires a one-layer closure. Intermediate repair requires some layered closure of deeper layers of subcutaneous tissue and superficial fascia. Complex repair involves more layered closure, debridement, extensive undermining, stints, or retention sutures.

Wound closures may use sutures, staples, or tissue adhesives. Wounds closed with adhesive strips must not be reported using wound closure procedure codes. When adhesive strips are the only wound closure material used, providers must report the most appropriate E/M visit procedure code on their claim.

Simple exploration of nerves, blood vessels, or tendons exposed in an open wound is considered inclusive to the wound closure and will not be reimbursed separately.

The lengths of multiple closures of wounds must be added together and billed as one procedure code if they meet at least one of the following criteria:

- The closures have the same CPT classification (see “Repair [Closure]” in the CPT manual).
- The closures are in anatomic sites that are grouped together in the same procedure code descriptor.

Providers must submit the procedure code that represents the total length of the repairs. Lengths of repairs from different CPT classifications or groupings of anatomic sites must be billed as separate procedure codes.

Wound closures must be billed using the following procedure codes:

<table>
<thead>
<tr>
<th>Procedure Codes</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>Repair Simple</strong></td>
</tr>
<tr>
<td>12001</td>
</tr>
<tr>
<td>12016</td>
</tr>
<tr>
<td><strong>Repair Intermediate</strong></td>
</tr>
<tr>
<td>12031</td>
</tr>
<tr>
<td>12046</td>
</tr>
<tr>
<td><strong>Repair Complex</strong></td>
</tr>
<tr>
<td>13100</td>
</tr>
<tr>
<td>13152</td>
</tr>
</tbody>
</table>

Multiple wounds on the same day will be paid the full allowed amount for the major (largest total length of the repair at the same anatomic site) wound and one-half the allowed amount for each additional laceration (total length of the repair at the same anatomic site).

No separate payment will be made for incision closures billed in addition to a surgical procedure when the closure is part of that surgical procedure.

No separate payment will be made for supplies in the office.
When the debridement is carried out separately without immediate primary closure, when gross contamination requires prolonged cleansing, or when large amounts of devitalized or contaminated tissue are removed, debridement may be reimbursed separately. Debridement rendered during the same surgical session as wound closure is considered inclusive to the closure and is not reimbursed separately.

Refer to: Subsection 9.2.69.11, “Supplies, Trays, and Drugs” in this handbook for the hospital-based emergency department.

Wound suture and wound closure are considered part of any surgical procedure performed on the same area, except for excision of benign or malignant lesion procedure codes that require more than simple closure. Providers may be reimbursed for the appropriate intermediate or complex closure procedure code. Multiple surgery guidelines apply.

The exceptions listed above apply to the following excision and closure procedure codes:

<table>
<thead>
<tr>
<th>Excision of Benign Lesion Procedure Code</th>
</tr>
</thead>
<tbody>
<tr>
<td>11400 11401 11402 11403 11404 11406 11420 11421 11422 11423</td>
</tr>
<tr>
<td>11424 11426 11440 11441 11442 11443 11444 11446</td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th>Excision of Malignant Lesion Procedure Codes</th>
</tr>
</thead>
<tbody>
<tr>
<td>11600 11601 11602 11603 11604 11606 11620 11621 11622 11623</td>
</tr>
<tr>
<td>11624 11626 11640 11641 11642 11643 11644 11646</td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th>Intermediate Closure Procedure Codes</th>
</tr>
</thead>
<tbody>
<tr>
<td>12031 12032 12034 12035 12036 12037 12041 12042 12044 12045</td>
</tr>
<tr>
<td>12046 12047 12051 12052 12053 12054 12055 12056 12057</td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th>Complex Closure Procedure Codes</th>
</tr>
</thead>
<tbody>
<tr>
<td>13100 13101 13102 13120 13121 13122 13131 13132 13133 13151</td>
</tr>
<tr>
<td>13152 13153 13160</td>
</tr>
</tbody>
</table>

### 9.2.22 Cochlear Implants

Cochlear implants, when medically indicated, are benefits of Texas Medicaid with prior authorization. A cochlear implant device (procedure code 69930) is an electronic instrument, part of which is implanted surgically to stimulate auditory nerve fibers, and part of which is worn externally to capture and amplify sound. These devices are available in single and multichannel models. Cochlear implants are used to provide awareness and identification of sound and to facilitate communication for persons who are profoundly hearing impaired.

Refer to: Subsection 3.2.1, “Cochlear Implants” in the Vision and Hearing Services Handbook (Vol. 2, Provider Handbooks) for additional information on benefit and authorization requirements for cochlear implants.

### 9.2.23 Continuous Glucose Monitoring (CGM)

CGM (procedure codes 95250 and 95251) is a benefit of Texas Medicaid with prior authorization. Procedure codes 95250 and 95251 are limited to once per 12 calendar months by any provider.

The rental or purchase of a continuous glucose monitoring system (CGMS) is considered part of the CGM and is not reimbursed separately.
9.2.23.1 Prior Authorization for Continuous Glucose Monitoring

CGM requires prior authorization and must be prescribed by a physician performing the glucose monitoring.

CGM may be prior authorized for clients with Type I diabetes or diabetes during pregnancy, including gestational diabetes. The client must be compliant with his or her current medical regimen, use insulin injections three or more times per day or be on an insulin pump, and have documented self-blood glucose monitoring at least four times per day. At least one or more of the following conditions must also be present:

- Frequent unexplained hypoglycemic episodes
- Unexplained large fluctuations in daily, preprandial blood glucose
- Episodes of ketoacidosis or hospitalization for uncontrolled glucose

Additional CGM services may be considered with documentation of medical necessity that indicates the client meets the criteria above and has a change in condition that would warrant a second procedure within 12 calendar months.

To avoid unnecessary denials, the physician must provide correct and complete information, including documentation of medical necessity for the requested services. The physician must maintain documentation of medical necessity in the client’s medical record. The requesting provider may be asked for additional information to clarify or complete a request for the use of CGM.

9.2.24 Developmental Screening and Testing and Aphasia Assessment

The following types of developmental screening and testing and aphasia assessment are benefits of Texas Medicaid when medically necessary:

- Developmental screening when performed outside of a Texas Health Steps (THSteps) medical checkup (procedure code 96110)
- Developmental testing (procedure codes 96112 and 96113 [add-on procedure code must be submitted with primary procedure code 96112])
- Assessment of aphasia (procedure code 96105)

Re-evaluations are a benefit of Texas Medicaid only to address a clinical need, to provide the documentation needed to measure a client’s status over time, and to direct the plan of care.

Procedure codes 96105, 96110, 96112, and 96113 are used to report medically necessary aphasia assessment, developmental screening, and testing.

Prior authorization is not required for developmental screening, developmental testing, and aphasia assessment.

9.2.24.1 Developmental Screening

Developmental screening requiring the use of a standardized, validated screening tool (procedure code 96110) is a benefit of Texas Medicaid for clients who are birth through 6 years of age.

Developmental screening is limited to once per rolling year, any provider, outside of a THSteps medical checkup when medically necessary. This screening should only be completed for a diagnosis of suspected developmental delay or to evaluate a change in the client’s developmental status outside of a THSteps medical checkup.

Developmental screening should be used to identify clients who are birth through 6 years of age and who may need a more comprehensive evaluation. Results of developmental screening may guide or identify the need for further testing. Clients with abnormal screening results must be referred to an appropriate
provider for further testing. Clients who are birth through 35 months of age who have suspected developmental delay must be referred to Texas Early Childhood Intervention (ECI) within 7 days after the child has been identified.

Refer to: Subsection 2.8, “Early Childhood Intervention (ECI) Services” in the Children’s Services Handbook (Vol. 2, Provider Handbooks) for additional information on the Texas ECI program.

Subsection 5.3.11.1.2, “Developmental Surveillance or Screening” in the Children’s Services Handbook (Vol. 2, Provider Handbooks) for additional information on developmental screening for THSteps checkups.

9.2.24.2 Developmental Testing

Developmental testing (procedure codes 96112 and 96113) is a benefit of Texas Medicaid for clients who are birth through 20 years of age.

Developmental testing consists of an extended evaluation and requires the use of a standardized norm-referenced tool. Developmental testing is medically necessary when there is suspected developmental delay supported by clinical evidence. Developmental testing is only medically indicated when clinical evidence suggests the following:

- Suspected developmental delay or atypical development when the diagnosis cannot be clearly identified through clinical interview or standardized screening tool alone.
- Retesting of a client to evaluate a change in developmental status that results in a change of treatment plan.

Procedure codes 96112 and 96113 are limited to two services per rolling year, same provider.

Developmental testing performed when a development delay or a change in the client’s developmental status is not suspected is not a benefit of Texas Medicaid.

Developmental testing is not a benefit when completed for the purposes of entering day care, Head Start, or a school setting.

Providers cannot bill the client for developmental testing that better fits the description of developmental screening.

The physician must maintain documentation of medical necessity in the client’s medical record. Retrospective review may be performed to ensure that the documentation supports the medical necessity of the service. The following information is required at least every six months to establish medical necessity:

- The physician’s prescription that includes a description of the specific service being prescribed
- The treatment plan that includes a copy of the current evaluation and documented age of the child at the time of the evaluation

9.2.24.3 Assessment of Aphasia

Aphasia assessment (procedure code 96105) is a benefit of Texas Medicaid when medically necessary and is limited to the following diagnosis codes:

<table>
<thead>
<tr>
<th>Diagnosis codes</th>
</tr>
</thead>
<tbody>
<tr>
<td>R4701</td>
</tr>
</tbody>
</table>

Procedure code 96105 is limited to two services per rolling year, same provider.
### 9.2.24.4 12-Hour Limitation for Procedure Codes 96110, 96112, and 96113

APRNs, PAs, and psychologists are limited to a maximum, combined total of 12 hours per day for developmental screening and testing, and inpatient and outpatient mental health services.

Because physicians (M.D. and D.O.) can delegate and may submit claims for services in excess of 12 hours per day, they are not subject to the 12-hour system limitation.

Developmental screening and testing are included in the 12-hour per day, per provider, system limitation. The following table lists the procedure codes that are included in the 12-hour per day system limitation, along with the time increments the system will apply based on the billed procedure code. The time increments applied will be used to calculate the 12-hour per day system limitation.

<table>
<thead>
<tr>
<th>Procedure Code</th>
<th>Time Applied by System</th>
</tr>
</thead>
<tbody>
<tr>
<td>96110</td>
<td>30 Minutes</td>
</tr>
<tr>
<td>96112</td>
<td>60 Minutes</td>
</tr>
<tr>
<td>96113</td>
<td>30 minutes</td>
</tr>
</tbody>
</table>

**Refer to:** Subsection 4.5, “Twelve Hour System Limitation” in the Behavioral Health and Case Management Services Handbook (Vol. 2, Provider Handbooks) for more information about procedure codes included in the 12-hour system limitation.

All providers, including physicians and all providers to whom they delegate services, are subject to retrospective review. HHSC and TMHP routinely perform retrospective reviews of all providers. All providers are subject to retrospective review for the total hours of services performed and billed in excess of 12 hours per day. Retrospective review may include:

- All E/M procedure codes, including those listed in the Evaluation and Management Section of the CPT Manual, billed with a diagnosis listed in the diagnosis table above under Neurobehavioral Testing
- All developmental screening and testing procedure codes included in the 12-hour system limitation

**Note:** Developmental screening and testing procedure codes and mental health procedure codes are included in the review. If a provider provides developmental and neurological assessment and testing at more than one location, any of these services may be retrospectively reviewed.

### 9.2.25 Diagnostic Tests

#### 9.2.25.1 Blood Pressure Monitoring

Blood pressure monitoring by either self-measured blood pressure monitoring or ambulatory blood pressure monitoring is a benefit of Texas Medicaid when used as a diagnostic tool to assist a physician in diagnosing hypertension in individuals whose blood pressure is either elevated, or inconclusive when evaluated in the office alone.

Self-measured blood pressure monitoring and ambulatory blood pressure monitoring may also be used for the following:

- Clients who are under treatment for established hypertension
- Evaluating refractory or treatment-resistant blood pressure
- Evaluating symptoms such as light-headedness corresponding with blood pressure changes
- Evaluating nighttime blood pressure
- Examining diurnal patterns of blood pressure
Self-measured blood pressure monitoring and ambulatory blood pressure monitoring are indicated for the evaluation of one of the following conditions:

- White coat hypertension, which is defined as the following:
  - Blood pressure measurements taken in the clinic or office are greater than 140/90 mm Hg on at least three separate visits, with two separate measurements made at each visit.
  - At least two separately documented blood pressure measurements taken outside of the clinic or office that are less than 140/90 mm Hg.
  - There is no evidence of end-organ damage.
- Resistant hypertension
- Hypotensive symptoms as a response to hypertension medications
- Nocturnal angina
- Episodic hypertension
- Syncope

Self-measured blood pressure monitoring and ambulatory blood pressure monitoring are indicated for initial diagnosis of hypertension and should not be used for maintenance monitoring.

Self-measured blood pressure monitoring may also be indicated for re-evaluation of clients previously diagnosed with hypertension.

Providers must document that the self-measured blood pressure monitoring was performed for at least 24 hours.

Procedure code 99473 is limited to one service per year, any provider. Procedure code 99473 may be considered for reimbursement more than once per year when the following documentation of medical necessity is submitted with the claim:

- Documentation of erroneous blood pressure readings—excessively high or low blood pressure, blood pressure readings excessively inconsistent with those measured professionally
- Documentation of erroneous blood pressure logs—day of the week, time of day, setting or location, or timing of medication administration inconsistent with prior professional instruction
- Documentation of poor health literacy, developmental, or intellectual challenges that may require repeated client education
- Client purchase or receipt of new blood pressure device

Procedure code 99474 is limited to four services per year, any provider, and may be reimbursed only if a claim for procedure code 99473 has been submitted within 12 rolling months.

Only one method of blood pressure monitoring (self-measured or ambulatory) may be reimbursed within a rolling 12-month period. Self-measured blood pressure monitoring submitted within the same rolling 12-month period as ambulatory blood pressure monitoring will be denied.

Use procedure codes 93784, 93786, 93788, and/or 93790 to bill in 24-hour increments for ambulatory blood pressure monitoring. Ambulatory blood pressure monitoring is limited to two services per lifetime, any provider. Ambulatory blood pressure monitoring performed more than twice per lifetime may be considered when documentation of medical necessity is submitted with the claim.

**9.2.25.2 Ambulatory and Long-Term Electroencephalogram (Ambulatory EEG)**

Ambulatory EEG monitoring is a covered benefit for clients in whom a seizure diathesis is suspected but not defined by history, physical, and resting EEG.
The EEG technical component procedure codes are limited to 3 studies for each physician for the same client per 6 months when medically necessary.

The following procedure codes should be submitted when billing for the EEG technical component:

<table>
<thead>
<tr>
<th>Procedure Codes</th>
</tr>
</thead>
<tbody>
<tr>
<td>95705</td>
</tr>
<tr>
<td>95715</td>
</tr>
</tbody>
</table>

Procedure code 95700 will be limited to three units per six months for each physician for the same client. Professional component procedure codes are limited to three studies per six months for each physician for the same client, when medically necessary.

Technical component procedure codes are limited to three studies per six months for each physician for the same client, when medically necessary.

Note: A study includes one unit of procedure code 95700 (set-up, education, and takedown) and any appropriate combination of the corresponding technical and professional procedure codes.

The following procedure codes should be submitted when billing for the EEG professional component:

<table>
<thead>
<tr>
<th>Procedure Codes</th>
</tr>
</thead>
<tbody>
<tr>
<td>95717</td>
</tr>
</tbody>
</table>

The procedure codes in the tables above may be reimbursed when they are submitted with the following diagnosis codes:

<table>
<thead>
<tr>
<th>Diagnosis Codes</th>
</tr>
</thead>
<tbody>
<tr>
<td>G0009</td>
</tr>
<tr>
<td>G0209</td>
</tr>
<tr>
<td>G0409</td>
</tr>
<tr>
<td>G0484</td>
</tr>
<tr>
<td>G0491</td>
</tr>
<tr>
<td>G0912</td>
</tr>
<tr>
<td>R410</td>
</tr>
<tr>
<td>Z052</td>
</tr>
</tbody>
</table>

Other diagnosis codes may be considered on appeal with supporting medical documentation to the TMHP Medical Director.

### 9.2.25.3 Bone Marrow Aspiration, Biopsy

Physicians may bill procedure code 85097 if interpretation is for smear interpretation, or procedure code 88305 if interpretation is for preparation and interpretation of cell block. If both procedure codes 85097 and 88305 are billed, procedure code 88305 is paid and procedure code 85097 is denied.

Physicians may bill procedure code 85097 or 88305 for preparation and interpretation of the specimen.

### 9.2.25.4 Cytopathology Studies—Other Than Gynecological

Procurement and handling of the specimen for cytopathology of sites other than vaginal, cervical, or uterine is considered part of the client’s E/M and will not be reimbursed separately.
Procedure codes 88160, 88161, and 88162 are reimbursed according to the POS where the cytopathology smear is interpreted.

Procedure code 88177 is limited to three services per day by the same provider.

### 9.2.25.5 Echoencephalography

Echoencephalography (procedure code 76506) is medically indicated for the following conditions or diagnosis codes:

<table>
<thead>
<tr>
<th>Diagnosis Codes</th>
</tr>
</thead>
<tbody>
<tr>
<td>A066  A170  A171  A1781  A1782  A1789  C410  C6961</td>
</tr>
<tr>
<td>C6962  C700  C710  C711  C712  C713  C714  C715</td>
</tr>
<tr>
<td>C716  C717  C718  C719  C7221  C7222  C7231  C7232</td>
</tr>
<tr>
<td>C7241  C7242  C7259  C729  C751  C752  C768  C7931</td>
</tr>
<tr>
<td>C7932  C7940  C7949  C7951  C7952  C7989  D075  D098</td>
</tr>
<tr>
<td>D164  D3161  D3162  D320  D329  D330  D331  D332</td>
</tr>
<tr>
<td>D333  D3500  D3501  D3502  D420  D421  D429  D432</td>
</tr>
<tr>
<td>D433  D434  D438  D439  D47Z1  D47Z2  D480  D487</td>
</tr>
<tr>
<td>D492  D496  D497  F0390  G060  G062  G07  G08</td>
</tr>
<tr>
<td>G132  G138  G232  G300  G301  G308  G309  G3101</td>
</tr>
<tr>
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</tr>
<tr>
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</tr>
<tr>
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<tr>
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</tr>
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</tr>
<tr>
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<td>I6010  I6011  I6012  I602  I6030  I6031  I6032  I604</td>
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<tr>
<td>I611  I612  I613  I614  I615  I616  I618  I619</td>
</tr>
<tr>
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</tr>
<tr>
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</tr>
<tr>
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<td>I63422  I63429  I63431  I63432  I63439  I6349  I6350  I63511</td>
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</tr>
<tr>
<td>I63532  I63533  I63539  I63543  I6381  I6389  I6601  I6602</td>
</tr>
<tr>
<td>Diagnosis Codes</td>
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<tr>
<td>I6603</td>
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<tr>
<td>I6623</td>
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<tr>
<td>I67850</td>
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<tr>
<td>I69212</td>
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<tr>
<td>Q283</td>
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<tr>
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<td>S060X9S</td>
</tr>
<tr>
<td>S061X2D</td>
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<td>S061X5A</td>
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<tr>
<td>S06357A</td>
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<tr>
<td>S06361A</td>
</tr>
</tbody>
</table>
9.2.25.6 Electrocardiogram (ECG)

Electrocardiograms (ECG) are a benefit of Texas Medicaid when used for the evaluation and management (E/M) of a confirmed or suspected primary disease of the heart, pericardium, and coronary arteries or when necessary for management of diseases that are not primarily cardiac, but can affect the heart directly or indirectly.

ECGs are limited to six treatments for each client, by any provider per benefit period.

For ECGs, a benefit period is defined as 12 consecutive months, beginning with the month the client receives the first ECG.

The following procedure codes may be reimbursed for ECGs: 93000, 93005, 93010, 93040, 93041, and 93042.

Claims that are denied for exceeding the six-ECG limitation may be appealed with documentation supporting medical necessity. The documentation must include the following:

- Diagnosis
- Treatment history
- Documentation of why additional ECGs are needed
The report of the professional component (the interpretation) for the ECG must be a complete written report that includes relevant findings and appropriate comparisons.

The interpretation may appear on the actual tracing.

When the ECG is performed in conjunction with the performance of an evaluation and management (E/M) service, the interpretation may appear with a progress note or other report of the E/M service; however, if the ECG is billed as a separate service from the E/M service, the interpretation should contain the same information as a report made upon the tracing itself.

A simple notation of “ECG/EKG normal” without an accompanying tracing will not suffice as documentation of a separately payable interpretation.

Appropriate documentation, which includes a copy of the ECG tracing, must be kept in the client’s medical record. Documentation must support the medical necessity of the ECG. Documentation may appear on the actual tracing or with a progress note or report. Documentation is subject to retrospective review.

Only an ECG interpretation that directly contributes to the diagnosis and treatment of a client may be considered for reimbursement. Services, such as routine admission ECGs performed without medical indications, that do not directly contribute to the diagnosis and treatment of an individual client are not considered medically necessary.

9.2.25.6.1 Prior Authorization for ECG

Prior authorization is not required for ECGs performed in the emergency room or inpatient hospital setting.

Prior authorization is required for more than six ECGs in a rolling 12-month period.

Requests for additional ECGs must be submitted on the Special Medical Prior Authorization (SMPA) Request Form along with documentation of medical necessity.

Providers may request a prior authorization up to 12 months in advance. When requesting retroactive authorization, a provider must submit the request no later than 14 calendar days after the ECG is completed.

Before submitting a prior authorization request for an ECG, a provider must have a completed SMPA Request Form that has been signed and dated by a physician who is familiar with the client. The completed SMPA Request Form must include the procedure codes and numerical quantities for the services requested. The completed SMPA Request Form with the original dated signature must be maintained by the prescribing physician in the client’s medical record.

The SMPA Request Form must include all of the following information, which is related to medical necessity:

- Procedure requested (CPT)
- Diagnosis
- Treatment history
- Treatment plan
Prior authorization requests submitted by paper, must be faxed or mailed with the completed SMPA Request Form to the SMPA department and a copy of the signed and dated form must be retained in the client’s medical record at the provider’s place of business. Requests may be faxed or mailed to the following address:

Texas Medicaid & Healthcare Partnership
Special Medical Prior Authorization
12357-B Riata Trace Parkway
Austin, TX 78727
Fax: 1-512-514-4213

Requests for prior authorization can also be submitted online through the TMHP website at www.tmhp.com.

9.2.25.7 Esophageal pH Probe Monitoring

Esophageal pH monitoring uses an indwelling pH microelectrode positioned just above the esophageal sphincter. The pH electrode and skin reference electrode are connected to a battery-powered pH meter and transmitter worn as a shoulder harness. The esophageal pH is monitored continuously and a strip chart is used to record the pH determinations. The patient is usually monitored for a 24-hour period. Esophageal pH monitoring is a medically appropriate adjunct procedure to help establish the presence or absence of gastroesophageal reflux.

Esophageal pH probe monitoring should be coded with procedure codes 91034, 91035, and 78262. Esophageal pH probe testing (procedure codes 78262, 91034, and 91035) are limited to two services per rolling year, same procedure, any provider.

Claims that are denied for exceeding two services per rolling year may be considered on appeal with documentation of one of the following:

- The client is new and the provider has been unsuccessful in obtaining the client’s previous records from a different provider.
- The provider is not aware that the client received previous esophageal testing.

Only one appeal will be considered per client, for the same provider. Providers must request prior authorization for any additional esophageal testing performed after the appealed service.

9.2.25.7.1 Prior Authorization

Esophageal pH probe testing (procedure codes 78262, 91034, and 91035) require prior authorization for services that exceed two per rolling year.

Requests for additional testing may be considered when submitted with documentation of medical necessity that supports, but is not limited to, the following:

- Adult’s unintentional weight loss is more than 5 percent of their normal body weight in a span of 12 months or less
- Child’s weight loss is 3 to 5 percent of their body mass in less than 30 days
- Symptoms of gastroesophageal reflux disease (GERD) that include heartburn and regurgitation that do not respond to treatment with medication
- Atypical symptoms of GERD, such as chest pain, coughing, wheezing, hoarseness, and sore throat

Prior authorization requests must be submitted to the Special Medical Prior Authorization Department using the Special Medical Prior Authorization (SMPA) Request Form. The completed prior authorization request form must be maintained by the requesting provider and the prescribing physician. The original, signed copy must be kept by the physician in the client’s medical record.
9.2.25.8 Helicobacter Pylori (H. pylori)

Initial testing for H. pylori may be performed using the following tests:

- Serology testing (procedure codes 83009 and 86677)
- Stool testing (procedure code 87338 with modifier QW)
- Breath testing (procedure codes 78267, 78268, 83013, and 83014)

Serology testing for H. pylori is a noninvasive diagnostic procedure that is preferred for initial diagnosis but is not indicated after a diagnosis has been made. Serology testing is not indicated or covered for monitoring a response to therapy.

Procedure codes 83009 and 86677 are allowed once per lifetime when submitted by any provider. A second test may be considered on appeal with documentation that indicates the original test result was negative for H. pylori.

Urea breath tests (UBTs) and fecal antigen tests provide reliable means of identifying active H. pylori infection before antibiotic therapy. UBTs are the most reliable non-endoscopic test to document eradication of H. pylori infection.

H. pylori is accepted as an etiologic factor in duodenal ulcers, peptic ulcer disease, gastric carcinoma, and primary B cell gastric lymphoma. H. pylori testing may be indicated for symptomatic clients who have a documented history of chronic/recurrent duodenal ulcer, gastric ulcer, or chronic gastritis. The history must delineate the failed conservative treatment for the condition.

H. pylori testing is not indicated or covered for any of the following:

- New onset uncomplicated dyspepsia.
- New onset dyspepsia responsive to conservative treatment (e.g., withdrawal of nonsteroidal anti-inflammatory drugs [NSAID] and/or use of antisecretory agents). If the treatment does not prove successful in eliminating the symptoms, further testing may be indicated to determine the presence of H. pylori.
- Screening for H. pylori in asymptomatic clients.
- Dyspeptic clients requiring endoscopy and biopsy.

H. pylori testing is not indicated under the following circumstances:

- There has been a negative endoscopy in the previous 90 days.
- An endoscopy is planned.
- H. pylori is of new onset and still being treated.

H. pylori testing will be denied if it is performed within 90 days of an upper gastrointestinal endoscopy. Procedure codes 87338 (with modifier QW), 78267, 78268, 83013, and 83014 may be reimbursed within the 90 days if the provider submits documentation that indicates the client was tested for eradication after treatment.

If a follow-up breath or stool test is used to document eradication of H. pylori, the medical record documentation must verify the history of the following previous complication(s):

- The client remains symptomatic after a treatment regimen for H. pylori.
- The client is asymptomatic after H. pylori eradication therapy but has a history of hemorrhage, perforation, or outlet obstruction from peptic ulcer disease.
- The client has a history of ulcer on chronic NSAID or anticoagulant therapy.
Testing for H. pylori eradication after the completion of antibiotic therapy (procedure codes 87338 [with modifier QW], 78267, 78268, 83013, and 83014) will be denied if billed less than 35 days after the initial test.

Procedure code 87339 is not a benefit of Texas Medicaid.

9.2.25.9 Myocardial Perfusion Imaging


9.2.25.10 Pediatric Pneumogram

A pediatric pneumogram (procedure code 94772) is a 12-hour to 24-hour recording of breathing effort, heart rate, oxygen level, and airflow to the lungs during sleep. The study is useful in identifying abnormal breathing patterns, with or without bradycardia, especially in premature infants.

The following diagnosis codes may be reimbursed for a pediatric pneumogram in infants from birth through 11 months of age:

<table>
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<th>Diagnosis Codes</th>
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<tr>
<td>P284</td>
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<td>R063</td>
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A pediatric pneumogram is limited to two services per lifetime without prior authorization when submitted with one of the diagnosis codes listed above. Additional studies may be considered under CCP with documentation of medical necessity, and will require prior authorization.

Referto: Section 2, “Medicaid Children’s Services Comprehensive Care Program (CCP)” in the Children’s Services Handbook (Vol. 2, Provider Handbooks).

EMGs, polysomnography, EEGs, and ECGs are denied when billed on the same day as a pediatric pneumogram.

Pediatric pneumograms are reimbursed on the same day as an apnea monitor (rented monthly) if documentation supports the medical necessity.

Pneumogram supplies are considered part of the technical component and are denied if billed separately.

9.2.26 Diagnostic Doppler Sonography

Diagnostic Doppler sonography is a benefit of Texas Medicaid when treatment decisions depend on the results. Authorization is not required for diagnostic Doppler services.

Doppler sonography uses a transducer that transmits and receives the returned sound waves as vibrations. The transducer turns the vibrations into electrical pulses that travel to the ultrasonic scanner where they are processed and transformed into a digital image. It is used to study blood flow throughout the body.

A vascular diagnostic study may be personally performed by a physician or by a technologist. The accuracy of noninvasive vascular diagnostic studies depends on the knowledge, skill, and experience of the technologist and physician performing and interpreting the study. Consequently, the physician who performs and/or interprets the study must be able to document training through recent residency training or post-graduate continuing medical education and experience and must maintain that documentation for post-payment review.
If noninvasive vascular diagnostic studies are performed by a technologist, the technologist must have demonstrated competency in ultrasound by receiving one of the following credentials in vascular ultrasound technology:

- Registered Vascular Specialist (RVS) provided by Cardiovascular Credentialing International (CCI)
- Registered Vascular Technologist (RVT) provided by the American Registry of Diagnostic Medical Sonographers (ARDMS)
- Vascular Sonographer (VS) provided by the American Registry of Radiologic Technologists (ARRT), Sonography

Alternately, such studies must be performed in a facility or vascular laboratory accredited by one of the following nationally recognized accreditation organizations. If a vascular laboratory or facility is accredited, the technologists performing noninvasive cerebrovascular arterial studies in that laboratory are considered to have demonstrated competency in cerebrovascular ultrasound:

- American College of Radiology (ACR) Vascular Ultrasound Accreditation Program
- Intersocietal Commission for the Accreditation of Vascular Laboratories (ICAVL)

### 9.2.26.1 Cerebrovascular Doppler Studies

Cerebrovascular Doppler sonography includes both extracranial and transcranial (intracranial) studies. This group of Doppler studies is used to investigate cerebral hemodynamics (e.g., blood flow, vasculitis, cerebral fluid collection/hydrocephalus, cerebral vascular disorders). Cerebrovascular Doppler sonography should not be used when treatment decisions will not be affected by the findings.

Cerebrovascular Doppler studies for the diagnosis of migraine are considered experimental and are not a benefit of Texas Medicaid.

Extracranial arterial Doppler (procedure codes 93880 and 93882) are limited to the following diagnosis codes:

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* Use R55 when symptomatology indicates a strong clinical suspicion of vertebrobasilar insufficiency
** Use R221 to report pulsatile neck mass
*** Use R0989 to report carotid bruit
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</table>

* Use R55 when symptomatology indicates a strong clinical suspicion of vertebrobasilar insufficiency
** Use R221 to report pulsatile neck mass
*** Use R0989 to report carotid bruit
Transcranial Doppler (procedure codes 93886, 93888, 93890, 93892, and 93893) are limited to the following diagnosis codes:

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* Use R55 when symptomatology indicates a strong clinical suspicion of vertebrobasilar insufficiency
** Use R221 to report pulsatile neck mass
*** Use R0989 to report carotid bruit

* Use G9389 to identify assessment of suspected brain death
** Use 1749 to report paradoxical cerebral embolism
*** Use R55 when symptomatology indicates a strong clinical suspicion of vertebrobasilar insufficiency
In addition to the diagnosis codes listed in the table above, procedure codes 93886 and 93888 are benefits for clients who are 2 through 16 years of age with sickle cell disease to evaluate the risk of stroke when submitted with the following diagnosis codes:

### Diagnosis Codes

- T82848D
- T82848S
- T82858A
- T82858D
- T82858S
- T82868A
- T82868D
- T82868S
- Z09

* Use G9389 to identify assessment of suspected brain death
** Use I749 to report paradoxical cerebral embolism
*** Use R55 when symptomatology indicates a strong clinical suspicion of vertebrobasilar insufficiency

9.2.26.2 Peripheral Doppler Studies

Peripheral Doppler sonography is used to determine vascular impedance and evaluate peripheral masses and peripheral nerve continuity.

9.2.26.3 Peripheral Arterial Doppler Studies

Noninvasive peripheral arterial examinations that are performed to establish the level and degree of arterial occlusive disease are reasonable and necessary if significant signs or symptoms of possible limbischemia are present, and the client is a candidate for invasive therapeutic procedures.

Peripheral arterial Doppler (procedure codes 93922, 93923, 93924, 93925, 93926, 93930, and 93931) are limited to the following diagnosis codes (unless otherwise indicated):

### Diagnosis Codes

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MEDICAL AND NURSING SPECIALISTS, PHYSICIANS, AND PHYSICIAN ASSISTANTS HANDBOOK  MAY 2021

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9.2.26.4 Peripheral Venous Doppler Studies

Non-invasive vascular diagnostic studies utilize ultrasonic Doppler and physiologic principles to assess irregularities in blood flow in the venous system.

Peripheral venous Doppler (procedure codes 93970 and 93971) are limited to the following diagnosis codes:

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In addition to the diagnosis codes listed in the table above, procedure code 93971 is also a benefit when submitted with diagnosis code Z01810, Z01818, or Z09.

Doppler echocardiography color flow velocity mapping (procedure code 93325) must be billed with one of the corresponding procedure codes in column B to be considered for reimbursement:

<table>
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<th>Column A Procedure Code</th>
<th>Column B Procedure Codes</th>
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9.2.26.5 **Limitations for Diagnostic Doppler Sonography**

Documentation of medical necessity for the diagnostic Doppler study must be maintained by the ordering provider in the client’s medical record.

Procedure codes described as complete bilateral studies are inclusive codes, and right and left studies billed on the same day will be reimbursed at a quantity of one.

Diagnostic Doppler procedure codes are limited to one study per day, same provider.

When medically necessary, multiple Doppler procedures (e.g., studies of extracranial arteries and intracranial arteries) billed on the same day by the same provider will be reimbursed at full fee for the first study and one-half fee for each additional study, regardless of the number of services billed.

The use of transcranial Doppler studies performed for the assessment of stroke risk in clients who are 2 through 16 years of age who have sickle cell anemia should be limited to once every 6 months.

The use of a simple hand-held or other Doppler device that does not produce hard copy output or that does not permit analysis of bidirectional vascular flow is considered part of the physical examination of the vascular system and is not separately reported.

9.2.27 **Evoked Response Tests and Neuromuscular Procedures**

The following services are a benefit of Texas Medicaid:

- Autonomic function test (AFT)
- Electromyography (EMG)
- Nerve conduction studies (NCS)
- Evoked potential (EP) testing
- Motion analysis studies

9.2.27.1 **Autonomic Function Tests**

AFTs are a benefit of Texas Medicaid when submitted with procedure codes 95921, 95922, 95923, 95924, and 95943.

Procedure codes 95921, 95922, 95923, 95924, and 95943 are limited to once per date of service, by the same provider.

Autonomic disorders may be congenital or acquired (primary or secondary). Some of the conditions under which autonomic function testing may be appropriate include, but are not limited to, the following:

- Amyloid neuropathy
- Diabetic autonomic neuropathy
- Distal small fiber neuropathy
- Excessive sweating
- Gastrointestinal dysfunction
- Idiopathic neuropathy
- Irregular heart rate
- Multiple system atrophy
- Orthostatic symptoms
- Pure autonomic failure
- Reflex sympathetic dystrophy or causalgia (sympathetically maintained pain)
• Sjogren’s syndrome

The reason for the referral, the specific autonomic function being tested, and a clear diagnostic impression must be documented in the client’s medical record for each AFT performed.

The client’s medical records must clearly document the medical necessity for the AFT. The medical record documentation must reflect the actual results of specific tests (such as latency and amplitude).

Medical necessity for reevaluation of a client (beyond the initial consultation and testing) must be clearly documented in the client’s medical record. Supporting documentation includes, but is not limited to, the following:

• The client has new symptoms unrelated to those previously evaluated, suggestive of a new diagnosis.
• Evidence that the client’s condition is changing rapidly, supported by the following:
  • Diagnosis
  • Current clinical signs and symptoms
  • Prior clinical condition
  • Expected clinical disease course
• Clinical benefit of additional studies.

The client’s medical records are subject to retrospective review. Wave form recordings obtained during the testing will aid documentation requirements in cases where a review becomes necessary.

9.2.27.2 Electromyography and Nerve Conduction Studies

Electromyography (EMG) and nerve conduction studies (NCS), collectively known as electrodiagnostic (EDX) testing, must be medically indicated and may be reimbursed with the diagnosis codes listed below. Testing must be performed using EDX equipment that provides assessment of all parameters of the recorded signals. Studies performed with devices designed only for screening purposes rather than diagnoses are not a benefit of Texas Medicaid.

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Any EDX testing procedures may be reimbursed up to four different dates of service per calendar year, same provider. Any E/M service will be denied as part of another service when billed for the same date of service as EMG or NCS service by the same provider.

Claims for nerve conduction studies that are denied for exceeding the maximum number of studies allowed per day, may be appealed with supporting medical record documentation.

The reason for the referral, the specific site(s) tested, and a clear diagnostic impression must be documented in the client’s medical record for each NCS or EMG study performed.

The client’s medical records must clearly document the medical necessity for the NCS and EMG testing. The medical record documentation must reflect the actual results of specific tests (such as latency and amplitude).

Medical necessity for re-evaluation of a client (beyond the initial consultation and testing) must be clearly documented in the client’s medical record. Supporting documentation includes, but is not limited to, the following:

- The client has new symptoms unrelated to those previously evaluated, suggestive of a new diagnosis. Examples may include suspected:
  - Peripheral nerve entrapment syndromes
  - Other neuropathies (traumatic, metabolic, or demyelinating)
  - Neuromuscular junction disorders (myasthenia gravis, botulism)
• Myopathies (dermatomyositis, congenital myopathies)
• Unexplained symptoms suggestive of peripheral nerve, muscle or neuromuscular junction pathology, manifested by muscle weakness, muscle atrophy, loss of dexterity, spasticity, sensory deficits, swallowing dysfunction, diplopia, or dysarthria
• The client’s diagnosis could not be confirmed on previous studies, although suspected.
• Evidence exists that the client’s condition is changing rapidly, supported by the following:
  • Diagnosis
  • Current clinical signs and symptoms
  • Prior clinical condition
  • Expected clinical disease course
• There is clinical benefit of additional electrodiagnostic studies.

The client’s medical records are subject to retrospective review. NCS hard copies of the waveform recordings obtained during the testing will aid documentation requirements in cases where a review becomes necessary.

9.2.27.2.1 EMG

The following EMG procedure codes may be reimbursed for one service per day, each procedure, by the same provider:

<table>
<thead>
<tr>
<th>Procedure Codes</th>
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<tbody>
<tr>
<td>01784 51785 95860 95861 95863 95864 95865 95867 95868 95869</td>
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<tr>
<td>95872 95875</td>
</tr>
</tbody>
</table>

Procedure code 95866 may be reimbursed up to two services per day, same provider. Procedure code 95870 may be reimbursed in multiple quantities if specific muscles are documented.

The needle EMG examination must be performed by a physician specially trained in electrodiagnostic medicine, as these tests are simultaneously performed and interpreted.

Surface or macro-EMG testing is considered experimental and is not a benefit of the Texas Medicaid.

9.2.27.2.2 NCS

NCS are reimbursed by Texas Medicaid with documentation of medical necessity using the following procedure codes:

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<th>Procedure Codes</th>
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<td>95885 95886 95887 95905 95907 95908 95909 95910 95911 95912</td>
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<tr>
<td>95913 95933 95937</td>
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</tbody>
</table>

NCS must be performed by one of the following:
• A physician
• A trained individual under the direct supervision of a physician. (Direct supervision means that the physician is in close physical proximity to the electrodiagnostic laboratory while testing is underway, immediately available to provide the trained individual with assistance and direction, and responsible for selecting the appropriate NCS to be performed.)
When the same studies are performed on unique sites by the same provider for the same date of service, studies for the first site must be billed without a modifier and studies for each additional site must be billed with modifier XE, XP, XS, or XU, indicating a distinct procedural service. Modifier 59 should be used when modifier XE, XP, XS, or XU is not appropriate.

Procedure codes 95907, 95908, 95909, 95910, 95911, 95912, and 95913 may be reimbursed only once when multiple sites on the same nerve are stimulated or recorded.

Prior authorization is required when the anticipated number of nerve conduction studies planned for an evaluation exceeds the following maximum number of studies:

<table>
<thead>
<tr>
<th>Procedure Code</th>
<th>Limitation</th>
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<tbody>
<tr>
<td>95885, 95886</td>
<td>Reimbursed once per extremity up to 4 units, using any combination of procedure codes, per day, any provider.</td>
</tr>
<tr>
<td>95885, 95886, 95887</td>
<td>Must be billed with one of the primary procedure codes 95907, 95908, 95909, 95910, 95911, 95912, or 95913.</td>
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<tr>
<td>95937</td>
<td>Up to 3 studies per day, per procedure, same provider without prior authorization.</td>
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</table>

Requests for prior authorization must be submitted to the Special Medical Prior Authorization department (SMPA) using the Special Medical Prior Authorization (SMPA) Request Form.

**Note:** An advanced practice registered nurse (APRN) or a physician assistant (PA) may sign all documentation related to the provision of evoked response tests and neuromuscular procedures on behalf of the client's physician when the physician delegates this authority to the APRN or PA. The APRN or PA provider's signature and license number must appear on the forms where the physician signature and license number blocks are required.

Requests must include documentation supporting medical necessity for the number of studies requested, and they must be received on or before the requested DOS. Requests received after the services are performed will be denied for DOS that occurred before the date the request was received.

Medical record documentation must establish medical necessity for the additional studies, including one or more of the following:

- Other diagnosis in the differential that require consideration should include provider notes about both of the following:
  - The additional diagnoses considered.
  - The clinical signs, symptoms, or electrodiagnostic findings that necessitated the inclusion.
- If multiple diagnoses have been established by nerve conduction studies and the recommendations in the table above for a single diagnostic category do not apply, then the provider should document all diagnoses established as a result of EDX testing.
- Testing of an asymptomatic contralateral limb to establish normative values for an individual client (particularly the elderly, diabetic, and clients with a history of ethyl alcohol [ETOH] usage).
- Comorbid clinical conditions are identified. The clinical condition must be one that may cause sensory or motor symptoms, for example:
  - Underlying metabolic disease (such as thyroid condition or diabetes mellitus)
  - Nutritional deficiency (alcoholism)
  - Malignant disease
  - Inflammatory disorder (including but not limited to lupus, sarcoidosis or Sjögren’s syndrome)
9.2.27.3 Evoked Potential Testing

Evoked potential (EP) tests are a benefit of Texas Medicaid when medically necessary. The most common EP tests are:

- Brainstem auditory evoked potentials (BAEPs)
- Motor evoked potentials (MEPs)
- Somatosensory evoked potentials (SEPs)
- Visual evoked potentials (VEPs)

Each EP test (procedure codes 92650, 92651, 92652, 92653, 95925, 95926, 95927, 95928, 95929, 95930, 95938, or 95939) is considered a bilateral procedure and is limited to once per date of service any provider regardless of modifiers that indicate multiple sites were tested.

EP tests may be reimbursed up to four services per rolling year, any combination of services by any provider. Claims that exceed the limitation of four services per rolling year may be considered for reimbursement on appeal with documentation that supports the medical necessity.

Intraoperative neurophysiology monitoring (procedure codes 95940 and 95941) is a benefit when performed in addition to each evoked potential test on the same day.

The documentation for the intraoperative neurophysiology monitoring must include the time for which each test is performed.

Procedure codes 95940 and 95941 are limited to a maximum of two hours per date of service, per client, same procedure, any provider.

Procedure codes 95940 and 95941 must be billed in conjunction with one of the following procedure codes or the service will be denied:

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<tr>
<th>Procedure Codes</th>
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<tbody>
<tr>
<td>95822 95860 95861 95863 95864 95865 95866 95867 95868 95869</td>
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Procedure codes 95940 and 95941 cannot be reported by the surgeon or anesthesiologist.

The reason for the referral, the specific nerve evoked potential being tested, and a clear diagnostic impression must be documented in the client’s medical record for each EP study performed.

The client’s medical records must clearly document the medical necessity for the EP testing. The medical record documentation must reflect the actual results of specific tests (such as latency and amplitude).

Medical necessity for re-evaluation of a client (beyond the initial consultation and testing) must be clearly documented in the client’s medical record. Supporting documentation includes, but is not limited to, the following:

- The client has new symptoms unrelated to those previously evaluated, suggestive of a new diagnosis.
- Evidence exists that the client’s condition is changing rapidly, supported by the following:
  - Diagnosis
  - Current clinical signs and symptoms
  - Prior clinical condition
  - Expected clinical disease course
- There is clinical benefit of additional studies.
The client’s medical records are subject to retrospective review. Wave form recordings obtained during
the testing will aid documentation requirements in cases where a review becomes necessary.

9.2.27.3.1 Visual Evoked Potentials

Some of the conditions under which VEP testing (procedure code 95930) may be appropriate include,
but are not limited to, the following:

- Identification of persons at increased risk for developing clinically definite multiple sclerosis.
- Diagnosing, monitoring, and assessing treatment response in multiple sclerosis.
- Localizing the cause of a visual field defect not explained by lesions seen on CT or MRI, or by
  metabolic disorders or infectious disease.
- Evaluating the signs and symptoms of visual loss in persons who are unable to communicate (e.g.,
  unresponsive persons, non-verbal persons).
- Evaluating clients who experience double vision, blurred vision, loss of vision, eye injuries, head
  injuries, or weakness of the eyes, arms, or legs.

9.2.27.4 Motion Analysis Studies

Motion analysis studies (procedure codes 96000, 96001, 96002, and 96003) are a benefit of Texas
Medicaid for clients who are 3 through 20 years of age.

Procedure codes 96000, 96001, 96002, and 96003 are limited to one per date of service by the same
provider and two per rolling year, any provider.

In the following table, the procedure codes in Column A will be denied when they are submitted on the
same date of service by the same provider as the procedure codes in Column B:

<table>
<thead>
<tr>
<th>Column A (Denied)</th>
<th>Column B</th>
</tr>
</thead>
<tbody>
<tr>
<td>96000</td>
<td>96001</td>
</tr>
<tr>
<td>95860, 95861, 95863, 95864, 95865, 95866, 95869, 95870, 95872</td>
<td>96002 or 96003</td>
</tr>
</tbody>
</table>

Documentation must include the following information that indicates the client meets all the require-
ments for motion analysis studies. The client must be:

- Ambulatory for a minimum of ten consecutive steps, with or without assistive devices.
- At least three years of age.
- Physically able to tolerate up to three hours of testing.

The reason for the referral and a clear diagnostic impression must be documented in the client's medical
record for each motion analysis study performed.

The client’s medical records must clearly document the medical necessity for the motion analysis study.
The medical record documentation must reflect the actual results of specific tests.

Medical necessity for re-evaluation of a client (beyond the initial consultation and testing) must be
clearly documented in the client’s medical record. Supporting documentation includes, but is not
limited to, the following:

- The client has new symptoms unrelated to those previously evaluated, suggestive of a new diagnosis.
- Evidence exists that the client’s condition is changing rapidly, supported by the following:
  - Diagnosis
  - Current clinical signs and symptoms
• Prior clinical condition
• Expected clinical disease course
• There is clinical benefit of additional studies.

The client’s medical records are subject to retrospective review.

9.2.28 Extracorporeal Membrane Oxygenation (ECMO)

ECMO may be effective on a short-term basis for clients with life-threatening respiratory and/or cardiac insufficiency.

ECMO may be reimbursed for clients who have the following clinical indications (this is not an all-inclusive list):

• Persistent pulmonary hypertension
• Meconium aspiration syndrome
• Respiratory distress syndrome
• Adult respiratory distress syndrome
• Congenital diaphragmatic hernia
• Sepsis
• Pneumonia
• Preoperative and postoperative congenital heart disease or heart transplantation
• Reversible causes of cardiac failure
• Cardiomyopathy
• Myocarditis
• Aspiration pneumonia
• Pulmonary contusion
• Pulmonary embolism

The following procedure codes may be used when billing ECMO:

<table>
<thead>
<tr>
<th>Procedure Codes</th>
</tr>
</thead>
<tbody>
<tr>
<td>33946 33947 33948 33949 33951 33952 33953 33954 33955 33956</td>
</tr>
<tr>
<td>33957 33958 33959 33962 33963 33964 33965 33966 33969 33984</td>
</tr>
<tr>
<td>33985 33986 33987 33988 33989</td>
</tr>
</tbody>
</table>

Terminal disease with expectation of short survival, advanced multiple organ failure syndrome, irreversible central nervous system injury and severe immunosuppression are contraindications to ECMO. Claims for ECMO services may be recouped if the services are provided in the presence of these conditions.

The initial 24 hours of veno-venous (VV) ECMO should be submitted using procedure code 33946. Procedure code 33948 should be used for each additional 24 hours. Procedure code 33946 is denied as part of procedure code 33948 if submitted with the same date of service. Procedure codes 33946 and 33948 are limited to one per day when billed by any provider.
The initial 24 hours of veno-arterial (VA) ECMO should be submitted using procedure code 33947. Procedure code 33949 should be used for each additional 24 hours. Procedure code 33947 is denied as part of procedure code 33949 if submitted with the same date of service. Procedure codes 33947 and 33949 are limited to one per day when billed by any provider.

If insertion of VV cannula (procedure codes 33951, 33952, 33953, 33954, 33955, and 33956) for prolonged extracorporeal circulation for cardiopulmonary insufficiency is submitted by the same provider with the same date of service as procedure code 33946 or 33948, the insertion of the cannula is denied, and the ECMO (procedure code 33946 or 33948) is considered for reimbursement.

If insertion of VA cannula (procedure codes 33951, 33952, 33953, 33954, 33955, and 33956) for prolonged extracorporeal circulation for cardiopulmonary insufficiency is submitted by the same provider with the same date of service as procedure code 33947 or 33949, the insertion of the cannula is denied, and the ECMO (procedure code 33947 and 33949) is considered for reimbursement.

9.2.29 Family Planning

Physicians, PAs, NPs, CNSs, and CNMs are encouraged to provide family planning services to Texas Medicaid clients, especially pregnant and postpartum clients. No separate enrollment is required. Providers are reimbursed for family planning services through Texas Medicaid (Title XIX) or through the DSHS Family Planning Program.

Refer to:
Section 2, “Medicaid Title XIX Family Planning Services” in the Gynecological, Obstetrics, and Family Planning Title XIX Services Handbook (Vol. 2, Provider Handbooks).

9.2.30 Gynecological Health Services

Gynecological examinations, surgical procedures, and treatments are benefits of Texas Medicaid.

Refer to: Section 6, “Gynecological Health Services” in the Gynecological, Obstetrics, and Family Planning Title XIX Services Handbook (Vol. 2, Provider Handbooks) for information about contraception, sterilizations, and family planning annual examinations.

9.2.31 Hospital Visits

Refer to: Subsection 9.2.56, “Physician Evaluation and Management (E/M) Services” in this handbook.

9.2.32 Hyperbaric Oxygen Therapy (HBOT)

Physicians who bill for the professional component of HBOT must use procedure code 99183. Hospital providers who bill for the chamber time must use procedure code G0277 with revenue code 413.

Note: Although oxygen may be administered by mask, cannula, or tube in addition to the hyperbaric treatment, the use of oxygen by mask, or other device, or applied topically is not considered hyperbaric treatment in itself.

Texas Medicaid recognizes the following indications for HBOT, as approved by the Undersea and Hyperbaric Medical Society (UHMS):

- Air or gas embolism
- Carbon monoxide poisoning
- Central retinal artery occlusion
- Compromised skin grafts and flaps
• Crush injuries, compartment syndrome, and other acute traumatic ischemias
• Decompression sickness
• Delayed radiation injury (soft tissue and bony necrosis)
• Diabetic foot ulcer
• Severe anemia
• Clostridial myositis and myonecrosis (gas gangrene)
• Intracranial abscess
• Necrotizing soft tissue infections
• Refractory osteomyelitis
• Acute thermal burn injuries

HBOT is not a replacement for other standard successful therapeutic measures.

Texas Medicaid considers HBOT experimental and investigational for any indications other than the ones approved by UHMS and outlined in this section. Non-covered indications include, but are not limited to, autism and traumatic brain injury.

Oxygen administered outside of a hyperbaric chamber, by any means, is not considered hyperbaric treatment.

The physician must be in constant attendance of hyperbaric oxygen therapy during compression and decompression of the chamber and may not delegate the rendering of the service. Both the facility’s medical record and the client’s medical record must contain documentation to support that there was a physician in attendance who provided direct supervision of the compression and decompression phases of the HBOT treatment. All documentation pertaining to HBOT is subject to retrospective review.

9.2.32.1 Prior Authorization for HBOT

HBOT procedure codes 99183 and G0277 require prior authorization. Prior authorization requests submitted for procedure code G0277 must also include revenue code 413. When requesting prior authorization, providers should use the Special Medical Prior Authorization (SMPA) Request Form on the TMHP website at www.tmhp.com.

Refer to: “Section 5: Fee-for-Service Prior Authorizations” (Vol. 1, General Information) for detailed information about prior authorization requirements.

The prior authorization request must include documentation that supports medical necessity and is specific to each appropriate covered indication as listed in the following table:

<table>
<thead>
<tr>
<th>Covered Indication</th>
<th>Total 30-Minute Intervals Allowed for Procedure Code G0277</th>
<th>Total Professional Sessions Allowed for Procedure Code 99183</th>
<th>Medical Necessity Documentation of the Following is Required</th>
</tr>
</thead>
<tbody>
<tr>
<td>Air or gas embolism</td>
<td>6</td>
<td>2</td>
<td>Evidence that gas bubbles are detectable by ultrasound, Doppler or other diagnostics</td>
</tr>
</tbody>
</table>

Note: The following Wagner wound classification grades apply only to the diabetic foot ulcer indications:
Grade 1: Superficial diabetic ulcer
Grade 2: Ulcer extension - involves ligament, tendon, joint capsule or fascia (No abscess or osteomyelitis)
Grade 3: Deep ulcer with abscess or osteomyelitis
Grade 4: Gangrene to portion of forefoot
Grade 5: Extensive gangrene of foot
<table>
<thead>
<tr>
<th>Covered Indication</th>
<th>Total 30-Minute Intervals Allowed for Procedure Code G0277</th>
<th>Total Professional Sessions Allowed for Procedure Code 99183</th>
<th>Medical Necessity Documentation of the Following is Required</th>
</tr>
</thead>
<tbody>
<tr>
<td>Carbon monoxide poisoning - initial authorization</td>
<td>15</td>
<td>5</td>
<td>Persistent neurological dysfunction secondary to carbon monoxide inhalation</td>
</tr>
<tr>
<td>Carbon monoxide poisoning - one subsequent authorization</td>
<td>9</td>
<td>3</td>
<td>Evidence of continuing improvement in cognitive functioning</td>
</tr>
<tr>
<td>Central retinal artery occlusion</td>
<td>36</td>
<td>6</td>
<td>Evidence of central retinal artery occlusion with treatment initiated within 24 hours of the occlusion</td>
</tr>
<tr>
<td>Compromised skin grafts and flaps - initial authorization</td>
<td>80</td>
<td>10</td>
<td>Evidence the flap or graft is failing because tissue is/has been compromised by irradiation or there is decreased perfusion or hypoxia</td>
</tr>
<tr>
<td>Compromised skin grafts and flaps - one subsequent authorization</td>
<td>40</td>
<td>5</td>
<td>Evidence of stabilization of graft or flap</td>
</tr>
<tr>
<td>Crush injury, compartment syndrome and other acute traumatic ischernias</td>
<td>36</td>
<td>12</td>
<td>Adjunct to standard medical and surgical interventions</td>
</tr>
<tr>
<td>Decompression sickness</td>
<td>28</td>
<td>1</td>
<td>Diagnosis based on signs and/or symptoms of decompression sickness after a dive or altitude exposure</td>
</tr>
<tr>
<td>Diabetic foot ulcer - initial authorization</td>
<td>60</td>
<td>30</td>
<td>After at least 30 days of standard medical wound therapy, with a wound pO2 less than 40 mmHg AND wound classified as Wagner grade 3 or higher. *</td>
</tr>
<tr>
<td>Diabetic foot ulcer - two subsequent authorizations</td>
<td>60</td>
<td>20</td>
<td>Evidence of continuing healing and wound pO2 less than 40 mmHg</td>
</tr>
<tr>
<td>Severe anemia</td>
<td>50</td>
<td>10</td>
<td>Hgb less than 6.0 sustained secondary to hemorrhage, hemolysis, or aplasia, when the client is unable to be cross matched or refuses transfusion because of religious beliefs</td>
</tr>
<tr>
<td>Clostridial myositis and myonecrosis (gas gangrene)</td>
<td>39</td>
<td>13</td>
<td>Evidence of unsuccessful medical and/or surgical wound treatment and positive Gram-stained smear of the wound fluid</td>
</tr>
<tr>
<td>Necrotizing soft tissue infections - initial authorization</td>
<td>36</td>
<td>12</td>
<td>Evidence of unsatisfactory response to standard medical and surgical treatment and advancement of dying tissue</td>
</tr>
</tbody>
</table>

Note: The following Wagner wound classification grades apply only to the diabetic foot ulcer indications:
Grade 1: Superficial diabetic ulcer
Grade 2: Ulcer extension - involves ligament, tendon, joint capsule or fascia (No abscess or osteomyelitis)
Grade 3: Deep ulcer with abscess or osteomyelitis
Grade 4: Gangrene to portion of forefoot
Grade 5: Extensive gangrene of foot
<table>
<thead>
<tr>
<th>Covered Indication</th>
<th>Total 30-Minute Intervals Allowed for Procedure Code G0277</th>
<th>Total Professional Sessions Allowed for Procedure Code 99183</th>
<th>Medical Necessity Documentation of the Following is Required</th>
</tr>
</thead>
<tbody>
<tr>
<td>Necrotizing soft tissue infections - two subsequent authorizations</td>
<td>15</td>
<td>5</td>
<td>Evidence that advancement of dying tissue has slowed</td>
</tr>
<tr>
<td>Delayed radiation injury (soft tissue and bony necrosis) - initial authorization</td>
<td>40</td>
<td>10</td>
<td>Evidence of unsatisfactory clinical response to conventional treatment</td>
</tr>
<tr>
<td>Delayed radiation injury - one subsequent authorization</td>
<td>40</td>
<td>10</td>
<td>Evidence of improvement demonstrated by clinical response</td>
</tr>
<tr>
<td>Refractory osteomyelitis - initial authorization</td>
<td>40</td>
<td>10</td>
<td>Evidence of unsatisfactory clinical response to conventional multidisciplinary treatment</td>
</tr>
<tr>
<td>Refractory osteomyelitis - one subsequent authorization</td>
<td>15</td>
<td>5</td>
<td>Evidence of improvement demonstrated by clinical response</td>
</tr>
<tr>
<td>Acute thermal burn injury - initial authorization</td>
<td>45</td>
<td>15</td>
<td>Partial or full thickness burns covering greater than 20% of total body surface area OR with involvement of the hands, face, feet or perineum</td>
</tr>
<tr>
<td>Acute thermal burn injury - three subsequent authorizations</td>
<td>30</td>
<td>10</td>
<td>Evidence of continuing improvement demonstrated by clinical response</td>
</tr>
<tr>
<td>Intracranial abscess - initial authorization</td>
<td>15</td>
<td>5</td>
<td>Adjunct to standard medical and surgical interventions when one or more of the following conditions exist: Multiple abscesses Abscesses in a deep or dominant location Compromised host Surgery contraindicated or client is a poor surgical risk</td>
</tr>
<tr>
<td>Intracranial abscess - one subsequent authorization</td>
<td>15</td>
<td>5</td>
<td>Evidence of improvement demonstrated by clinical response and radiological findings</td>
</tr>
</tbody>
</table>

*Note: The following Wagner wound classification grades apply only to the diabetic foot ulcer indications:*

- **Grade 1**: Superficial diabetic ulcer
- **Grade 2**: Ulcer extension - involves ligament, tendon, joint capsule or fascia (No abscess or osteomyelitis)
- **Grade 3**: Deep ulcer with abscess or osteomyelitis
- **Grade 4**: Gangrene to portion of forefoot
- **Grade 5**: Extensive gangrene of foot

Procedure code 99183 is authorized according to the number of professional sessions (total HBOT treatments), and procedure code G0277 is authorized according to the number of 30-minute intervals of chamber time. The units in the columns for procedure codes 99183 and G0277 represent the maximum number of sessions and intervals that are allowed for that procedure code per authorization.

Limitations beyond those listed in the table above are considered experimental and investigational.
In emergency situations, the prior authorization request must be submitted no later than three business days after the date the service is rendered. Providers must not submit a claim until the prior authorization request has been approved. If the request has not been approved, the claim will be denied.

**9.2.33 Ilizarov Device and Procedure**

Providers must use procedure codes 20692, 20693, 20694, and 20999 when submitting claims for the Ilizarov procedure. A global fee payment methodology is applied to the Ilizarov device procedure codes. Procedure codes 20692, 20693, 20694, and 20999 include the preconstruction, surgical application, adjustments to the device for up to 6 months, and the removal of the device.

Providers who bill for other external fixator devices, such as the Monticelli device, should continue to use procedure codes 20690 or 20692, where applicable, when billing for the surgical applications.

**9.2.34 Immunization Guidelines and Administration**

Texas Medicaid reimburses immunizations (vaccines and toxoids) that the Advisory Committee on Immunization Practices (ACIP) recommends as routine.

Providers must follow the most current ACIP recommendations unless they conflict with guidelines from the Texas Vaccines for Children (TVFC) Program, in which case providers must follow TVFC guidelines. Providers must also provide the appropriate vaccine information statements (VISs) produced by the Centers for Disease Control and Prevention (CDC). VISs explain the benefits and risks of the vaccines and toxoids administered.

*Note:* Administered vaccines and toxoids must be reported to DSHS. After obtaining consent, DSHS submits all reported vaccines and toxoids to a centralized repository of immunization histories. This lifespan registry is known in Texas as ImmTrac2.

**9.2.34.1 Administration Fee**

An administration fee may be reimbursed for all covered vaccines and toxoids that are administered according to the ACIP. The following procedure codes may be reimbursed when billed for vaccine and toxoid administration:

<table>
<thead>
<tr>
<th>Procedure Code</th>
<th>90460</th>
<th>90461</th>
<th>90471</th>
<th>90472</th>
<th>90473</th>
<th>90474</th>
</tr>
</thead>
</table>

Procedure codes 90460 and 90461 are benefits for services rendered to clients who are birth through 18 years of age when counseling is provided for the immunization administered.

Procedure codes 90471, 90472, 90473, and 90474 are benefits when counseling is not provided for the immunization administered. Procedure codes 90471 and 90472 may be reimbursed for services rendered to clients of any age. Procedure codes 90473 and 90474 are restricted to clients who are 20 years of age and younger.

The administration fee may be reimbursed when the procedure code for the vaccine or toxoid administered (regardless of the source of the vaccine or toxoid) and the administration fee procedure code are billed on the same claim with the same date of service. Only one administration fee may be reimbursed to any provider for each vaccine or toxoid administered per day.
The following vaccines and toxoids procedure codes are a benefit of Texas Medicaid for clients who are 20 years of age and younger based on the number of recognized components as follows:

<table>
<thead>
<tr>
<th>Procedure Code</th>
<th>Number of Recognized Components**</th>
<th>Procedure Code</th>
<th>Number of Recognized Components**</th>
</tr>
</thead>
<tbody>
<tr>
<td>90620*</td>
<td>1</td>
<td>90621*</td>
<td>1</td>
</tr>
<tr>
<td>90630</td>
<td>1</td>
<td>90632</td>
<td>1</td>
</tr>
<tr>
<td>90633*</td>
<td>1</td>
<td>90636</td>
<td>2</td>
</tr>
<tr>
<td>90644</td>
<td>2</td>
<td>90647*</td>
<td>1</td>
</tr>
<tr>
<td>90648*</td>
<td>1</td>
<td>90651*</td>
<td>1</td>
</tr>
<tr>
<td>90655*</td>
<td>1</td>
<td>90654</td>
<td>1</td>
</tr>
<tr>
<td>90657*</td>
<td>1</td>
<td>90656*</td>
<td>1</td>
</tr>
<tr>
<td>90660*</td>
<td>1</td>
<td>90658*</td>
<td>1</td>
</tr>
<tr>
<td>90670*</td>
<td>1</td>
<td>90661</td>
<td>1</td>
</tr>
<tr>
<td>90673</td>
<td>1</td>
<td>90672*</td>
<td>1</td>
</tr>
<tr>
<td>90680*</td>
<td>1</td>
<td>90674</td>
<td>1</td>
</tr>
<tr>
<td>90682</td>
<td>1</td>
<td>90681*</td>
<td>1</td>
</tr>
<tr>
<td>90686*</td>
<td>1</td>
<td>90685*</td>
<td>1</td>
</tr>
<tr>
<td>90688*</td>
<td>1</td>
<td>90687*</td>
<td>1</td>
</tr>
<tr>
<td>90698*</td>
<td>5</td>
<td>90696*</td>
<td>4</td>
</tr>
<tr>
<td>90702*</td>
<td>2</td>
<td>90700*</td>
<td>3</td>
</tr>
<tr>
<td>90710*</td>
<td>4</td>
<td>90707*</td>
<td>3</td>
</tr>
<tr>
<td>90714*</td>
<td>2</td>
<td>90713*</td>
<td>1</td>
</tr>
<tr>
<td>90716*</td>
<td>1</td>
<td>90715*</td>
<td>3</td>
</tr>
<tr>
<td>90732*</td>
<td>1</td>
<td>90723*</td>
<td>5</td>
</tr>
<tr>
<td>90734*</td>
<td>1</td>
<td>90733</td>
<td>1</td>
</tr>
<tr>
<td>90744*</td>
<td>1</td>
<td>90743</td>
<td>1</td>
</tr>
<tr>
<td>90748*</td>
<td>2</td>
<td>90746</td>
<td>1</td>
</tr>
<tr>
<td>90756*</td>
<td>1</td>
<td>90749</td>
<td>1</td>
</tr>
</tbody>
</table>

* TVFC-distributed vaccine/toxoid  
** The number of components applies if counseling is provided and procedure codes 90460 and 90461 are submitted.

Each vaccine or toxoid and its administration must be submitted on the claim in the following sequence: the vaccine procedure code immediately followed by the applicable immunization administration procedure code(s). All of the immunization administration procedure codes that correspond to a single vaccine or toxoid procedure code must be submitted on the same claim as the vaccine or toxoid procedure code.

Each vaccine or toxoid procedure code must be submitted with the appropriate “administration with counseling” procedure code(s) (procedure codes 90460 and 90461) or the most appropriate “administration without counseling” procedure code (procedure code 90471, 90472, 90473, or 90474). If an “administration with counseling” procedure code is submitted with an “administration without counseling” procedure code for the same vaccine or toxoid, the second administration of the vaccine or toxoid will be denied.
Administration with Counseling

Providers must submit claims for immunization administration procedure codes 90460 or 90461 based on the number of components per vaccine. Providers must specify the number of components per vaccine by billing 90460 and 90461 as defined by the procedure code descriptions:

- Procedure code 90460 is submitted for the administration of the first component.
- Procedure code 90461 is submitted for the administration of each additional component identified in the vaccine.

Procedure code 90461 will be denied if procedure code 90460 has not been submitted on the same claim for the same vaccine or toxoid.

The necessary counseling that is conducted by a physician or other qualified health-care professional must be documented in the client’s medical record.

The following is an example of how to submit claims for immunization administration procedure codes when counseling is provided:

<table>
<thead>
<tr>
<th>Procedure Code</th>
<th>Quantity Billed</th>
</tr>
</thead>
<tbody>
<tr>
<td>Vaccine or toxoid procedure code with 1 component</td>
<td>1</td>
</tr>
<tr>
<td>90460 (1st component)</td>
<td>1</td>
</tr>
<tr>
<td>Vaccine or toxoid procedure code with 3 components</td>
<td>1</td>
</tr>
<tr>
<td>90460 (1st component)</td>
<td>1</td>
</tr>
<tr>
<td>90461 (2nd and 3rd components)</td>
<td>2</td>
</tr>
</tbody>
</table>

**Note:** The term “components” refers to the number of antigens that prevent disease(s) caused by one organism. Combination vaccines are those that contain multiple vaccine components.

Administration without Counseling

Procedure codes 90471, 90472, 90473, and 90474 may be reimbursed per vaccine based on the route of administration.

The following is an example of how to submit claims for injection administration procedure codes when counseling is not provided:

<table>
<thead>
<tr>
<th>Procedure Code</th>
<th>Quantity Billed</th>
</tr>
</thead>
<tbody>
<tr>
<td>Vaccine or toxoid procedure code</td>
<td>1</td>
</tr>
<tr>
<td>90471 (Injection administration)</td>
<td>1</td>
</tr>
<tr>
<td>Vaccine or toxoid procedure code</td>
<td>1</td>
</tr>
<tr>
<td>90472 (Injection administration)</td>
<td>1</td>
</tr>
<tr>
<td>Vaccine or toxoid procedure code</td>
<td>1</td>
</tr>
<tr>
<td>90472 (Injection administration)</td>
<td>1</td>
</tr>
</tbody>
</table>

**9.2.34.2 Documentation**

Providers must document the following information in the client’s medical record, which is subject to retrospective review to determine appropriate utilization and reimbursement of this service:

- The vaccine or toxoid given
- The date of the vaccine or toxoid administration (day, month, year)
- The name of the vaccine or toxoid manufacturer and the vaccine or toxoid lot number
• The signature and title of the person administering the vaccine or toxoid
• The organization’s name and address
• The publication date of the VIS issued to the client, parent, or guardian
• The site at which the vaccine was given (recommended)

9.2.34.3 Vaccine Adverse Event Reporting System (VAERS)
VAERS encourages providers to report any adverse event that occurs after the administration of any vaccine in the United States, even if it’s unclear whether a vaccine caused it. The National Childhood Vaccine Injury Act (NCVIA) requires health-care providers to report:
• Any adverse event listed by the vaccine manufacturer as a contraindication to subsequent doses of the vaccine.
• Any reaction listed in the VAERS Reportable Events Table that occurs within the specified time period after vaccination.

Clinically significant adverse events should be reported even if it is unclear whether a vaccine caused the event.

Documentation of the injection site is recommended but not required.

A copy of the Reportable Events Table can be obtained by calling VAERS at 1-800-822-7967 or by downloading it from vaers.hhs.gov/docs/VAERS_Table_of_Reportable_Events_Following_Vaccination.pdf.

9.2.35 Immunizations for Clients Birth through 20 Years of Age
Administration of vaccines and toxoids to clients who are birth through 20 years of age may be a benefit of THSteps when provided as part of a THSteps medical checkup. A THSteps provider who bills vaccines and toxoids with diagnosis or age restrictions is subject to those restrictions. In addition to the age appropriate diagnosis for the THSteps preventive care medical checkup, providers must bill the claim with the diagnosis code that indicates the condition that necessitates the vaccine or toxoid.

If an immunization is administered as part of the preventive care medical checkup, diagnosis code Z23 may also be included on the claim, in addition to the age-appropriate diagnosis.

If an immunization is the only service provided during an office visit, providers may submit only diagnosis code Z23 on the claim.

Administration of vaccines and toxoids to clients who are birth through 20 years of age may be a benefit of CCP when the vaccine or toxoid is provided as part of an acute medical visit outside of a THSteps medical checkup.

Refer to: Section 5, “THSteps Medical” in the Children’s Services Handbook (Vol. 2, Provider Handbooks) for more information on THSteps age related diagnosis codes.

9.2.35.1 Vaccine Coverage Through the TVFC Program
Providers may refer to the TVFC web site at www.dshs.texas.gov/immunize/tvfc/default.shtm for information about the program and for a list of vaccines available through the program.

Note: All vaccines and toxoids recommended by Advisory Committee on Immunization Practices (ACIP) are available from the TVFC Program to enrolled clinic sites. Clinics participating in the TVFC Program have agreed to administer all ACIP-recommended vaccines to the eligible populations that are served.

When a single antigen vaccine or toxoid or a comparable antigen vaccine or toxoid is available through TVFC, but the provider chooses to use a different ACIP-recommended product, the administration fee will be reimbursed but the vaccine or toxoid will not be reimbursed.
Although Texas Medicaid does not mandate that providers enroll in TVFC, Texas Medicaid will not reimburse providers when the vaccine is available through TVFC. Only the administration fee will be reimbursed through Texas Medicaid when the vaccine or toxoid procedure code is identified on the claim. Clients may not be billed for vaccines and toxoids that are available through TVFC.

If a vaccine or toxoid meets the definition of “not available” through TVFC, it may be separately reimbursed through CCP when billed with modifier U1. Modifier U1 may be used in the following situations:

- The TVFC, based on their federal resolution (distribution/guidelines), does not distribute an HHSC-approved vaccine or toxoid following the ACIP recommendation, and the provider purchases vaccine to administer to all ACIP-recommended ages or risk groups.
- A new vaccine or toxoid approved by the ACIP with established guidelines, but has not been negotiated or added to a TVFC contract
- Funding for new vaccine or toxoid has not been established by TVFC
- Insufficient vaccine and toxoid supply due to national supply or distribution issues, as reported to HHSC by TVFC

HHSC will notify providers if a vaccine or toxoid meets the definition of “not available” from TVFC and when the provider’s privately purchased vaccine or toxoid may be billed with modifier U1. Modifier U1 must not be used due to a provider’s failure to enroll in TVFC or to maintain sufficient TVFC vaccine or toxoid inventory.


### 9.2.35.2 Vaccine and Toxoid Procedure Codes

The following vaccine and toxoid procedure codes may be reimbursed for Texas Medicaid clients who are birth through 20 years of age:

<table>
<thead>
<tr>
<th>Procedure Codes</th>
<th>Bacillus Calmette-Guérin (BCG)</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>Refer to:</strong></td>
<td>Subsection 9.2.9, “Bacillus Calmette-Guérin (BCG) Intravesical for Treatment of Bladder Cancer” in this handbook.</td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th>Procedure Codes</th>
<th>Hepatitis A and B</th>
</tr>
</thead>
<tbody>
<tr>
<td>90630</td>
<td>90632</td>
</tr>
<tr>
<td>90740</td>
<td>90743</td>
</tr>
<tr>
<td>90748*</td>
<td>90744*</td>
</tr>
<tr>
<td>90746</td>
<td>90747</td>
</tr>
</tbody>
</table>

Providers must document in the client’s medical record the indication for the hepatitis B vaccine, for dialysis patients. These records are subject to retrospective review to determine appropriate utilization of and reimbursement for this service.

Procedure codes 96372 and 96374 may be reimbursed for the administration of hepatitis B vaccine procedure codes 90740 and 90747.

Providers are expected to follow the ACIP recommendations for administration.

<table>
<thead>
<tr>
<th>Procedure Codes</th>
<th>Hepatitis B Immune Globulin</th>
</tr>
</thead>
<tbody>
<tr>
<td>90371</td>
<td>96372</td>
</tr>
<tr>
<td>96374</td>
<td>J1571</td>
</tr>
<tr>
<td>J1573</td>
<td>* Indicates a vaccine or toxoid distributed through TVFC. Vaccines and toxoids available through TVFC for clients who are birth through 18 years of age will not be reimbursed through Texas Medicaid. These vaccines and toxoids will be processed as informational.</td>
</tr>
</tbody>
</table>
### Providers must document in the client’s medical record the indication for the immunoglobulin. These records are subject to retrospective review to determine appropriate utilization of and reimbursement for this service.

### Intramuscular hepatitis B immune globulin (HBIg) may be reimbursed when medically necessary to provide coverage for acute exposure to the hepatitis B virus. HBIg is not provided through TVFC.

- Procedure codes 90371, J1571, and J1573 must be billed with diagnosis code Z205, Z206, or Z20828.
- Only one HBIg procedure code will be paid if billed with the same date of service by any provider as any other HBIg procedure code.

- Procedure codes 96372 and 96374 may be reimbursed for HBIg administration. Providers are expected to follow the ACIP recommendations for administrations.

#### Hbi

- Procedure codes 90647, 90648, 90651, 90654, 90655, 90656, 90657, 90658, 90660, 90661, 90672, 90673, 90674, 90682, 90685, 90686, 90687, 90688, 90756

**Human Papilloma (HPV)**

- Procedure codes 90651

**Influenza**

- Procedure codes 90654, 90655, 90656, 90657, 90658, 90660, 90661, 90672, 90673, 90674, 90682, 90685, 90686, 90687, 90688, 90756

- Influenza vaccine is a benefit of Texas Medicaid for high-risk clients who are not covered by THSteps or TVFC or when the vaccine is not declared available through the TVFC.

- Texas Medicaid considers the influenza season in the United States to be October through the end of May.

- Procedure codes 90655, 90657, 90685, and 90687 are limited to clients who are 6 through 35 months of age.

- Procedure codes 90656 and 90658 are limited to clients who are 3 years of age and older.

- Procedure codes 90686 and 90688 are limited to clients who are 6 months of age and older.

- Procedure code 90682 is limited to clients who are 18 years of age and older.

- Procedure code 90756 is limited to clients who are 4 years of age and older.

#### Measles, Mumps, Rubella Vaccine (MMR)

- Procedure code 90707

#### Measles, Mumps, Rubella, and Varicella Vaccine (MMRV)

- Procedure code 90710

#### Pneumococcal and Meningococcal

- Procedure codes 90620, 90621, 90644, 90670, 90733, 90734

* Indicates a vaccine or toxoid distributed through TVFC. Vaccines and toxoids available through TVFC for clients who are birth through 18 years of age will not be reimbursed through Texas Medicaid. These vaccines and toxoids will be processed as informational.
9.2.36 Immunizations for Clients Who Are 21 Years of Age and Older

Vaccines and toxoids may be reimbursed through Texas Medicaid at a fee determined by HHSC when the vaccine is medically necessary. Providers are expected to follow the ACIP recommendations for administration.

The following immunizations are identified and recommended by the ACIP as medically-necessary for clients who are 21 years of age and older (this list is not all-inclusive):

### Procedure Codes

<table>
<thead>
<tr>
<th>Procedure Codes</th>
</tr>
</thead>
<tbody>
<tr>
<td>BCG, Refer to: Subsection 9.2.9, “Bacillus Calmette-Guérin (BCG) Intravesical for Treatment of Bladder Cancer” in this handbook.</td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th>Immunization Procedure Codes</th>
</tr>
</thead>
<tbody>
<tr>
<td>Hepatitis A and B</td>
</tr>
<tr>
<td>90632</td>
</tr>
</tbody>
</table>

Providers must document in the client’s medical record the indication for the hepatitis B vaccine, for dialysis patients. These records are subject to retrospective review to determine appropriate utilization of and reimbursement for this service.

Procedure codes 96372 and 96374 may be reimbursed for the administration of hepatitis B vaccine procedure codes 90740 and 90747.
### Immunization Procedure Codes

#### Hepatitis B Immune Globulin

<table>
<thead>
<tr>
<th>Code</th>
<th>90371</th>
<th>96372</th>
<th>96374</th>
<th>J1571</th>
<th>J1573</th>
</tr>
</thead>
</table>

Providers must document in the client’s medical record the indication for the immunoglobulin. These records are subject to retrospective review to determine appropriate utilization of and reimbursement for this service.

Intramuscular HBlg may be reimbursed when medically necessary to provide coverage for acute exposure to the hepatitis B virus. HBlg is not provided through TVFC.

Procedure codes 90371, J1571, and J1573 must be billed with diagnosis code Z205, Z206, or Z20828. Only one HBlg procedure code will be paid if billed with the same date of service by any provider as any other HBlg procedure code.

Procedure codes 96372 and 96374 may be reimbursed for HBlg administration.

#### Hepatitis A and B

<table>
<thead>
<tr>
<th>Code</th>
<th>90636</th>
</tr>
</thead>
</table>

#### Human Papilloma (HPV)

<table>
<thead>
<tr>
<th>Code</th>
<th>90651</th>
</tr>
</thead>
</table>

#### Influenza

<table>
<thead>
<tr>
<th>Code</th>
<th>90630</th>
<th>90654</th>
<th>90656</th>
<th>90658</th>
<th>90661</th>
</tr>
</thead>
</table>

<table>
<thead>
<tr>
<th>Code</th>
<th>90662</th>
<th>90673</th>
<th>90674</th>
<th>90682</th>
<th>90686</th>
</tr>
</thead>
</table>

<table>
<thead>
<tr>
<th>Code</th>
<th>90688</th>
<th>90756</th>
</tr>
</thead>
</table>

Influenza vaccine is a benefit of Texas Medicaid for all clients. Texas Medicaid considers the influenza season in the United States to be October through the end of May. The optimal time to receive influenza vaccine is as early in the season as it is available. However, clients should continue to receive influenza vaccine through March. The vaccine may be administered one time per influenza season.

Procedure code 90682 is limited to clients who are 18 years of age and older.

#### MMR

<table>
<thead>
<tr>
<th>Code</th>
<th>90707</th>
</tr>
</thead>
</table>

#### Pneumococcal and Meningococcal

<table>
<thead>
<tr>
<th>Code</th>
<th>90620</th>
<th>90621</th>
<th>90670</th>
<th>90732</th>
</tr>
</thead>
</table>

The initial pneumococcal polysaccharide vaccine is limited to one per client per lifetime. Revaccination is recommended five years (not interpreted to mean every five years) after the initial dose for high-risk individuals.

Revaccination after a second dose is not reimbursed.

#### Shingles

<table>
<thead>
<tr>
<th>Code</th>
<th>90736</th>
<th>90750</th>
</tr>
</thead>
</table>

Procedure code 90736 is limited to clients who are 60 years of age and older. Procedure code 90750 is limited to clients who are 50 years of age and older.

#### Tetanus

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<thead>
<tr>
<th>Code</th>
<th>90714</th>
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</thead>
</table>

#### Tetanus, Diphtheria, and Acellular Pertussis Vaccine (Tdap)

<table>
<thead>
<tr>
<th>Code</th>
<th>90715</th>
</tr>
</thead>
</table>

The specific diagnosis necessitating the vaccine or toxoid is required when billing the administration fee procedure code in combination with the appropriate vaccine procedure code. Diagnosis code Z23 may also be included. The type of immunization given will be identified by the procedure code.
9.2.37 Postexposure Prophylaxis for Rabies

Postexposure prophylaxis for rabies procedure codes 90375, 90376, 90377, and 90675 is a benefit of Texas Medicaid. Rabies vaccine for pre-exposure procedure code 90676 is not a benefit of Texas Medicaid.

Postexposure rabies vaccine is limited to clients with diagnosis code Z203.

Animal bites to people must be reported as soon as possible to the Local Rabies Control Authority (LRCA).

Postexposure prophylaxis for rabies is not necessary following exposure to an animal that tests negative for the rabies virus.

An exposed person who has never received a complete pre- or postexposure rabies vaccine series will first receive a dose of rabies immune globulin (HRIG). This is a blood product that contains antibodies against rabies and gives immediate, short-term protection. The injection should be given in or near the wound area.

HRIG that is not administered when vaccination begins can be administered up to seven days after the administration of the first dose of vaccine. Beyond the seventh day, HRIG is not recommended since an antibody response to the vaccine is presumed to have occurred, and HRIG may inhibit the immune response to the vaccine.

The recommended dose of HRIG is 20 IU/kg body weight. This formula is applicable to all age groups, including children.

The postexposure treatment will also include five doses of rabies vaccine (1.0 ml. intramuscular). The first dose should be given as soon as possible after the exposure (day 0). Additional doses should be given on days 3, 7, 14, and 28 after the first shot. For an exposed person who has previously been vaccinated with a complete pre- or postexposure vaccine series, two doses of rabies vaccine should be given on days 0 and 3.

Health care providers, who determine their client requires the preventative rabies vaccination series after valid rabies exposure, may obtain the biologicals directly from the manufacturer or through one of the DSHS depots around the state.

Injection administration is a benefit for administration of rabies vaccine for post exposure.

9.2.37.1 Prior Authorization for Postexposure Rabies Vaccine

Prior authorization is not required for postexposure rabies vaccine. The physician must maintain documentation of the exposure in the client’s medical record.

9.2.37.2 Limitations for Postexposure Rabies Vaccine

Reimbursement for postexposure rabies vaccine is limited to one per client per day, by any provider.

Reimbursement for postexposure rabies vaccine is limited to 5 occurrences per 90 rolling days. Claims billed for any vaccine given beyond 90 rolling days will be denied.

9.2.37.2.1 Obtaining Rabies Vaccine and HRIG from DSHS for PEP Use

Providers may obtain the vaccine and HRIG directly from the manufacturer. If a provider is not able to obtain the vaccine and/or HRIG directly, providers may contact DSHS local or state public health professionals.

For each potential rabies exposure, providers must consult with their local health department or the DSHS regional ZC program office that serves their area. Requests for consultations made to DSHS after-hours or on holidays should be directed to the DSHS On-Call Physician at 1-888-963-7111.
Local public health professionals or regional ZC staff will help providers determine whether or not the exposure situation warrants PEP. If the exposure situation is determined to be valid, providers will be given detailed information about how to obtain rabies vaccine and HRIG for the patient.

Providers can refer to the following DSHS web pages for the contact information of local public health professionals:

- Full Service Local Health Departments and Districts of Texas at [www.dshs.texas.gov/regions/lhds.shtm](http://www.dshs.texas.gov/regions/lhds.shtm)
- Zoonosis Control Branch at [www.dshs.texas.gov/idcu/health/zoonosis/contact/](http://www.dshs.texas.gov/idcu/health/zoonosis/contact/)
- Regional DSHS ZC offices

9.2.38 Implantable Infusion Pumps

Implantable infusion pump (IIPs) are intended to provide long-term, continuous, or intermittent drug infusion. They may be medically necessary in the following circumstances:

- Administration of intrathecal or epidural antispasmodic drugs to treat refractory intractable spasticity
- Administration of Intrathecal, epidural, or central venous analgesic (opioid or non-opioid) drugs for treatment of severe chronic intractable pain
- Administration of intrahepatic chemotherapy for primary liver cancer or metastatic cancer with metastases limited to the liver
- Administration of intra-arterial chemotherapy in head and neck cancers

An implantable infusion pump is not a benefit for the following uses:

- Continuous insulin infusion for diabetes
- Continuous heparin infusion for recurrent thromboembolic disease
- Continuous intralesional infusion for severe chronic intractable pain
- Continuous intra-arterial infusion
- Continuous intra-articular infusion for severe chronic intractable pain
- Administration of antibiotics for osteomyelitis

All supplies associated with an IIP are included with the reimbursement for the surgery to implant the infusion pump and are not reimbursed separately.

Providers may be reimbursed for implantable infusion pumps using procedure codes E0782, E0783, and E0786. If procedure codes E0782 and E0783 are billed with the same date of service, only one may be reimbursed.

9.2.38.0.1 Prior Authorization for Implantable Infusion Pumps

Implantable infusion pumps (procedure codes E0782, E0783, and E0786) require prior authorization.
Prior authorization is not required for the physician services associated with the insertion, revision, removal, refilling, or maintenance of the IIP.

Providers must request prior authorization through the Special Medical Prior Authorization (SMPA) department. The ASC or DME provider may submit a request for prior authorization using the Special Medical Prior Authorization (SMPA) Form, which must be completed and signed by a physician.

The completed, signed and dated SMPA form must be maintained by the provider and the prescribing physician in the client’s medical record.

The completed SMPA Form must include the procedure code and quantity for the services that are requested. Documentation that is submitted with the prior authorization request must indicate whether the IIP will be provided by the ASC or the DME provider.

To avoid unnecessary denials, the physician must provide correct and complete information, including documentation of medical necessity for the requested IIP. The requesting provider may be asked for additional information to clarify or complete a request for the IIP.

Documentation submitted with the prior authorization request must indicate the client or caregiver has:

- The ability to provide a return demonstration performance.
- The attention, desire, interest, flexibility, and independence.
- An understanding of cause and effect and object permanence.

As indicated in the following sections, supporting documentation that is based on the type of IIP requested must be included with the request for prior authorization. All of the documentation listed under the specific type of IIP must be included with the request for prior authorization.

### 9.2.38.0.2 IIP for Administration of Anti-spasmodic Drug to Treat Severe Refractory Spasticity

The following documentation is required for prior authorization:

- Initial evaluation
- Type of surgical implantation and description of IIP requested
- Symptoms:
  - Degree of spasticity
  - Affected muscle groups
  - Functional impact
  - Duration of symptoms
- Any recent hospitalizations (within past 12 months)
- Comorbid conditions
- All pertinent laboratory and radiology results
- Treatment history of self-administration with evidence of:
  - A minimum of six weeks of non-invasive methods of spasticity control, including, but not limited to, oral antispasmodics, that either:
    - Failed to adequately control the spasticity, or
    - Produced intolerable side effects
  - The role, participation, and compliance of the family or client that demonstrate the following:
    - The ability to provide a return demonstration performance
• Attentiveness, desire, interest, flexibility, and independence
• An understanding of cause and effect and object permanence
• Favorable response to a trial intrathecal dose of the antispasmodic
• No contraindications to implantation exist, including, but not limited to, the following:
  • Coagulopathy
  • Infection
  • Other implanted devices where the “crosstalk” between devices may inadvertently change the prescription
  • Allergy or hypersensitivity to the drug being administered
• Treatment plan, including the following:
  • Antispasmodic to be infused
  • Follow-up, including pump refilling, maintenance, and monitoring of changes in infusion rate
  • Expected outcome
  • Treatment goals

9.2.38.1 IIP for Administration of Analgesic (Opioid or Nonopioid) Drug for Treatment of Severe Intractable Pain

The following documentation is required for prior authorization:
• The initial evaluation
• Type of surgical implantation and description of IIP requested
• Symptoms:
  • Severity of pain
  • Functional impact
• Source of pain or location, including whether pain is malignant or non-malignant
• Duration of symptoms
• Any recent hospitalizations (within the past 12 months)
• Comorbid conditions
• All pertinent laboratory and radiology results
• A life expectancy of at least three months

  Note: The standard of care for treatment of severe intractable pain for a client with a life expectancy of less than three months is to use less invasive techniques such as an external infusion pump.

For malignant pain, the following documentation is required for prior authorization:
• Treatment history with evidence of a favorable response to a trial intrathecal dose of the analgesic drug, defined as a minimum of 50 percent reduction in pain
• Failure of more conservative methods of pain control, including, but not limited to, oral analgesics, surgery, or therapy, that were ineffective due to one of the following:
• Failed to adequately control the pain, or
• Produced intolerable side effects

Note: The standard of care for treatment of severe intractable pain for a client with a life expectancy of less than three months is to use less invasive techniques such as an external infusion pump.

For nonmalignant pain, the following documentation is required for prior authorization:

• A minimum of six months of more conservative methods of pain control, including but not limited to oral analgesics, surgery, attempts to eliminate physical and behavioral abnormalities that may cause an exaggerated pain reaction, that were ineffective due to one of the following:
  • Failed to adequately control the pain, or
  • Produced intolerable side effects

Examples of non-malignant severe intractable pain include, but are not limited to, the following:

• Complex regional pain syndrome I & II (causalgia/RSD) refractory to other treatments.
• Post herpetic neuralgia
• Failed back syndrome
• Phantom limb pain
• Arachnoiditis (proven with MRI/increased CSF protein levels)
• Spinal cord myelopathy (refractory to conservative measurements)
• The role, participation, and compliance of the family or client that demonstrate the following:
  • The ability to provide a return demonstration performance
  • Attentiveness, desire, interest, flexibility, and independence
  • An understanding of cause and effect and object permanence
• No contraindications to implantation exist, including, but not limited to, the following:
  • Coagulopathy
  • Infection
  • Other implanted devices where the “crosstalk” between devices may inadvertently change the prescription
  • Tumor encroachment on the thecal sac
  • Allergy or hypersensitivity to the drug being administered
  • Treatment plan, including the following:
    • Analgesic to be infused
    • Follow-up including pump refilling, maintenance, and monitoring of changes in infusion rate
    • Expected outcome
    • Treatment goals

9.2.38.2 IIP for Administration of Intrahepatic Chemotherapy in Primary Liver Cancer or Colorectal Cancer with Liver Metastases

The following documentation is required for prior authorization:

• The initial evaluation
• Type of surgical implantation and description of IIP requested
• Diagnosis of one of the following:
• Primary liver cancer
• Metastatic cancer with metastases limited to the liver
• Any recent hospitalizations (within the past 12 months)
• Comorbid conditions
• All pertinent laboratory and radiology results
• The role, participation, and compliance of the family and/or client demonstrating:
  • The ability to provide a return demonstration performance
  • Attentiveness, desire, interest, flexibility, and independence
  • An understanding of cause and effect and object permanence
• No contraindications to implantation exist, including, but not limited to, the following:
  • Coagulopathy
  • Infection
  • Other implanted devices where the “crosstalk” between devices may inadvertently change the prescription
  • Allergy or hypersensitivity to the drug being administered
• Treatment plan, including the following:
  • Chemotherapeutic agent to be infused. The prescribed drug must be approved by the U.S. Food and Drug Administration (FDA) for the intended use and must be compatible with the implantable device (such as floxuridine or methotrexate)
  • Follow-up, including pump refilling, maintenance, and monitoring of changes in infusion rate
  • Expected outcome
  • Treatment goals

9.2.38.3 IIP for Administration of Intra-Arterial Chemotherapy in Head and Neck Cancers

The following documentation is required for prior authorization:
• Initial evaluation
• Type of surgical implantation and description of IIP requested
• Diagnosis and site(s) of any metastases
• Any hospitalizations (within the past 12 months) and all other diagnoses
• All pertinent laboratory and radiology results
• The role, participation, and compliance of the family or client that demonstrates the following:
  • The ability to provide a return demonstrate performance
  • Attentiveness, desire, interest, flexibility, and independence
  • An understanding of cause and effect and object permanence
• No contraindications to implantation exist, including, but not limited to, the following:
  • Coagulopathy
  • Infection
• Other implanted devices where the “crosstalk” between devices may inadvertently change the prescription
• Allergy or hypersensitivity to the drug being administered
• Treatment plan, including the following:
  • Chemotherapeutic agent to be infused
  • Follow-up, including pump refilling, maintenance, and monitoring of changes in infusion rate
  • Expected outcome
  • Treatment goals

9.2.38.4 Replacement of an IIP

An IIP is expected to last a minimum of five years. Prior authorization for replacement of an IIP is considered within five years when one of the following occurs:

• There has been a significant change in the client’s condition and the current equipment no longer meets the client’s needs.
• The equipment is no longer functional and either cannot be repaired or it is not cost-effective to repair.
• Loss or irreparable damage to the IIP has occurred. The following must be submitted with the prior authorization request:
  • A copy of the police or fire report, when appropriate
  • A statement about the measures to be taken in order to prevent reoccurrence

Replacement of an IIP for a client who is birth through 20 years of age that does not meet the criteria above may be considered for prior authorization through CCP.

The DME Certification and Receipt Form is required and must be completed before reimbursement can be made for any DME delivered to a client. The certification form must include the name of the item, the date the client received the DME, and the signatures of the provider and the client or primary caregiver.

The DME provider must maintain the signed and dated form in the client’s medical record.

Refer to: Subsection 2.7.3.5, “DME Certification and Receipt Form” in the Children’s Services Handbook (Vol. 2, Provider Handbooks) for more information about this form.

9.2.38.5 Implantation of Catheters, Reservoirs, and Pumps

The following procedure codes may be used to bill the implantation of catheters and infusion pumps or devices for long term medication administration:

<table>
<thead>
<tr>
<th>Procedure Codes</th>
</tr>
</thead>
<tbody>
<tr>
<td>62350</td>
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</tbody>
</table>

Procedure code 62350 or 63251 may be reimbursed when billed for the same date of service as procedure code 62360, 62361, or 62362.

Procedure codes 62355 and 62365 do not require prior authorization.
The following procedure codes are denied as included in the total anesthesia time when billed with the same date of service as an anesthesia procedure by the same physician:

<table>
<thead>
<tr>
<th>Procedure Codes</th>
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<tbody>
<tr>
<td>62350</td>
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<tr>
<td>62351</td>
</tr>
<tr>
<td>62355</td>
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<tr>
<td>62360</td>
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<tr>
<td>62361</td>
</tr>
<tr>
<td>62362</td>
</tr>
<tr>
<td>62365</td>
</tr>
</tbody>
</table>

These procedure codes are considered for reimbursement according to multiple surgery guidelines when billed with the same date of service as another surgical procedure performed by the same physician.

Procedure codes 95990, 96521, and 96522 are considered for reimbursement when used for refilling an implantable pump.

Procedure codes 62367, 62368, 62369, and 62370 may be used to bill for electronic analysis of an implantable infusion pump.

Procedure codes 62369 and 62370 will be denied when billed for the same date of service by the same provider as procedure code 62362.

The following procedure codes may be used to bill the insertion, revision, removal, or repair associated with implantable infusion pumps:

<table>
<thead>
<tr>
<th>Procedure Codes</th>
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</thead>
<tbody>
<tr>
<td>36260</td>
</tr>
<tr>
<td>36261</td>
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<tr>
<td>36262</td>
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### 9.2.38.6 Drug Monitoring Services

Providers must use the most appropriate procedure codes when submitting claims for drug monitoring services that monitor prescribed medications that can be abused when used for the treatment of chronic pain. These claims are subject to retrospective review. Claims may be reprocessed and recouped if they are submitted for these drug monitoring services in the office setting using a procedure code for a quantitative test rather than a qualitative or semiquantitative test.

An enzyme immunoassay (EIA) device can be used to provide preliminary qualitative or semiquantitative test results for point-of-care monitoring purposes. EIA devices and the reagents used to perform in-office drug testing are cleared by the FDA only to obtain qualitative or semiquantitative initial screen or preliminary results.

Immunoassay and enzyme assay are tests that produce qualitative and semiquantitative results, so these tests must not be reported with procedure codes for quantitative tests. A qualitative or semiquantitative test is not a quantitative test and must not be billed as such.

The initial drug screen or preliminary result testing yields qualitative and semiquantitative results, which must be reported with an appropriate drug testing procedure code, as categorized in the CPT manual as “Drug Testing.” Only those procedure codes that are a benefit of Texas Medicaid may be reimbursed.

CPT-categorized “Chemistry” and “Therapeutic Drug Assay” procedure codes are for quantitative tests and must not be reported for an initial screen or preliminary result that was performed in the point-of-care setting.

**Refer to:** The CPT manual for drug testing, chemistry, and therapeutic drug assay procedure codes, and to the Texas Medicaid fee schedule for procedure codes that may be reimbursed by Texas Medicaid.

Using procedure codes for quantitative tests to report preliminary qualitative or semiquantitative test results is considered systematic upcoding and may lead to administrative sanctions, civil monetary penalties, and criminal prosecution.
Providers may refer to the CMS website for more information about laboratory tests that may be rendered in the office setting. For tests that require a CLIA certificate of waiver, CMS publishes a list of all waived tests. The list is updated quarterly and includes the procedure code to use when billing a test.

9.2.39 Laboratory Services

Texas Medicaid benefits are provided for professional and technical services ordered by a physician and provided under the supervision of a physician in a setting other than a hospital (inpatient or outpatient). All laboratory services must be documented in the client’s medical record as medically necessary and referenced to an appropriate diagnosis. Texas Medicaid does not reimburse baseline or screening laboratory studies.

Providers may bill only for laboratory tests that are actually provided in their office. Any test sent to an outside laboratory must not be billed on the provider’s claim. Laboratories bill Texas Medicaid directly for the tests they perform.

Unless otherwise noted, interpretation of laboratory tests is considered part of the provider’s professional services (hospital, office, or emergency room visits) and must not be billed separately. Modifier Q4 is required for laboratory, radiology, and ultrasound interpretations by any provider other than the attending physician.

Laboratory tests that are generally considered part of a laboratory panel (e.g., chemistries, CBCs, urinalyses [UAs]) and that are performed on the same day must be billed as a panel regardless of the method used to perform the tests (automated or manual).

Physician interpretations that are requested of a consulting pathologist and require professional reading and reporting of results may be billed to Texas Medicaid separately as a professional charge.

All providers of laboratory services must comply with the rules and regulations of CLIA. Providers not complying with CLIA cannot be reimbursed for laboratory services.

Texas Medicaid follows the Medicare categorization of tests for CLIA certificate holders.

Refer to: The CMS website at www.cms.gov/CLIA/10_Categorization_of_Tests.asp for information about procedure code and modifier QW requirements.

Subsection 2.2.5, “Automated Laboratory Tests and Laboratory Paneling” in the Radiology and Laboratory Services Handbook (Vol. 2, Provider Handbooks) for claims processing instructions.


Subsection 3.4.2, “Reimbursement” in the Radiology and Laboratory Services Handbook (Vol. 2, Provider Handbooks) for claims processing instructions.

Subsection 2.2, “Fee-for-Service Reimbursement Methodology” in “Section 2: Texas Medicaid Fee-for-Service Reimbursement” (Vol. 1, General Information) for more information about reimbursement.

9.2.39.1 THSteps Laboratory Services

Refer to: Subsection 5.3.11.6, “Laboratory Test” in the Children’s Services Handbook (Vol. 2, Provider Handbooks).

9.2.39.2 Laboratory Handling Charge

The laboratory handling charge covers the expense of obtaining and packaging the specimen and sending it to a reference laboratory.
A laboratory handling charge (procedure code 99000) may be billed if the specimen is obtained by venipuncture or catheterization and sent to an outside lab. The reference laboratory name and address or provider identifier must be listed in Block 32 of the CMS-1500 claim form, and Block 20 must be completed.

The provider is required to forward the client’s name, address, Medicaid ID number, and diagnosis, if appropriate, with the specimen to the reference laboratory so the laboratory may bill Texas Medicaid for its services.

A provider may bill only one laboratory handling charge per client visit unless the specimen is divided and sent to different laboratories or different specimens are collected and sent to different labs. The claim must indicate the name and/or address of each laboratory to which a specimen is sent for more than one laboratory handling fee to be paid. This laboratory handling benefit does not apply to THSteps medical checkup providers who must submit specimens to the DSHS Laboratory.

9.2.39.3 Blood Counts

Texas Medicaid considers a baseline CBC appropriate for the evaluation and management of existing and suspected disease processes. CBCs should be individualized and based on client history, clinical indications, or proposed therapy and will not be reimbursed for screening purposes.

Refer to: Subsection 2.2.7, “Complete Blood Count (CBC)” in the Radiology and Laboratory Services Handbook (Vol. 2, Provider Handbooks) for more information about blood counts.

9.2.39.4 Clinical Lab Panel Implementation

Refer to: Subsection 2.2.5, “Automated Laboratory Tests and Laboratory Paneling” in the Radiology and Laboratory Services Handbook (Vol. 2, Provider Handbooks) for more information about laboratory panels.

9.2.39.5 Clinical Pathology Consultations

Clinical pathology consultations (procedure code 80500 or 80502) are a benefit of Texas Medicaid for services rendered by a consultant who is either a clinical pathologist or a geneticist. In a clinical pathology consultation, the consultant may also help the ordering physician determine whether further study is appropriate, based on test results.

Providers may be reimbursed for clinical pathology consultations when the claim indicates the following information:

- The name and address or provider identifier of the physician who requested the consultation.
- A written narrative report describing the findings of the consultation, which will also be included in the client’s medical record.

Note: To submit claims for interpretation, the provider must document an interaction that clearly shows that the consultant interpreted the test results and made specific recommendations to the attending physicians.

If the claim does not include all of this information, the clinical pathology consultation will be denied.

Note: Geneticists who provide a pathology consultation must submit claims using their acute care provider identifier.

Routine conversations held between a consultant and attending physicians about test orders or results are not consultations. Information that can be furnished by a non-physician laboratory specialist does not qualify as a consultation service.

9.2.39.6 Cytogenetics Testing

Cytogenetics testing is a group of laboratory tests involving the study of chromosomes.
Clinical evidence supports the significance of cytogenetics evaluation in the diagnosis, prognosis, and treatment of acute leukemias and lymphomas, especially in children. The detection of the well-defined recurring genetic abnormalities often enables a correct diagnosis with important prognostic information that affects the treatment protocol.

Reimbursement for cytogenetics testing is limited to the following diagnosis codes:

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Cytogenetics testing may be reimbursed with the following procedure codes and limitations:

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### 9.2.39.7 Maternal Serum Alpha-Fetoprotein (MSAFP)

MSAFP may be reimbursed once per pregnancy per provider for all pregnant women eligible for Medicaid. For additional services, payment is allowed with documentation attached to the claim. Procedure code 82105 should be used for MSAFP.

### 9.2.40 Pharmacogenetics

Pharmacogenetic testing of cytochrome p450 (CYP450) metabolic pathway may be considered medically necessary only if the results of the testing are necessary to differentiate between treatment options.
The use of pharmacogenetics may be considered medically necessary once in a lifetime to determine effective response to drug therapy for the following:

### 9.2.40.1 Testing of Polymorphic 2C19

Pharmacogenetics testing of polymorphic 2C9 (procedure code 81227) may be considered for clopidogrel treatment and requires prior authorization and may be considered medically necessary when all of the following conditions are met:

- The client has never received genetic testing of the 2C19 alleles.
- The client has never received clopidogrel treatment.
- The clopidogrel treatment will be used for one of the following diseases or conditions:
  - ST elevated and non-ST elevated myocardial infarction (STEMI and NSTEMI)
  - Subsequent STEMI and NSTEMI
  - Dressler’s syndrome
  - Unstable angina
  - Cerebral infarction due to embolism of cerebral arteries
  - Occlusion and stenosis of cerebral arteries, not resulting in cerebral infarction
  - Peripheral vascular disease, including unspecified

**Note:** The routine use of genetic testing to screen patients treated with clopidogrel who are undergoing percutaneous coronary intervention (PCI) is not a benefit of Texas Medicaid.

### 9.2.40.2 Testing of Polymorphic 2D6

Pharmacogenetics testing of polymorphic 2D6 (procedure code 81226) may be considered medically necessary when all of the following conditions are met:

- Group one:
  - The client has never received genetic testing of the 2D6 alleles
  - The client has a diagnosis of Gaucher disease type 1
  - Treatment with eliglustat (Cerdelga®) is being considered
- Group two:
  - The client has never received genetic testing of the 2D6 alleles
  - The client has a diagnosis of Huntington’s disease
  - Treatment with tetrabenazine (Xenazine®) is being considered in a dosage greater than 50mg per day.

Prior authorization is not required for the initial pharmacogenetic testing of polymorphic 2D6 (procedure code 81226) that is performed on a client. Prior authorization is required for repeat testing.
9.2.40.3 Testing of Polymorphic 2C9
Pharmacogenetics testing of polymorphic 2C9 (procedure code 81227) requires prior authorization and may be considered for warfarin treatment and may be considered medically necessary when all of the following conditions are met:

- The client has never received genetic testing of the 2C9 alleles
- The client has never received warfarin (vitamin K antagonists) treatment
- The warfarin treatment will be used for one of the following diseases or conditions:
  - Irregular heartbeat or rhythm
  - Prosthetic (replacement or mechanical) heart valves
  - Myocardial infarction
  - Risk of venous thrombosis (swelling and blood clot in a vein)
  - Risk of pulmonary embolism (a blood clot in the lung)

9.2.40.4 Prior Authorization Requirements
Prior authorization is required for requests for pharmacogenetic testing for more than once in a lifetime. Prior authorization requests must be submitted on the Special Medical Prior Authorization (SMPA) Request Form. The form must be completed, signed, dated, and submitted by the prescribing or ordering provider.

Prior authorization requests from laboratories will not be processed. The requesting provider must share the prior authorization number with the laboratory submitting the claim.

The prior authorization request must include the following:

- Laboratory TPI in section D of the SMPA Request Form
- Proposed or current treatment plan, including the drug name, dosage, and frequency that support the medical necessity of the service requested
  - This information may be documented in the “Statement of medical necessity” field under Section C of the SMPA Request Form or submitted separately with the prior authorization request.
- For prior authorization of procedure code 81225, the ordering provider must include a statement on the SMPA Request Form attesting that the client has never received clopidogrel treatment.
- For prior authorization of procedure code 81227, the ordering provider must include a statement on the SMPA Request Form attesting that the client has never received warfarin treatment.

Prior authorization requests to repeat the same test (procedure code 81225, 81226, or 81227) will be reviewed by the medical director when one of the following criteria is met:

- The client has Huntington’s disease and a history of pharmacogenetic testing of 2D6 (procedure code 81226) for tetrabenzine treatment, and the new request is for the same testing of 2D6 but for eliglustat to treat Gaucher disease type 1.
- The client has Gaucher disease type 1 and a history of pharmacogenetic testing of 2D6 (procedure code 81226) for eliglustat treatment, and the new request is for the same testing of 2D6 but for tetrabenzine to treat Huntington’s disease.

Previous test results are unavailable. Every reasonable effort must be made to obtain the test results from the client’s provider or laboratory who previously ordered or conducted testing. Documentation of these efforts must be submitted with the prior authorization request.
9.2.40.5  Exclusions

The following services are not a benefit of Texas Medicaid:

- Pharmacogenetics tests of polymorphisms in a p450 superfamily other than 2D6, 2C19, or 2C9, which are performed for the purpose of aiding in the choice of drug or dose to increase efficacy or avoid toxicity, as they are considered experimental and investigational

- The routine clinical use of genetic testing to screen patients treated with clopidogrel who are undergoing percutaneous coronary intervention (PCI)

- The use of any of the 2D6, 2C19, or 2C9 tests for the following conditions, drugs, or treatments:
  - Opioid pain medicines (codeine, oxycodone, hydrocodone, tramadol, fentanyl, and methadone)
  - Selective serotonin reuptake inhibitors (SSRIs)
  - Selective norepinephrine reuptake inhibitors (SNRIs)
  - Beta blockers
  - Selective tricyclic antidepressants
  - Selective antipsychotic drugs
  - Efavirenz and other antiretroviral therapies for human immunodeficiency virus (HIV) infection
  - Immunosuppressants for organ transplantation
  - Aricept® (donepezil) for individuals with Alzheimer’s disease
  - p450 polymorphisms test panels for any of the 3 alleles 2C19, 2D6, or 2C9

9.2.41  Lung Volume Reduction Surgery (LVRS)

LVRS is a benefit for clients who are not high risk but have a presence of severe, upper-lobe emphysema (as defined by radiologist assessment of upper-lobe predominance on CT scan) or who are not high risk but have a presence of severe, non-upper-lobe emphysema with low exercise capacity.

*Note:* Clients who have low exercise capacity are those whose maximal exercise capacity is at or below 25 watts for women and 40 watts for men after completion of the pre-operative therapeutic program in preparation for LVRS. Exercise capacity is measured by incremental, maximal, symptom-limited exercise with a cycle ergometer utilizing a 5- or 10-watt-per-minute ramp on 30-percent oxygen after 3 minutes of unloaded pedaling.

LVRS must be performed in a facility that meets at least one of the following requirements:

- Certified under the Disease Specific Care Certification Program for LVRS by the Joint Commission on Accreditation of Health Care Organization

- Approved by Medicare as a lung or heart-lung transplant facility

The surgery must be both preceded and followed by a program of diagnostic and therapeutic services that are consistent with those provided in the National Emphysema Treatment Trial (NETT) and designed to maximize the client’s potential to successfully undergo and recover from surgery. The program must meet all of the following requirements:

- Include a 6- to 10-week series of at least 16, and no more than 20, pre-operative sessions, each lasting a minimum of 2 hours

- Include at least 6, and no more than 10, post-operative sessions, each lasting a minimum of 2 hours, within 8 to 9 weeks after the LVRS

- Be consistent with the care plan that was developed by the treating physician following the performance of a comprehensive evaluation of the client’s medical, psychosocial, and nutritional needs
• Be arranged, monitored, and performed under the coordination of the facility where the surgery takes place

Clients must have surgical clearance by a licensed cardiologist for any of the following conditions:
• Unstable angina
• Left ventricular ejection fraction (LVEF) cannot be estimated from the echocardiogram
• LVEF less than 45 percent
• Dobutamine-radionuclide cardiac scan indicates coronary artery disease or ventricular dysfunction
• Arrhythmia (more than 5 premature ventricular contractions (PVC) per minute)
• Cardiac rhythm other than sinus
• PVCs on electrocardiogram (EKG) at rest

For clients with cardiac ejection fraction less than 45 percent, there must be no history of congestive heart failure or myocardial infarction within six months of consideration for surgery.

Clients must have surgical clearance by a licensed pulmonologist, thoracic surgeon, and anesthesiologist after completion of pre-operative rehabilitation.

Procedure codes 32491, G0302, G0303, G0304, and G0305 are limited to one per rolling year per client for any provider.

Pre-operative pulmonary rehabilitation services for preparation for LVRS (procedure codes G0302, G0303, and G0304) and post-discharge pulmonary surgery services LVRS (procedure code G0305) will be restricted to diagnosis codes J430, J431, J432, J438, and J983.

Procedure code G0305 may be reimbursed only if a claim for LVRS (procedure code 32491) has been submitted within the past 12 months.

9.2.41.1 Prior Authorization for Lung Volume Reduction Surgery

LVRS must be prior authorized and is limited to clients who have severe emphysema, disabling dyspnea, and evidence of severe air trapping. The following documentation must be submitted with the request for prior authorization:
• The client’s history and physical examination is consistent with emphysema
• BMI less than 31.1 kg/m2 (men) or less than 32.3 kg/m2 (women)
• Pulmonary status that is stable with less than 20 mg prednisone (or equivalent) per day
• A radiographic high resolution computer tomography (HRCT) scan has been conducted that shows evidence of bilateral emphysema.
• The forced expiratory volume in one second (FEV1) (maximum of pre- and postbronchodilator values) is less than or equal to 45 percent of the predicted value. If the client is 70 years of age and older, FEV1 is 15 percent of the predicted value or more.
• The total lung capacity (TLC) greater than 100 percent predicted postbronchodilator
• Residual volume (RV) greater than 150 percent predicted postbronchodilator found on prerrehabilitation pulmonary function study.
• Arterial blood gas level (pre-rehabilitation):
  • Partial pressure of carbon dioxide (PaCO2) less than or equal to 60 mm Hg (PaCO2 less than or equal to 55 mm Hg if one mile above sea level)
  • Partial pressure of oxygen (PaO2) greater than or equal to 45 mm Hg on room air (PaO2 greater than or equal to 30 mm Hg if one mile above sea level)
• The plasma cotinine is less than or equal to 13.7 ng/ml (if the client is not using nicotine products) or the carboxyhemoglobin is less than or equal to 2.5 percent (if the client is using nicotine products).

• Nonsmoking for four months prior to initial interview and throughout evaluation for surgery

• Successful 6-minute walk test equal to or greater than 140 meters following pre-operative rehabilitation

• Successful completion of three minute unloaded pedaling in an exercise tolerance test both before and after pre-operative rehabilitation

To complete the prior authorization process, a provider must mail or fax the request to the TMHP Special Medical Prior Authorization Unit and include documentation of medical necessity.

• Requisition forms from the laboratory are not sufficient for verification of the personal and family history.

• Medical documentation that is submitted by the physician must verify the client’s diagnosis or family history.

Prior authorization is not required for the associated preoperative pulmonary surgery services for preparation for LVRS (procedure codes G0302, G0303, and G0304) or the associated postdischarge pulmonary surgery services after LVRS (procedure code G0305).

9.2.41.1.1 Noncovered Conditions

LVRS is not a benefit in any of the following clinical circumstances:

• A client with characteristics that carry a high risk for perioperative morbidity and/or mortality

• A disease that is unsuitable for LVRS

• A medical condition or other circumstance that makes it likely that the client will be unable to complete the preoperative and postoperative pulmonary diagnostic and therapeutic program required for surgery

• The client presents with FEV1 less than or equal to 20 percent of predicted value, and either a homogeneous distribution of emphysema on the CT scan or a carbon monoxide diffusing capacity of less than or equal to 20 percent of predicted value (a high-risk group identified in October 2001 by the NETT)

• The client satisfies the criteria outlined above and has severe, non-upper-lobe emphysema with a high-exercise capacity. High-exercise capacity is defined as a maximal workload at the completion of the preoperative diagnostic and therapeutic program that is above 25 watts for women or 40 watts for men (under the measurement conditions for cycle ergometry).

• A previous LVRS (laser or excision) on the same lung

• A pleural or interstitial disease which precludes surgery

• A giant bulla (greater than 1/3 the volume of the lung in which the bulla is located)

• A clinically significant bronchiectasis

• A pulmonary nodule requiring surgery

• A previous lobectomy

• Uncontrolled hypertension (systolic greater than 200 mm Hg or diastolic greater than 110 mm Hg)

• Oxygen requirement greater than 6 liters per minute during resting to keep oxygen saturation greater than or equal to 90 percent

• A history of recurrent infections with clinically significant production of sputum
• Unplanned weight loss greater than 10 percent within 3 months before the consideration of surgery
• Pulmonary hypertension, defined as the mean pulmonary artery pressure of 35 mmHg or greater on the right heart catheterization or peak systolic pulmonary artery pressure of 45 mmHg or greater. Right heart catheterization is required to rule out pulmonary hypertension if the peak systolic pulmonary artery pressure is greater than 45 mmHg on an echocardiogram
• Resting bradycardia (less than 50 beats per minute)
• Frequent multifocal premature ventricular contractions (PVCs) of complex ventricular arrhythmia or sustained supraventricular tachycardia (SVT)
• Evidence of a systemic disease or neoplasia that is expected to compromise survival

9.2.42  Diagnostic and Therapeutic Breast Procedures

Diagnostic, mastectomy, and breast reconstruction procedures are benefits of Texas Medicaid. These are physician-directed services including, but not limited to diagnostic and surgical breast procedures provided by physicians in the office, outpatient, or inpatient hospital settings, and external breast prostheses provided by durable medical equipment (DME) providers in the home setting.

Categories of service include:
• Diagnostic breast procedures
• Mastectomy
• Reconstructive breast procedures
• Treatment of complications of breast reconstruction
• External breast prostheses

9.2.42.1  Diagnostic Procedures

Diagnostic breast procedures for a condition or malignancy of the breast may include:
• Puncture aspiration
• Mastotomy
• Injection procedure for ductogram or galactogram
• Percutaneous biopsy, with or without imaging guidance
• Incisional biopsy
• Nipple exploration
• Excision of the following:
  • Lactiferous duct fistula
  • Benign or malignant breast lesion
  • Chest wall tumor

The following procedure codes may be reimbursed for diagnostic breast procedures:

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<td>19284</td>
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The following services are not benefits of Texas Medicaid:

- Mastectomy for a diagnosis of fibrocystic disease in the absence of documented risk factors.
- Cosmetic services performed primarily to improve appearance.
- Commercial or “decorative” tattooing.
- Replacement of external breast prostheses when the damage is due to abuse or neglect by the client, client’s family, or the caregiver.

### 9.2.42.2 Therapeutic Procedures

#### 9.2.42.2.1 Mastectomy Procedures

Mastectomy and partial mastectomy (e.g., lumpectomy, tylectomy, quadrantectomy, and segmentectomy) are benefits when it is medically necessary to remove a breast or portion of a breast for conditions including, but not limited to:

- Developmental abnormality
- Congenital defect
- Trauma or injury to chest wall
- Primary or secondary malignancy of the breast
- Carcinoma in situ of the breast

The following procedure codes for mastectomy are benefits of Texas Medicaid:

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Procedure codes 19301, 19302, 19303, 19305, 19306, and 19307 may be reimbursed without prior authorization for services rendered to male or female clients who are 18 years of age and older.

Prior authorization is required for services rendered to clients who are 17 years of age and younger.

Procedure codes 19303, 19305, 19306, and 19307 are limited to 1 service per breast per lifetime.

#### 9.2.42.2.2 Prophylactic Mastectomy

Prophylactic mastectomy is a benefit after a thorough assessment of a client’s unique risk factors, health, and the level of concern. Prophylactic mastectomy is limited to clients who are at moderate- to high-risk for the development of breast cancer.

Moderate- to high-risk clients are those who meet one or more of the following criteria for development of breast cancer:

- Current or previous diagnosis of breast cancer
- Family history of breast cancer in mother, sister, or daughter, especially before the age of 50
- Presence of any of the following genetic mutations:
  - Breast cancer gene 1 (BRCA1)
  - Breast cancer gene 2 (BRCA2)
  - Tumor protein 53 (TP 53)
  - Phosphatase and tensin homolog (PTEN)
  - Lobular carcinoma in situ (LCIS)
- Radiation therapy to the chest before a client reaches 30 years of age

Documentation that supports medical necessity for prophylactic mastectomy must include the information listed above.

Documentation that as a candidate for prophylactic mastectomy, the client has undergone counseling regarding cancer risks. Counseling must include assessment of all of the following:

- The client’s ability to understand the risks and long-term implications of the surgical procedure, and
- The client’s informed choice to proceed with the surgical procedure.

9.2.42.2.3 Mastectomy for Pubertal Gynecomastia

Mastectomy for pubertal gynecomastia is a benefit with prior authorization for males who are 20 years of age and younger. Procedure code 19300 may be reimbursed for mastectomy for pubertal gynecomastia.

The following documentation must be submitted with the prior authorization request for procedure code 19300:

- The gynecomastia classification (grade II, III, or IV) as defined by the American Society of Plastic Surgeons classification.
- Evidence that puberty is near completion, as indicated by the following:
  - 95 percent of adult height based on bone age, and
  - Tanner stage V has been achieved.
- Evidence that the client has been off gynecomastia inducing drugs or other substances for a minimum of one year when this is identified as the cause of the gynecomastia.
- Evidence of resolution as supported by appropriate test results and treatment for hormonal causes, including hyperthyroidism, estrogen excess, prolactinomas, and hypogonadism, for a minimum of one year when identified as the cause of the gynecomastia.
- Evidence of a psychiatric assessment performed by a psychiatrist or psychologist.
- Client’s history and treatment plan including planned surgical procedure and timelines.
- Identification of which breast or breasts, require mastectomy.

Documentation that supports medical necessity for mastectomy for pubertal gynecomastia must be maintained in the client’s medical record, and must include the following:

- A complete medical and family history, including:
  - Gynecomastia classification
  - Bone age
  - Tanner stage
  - Use of any gynecomastia inducing drugs or substances and date last ingested
  - Hormonal causes of gynecomastia, treatment, and length of treatment
  - Psychiatric assessment performed by a psychiatrist or psychologist and outcome
  - Affected breast or breasts
  - A thorough physical examination
  - Medically indicated laboratory testing and any other testing including results
9.2.42.3 **Breast Reconstruction**

Breast reconstruction may be performed in a single stage or several stages. Breast reconstruction is a benefit when all of the following criteria are met:

- The client has a documented history of one or more of the following:
  - Mastectomy
  - Congenital defect
  - Developmental abnormality
  - Trauma or injury to the chest wall
- The client meets age and gender criteria for the requested procedure.
- The physician has documented a treatment plan in the client's medical record that addresses the recommended breast reconstruction.
- Reconstruction to attain symmetry is required and may include a surgical procedure to the contralateral breast and may be either a reduction or an augmentation.

Procedure options for breast reconstruction following a mastectomy include, but are not limited to the following:

- Superficial inferior epigastric artery (SIEA) flap
- Deep inferior epigastric artery (DIEP) flap
- Transverse rectus abdominis myocutaneous (TRAM) flap
- Breast implants (saline or silicone)
- Reduction mammoplasty
- Mastopexy
- Reconstruction of the nipple or areola (small flaps)
- Tattooing to correct color defects of the skin
- Treatment for complications of breast reconstruction

Documentation that supports medical necessity for breast reconstruction, including tattooing, must include the following:

- Diagnosis resulting in the need for breast reconstruction,
- Date of mastectomy, when appropriate,
- Date of any previous breast reconstruction procedures, when appropriate,
- Treatment plan to include planned surgical procedures and timeline for completion, and
- When appropriate, identification of the complication.

All Medicaid services, including breast reconstruction after breast cancer surgery, are covered for Medicaid Breast and Cervical Cancer (MBCC) clients who are receiving active cancer treatment. "Active treatment" is defined as medical treatment following a cancer diagnosis that is intended to cure or otherwise treat a diagnosed cancer.

Active treatment may include some or all of the following:

- Surgery
- Chemotherapy
- Radiotherapy
• Medication (e.g., ongoing hormonal treatments for estrogen and progesterone breast cancer)
• Active disease surveillance for triple negative receptor breast cancer

Reconstructive surgery (e.g., breast reconstruction) is considered “active treatment” if it is intended to permanently correct a physical condition resulting from either the diagnosed cancer or the treatment of the diagnosed cancer.

Ongoing treatment of a persistent condition resulting from a diagnosed cancer or treatment of a diagnosed cancer is not considered “active treatment” if cancer is no longer present or in need of treatment.

The following breast reconstruction procedure codes may be reimbursed without prior authorization for services rendered to clients who are 18 years of age and older:

<table>
<thead>
<tr>
<th>Procedure Codes</th>
</tr>
</thead>
<tbody>
<tr>
<td>11970</td>
</tr>
<tr>
<td>19364</td>
</tr>
</tbody>
</table>

*Procedure codes are limited to females only.

Prior authorization is required for services rendered to clients who are 17 years of age and younger or when the client does not meet gender or age criteria.

Procedure codes 11920, 11921, and 11922 may be reimbursed when performed as part of breast reconstruction.

Breast reconstruction claims denied for no history of previous mastectomy may be appealed with supporting documentation indicating the date of mastectomy, or the identified trauma, injury, or congenital or developmental abnormality.

9.2.42.3.1 Tattooing to Correct Color Defects of the Skin

Tattooing to correct color defects of the skin (procedure codes 11920, 11921, and 11922) are limited to two services per lifetime.

Tattooing claims denied for no history of breast reconstruction may be appealed with supporting documentation indicating the date of breast reconstruction, or the identified trauma, injury, or congenital or developmental abnormality.

9.2.42.3.2 Treatment for Complications of Breast Reconstruction

The treatment of complications related to breast reconstruction may be reimbursed using procedure codes 19328, 19330, 19370, 19371, and 19380.

Procedure codes 19328, 19330, 19370, and 19371 may be reimbursed for services rendered to female clients only.

9.2.42.3.3 Chest Wall Procedures

Excision of chest wall tumors may be reimbursed using procedure codes 21601, 21602, and 21603.

Procedure code 21603 is limited to once per lifetime.

9.2.42.3.4 External Breast Prostheses

External breast prostheses are available through a durable medical equipment (DME) provider for a female client with a history of a medically necessary mastectomy procedure.
The following procedure codes may be reimbursed for external breast prosthesis services rendered to female clients of any age:

<table>
<thead>
<tr>
<th>Procedure Code</th>
<th>Limitation</th>
</tr>
</thead>
<tbody>
<tr>
<td>L8000</td>
<td>4 per rolling year</td>
</tr>
<tr>
<td>L8001</td>
<td>4 per rolling year, per modifier</td>
</tr>
<tr>
<td>L8002</td>
<td>4 per rolling year</td>
</tr>
<tr>
<td>L8010</td>
<td>8 per rolling year</td>
</tr>
<tr>
<td>L8015</td>
<td>2 per rolling year</td>
</tr>
<tr>
<td>L8020</td>
<td>1 per 6 rolling months</td>
</tr>
<tr>
<td>L8030</td>
<td>per 2 rolling years</td>
</tr>
<tr>
<td>L8031</td>
<td>per 2 rolling years</td>
</tr>
<tr>
<td>L8032</td>
<td>8 per rolling year</td>
</tr>
<tr>
<td>L8033</td>
<td>8 per rolling year</td>
</tr>
<tr>
<td>L8035</td>
<td>Requires prior authorization</td>
</tr>
<tr>
<td>L8039</td>
<td>Requires prior authorization</td>
</tr>
</tbody>
</table>

Replacement of external breast prostheses may be considered at any time, through the prior authorization with documentation.

For a new or replacement external breast prosthesis procedure code outside the limitations, all of the following documentation must be submitted with the prior authorization request:

- The client’s diagnosis
- Documentation of medical necessity for the requested prosthesis
- Documentation indicating the reason for recommending the requested prosthesis

When requesting a prior authorization for procedure code L8035 (custom prosthesis), all of the following documentation must be submitted with the prior authorization request:

- The client’s diagnosis
- Documentation of medical necessity for the requested prosthesis
- Documentation indicating the reason for recommending the requested prosthesis

When requesting a prior authorization for procedure code L8039 (other prosthesis), all of the following documentation must be submitted with the prior authorization request:

- A clear, concise description of the breast prosthesis requested
- Reason for recommending the requested prosthesis
- A CPT or HCPCS procedure code, which is comparable to the procedure being requested
• Documentation that this breast prosthesis is not investigational or experimental
• The provider’s intended fee for the requested prosthesis

### 9.2.42.4 Prior Authorization Requirements for Diagnostic and Therapeutic Breast Procedures

Prior authorization is not required for the following when all of the following criteria are met:

- The procedure is a mastectomy or breast reconstruction for clients who are 18 year of age or older.
- The request is for one of the following external breast prosthesis procedure codes: L8000, L8001, L8002, L8010, L8015, L8020, or L8030.
- The procedure is for partial mastectomy procedure codes 19301 and 19302 for clients of any age.

Prior authorization is required for the following:

- Mastectomy or breast reconstruction when the client is 17 years of age or younger, or does not meet gender criteria
- Mastectomy for pubertal gynecomastia
- Procedure code 19499 (unlisted procedure)
- External breast prosthesis procedure codes L8035 (custom prosthesis) and L8039 (other prosthesis)
- Any request for new or replacement external breast prosthesis outside of the limitations

### 9.2.42.4.1 Unlisted Breast Procedure

All of the following documentation must be submitted for procedure code 19499 with the prior authorization request:

- A clear, concise description of the procedure to be performed
- Reason for recommending this particular procedure
- A Current Procedural Terminology (CPT) or Healthcare Common Procedure Coding System (HCPCS) procedure code, which is comparable to the procedure being requested
- Documentation that this procedure is not investigational or experimental
- Place of service the procedure is to be performed
- The provider’s intended fee for this procedure

### 9.2.42.4.2 Documentation Requirements

In addition to documentation requirements outlined in the “Prior Authorization Requirements” section above, the following requirements apply:

- All services are subject to retrospective review. Documentation in the client’s medical record must be maintained by the physician and must support the medical necessity for the services provided.
- Services not supported by documentation are subject to recoupment.

### 9.2.43 Neurostimulators

Neurostimulator and neuromuscular stimulator procedures and the rental or purchase of devices and associated supplies, such as leads and form fitting conductive garments are a benefit of Texas Medicaid when medically necessary.

Neurostimulator devices are considered DME, so providers must complete both the Home Health (Title XIX) DME/Medical Supplies Physician Order Form (Title XIX Form) to prescribe the DME and the DME Certification and Receipt Form to show receipt of the DME by the client. Both forms must be maintained in the client’s medical record.
9.2.43.1 Prior Authorization for Neurostimulators

All devices and related procedures for the initial application or surgical implantation of the stimulator or neuromuscular stimulator device require prior authorization.

Requests for prior authorization must be submitted to the Special Medical Prior Authorization (SMPA) department with documentation supporting the medical necessity of the requested device. Providers may use the Special Medical Prior Authorization (SMPA) Request Form when they submit requests to the SMPA department.

To avoid unnecessary denials, the physician must provide correct and complete information including documentation for medical necessity of the equipment and/or supplies requested. The physician must maintain documentation of medical necessity in the client’s medical record. The requesting provider may be asked for additional information to clarify or complete a request for the equipment and/or supplies. Prior authorization requests for all neurostimulators and related procedures must include the provider identifiers for both the surgeon and the facility.

A neurostimulator device that has been purchased is anticipated to last a maximum of five years and may be considered for replacement when five years have passed and/or the equipment is no longer repairable. At that time, replacement of the device will be considered. Replacement devices require prior authorization. Replacement of equipment may also be considered when loss or irreparable damage has occurred. A copy of the police or fire report when appropriate, and the measures to be taken to prevent reoccurrence must be submitted.

9.2.43.2 Neuromuscular Electrical Stimulation (NMES)

NMES application and the rental or purchase of devices and conductive garments are a benefit of Texas Medicaid when medically necessary and prior authorized. Prior authorization requests for NMES must include documentation of a spinal cord injury or disuse atrophy that is refractory to conventional therapy.

NMES may be reimbursed using the following procedure codes:

<table>
<thead>
<tr>
<th>Procedure Codes</th>
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</thead>
<tbody>
<tr>
<td>64580</td>
</tr>
</tbody>
</table>

9.2.43.2.1 NMES Rental

The rental of a NMES device may be considered before purchase and is limited to a one-month trial period with consideration for one additional month’s trial with documentation of medical necessity. Supplies are considered to be part of the rental and will not be separately reimbursed. Garments may be considered for reimbursement during the rental period when medically necessary.
9.2.43.2.2 NMES Purchase

The purchase of a NMES device is limited to once per five years, and may be reimbursed when there is documentation of successful test stimulation (during rental or other therapeutic period) that showed improvement as measured by the following:

- A demonstrated increase in range of motion.
- The client’s improved ability to complete activities of daily living or perform activities outside the home.

Garments may be considered for reimbursement during the purchase period when medically necessary.

9.2.43.2.3 NMES for Muscle Atrophy

NMES may be reimbursed when used to treat muscle disuse atrophy when brain, spinal cord, and peripheral nerve supply to the muscle is intact, as well as other non-neurological conditions. Examples of NMES treatment for non-neurological conditions include, but are not limited to, casting or splinting of a limb, contracture due to scarring of soft tissue as in burn lesions, and hip replacement surgery until orthotic training begins.

9.2.43.2.4 NMES for Walking in Clients with Spinal Cord Injury (SCI)

The type of NMES that is used to enhance the ability to walk of SCI clients is commonly referred to as functional electrical stimulation (FES). These devices are surface units that use electrical impulses to activate paralyzed or weak muscles in precise sequence.

The use of NMES/FES is limited to SCI clients who have completed a training program which consists of at least 32 physical therapy sessions with the device over a period of three months.

The trial period of physical therapy will enable the treating physician to properly evaluate the client’s ability to use NMES/FES devices frequently and for the long term.

Physical therapy necessary to perform this training must be directly performed by the physical therapist as part of a one-on-one training program. The goal of physical therapy must be to train SCI clients on the use of NMES/FES devices to achieve walking, not to reverse or retard muscle atrophy.

NMES/FES is a benefit for SCI clients who have all of the following characteristics:

- Clients with intact lower motor unit (L1 and below) (both muscle and peripheral nerve).
- Clients with muscle and joint stability for weight bearing at upper and lower extremities that can demonstrate balance and control to maintain an upright posture while standing independently for at least three minutes.
- Clients who demonstrate brisk muscle contraction to NMES and have sensory perception electrical stimulation sufficient for muscle contraction.
- Clients who possess high motivation, commitment, and cognitive ability to use such devices for ambulation, as established by provider interview and documentation.
- Clients who can transfer independently.
- Clients who can demonstrate hand and finger function to manipulate controls.
- Clients with at least six-month post recovery spinal cord injury and restorative surgery.
- Clients with hip and knee degenerative disease and no history of long bone fracture secondary to osteoporosis.

NMES and FES used for walking is not a benefit in SCI clients with any of the following:

- Cardiac pacemakers
- Severe scoliosis or severe osteoporosis
- Skin disease or cancer at area of stimulation
- Irreversible contracture
- Autonomic dyslexia

9.2.43.3 Transcutaneous Electrical Nerve Stimulation (TENS)

TENS involves the attachment of a transcutaneous nerve stimulator to the surface of the skin over the peripheral nerve to be stimulated.

TENS may be reimbursed for the treatment of acute postoperative pain or chronic pain that is refractory to conventional therapy.

TENS may be reimbursed using the following procedure codes:

<table>
<thead>
<tr>
<th>Procedure Codes</th>
</tr>
</thead>
<tbody>
<tr>
<td>E0720</td>
</tr>
</tbody>
</table>

9.2.43.3.1 TENS Rental

Rental of a TENS device will be considered for prior authorization when there is documentation of a condition that indicates acute postoperative pain or chronic pain that is refractory to conventional therapy.

The rental of a TENS device is limited to one-month trial period with consideration for one additional month’s trial with documentation of medical necessity. Supplies, such as lead wires and electrodes, are considered to be part of the rental and will not be separately reimbursed. Garments may be considered during the rental period when medically necessary.

When the TENS device is rented for a trial period rather than supplied by the provider, the combined payment made for professional services and the rental of the stimulator must not exceed the amount which would be reimbursed for the total service, including the stimulator, if furnished by the provider alone.

9.2.43.3.2 TENS Purchase

The purchase of a TENS device is limited to once every five years and may be reimbursed with prior authorization when there is documentation of the following:

- A condition that indicates chronic pain that is refractory to conventional therapy.
- A successful test stimulation (during rental or other therapeutic period) that showed improvement as measured by demonstrated increase in range of motion.
- The client’s improved ability to complete activities of daily living or perform activities outside the home.

9.2.43.4 NMES and TENS Garments

The rental of the NMES/TENS garment is not covered during the trial rental period unless the client has a documented skin problem prior to the start of the trial period, and HHSC or its designee determines that use of such an item is medically necessary for the client based on the documentation submitted.

The purchase of conductive garments for NMES/TENS devices may be considered when:

- The garment has been prescribed by a physician for use in providing covered NMES/TENS treatment.
- A NMES/TENS device has been purchased for the client’s use.
- The conductive garment is necessary for one of the medical indications outlined below:
• The client cannot manage without the conductive garment because there is such a large area or so many sites to be stimulated and the stimulation would have to be delivered so frequently that it is not feasible to use conventional electrodes, adhesive tapes, and lead wires.

• The client cannot manage the treatment for chronic intractable pain without the conductive garment because the areas or sites to be stimulated are inaccessible with the use of conventional electrodes, adhesive tapes, and lead wires.

• The client has a documented medical condition such as skin problems that preclude the application of conventional electrodes, adhesive tapes, and lead wires.

9.2.43.5 NMES and TENS Supplies
Supplies for purchased devices are limited as follows:
• If additional electrodes are required, procedure code A4556 may be considered for reimbursement at a maximum of 15 per month.
• If additional lead wires are required, procedure code A4557 may be considered for reimbursement at a maximum of 2 per month.
• Procedure code A4595 is limited to 1 per month.

Supplies are included in the rental and will not be reimbursed separately.
Supply procedure codes A4556, A4557, or A4595 may be reimbursed for clients with a purchased device and a claims history of an NMES/TENS procedure within the past five years. Providers must maintain documentation in the client’s medical record that a device has been purchased. Additional documentation such as the purchase date, serial number, and purchasing entity of the device may be required.

9.2.43.6 Diaphragm-Pacing Neuromuscular Stimulation
Diaphragm-pacing neuromuscular stimulation is a benefit of Texas Medicaid when medically necessary and prior authorized.

Diaphragm-pacing neuromuscular stimulation is the electrical stimulation to one or both of the phrenic nerves or to the phrenic motor point regions of the diaphragm muscles that cause contraction of one or both of the two hemidiaphragms rhythmically to produce inspiration.

Diaphragm-pacing neuromuscular stimulation may be reimbursed when billed with procedure codes 64575 and 64590.

9.2.43.6.1 Prior Authorization for Diaphragm-Pacing Neuromuscular Stimulation
The surgical implantation of the diaphragm-pacing neuromuscular stimulator and purchase of a device are considered for prior authorization when medically necessary for individuals with severe, chronic respiratory failure that requires mechanical ventilation for any of the following reasons:
• Improvement of ventilatory function in stable, non-acute members with spinal cord injury (SCI) with high quadriplegia at or above C-3
• Alveolar hypoventilation, either primary or secondary to brainstem disorder
• Amyotrophic lateral sclerosis

All of the following criteria must be met:
• The phrenic nerves are viable
• Diaphragmatic function is sufficient to accommodate chronic stimulation
• Pulmonary function is known to be adequate
• The client has normal chest anatomy, a normal level of consciousness, and has the ability to participate in and complete the training and rehabilitation associated with the use of the device
9.2.43.7 Dorsal Column Neurostimulator (DCN)

DCN involves the surgical implantation of neurostimulator electrodes within the dura mater (endodural) or the percutaneous insertion of electrodes in the epidural space. The neurostimulator system stimulates pain-inhibiting nerve fibers, masking the sensation of pain with a tingling sensation (paresthesia).

DCN implantation may be reimbursed using procedure codes 63650, 63655, or 63685.

Conditions that may indicate chronic intractable pain include, but are not limited to, the following:

- Post-amputation “ghost” pain
- Cancer with bone metastasis
- Causalgia of upper/lower limb
- Herniated disc
- Radiculitis
- Spinal stenosis
- Spinal surgery
- Tic douloureux (trigeminal neuralgia)

9.2.43.7.1 Prior Authorization for Dorsal Column Neurostimulators

DCN electrode implantation and the purchase of devices is a benefit of Texas Medicaid when medically necessary and prior authorized.

The surgical implantation of DCN device may be considered for prior authorization for clients who have chronic intractable pain with documentation that indicates the following:

- Other treatment modalities, including pharmacological, surgical, physical, and/or psychological therapies, have been tried and shown to be unsatisfactory, unsuitable, or contraindicated for the client.
- The client has undergone careful screening, evaluation, and diagnosis by a multidisciplinary team prior to implantation.
- There has been evidence of pain relief during a trial period for DCN with a temporarily implanted electrode or electrodes preceding the permanent implantation.

Note: A trial period including device and supplies is considered part of DCN procedures and will not be separately reimbursed.

- All the facilities, equipment, and professional and support personnel required for the proper diagnosis, treatment, training, and the client’s follow-up are available.

9.2.43.8 Gastric Electrical Stimulation (GES)

GES involves electrical stimulation of the lower stomach (antrum) with a fully implantable system that consists of two unipolar intramuscular leads (thin wires) and a neurostimulator device.

GES is a benefit of Texas Medicaid when medically necessary and prior authorized for the treatment of chronic intractable nausea and vomiting that is secondary to gastroparesis that has proven to be refractory to medical management.

GES may be reimbursed with procedure codes 43647, 43881, and 64590.

GES is a benefit for Texas Medicaid clients with the following conditions:

- Organic obstruction or pseudo-obstruction
- A primary eating or swallowing disorder
• Chemical dependency
• Pregnancy

9.2.43.8.1 Prior Authorization for GES

The surgical implantation of a GES and purchase of a device are considered for prior authorization for chronic intractable nausea and vomiting secondary to gastroparesis of diabetic or idiopathic etiology when all of the following criteria are met:

• Gastric emptying is significantly delayed as documented by standard scintigraphic imaging of solid food.

• Patient is refractory or intolerant of two out of three classes of prokinetic medications and two out of three antiemetic medications.

• The client’s nutritional status is sufficiently low that all of the following criteria for total parenteral nutrition are met:
  • Adequate trials of dietary adjustment, oral supplements, or tube enteral nutrition have demonstrated that the patient can receive no more than 30 percent of his/her caloric needs orally and/or by tube.
  • The patient must be in a stage of wasting as indicated by all of the following:
    • Weight is significantly less than normal body weight for a patient’s height and age in comparison with pre-illness weight.
    • Serum albumin is less than 3.4 grams.
    • BUN is less than 10 mg.
    • Phosphorus level is less than 2.5 mg.

9.2.43.9 Intracranial Neurostimulators

The surgical implantation, revision, and removal of intracranial deep brain stimulators (DBS) are a benefit for the relief of chronic intractable pain when more conservative methods, such as TENS, PENS, or pharmacological management have failed or were contraindicated.

Intracranial neurostimulation may be reimbursed using the following procedure codes:

<table>
<thead>
<tr>
<th>Procedure Codes</th>
</tr>
</thead>
<tbody>
<tr>
<td>61781</td>
</tr>
</tbody>
</table>

9.2.43.9.1 Prior Authorization for Intracranial Neurostimulators

Intracranial neurostimulation involves the stereotactic implantation of electrodes in the brain and is a benefit of Texas Medicaid when medically necessary and prior authorized.

The surgical implantation and purchase of an intracranial neurostimulation device may be considered for prior authorization for chronic intractable pain or treatment of intractable tremors.

Requests for prior authorization must include documentation of the following:

• Other treatment modalities, including pharmacological, surgical, physical, and psychological therapies, have been tried and shown to be unsatisfactory, unsuitable, or contraindicated for the client.

• The client has undergone careful screening, evaluation, and assessment by a multidisciplinary team prior to implantation.

• The client has reported pain relief with a temporarily implanted electrode preceding the permanent implantation.
• All the facilities, equipment, and support personnel required for the proper assessment, treatment, training, and client’s follow-up are available.

Prior authorization will not be given for the treatment of motor function disorders such as multiple sclerosis; however, the implantation, revision, and removal of deep brain stimulators may be reimbursed for the treatment of intractable tremors due to the following:

• Idiopathic Parkinson’s disease
• Essential tremor

9.2.43.10 Pelvic Floor Stimulation

Purchase of a non-implantable pelvic floor stimulator (procedure code E0740) is a benefit of Texas Medicaid for the treatment of stress or urge incontinence in clients who have failed conservative treatment, such as Kegel exercises, behavior management, bladder training, or medication.

Purchase of the pelvic floor stimulator device is limited to once per five years. All accessories and supplies are considered part of the purchase price and are not reimbursed separately.

9.2.43.10.1 Prior Authorization for Pelvic Floor Stimulation

Prior authorization is required for the purchase of a pelvic floor stimulator device.

Documentation submitted with the prior authorization request must demonstrate that the client:

• Has a diagnosis of stress or urge incontinence.
• Has completed a six-month trial of pelvic muscles exercises with no significant clinical improvement.

9.2.43.11 Percutaneous Electrical Nerve Stimulation (PENS)

PENS is a benefit of Texas Medicaid when medically necessary and prior authorized. Devices and supplies are considered a part of the service and are not separately reimbursable.

PENS is a diagnostic procedure for the treatment of chronic pain involving the stimulation of peripheral nerves by a needle electrode inserted through the skin.

9.2.43.11.1 Prior Authorization for PENS

PENS services may be reimbursed with prior authorization for clients who meet the following criteria:

• The client has a diagnosis that indicates chronic pain, which is refractory to conventional therapy.
• Treatment with TENS has failed or is contraindicated for the client.

PENS may be reimbursed using the following procedure codes: 64553, 64555, or 64590. The revision or removal of a peripheral neurostimulator used in PENS therapy may be reimbursed without prior authorization using procedure code 64595.

9.2.43.12 Sacral Nerve Stimulators (SNS)

SNS are a benefit of Texas Medicaid when medically necessary and prior authorized. SNS implantation may be reimbursed using procedure code 64561, 64581, or 64590.

SNS involves the use of pulse generators that transmit electrical impulses to the sacral nerves through a surgically implanted wire for treatment of urinary retention, urinary frequency, and urinary/fecal incontinence.

9.2.43.12.1 Prior Authorization for SNS

The surgical implantation of SNS and purchase of a device may be considered for prior authorization with the following:

• Urinary incontinence secondary to urethral instability and/or detrusor muscle instability.
• Chronic voiding dysfunction.
• Non-obstructive urinary retention.
• Fecal incontinence.

Additionally, the medical record of the client must have documentation of the following:

• The urinary retention, urinary frequency, and urinary/fecal incontinence are refractory to conventional therapy (documented behavioral, pharmacological, and/or surgical corrective therapy).
• The client is an appropriate surgical candidate such that implantation with anesthesia can occur.

9.2.43.13 Vagal Nerve Stimulators (VNS)

VNS are a benefit of Texas Medicaid when medically necessary and prior authorized, for the treatment of intractable partial onset seizures.

VNS involves the use of devices that deliver electrical pulses to the cervical portion of the vagus nerve by an implanted generator.

9.2.43.13.1 Prior Authorization for VNS

The surgical implantation and purchase of VNS devices may be considered for prior authorization for clients with partial onset intractable seizures when there is failure, contraindication, or intolerance to all suitable medical and pharmacological management.

The surgical implantation of VNS may be reimbursed using procedure code 61885, 61886, 64553, or 64568.

VNS are not a benefit of Texas Medicaid in the following cases:

• For the treatment of clients with an absent left vagus nerve
• For the treatment of clients with depression
• For the treatment of clients with diseases or conditions with a poor prognosis or are progressively terminal in nature

Incapacities due to intellectual disabilities or cerebral palsy may confound the assessment of benefits resulting from VNS. When a diagnosis of intellectual disabilities or cerebral palsy exists, the treating physician must document in the client's medical record how VNS will measurably benefit the client in spite of intellectual disabilities or cerebral palsy.

9.2.43.14 Prior Authorization of Neurostimulator Devices Procedure Codes

The following device procedure codes may be reimbursed with prior authorization:

<table>
<thead>
<tr>
<th>Procedure Codes</th>
</tr>
</thead>
<tbody>
<tr>
<td>L8681</td>
</tr>
</tbody>
</table>

Neuromuscular devices and the implantation codes must be billed on the same day by any provider.

To identify the service as a VNS device, procedure code L8686 must be submitted with modifier TG. Only one similar device code may be reimbursed per date of service for any provider.

9.2.43.15 Supplies for Neurostimulators

Supply procedure codes A4290, C1883, C1897, L8680, and L8696 may be reimbursed for clients with a purchased device and a claims history of a prior neurostimulator or neuromuscular stimulator implantation within the past five years. Providers must maintain documentation in the client’s medical record that a device has been purchased. Additional documentation such as the purchase date, serial number and purchasing entity of the initial implantable device may be required. Supplies for implantable devices
may be considered for reimbursement on appeal with documentation of a prior neurostimulator or a neuromuscular stimulator implantation procedure for clients with a history that is more than five years or for those who have a neurostimulator that was not received through Texas Medicaid.

To identify the service as a VNS implantable electrode, procedure code L8680 must be submitted with modifier TG.

9.2.43.16  Electrocorticogram of Implanted Neurostimulator
Electrocorticogram (procedure code 95836) is a benefit of Texas Medicaid and may be reported only once per each 30 day period.

9.2.43.17  Electronic Analysis for Neurostimulators
The following procedure codes may be reimbursed without prior authorization for the electronic analysis of the implanted neurostimulator and neuromuscular stimulation:

<table>
<thead>
<tr>
<th>Procedure Codes</th>
</tr>
</thead>
<tbody>
<tr>
<td>95970 95971 95972 95976 95977 95980 95981 95982 95983 95984</td>
</tr>
</tbody>
</table>

9.2.43.18  Revision or Removal of Neurostimulator Devices
The revision or removal of implantable neurostimulators may be reimbursed without prior authorization using the following procedure codes:

<table>
<thead>
<tr>
<th>Procedure Codes</th>
</tr>
</thead>
<tbody>
<tr>
<td>43648 43882 61781 63661 63662 63663 63664 63688 61880 61888</td>
</tr>
<tr>
<td>64569 64570 64585 64595</td>
</tr>
</tbody>
</table>

9.2.43.19  Noncovered Neurostimulator Services
The following services are not a benefit of Texas Medicaid:

- VNS is not a benefit when provided for the treatment of depression.
- Neurostimulation and neuromuscular stimulation services for indications other than those outlined above.

9.2.44  Newborn Services
The newborn period is defined as the time from birth through 28 days of life. This section addresses routine newborn care, attendance at delivery, newborn resuscitation, neonatal critical care, and intensive (noncritical) low birth weight services.

Retrospective review may be performed to ensure documentation supports the medical necessity of the service and any modifier used when billing a claim.

All newborn E/M procedure codes must have a newborn outcome diagnosis code included on the claim. Modifier 25 may be used to identify a significant separately identifiable E/M provided on the same day by the same physician as a procedure or other service. Documentation that supports the provision of a significant, separately identifiable E/M service must be maintained in the client’s medical record and made available to Texas Medicaid upon request.

Physician standby (procedure code 99360) is not a benefit.

**Note:** Some of the services addressed in this section may also be used for care beyond 28 days of life.

**Refer to:** Subsection 9.2.56, “Physician Evaluation and Management (E/M) Services” in this handbook.
Refer to: Subsection 2.2.23.13, “Cardiorespiratory Monitor (CRM)” in the *Durable Medical Equipment, Medical Supplies, and Nutritional Products Handbook (Vol. 2, Provider Handbooks)* for authorization of cardiorespiratory monitors.

### 9.2.44.1 Circumcisions for Newborns

Texas Medicaid may provide reimbursement for circumcisions billed with procedure code 54150 or procedure code 54160.

### 9.2.44.2 Hospital Visits and Routine Care

The following procedure codes may be reimbursed for neonatal care and intensive care services:

<table>
<thead>
<tr>
<th>Service</th>
<th>Procedure Code(s)</th>
<th>Benefit(s) and Limitation(s)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Initial hospital E/M admission</td>
<td>99221 99222 99223</td>
<td>If the client is readmitted within the first 28 days of life, the provider must bill an initial hospital evaluation and management (E/M) admission. Reimbursed one per day, any provider.</td>
</tr>
<tr>
<td>Hospital discharge</td>
<td>99238 99239</td>
<td>Reimbursed for the client’s discharge from the hospital.</td>
</tr>
<tr>
<td>Subsequent hospital and hospital consultation services</td>
<td>99251 99252 99253 99254 99255</td>
<td>Services for a client who is not critically ill and unstable but who happens to be in a critical care unit must be reported using subsequent hospital codes (99478, 99479, and 99480) or hospital consultation codes (99251, 99252, 99253, 99254, and 99255).</td>
</tr>
<tr>
<td>Initial newborn care</td>
<td>99460*</td>
<td>May be reimbursed once per lifetime, any provider.</td>
</tr>
<tr>
<td>Normal newborn care</td>
<td>99461*</td>
<td>May be reimbursed once per lifetime, any provider. Subsequent visits must be billed using an appropriate visit code based on the place of service.</td>
</tr>
<tr>
<td>Subsequent hospital care</td>
<td>99462</td>
<td>Reimbursable once per day in the hospital and limited to a total of seven days. Restricted to clients who are birth through seven days of age. If the client is diagnosed with a condition that requires more complex care and/or must stay more than 8 days, the provider must bill subsequent neonatal and pediatric care critical or intensive care (procedure codes 99469, 99478, 99479, or 99480). If the client is readmitted, the provider must bill an initial hospital E/M admission (procedure code 99221, 99222, 99223, or 99468) and the appropriate code for inpatient neonatal critical care (procedure code 99469). Procedure code 99462 is not reimbursable in the birthing center.</td>
</tr>
<tr>
<td>Newborn admission and discharge, same date</td>
<td>99463**</td>
<td>May be reimbursed once per lifetime when submitted by any provider. Reimbursed for newborns who are admitted and discharged on the same day from the hospital or birthing room setting (either hospital or birthing center).</td>
</tr>
<tr>
<td>Attendance at delivery</td>
<td>99464</td>
<td>May be reimbursed once, and only on the day of delivery, when billed by a physician other than the delivering physician.</td>
</tr>
</tbody>
</table>

* Newborn examinations billed with procedure codes 99460, 99461, and 99463 may be counted as a THSteps periodic medical checkup when all necessary components are completed and documented in the medical record.

** If the client is readmitted within the first 28 days of life, the provider must bill an initial hospital evaluation and management (E/M) admission (procedure code 99221, 99222, or 99223).
Note: Services for a newborn’s unsuccessful resuscitation may be billed under the mother’s Texas Medicaid number using procedure code 99499.

Refer to: Section 5, “THSteps Medical” in the Children’s Services Handbook (Vol. 2, Provider Handbooks).

Subsection 5.3.9, “Newborn Examination” in the Children’s Services Handbook (Vol. 2, Provider Handbooks) for a list of the required components for an initial THSteps exam.

Retrospective review may be performed to ensure documentation supports the medical necessity of the service and any modifier used when billing a claim.

In the following table, procedure codes in Column A will be denied when billed with the same date of service by the same provider as a procedure code in Column B:

<table>
<thead>
<tr>
<th>Column A (Denied)</th>
<th>Column B</th>
</tr>
</thead>
<tbody>
<tr>
<td>99238, 99239</td>
<td>99460, 99461, 99463</td>
</tr>
<tr>
<td>99462</td>
<td>99238, 99239</td>
</tr>
<tr>
<td>36410, 96361, 99292, 99307, 99354, 99355, 99356, 99357</td>
<td>99468, 99469</td>
</tr>
</tbody>
</table>

* Newborn examinations billed with procedure codes 99460, 99461, and 99463 may be counted as a THSteps periodic medical checkup when all necessary components are completed and documented in the medical record.

** If the client is readmitted within the first 28 days of life, the provider must bill an initial hospital evaluation and management (E/M) admission (procedure code 99221, 99222, or 99223).
9.2.44.3 Newborn Hearing Screening

The newborn hearing screening procedure is a screening procedure, not diagnostic, and will not be reimbursed separately from the usual inpatient newborn delivery payment. Special investigations and examination codes are not appropriate for use with hearing screening of infants. For more information on newborn hearing screening, providers may contact:

Texas Early Hearing Detection and Intervention
PO Box 149347, MC-1918
Austin, TX. 78714-9347
1-512-458-7111, Ext. 2600
www.dshs.texas.gov/tehdi

Subsection 5.3.11.2.3, “Hearing Screening” in the Children's Services Handbook (Vol. 2, Provider Handbooks) for additional information about hearing screenings.

9.2.45 Occupational Therapy (OT) Services

Occupational therapy (OT) is a payable benefit to physicians.

Refer to: Section 4, “Therapy Services Overview” in the Physical Therapy, Occupational Therapy, and Speech Therapy Services Handbook (Vol. 2, Provider Handbooks) for information about occupational therapy services provided by a physician.

9.2.46 Ophthalmology

When an ophthalmologist sees a client for a minor condition that does not require a complete eye exam, such as conjunctivitis, providers are to use the appropriate office E/M code.

Providers are to use the eye exam procedure codes with a diagnosis of ophthalmological disease or injury.


9.2.46.1 Corneal Transplants

Corneal transplants are benefits of Texas Medicaid. Corneal transplants are subject to global surgery fee guidelines. Procedure codes 65710, 65730, 65750, 65755, 65756, and 65757 are used for this surgery.

Bioengineered cornea transplants remain investigational at this time and are not considered for reimbursement under Texas Medicaid.

Procurement of the cornea is not reimbursed separately.
9.2.46.2 Eye Surgery by Laser

Eye surgery by laser is a benefit of Texas Medicaid when medically necessary and meets the conditions and limitations stated in this section.

Authorization is not required for eye surgery by laser.

All procedure codes in this section are subject to multiple surgery guidelines. For bilateral procedures, the following modifiers must be added to the claim to indicate that the procedures were performed on the right and left eyes:

- Modifier RT to indicate the right eye
- Modifier LT to indicate the left eye

All procedures may be reimbursed only to physicians and are limited to reimbursement once every 90 days for the same eye with the exception of infants from birth through 23 months of age. Procedures performed on infants from birth through 23 months of age are not subject to any frequency restrictions.

9.2.46.2.1 Other Eye Surgery Procedures

Anterior Segment of the Eye–The Cornea

Laser surgery to the cornea by laser-assisted in situ keratomileusis (LASIK) or photorefractive keratectomy (PRK) for the purpose of correcting nearsightedness (myopia), farsightedness (hyperopia), or astigmatism is not a benefit of Texas Medicaid.

Reimbursement for laser surgery to the cornea, procedure codes 65450, 65855, and 65860 is limited to once every 90 days for the same eye.

Anterior Segment of the Eye–The Iris, Ciliary Body

Laser surgery to the anterior segment of the eye–the iris, ciliary body may be reimbursed only when billed with one of the following procedure codes:

<table>
<thead>
<tr>
<th>Procedure Codes</th>
</tr>
</thead>
<tbody>
<tr>
<td>66600</td>
</tr>
</tbody>
</table>

Reimbursement for procedure codes 66600, 66605, 66710, 66711, 66761, 66762, and 66770 is limited to once every 90 days for the same eye.

Claims for iridectomy (66600, 66605, 66625, 66630, or 66635) or iridotomy (66500 or 66505) are not reimbursed when billed for the same date of service as a trabeculectomy (66170 or 66172). These claims are considered for review when filed on appeal with documentation of medical necessity. The iridectomy is considered part of a trabeculectomy. An iridectomy billed with any other eye surgery on the same day suspends for review.

An iridectomy is also considered part of certain types of cataract extractions. An iridectomy (66600 or 66605) is not reimbursed when billed for the same date of service as the cataract surgeries listed in the following table. The iridectomy is considered part of the cataract surgery. These claims are considered for review when filed on appeal with documentation of medical necessity.

<table>
<thead>
<tr>
<th>Procedure Codes</th>
</tr>
</thead>
<tbody>
<tr>
<td>65920</td>
</tr>
<tr>
<td>66986</td>
</tr>
</tbody>
</table>
Posterior Segment of the Eye–Retina or Choroid

Laser surgery to the retina or choroid may be reimbursed only when billed with one of the following procedure codes:

<table>
<thead>
<tr>
<th>Procedure Codes</th>
</tr>
</thead>
<tbody>
<tr>
<td>67105 67107 67108 67110 67113 67145 67210 67220 67221 67225</td>
</tr>
<tr>
<td>67228 67229 G0186</td>
</tr>
</tbody>
</table>

Procedure code 67229 is restricted to clients who are birth through 1 year of age.

When billed for the same date of service, same eye, any provider, procedure code 67031 will be denied as part of any of the following procedure codes:

<table>
<thead>
<tr>
<th>Procedure Codes</th>
</tr>
</thead>
<tbody>
<tr>
<td>67036 67108 67110 67120 67121 67141 67142 67208 67210 67218</td>
</tr>
<tr>
<td>67227 67228</td>
</tr>
</tbody>
</table>

When billed for the same date of service, same eye, any provider, only one of the following procedure codes may be reimbursed: 67220, 67221, 67225, or G0186.

When billed for the same date of service, same eye, by any provider, procedure codes 67025, 67028, 67031, 67036, 67040, and 67105 will be denied as part of 67108.

Posterior Segment of the Eye, Vitreous–Vitrectomy

Laser surgery to the vitreous may be reimbursed only when billed with one of the following procedure codes: 67031, 67039, 67040, and 67043.

Reimbursement for procedure codes 67031, 67039, 67040, and 67043 is limited to once every 90 days for the same eye.

When billed for the same date of service, same eye, any provider procedure codes 67500 and 69990 are denied as part of 66821.

Procedure code 66821 is denied as part of 66830, 67031, and 67228.

Procedure codes 66820, 66984, 66985, and 67036 will pay according to multiple surgery guidelines when billed with procedure code 66821.

When billed for the same date of service, same eye, different provider procedure codes 66821, 67005, 67010, and 69990 will be denied as part of 67031.

When billed for the same date of service, same eye, any provider procedure code 67031 will be denied as part of any of the following procedure codes: 67036, 67108, 67110, 67120, 67121, 67208, 67218, 67227, and 67228.

9.2.46.3 Eye Surgery by Incision

The following restrictions apply to vitrectomy and cataract surgeries:

- Procedure codes 66500, 66505, 66605, 66625, 66630, and 66635 are denied as part of another procedure when billed with the following cataract surgeries: 65920, 66840, 66850, 66852, 66920, 66930, 66940, 66983, 66984, 66985, and 66986. Claims may be appealed with additional documentation to demonstrate the medical necessity.
- Procedure code 66020 is denied as part of another procedure when billed with any related eye surgery procedure code.
- Procedure code 67036 may be reimbursed when billed alone.
• Procedure code 67036 is denied as part of another procedure when billed with procedure codes 67039, 67040, 67041, 67042, 67043, or 67108.

• Procedure codes 67039 and 67040 are combined and reimbursed as procedure code 67108 when billed by the same provider for the same date of service.

• For clients who are 8 years of age and younger, the following cataract extraction and vitrectomy procedure codes, performed on the same eye, will be considered for payment per multiple surgery guidelines:

<table>
<thead>
<tr>
<th>Procedure Codes</th>
</tr>
</thead>
<tbody>
<tr>
<td>66840</td>
</tr>
<tr>
<td>67015</td>
</tr>
<tr>
<td>67042</td>
</tr>
</tbody>
</table>

• For clients who are nine years of age and older, the following procedure codes will be paid when performed on the same eye as a cataract extraction:

<table>
<thead>
<tr>
<th>Procedure Codes</th>
</tr>
</thead>
<tbody>
<tr>
<td>67005</td>
</tr>
<tr>
<td>67040</td>
</tr>
</tbody>
</table>

• For clients who are nine years of age and older, the following procedure codes will be denied as part of the codes listed above, when performed on the same eye:

<table>
<thead>
<tr>
<th>Procedure Codes</th>
</tr>
</thead>
<tbody>
<tr>
<td>66840</td>
</tr>
</tbody>
</table>

Reimbursement for procedure codes 67041, 67042, and 67043 is limited to once every 90 days for the same eye.

9.2.46.4 Intraocular Lens (IOL)
An IOL (V2630, V2631, and V2632) may be reimbursed only to physicians in the office setting (POS 1). Providers must submit a copy of the manufacturer’s invoice for procedure code V2631 to TMHP with their claim. Reimbursement for the lens is limited to the actual acquisition cost for the lens (taking into account any discount) plus a handling fee not to exceed five percent of the acquisition cost.

Medicaid does not reimburse physicians who supply IOLs to ASCs/HASCs.

Reimbursement for the surgical procedure necessary to implant an IOL remains unchanged.

9.2.46.5 Intravitreal Drug Delivery System
Procedure codes 67027 and 67121 pertain to the procurement, implantation, and removal of an intravitreal drug delivery system (e.g., a ganciclovir implant). They are set to deny when billed concurrently.

9.2.46.6 Other Eye Surgery Limitations
The following procedure codes require modifier LT or RT to identify the eye for which the surgery is being performed:

<table>
<thead>
<tr>
<th>Procedure Codes</th>
</tr>
</thead>
<tbody>
<tr>
<td>65205</td>
</tr>
<tr>
<td>67345</td>
</tr>
</tbody>
</table>
In the following table, the procedure codes in Column A may be reimbursed only when at least one corresponding procedure code from Column B has been paid to the same provider for the same date of service:

<table>
<thead>
<tr>
<th>Column A Procedure Codes</th>
<th>Column B Procedure Codes</th>
</tr>
</thead>
<tbody>
<tr>
<td>66990</td>
<td>65820, 65875, 65920, 66985, 66986, 67036, 67039, 67040, 67041, 67042, 67043, or 67113</td>
</tr>
<tr>
<td>67320, 67331, 67332, 67334</td>
<td>67311, 67312, 67314, 67316, or 67318</td>
</tr>
<tr>
<td>67335, 67340</td>
<td>67311, 67312, 67314, 67316, 67318, 67320, 67331, 67332, or 67334</td>
</tr>
<tr>
<td>V2790</td>
<td>65780</td>
</tr>
</tbody>
</table>

### 9.2.47 Organ/Tissue Transplants

Organ/tissue transplants that include bone marrow, peripheral stem cell, heart, intestine, lung, liver, kidney, or pancreas are a benefit of Texas Medicaid.

Solid organ transplants are a benefit of Texas Medicaid when medically necessary based on safety and efficacy, as demonstrated by scientific evidence and by controlled clinical studies, in accordance with the Texas Administrative Code (TAC). Solid organ transplants are limited to clients with a critical medical condition who are expected to have a successful clinical outcome that will result in a return to improved functional independence. Benefits are not available for the following experimental or investigational services:

- Artificial and bioartificial livers
- Xenotransplantation of solid organs
- Thymus transplant

Coverage is limited to one transplant per organ system (or organ systems for combined transplants) per lifetime except for one subsequent transplant because of organ rejection.

Solid organ transplants require prior authorization and may be reimbursed only when performed in a Medicaid-enrolled facility that is a designated children’s hospital with a transplant unit or program, or certified for the procedure by the United Network for Organ Sharing (UNOS).

The facility must be in Texas, unless there are no Texas facilities certified by UNOS or designated as a Children’s Hospital with a transplant unit or program for the requested procedure.

All requests for out-of-state (OOS) services, whether for pre-transplant evaluation, transportation, or post-transplant monitoring, must be sent to the medical director for prior authorization review. Texas Medicaid will consider authorizing OOS services when the following criteria are met:

- The client does not leave Texas to receive care that can be received in Texas.
- An in-state facility approved for the procedure has declined to accept the client and documentation is submitted to explain why the in-state team cannot perform the procedure.
- There is no physician provider or facility with the level of expertise required to perform the necessary procedure available in Texas, or the client has received an initial transplant at the OOS facility and requires additional transplant services due to complications or graft loss.
- There is reasonable assurance that the client meets the clinical criteria required by Texas Medicaid for transplant approval.
- The service is necessary, reasonable, and federally allowable, and the facility and physicians agree to accept Medicaid reimbursement for these services.
• The OOS facility must be certified by UNOS or designated as a Children’s Hospital with a transplant unit or program.

When requesting an OOS prior authorization for a pre-transplant evaluation, the provider must submit a copy of the transplant evaluation performed by a Texas facility to support the need for an OOS solid organ pre-transplant evaluation.

When requesting an OOS prior authorization for transplant of a solid organ, the provider must submit a copy of the transplant evaluation performed by a Texas facility and a copy of the transplant evaluation performed by the OOS facility to support the need for an OOS solid organ transplant.

When requesting an OOS prior authorization for post-transplant monitoring or other post-transplant services, the provider must submit documentation that the client received the initial transplant at the same OOS facility to include complications or graft loss if present, in order to support the need for OOS solid organ post-transplant monitoring or other post-transplant services.

Expenses incurred for the procurement of a living donor’s organ are not a benefit of Texas Medicaid.

Refer to: Subsection 3.2.5, “Organ and Tissue Transplant Services” in the Inpatient and Outpatient Hospital Services Handbook (Vol. 2, Provider Handbooks) for more information about the transplant facility approval criteria.

Subsection 3.2.5.2, “Transplant Benefits and Limitations” in the Inpatient and Outpatient Hospital Services Handbook (Vol. 2, Provider Handbooks) for more information about organ/tissue transplant program limitations.

9.2.47.1 General Prior Authorization Requirements

Solid organ transplant prior authorization requests must include all of the following:

• A complete history and physical
• A statement of the current medical conditions and status of the transplant recipient
• Documentation of how the client meets the prior authorization criteria specified for the transplant requested
• Documentation of the absence of co-morbidities or contraindications such as the following:
  • Severe pulmonary hypertension
  • End-stage cardiac, renal, hepatic, or other organ dysfunction unrelated to the primary disorder
  • Uncontrolled HIV infection or AIDS defining illness
  • Multiple organ compromise secondary to infection, malignancy, or condition with no known cure
  • Ongoing or recurrent active infections that are not effectively treated
  • Psychiatric instability severe enough to jeopardize incentive for adherence to medical regimen
  • Active alcohol or chemical dependency that might interfere with compliance to a medical regimen
• History of compliance with other medical treatments, regimen, and plan of care

Backbench procedures do not require prior authorization but may only be reimbursed when a corresponding transplant procedure has been paid for the same date of service.

Note: Clients who are birth through 20 years of age and who do not meet the criteria for coverage may be considered through the Comprehensive Care Program (CCP).

Additional prior authorization criteria, if applicable, specific to each type of transplant are outlined in the following sections.
If prior authorization is not obtained for a solid organ transplant, services directly related to the transplant within the three-day preoperative and six-week postoperative period are also denied regardless of who provides the services (e.g., laboratory services, status post visits, radiology services). However, coverage for other services needed as a result of complication of the transplant or for services unrelated to the transplant may be considered when medically necessary, reasonable, and federally allowable.

Claims for transplant clients are placed on active review when the transplant was not prior authorized so that the services related to the transplant can be monitored.

9.2.47.2 Heart Transplants

9.2.47.2.1 Prior Authorization for Heart Transplants

A heart transplant to a client for primary heart dysfunction must be documented as the client being unresponsive to more conventional and/or standard therapies to be considered for coverage.

Procedure code 33945 may be considered for prior authorization with medical necessity documentation that indicates a New York Heart Association (NYHA) Class III or IV cardiac disease with one of the following medical conditions:

- Congenital heart disease
- Valvular heart disease
- Viral cardiomyopathy
- Familial and restrictive cardiomyopathy

9.2.47.3 Intestinal Transplants

An intestinal transplant may be considered for clients who are dependent on parental nutrition and have compromised venous access, have had two or more episodes of central line sepsis, or who have begun to manifest progressive parental nutrition associated liver dysfunction. Procedure codes 44135 and 44136 must be prior authorized.

Small bowel transplantation is considered medically necessary in clients with irreversible intestinal failure including, but not limited to:

- Short bowel syndrome
- Pseudo-obstruction
- Microvillus inclusion
- Tumor

The prior authorization request must include documentation of irreversible intestinal failure with failed total parenteral nutrition (TPN) therapy. The client has experienced TPN failure if any one of the following criteria is met:

- Impending or overt liver failure due to TPN-induced liver injury. Clinical indicators include the following:
  - Increased serum bilirubin levels
  - Increased liver enzyme levels
  - Splenomegaly
  - Thrombocytopenia
  - Gastroesophageal varices
  - Coagulopathy
  - Stomal bleeding
• Hepatic fibrosis
• Cirrhosis
• Thrombosis of major central venous channels (subclavian, jugular, or femoral veins). Thrombosis of two or more of these vessels is considered a life-threatening complication and TPN failure.
• Frequent central line–related sepsis. Two or more episodes of central-line–induced systemic sepsis per year that require hospitalization are considered TPN failure. A single episode of central-line–related fungemia, septic shock, or acute respiratory distress syndrome is considered TPN failure.
• Frequent episodes of severe dehydration despite TPN and intravenous fluid supplement. Under certain medical conditions, such as secretory diarrhea and nonconstructable gastrointestinal tract, the loss of combined gastrointestinal and pancreatobiliary secretions exceed the maximum intravenous infusion rates that can be tolerated by the cardiopulmonary system.

Diagnoses that indicate intestinal failure include, but are not limited to, the following:
• Small bowel syndrome resulting from inadequate intestinal propulsion due to neuromuscular impairment
• Small bowel syndrome resulting from postsurgical conditions due to resections
• Intestinal cysts
• Mesenteric cysts
• Small bowel or other tumors involving small bowel
• Crohn’s disease
• Mesenteric thrombosis
• Volvulus
• Short-gut syndrome in which there is liver function impairment (usually secondary to TPN)

9.2.47.4 Kidney Transplants

9.2.47.4.1 Prior Authorization for Kidney Transplants
Procedure codes 50360 and 50365 must be prior authorized. Medical necessity documentation of one of the following is required:
• Hemodialysis or continuous ambulatory peritoneal dialysis (CAPD).
• Chronic renal failure with anticipated deterioration to end-stage renal disease.
• End-stage renal disease, evidenced by a creatinine clearance below 20 ml/min or development of symptoms of uremia.
• End-stage renal disease that requires dialysis or is expected to require dialysis within the next 12- to 18-month period.

9.2.47.4.2 Cytogam
Procedure code J0850 is reimbursable by Texas Medicaid. Cytogam is indicated for the attenuation of primary cytomegalovirus disease in seronegative kidney transplant recipients who receive a kidney from a seropositive donor. Payment of cytogam is limited to diagnosis code Z940, Z941, Z942, Z943, Z944, or Z9483. Cytogam is payable only in the office or outpatient setting.

Refer to: Subsection 3.2.5, “Organ and Tissue Transplant Services” in the Inpatient and Outpatient Hospital Services Handbook (Vol. 2, Provider Handbooks) for more information about the transplant facility approval criteria.
9.2.47.5 Liver Transplants

9.2.47.5.1 Prior Authorization for Liver Transplants

For a client to be considered for coverage of a liver transplant, the medical records for the client must include documentation showing the client is unresponsive to more conventional and/or standard therapies.

Authorization of procedure codes 47133 and 47135 requires medical necessity documentation of liver disease in one of the following categories:

- Primary cholestatic liver disease
- Other cirrhosis:
  - Alcoholic
  - Hepatitis C, non-A, non-B, and Hepatitis B
- Fulminant hepatic failure
- Metabolic diseases
- Malignant neoplasms
- Benign neoplasms
- Biliary atresia

9.2.47.6 Lung Transplants

9.2.47.6.1 Prior Authorization for Lung Transplants

A lung transplant to a client must be documented as unresponsive to more conventional and/or standard therapies to be considered for coverage.

Prior authorization of procedure codes 32851, 32852, 32853, 32854, and S2060 may be considered with medical necessity documentation of the following:

- Symptoms at rest directly related to chronic pulmonary disease and resultant severe functional limitation
- End-stage pulmonary diseases in one of these categories:
  - Obstructive lung disease
  - Restrictive lung disease
  - Cystic Fibrosis
  - Pulmonary hypertension

9.2.47.7 Pancreas Transplant

9.2.47.7.1 Prior Authorization for Pancreas Transplant

A pancreas/simultaneous kidney-pancreas transplant must be documented as the client being unresponsive to more conventional and/or standard therapies to be considered for coverage.

For prior authorization of procedure codes 48160 and 48554, medical necessity documentation must be submitted that shows the following:

- Recurrent, acute, and severe metabolic and potentially life-threatening complications requiring medical attentions such as:
  - Hypoglycemia
  - Hyperglycemia
• Ketoacidosis
• Failure of exogenous insulin-based management to achieve sufficient glycemic control (HbA1c of greater than 8.0) despite aggressive conventional therapy
• Insensitivity to hypoglycemia; or
• Satisfactory kidney function (creatinine clearance greater than 40mL/min), except for kidney-pancreas transplants; and
• Type 1 diabetes with secondary diabetic complications that are progressive despite the best medical management; and
• At least two of the following secondary complications:
  • Diabetic neuropathy
  • Retinopathy
  • Gastroparesis
  • Autonomic neuropathy
  • Extremely labile (brittle) insulin-dependent diabetes mellitus

9.2.47.8 Multi-Organ Transplants
Procedure codes 33935, S2053, and S2054 may be considered for prior authorization if medical necessity documentation meets the requirements for each organ.

Procedure code S2065 may be considered for prior authorization if medical necessity documentation indicates the client meets criteria for a pancreas transplant and has end-stage renal disease that requires dialysis or is expected to require dialysis within the next 12 months.

9.2.47.9 Nonsolid Organ Transplants
Nonsolid organ transplants covered by Texas Medicaid include allogeneic and autologous stem cell transplantation, allogeneic and autologous bone marrow transplantation, autologous islet cell transplantation, and hematopoietic progenitor cell (HPC) boost infusion.

9.2.47.9.1 Allogeneic and Autologous Bone Marrow and Stem Cell Transplantation
Stem cell transplantation is a process in which stem cells are obtained from either a client’s or donor’s bone marrow, peripheral blood, or umbilical cord blood for intravenous infusion. The transplant can be used to effect hematopoietic reconstitution following severely myelotoxic doses of chemotherapy and/or radiotherapy used to treat various malignancies, and also can be used to restore function in clients having an inherited or acquired deficiency or defect.

Benefits are not available for any experimental or investigational services, supplies, or procedures.

Coverage of bone marrow and stem cell transplantation is limited to the following procedure codes: 38206, 38230, 38232, 38240, 38241, 38242, and S2142.

Texas Medicaid recognizes the following covered indications for stem cell transplants:
• Allogeneic
  • Hematological malignancy
  • Lymphatic malignancy
  • Bone marrow disorders
  • Hemoglobinopathies
  • Platelet function disorders
• Immunodeficiency disorders
• Inherited metabolic disorders
• Multiple myeloma/plasma cell disorders

• Autologous
  • Hematological malignancy
  • Lymphatic malignancy
  • Germ cell tumors
  • Brain tumors
  • Small round blue cell tumors of childhood
  • Multiple myeloma/plasma cell disorders

• Indications for additional infusions
  • Infusion of stem cells for failure to graft (autologous)
  • Donor leukocyte infusion for persistent or relapsed malignant disease (allogeneic)

• Indications for re-transplantation
  • Relapse of disease
  • Failure to engraft or poor graft function

9.2.47.9.2 Autologous Islet Cell Transplantation
Autologous islet cell transplantation associated with the complete or partial removal of the pancreas (procedure code 48160) is a benefit of Texas Medicaid only for clients with a diagnosis of chronic pancreatitis.

Allogeneic islet cell transplantation is not a benefit.

9.2.47.9.3 HPC Boost Infusion
Prior authorization is required for HPC boost infusion procedure code 38243. The prior authorization request must include documentation of a prior stem cell transplant.

Requests for more than two boost procedures per lifetime requires medical necessity review and approval by the medical director.

9.2.47.9.4 Prior Authorization for Nonsolid Organ Transplants
All nonsolid organ transplants require mandatory prior authorization and must be performed in a Texas facility that is a designated children’s hospital or a facility in compliance with the criteria set forth by the Organ Procurement and Transportation Network (OPTN), the United Network for Organ Sharing (UNOS), or the National Marrow Donor Program (NMDP). Prior authorization is effective for the date span specified on the prior authorization approval letter. If the transplant has not been performed by the end of the authorization period, the physician must apply for an extension.

Documentation supplied with the prior authorization request must include the following:
• A complete history and physical.
• A statement of the client’s current medical condition and the expected long-term prognosis for the client from the proposed procedure.

Each subsequent transplant must be prior authorized separately.

Peripheral or umbilical cord blood stem cell transplantation may be authorized in lieu of bone marrow transplantation (BMT), but will not be approved when performed simultaneously.
If a stem cell transplant has been prior authorized for a client who is 21 years of age or older, a maximum of 30 days of inpatient hospital services during a Title XIX spell of illness may be covered beginning with the actual first day of the transplant. This coverage is in addition to covered inpatient hospital days provided before the actual first day of the transplant. This 30-day period is considered a separate inpatient hospital admission for reimbursement purposes, but is included under one hospital stay.

Bone marrow harvesting (38230) or peripheral stem cell harvesting (38206) for autologous bone marrow or stem cell transplants are a benefit of Texas Medicaid and require prior authorization.

Autologous harvesting of stem cells (single or multiple sessions) may be reimbursed to the facility when prior authorized by HHSC or its designee and performed in the outpatient setting (POS 5). Harvesting of stem cells performed in the inpatient setting (POS 3) is included in the DRG and will not be reimbursed separately.

Physician services for the storage of stem cells are not a benefit of Texas Medicaid.

Donor expenses are included in the global fee for the transplant recipient and are not reimbursed separately. Therefore, allogeneic bone marrow or stem cell harvesting procedures are not a benefit of Texas Medicaid.

Stem cell transplants for other conditions may be considered on a case by case basis. Documentation for prior authorization must be submitted to determine whether the transplant is medically necessary and appropriate.

9.2.47.10 Organ Procurement
The appropriate DRG reimbursement coverage to the approved institution for a prior authorized solid organ transplant procedure includes procurement of the organ and services associated with the organ procurement as specified by HHSC or its designee. Documentation of organ procurement must be maintained in the hospital medical records.

9.2.48 Orthognathic Surgery
Orthognathic surgery is a benefit of Texas Medicaid only when it is necessary for medical reasons, or when it is necessary as part of an approved plan of care in the Texas Medicaid Dental Program. Orthognathic surgery is administered and may be reimbursed as part of the medical/surgical benefit of Texas Medicaid and not as part of the Texas Medicaid Dental Program.

Treatment of malocclusion is a benefit of the Texas Medicaid Dental Program. Orthognathic surgery is a benefit when it is necessary as part of the approved dental benefit.

Maxillary and/or mandibular facial skeletal deformities are associated with clearly abnormal masticatory malocclusion.

Orthognathic surgery may be considered medically necessary for the following client conditions:

- Producing signs or symptoms of masticatory dysfunction
- Facial skeletal discrepancies associated with documented sleep apnea, airway defects, and soft tissue discrepancies
- Facial skeletal discrepancies associated with documented speech impairments
- Structural abnormalities of the jaws secondary to infection, trauma, neoplasia, or congenital anomalies

Orthognathic surgery may be considered for reimbursement when required for the client to access a dental service. Orthognathic surgery that is done primarily to improve appearance and not for reasons of medical necessity is considered cosmetic and is not a benefit of Texas Medicaid.
9.2.48.1 Prior Authorization for Orthognathic Surgery

The following orthognathic medical surgical services may be considered for reimbursement to oral and maxillofacial surgeons with prior authorization. A narrative explaining medical necessity must be provided with the authorization request.

<table>
<thead>
<tr>
<th>Procedure Codes</th>
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<tbody>
<tr>
<td>21010</td>
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9.2.49 Osteopathic Manipulative Treatment (OMT)

OMT, when performed by a physician (MD or DO), is a benefit of Texas Medicaid for the acute phase of the acute musculoskeletal injury or the acute phase of an acute exacerbation of a chronic musculoskeletal injury with a neurological component.

OMT is covered when it is performed with the expectation of restoring the patient’s level of function, which has been lost or reduced by injury or illness. Manipulations should be provided in accordance with an ongoing, written treatment plan that supports medical necessity. A model of documentation that supports medical necessity for the treatment plan includes the following:

- Specific modalities/procedures to be used in treatment
- Diagnosis
- Region treated
- Degree of severity
- Impairment characteristics
- Physical examination findings (X-ray or other pertinent findings)
- Specific statements of long- and short-term goals
- Reasonable estimate of when the goals will be reached (estimated duration of treatment)
- Frequency of treatment (number of times per week)
- Equipment and techniques used

The treatment plan must be updated as the client’s condition changes. Treatment plans must be maintained in the medical records and are subject to retrospective review.

Reimbursement is contingent on correct documentation of the condition. The acute modifier AT must be submitted with the claim for payment to be made. Paper claims submitted without modifier AT will be denied; electronic claims will be rejected. The AT modifier is described as representing treatment provided for an acute condition or an exacerbation of a chronic condition that persists less than 180 days from the start date of therapy. If the condition persists for more than 180 days from the start of therapy, the condition is considered chronic, and treatment is no longer considered acute. Providers may file an appeal for claims denied as being beyond the 180 days of therapy with supporting documentation that
the client’s condition has not become chronic and the client has not reached the point of plateauing. Plateauing is defined as the point at which maximal improvement has been documented and further improvement ceases.

The following procedure codes are payable when billing for OMT to the head, cervical, thoracic, lumbar, sacral, pelvic, lower extremities, upper extremities, rib cage, abdominal, and visceral regions: 98925, 98926, 98927, 98928, and 98929.

OMT will be denied when billed on the same date of service by the same provider as any of the following procedure codes:

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<tr>
<th>Procedure Codes</th>
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When multiples of procedure codes 98925, 98926, 98927, 98928, and 98929 are billed on the same day by the same provider, the most inclusive code is paid and the others are denied.

An E/M or initial or subsequent care visit or consultation may be paid in addition to OMT billed on the same day if the client’s condition requires a visit for a significant and separately identifiable service above and beyond the usual pre- and post-care associated with the OMT procedure, even if the visit and OMT are related to the same symptom or condition. Modifier 25 must be submitted with the E/M procedure code to identify a separate and distinct service rendered on the same day as OMT.

Documentation that supports the provision of a significant, separately identifiable E/M service must be maintained in the client’s medical record and made available to Texas Medicaid upon request.

Procedure code 97140 will be denied as part of another service if billed on the same date of service as procedure codes 98925, 98926, 98927, 98928, or 98929.

9.2.50 Pain Management

Pain management is a benefit of Texas Medicaid.

Procedure codes 62350, 62351, 62355, 62360, 62361, 62362, and 62365 billed on the same day as another surgical procedure performed by the same physician are paid according to multiple surgery guidelines.

Procedure codes 62350, 62351, 62355, 62360, 62361, 62362, and 62365 billed on the same day as an anesthesia procedure performed by the same physician are denied as included in the total anesthesia time.
Reimbursement to the physician for the surgical procedure is based on the assigned RVUs or maximum fee. Outpatient facilities are reimbursed at their reimbursement rate. Inpatient facilities are reimbursed under the assigned diagnosis-related group (DRG). No separate payment for the intrathecal pump is made.

Use the following procedure codes when billing for the implantation/revision/replacement of the pump/catheter:

<table>
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<tr>
<th>Procedure Codes</th>
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<td>62350</td>
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Procedure codes 62367 and 62368 do not require prior authorization and are payable as a medical service only.

Refer to: Subsection 9.2.38, “Implantable Infusion Pumps” in this handbook for more information about implanted pumps.

Acute pain is defined as pain caused by occurrences such as trauma, a surgical procedure, or a medical disorder manifested by increased heart rate, increased blood pressure, increased respiratory rate, shallow respirations, agitation or restlessness, facial grimace, or splinting.

Chronic pain is defined as persistent, often lasting more than six months; symptoms are manifested similarly to that of acute pain.

Postoperative refers to the time frame immediately following a surgical procedure in which a catheter is maintained in the epidural or subarachnoid space for the duration of the infusion of pain medication.

**9.2.50.1 Epidural and Subarachnoid Infusion (Not Including Labor and Delivery)**

Epidural and subarachnoid infusion for pain management is payable for acute, chronic, and postoperative pain management.

Procedure code 01996 is limited to once per day and is denied when billed on the same day as a surgical/anesthesia procedure. Procedure code 01996 billed longer than 30 days requires medical necessity documentation. Cancer diagnoses are excluded from the 30-day limitation.

Procedure code 01996 is payable to CRNAs and physicians.

**9.2.52 Panniculectomy and Abdominoplasty**

Procedure codes 15830 and 15847 are benefits of Texas Medicaid when prior authorized.

To avoid unnecessary denials, the physician must provide correct and complete information, including documentation establishing medical necessity of the service requested. This documentation must remain in the client’s medical record and is subject to retrospective review.

**9.2.52.1 Panniculectomy**

A panniculectomy (procedure code 15830) may be reimbursed with prior authorization for one of the following conditions when the panniculus hangs to or below the level of the pubis:

- A panniculus has recurrent non-healing ulcers.
• Client is insulin dependent with recurring infection and causing the prolapse of a ventral hernia.
• Panniculus directly causes significant clinical functional impairment.

Panniculectomy is not a benefit when one of following is the primary purpose:
• To remove excess skin and fat from the middle and lower abdomen in order to contour and alter the appearance of the abdominal area to improve appearance.
• Dissatisfaction with personal body image.
• To minimize the risk of ventral hernia formation of recurrence.
• For the sole purpose of treating neck or back pain.

Panniculectomy may be prior authorized when the client meets one of the following:
• Panniculectomy is planned and there is no history of significant weight loss or gastric bypass surgery.
• Panniculectomy is planned without history of gastric bypass surgery but with significant weight loss and the panniculus hangs to or below the level of the pubis.
• Panniculectomy is planned with history of gastric bypass surgery or abdominoplasty and the client is 12 months post-surgery.

If a panniculectomy is planned and there is no history of significant weight loss or gastric bypass surgery, or a panniculectomy is planned without history of gastric bypass surgery but with significant weight loss and the panniculus hangs to or below the level of the pubis, one of the following must be met:
• Documentation of recurrent episodes of infection or recurrent non-healing ulcers over three months that are non-responsive to treatment or appropriate medical therapy, such as oral or topical prescription.
• The client is insulin-dependent and has a serious infection control problem and the panniculus is causing the prolapse of a ventral hernia.
• Documentation by the treating physician that the panniculus directly causes significant clinical functional impairment. Clinical functional impairment may be indicated by associated musculoskeletal dysfunction or interference with activities of daily living and there is reasonable evidence to support that this surgical intervention will correct the condition.

If a panniculectomy is planned with a history of gastric bypass surgery or abdominoplasty and the client is 12 months post-surgery, the following must be met:
• Documentation that the panniculus hangs to or below the level of the pubis and the client has maintained a significant (100 pounds or more), stable weight loss for at least six months. Documentation must include the weight loss history, prior and current height, prior and current weight, and the history and physical including all previous surgeries.
• Documentation of recurrent episodes of infection or recurrent non-healing ulcers over three months that are non-responsive to treatment or appropriate medical therapy, such as oral or topical prescription. The 12-month post-gastric bypass requirement may be waived.
• The client is insulin-dependent and has a serious infection control problem and the panniculus is causing the prolapse of a ventral hernia. The 12-month post-gastric bypass requirement may be waived.
• Documentation by the treating physician that the panniculus directly causes significant clinical functional impairment. The 12-month post-gastric bypass requirement may be waived. Clinical functional impairment may be indicated by associated musculoskeletal dysfunction or interference with activities of daily living and there is reasonable evidence to support that this surgical intervention will correct the condition.
All medical record documentation pertinent to the client’s evaluation and treatment must support medical necessity of the panniculectomy. Documentation may include the following:

- Office records
- Consultation reports
- Operative reports
- Other hospital records (examples: pathology report, history and physical)

Documentation to support the panniculectomy must be submitted with the request for prior authorization. In addition to medical record documentation, the provider may also submit a letter of support or an explanation to substantiate medical necessity.

This service is typically expected to be limited to once per lifetime; however, repeat panniculectomies may be considered for prior authorization upon submission of supporting documentation as outlined above.

A panniculectomy provided as a secondary surgery may be considered for prior authorization when the panniculus interferes with a medically necessary intra-abdominal surgery (e.g., abdominal hernia repair or hysterectomy) or to facilitate an improved anatomical field in order to provide radiation treatment to the abdomen. Documentation of medical necessity must include:

- The comorbidity for the diagnosis of the primary surgery or for the nature of the condition undergoing radiation treatment.

- Documentation supporting the need for the panniculectomy as the panniculus hangs below the level of the pubis and will significantly interfere with a planned surgical procedure, or the abdominal structures identified as requiring radiation therapy will not be adequately treated due to the size of the panniculus.

A panniculectomy provided as a secondary surgery may be considered when the primary surgery was performed for an urgent condition defined as a symptom or condition that is not an emergency, but requires further diagnostic workup or treatment within 24 hours to avoid a subsequent emergent situation.

The need for the panniculectomy as a secondary surgery in conjunction with a primary urgent surgery must be supported by retrospective review of submission of all of the following documentation:

- History and physical and the operative report.

- The panniculus hangs below the level of the pubis and would have significantly interfered with the urgent primary surgical procedure.

9.2.52.2 Abdominoplasty

An abdominoplasty (procedure code 15847) is a benefit for clients who are birth through 20 years of age and may be reimbursed with prior authorization for one of the following conditions:

- Prune belly
- Diastasis recti in the presence of a true midline hernia (ventral or umbilical)

Abdominoplasty is not a benefit when one of the following is the primary purpose:

- To remove excess skin and fat and tighten abdominal wall from the middle and lower abdomen in order to contour and alter the appearance of the abdominal area to improve appearance.

- Dissatisfaction with personal body image.

- To repair diastases recti (unless prior authorization criteria has been met).
Abdominoplasty may be prior authorized when the client meets all of the following criteria:

- Documented diagnosis of prune belly (i.e., Eagle Barret syndrome) or repair of diastasis recti in the presence of a true midline hernia (ventral or umbilical).
- Documentation for reconstructive surgery that must include appropriate historical medical record documentation and may include any of the following:
  - Consultation reports
  - Operative reports or other applicable hospital records (examples: pathology report, history and physical)
  - Office records
  - Letters with pertinent information from provider (when medical records are requested, a letter of support or explanation may be helpful, but alone will not be considered sufficient documentation to make a medical necessity determination)
- For repair of diastasis recti with a true midline hernia, documentation must also include all of the following:
  - The size of the hernia
  - Whether it is reducible, painful, or other symptoms
  - Whether there is a defect rather than just thinning of the abdominal fascia

Consideration of other abdominal diagnoses may be considered for prior authorization with the submission of additional supporting documentation that may include the following:

- Consultation reports
- Operative reports or other applicable hospital records (examples: pathology report, history and physical)
- Office records
- Letters with pertinent information from provider (when medical records are requested, a letter of support or explanation may be helpful, but alone will not be considered sufficient documentation to make a medical necessity determination)

### 9.2.53 Penile and Testicular Prostheses

The following services are a benefit of Texas Medicaid for male clients:

- Removal of a penile prosthesis without replacement (procedure codes 54406 and 54415).
- Insertion of testicular prosthesis for the replacement of congenitally absent testes or testes lost due to disease, injury, or surgery (procedure code 54660)—prior authorization is required.

Procedure code 54660 is a benefit for clients who are birth through 20 years of age. Insertion of a testicular prosthesis may be prior authorized with the following criteria:

- The client has lost a testicle as a result of cancer or trauma or has congenital absence of a testicle.
- The loss of the testicle has resulted in detrimental psycho-social sequelae, as evidenced by a psychiatric evaluation.

Requests for prior authorization must be submitted by the physician to the Special Medical Prior Authorization (SMPA) department using the Special Medical Prior Authorization (SMPA) Request Form. The request must be submitted with documentation that supports medical necessity.
9.2.54 **Percutaneous Transluminal Coronary Interventions**

Percutaneous transluminal coronary interventions are a therapeutic option for clients who have arteriosclerotic heart disease.

When any of the following procedure codes are performed on the same date of service and on the same vessel as intracoronary vessel stenting, any provider, only the stenting procedure code will be considered for reimbursement: 92973, 92982, 92984, 92995, and 92996.

Angioplasty, atherectomy, or thrombectomy performed on different coronary vessels may be reimbursed separately. When different coronary vessels are not indicated, only the stenting procedure will be paid.

9.2.55 **Physical Therapy (PT) Services**

Physical therapy (PT) is a payable benefit to physicians.

Refer to: Section 4, “Therapy Services Overview” in the Physical Therapy, Occupational Therapy, and Speech Therapy Services Handbook (Vol. 2, Provider Handbooks) for information about physical therapy services provided by a physician.

9.2.56 **Physician Evaluation and Management (E/M) Services**

E/M is a benefit of Texas Medicaid. Providers must follow either the 1995 or 1997 Documentation Guidelines for Evaluation and Management Services published by CMS when selecting the level of service provided.

The following E/M services are benefits of Texas Medicaid:

- Domiciliary, rest home, or custodial care services
- Emergency department services
- Group clinical visits
- Home services
- Hospital services including inpatient, observation, critical care, discharge, and concurrent care services (includes consultation and prolonged services)
- Nursing facility services
- Office or other outpatient services for new and established patients (includes consultation and prolonged services)
- Preventive care visits
- Services outside of business hours
- Tobacco use cessation

Claims submitted to TMHP by physicians for services provided during an inpatient hospital stay must be received by TMHP within 95 days of each date of service, not 95 days of the discharge date.

Inpatient claims must indicate the facility’s provider identifier in Block 32 or in the appropriate field of electronic software.
9.2.56.1 Office or Other Outpatient Hospital Services

9.2.56.1.1 New and Established Patient Services

A new patient is one who has not received any professional services from a physician or from another physician of the same specialty who belongs to the same group practice, within the past three years. Providers must use procedure codes 99202, 99203, 99204, and 99205 when billing for new patient services provided in the office or an outpatient or other ambulatory facility. New patient visits are limited to one every three years, per client, per provider.

An established patient is one who has received professional services from a physician or from another physician of the same specialty within the same group practice, within the last three years. Providers must use procedure codes 99211, 99212, 99213, 99214, and 99215 when billing for established patient services provided in the office or an outpatient or other ambulatory facility.

Established office or outpatient care visits are limited to once per day, same provider. When a new patient checkup is billed for the same date of service as a new patient acute care visit, both new patient services may be reimbursed when billed by the same provider or provider group if no other acute care visits or preventive care medical checkups have been billed in the past three years.

Modifier 25 must be submitted when the services rendered are performed for a significant separately identifiable service by the same physician or physician group on the same date of service. Modifier 25 is required when the provider submits a claim with the following:

- A second office or outpatient visit on the same day as another office or outpatient visit
- An office or outpatient visit beyond the usual preoperative care associated with the procedure that was performed

**Note:** Office or outpatient visits provided on the same date of service as a planned procedure (minor or extensive) are included in the cost of the procedure and are not separately reimbursed.

Documentation that supports the provision of a significant, separately identifiable E/M service must be maintained in the client’s medical record and made available to Texas Medicaid upon request. The documentation must clearly indicate what the significant problem/abnormality was, including the important, distinct correlation with signs and symptoms to demonstrate a distinctly different problem that required additional work and must support that the requirements for the level of service billed were met or exceeded.

The date and time of both services performed must be outlined in the medical record and the time of the second service must be different than the time of the first service, although a different diagnosis is not required.

Examples of additional visits to which Modifier 25 must be appended include, but are not limited to:

- A second E/M service for the same date of service as a group visit with the required E/M visit.
- An established patient E/M service for the same date of service as a THSteps medical checkup.
- An E/M service for the same date of service as a scheduled procedure.

Office visits (procedure codes 99202, 99203, 99204, 99205, 99211, 99212, 99213, 99214, and 99215) provided on the same date of service as a planned procedure (minor or extensive) are included in the cost of the procedure and are not separately reimbursed.

Office visit procedure code 99211, 99212, 99213, 99214, or 99215 must be billed by the same provider with the same date of service as a group clinical visit.

9.2.56.2 Group Clinical Visits

Texas Medicaid may reimburse physicians for group clinical visits (procedure code 99078) providing clinical services and educational counseling to a group of clients with the same condition.
To be considered for reimbursement, procedure code 99078 must be billed for the same date of service by the same provider as E/M procedure code 99211, 99212, 99213, 99214, or 99215.

Group clinical visits may be reimbursed for established patients only. The client’s plan of care must be determined and documented in the medical record by the physician before attending group clinical visits.

Participation of established patients in a group clinical visit is optional. Informed consent must be obtained from the client and maintained in the medical record before rendering group clinical visit services.

The physician leading the group clinical visit is responsible for the effectiveness and content of the information provided during the group clinical visit.

Nationally approved curriculum on asthma and diabetes, such as that available through the American Association of Diabetic Educators and Asthma Education and Prevention Programs approved by the CDC must be incorporated into the educational portion of group clinical visits.

Group clinical visits must last at least 1 hour, but no longer than 2 hours, with a minimum of 2 clients and a maximum of 20.

To promote self-management of the chronic disease, the group visit must include a presentation that instructs and informs the client about clinical issues, including how to prevent disease exacerbation or complications, properly use medications and other therapeutic techniques, or live with chronic illness topics. Group visit presentations must include:

- **Diabetic education consisting of the following:**
  - What diabetes is
  - Nutrition
  - Exercise and physical activity
  - Prevention of acute complications
  - Prevention of chronic complications
  - Monitoring
  - Medication

- **Asthma education consisting of the following:**
  - What is asthma?
  - What are symptoms of asthma?
  - What happens during an episode of asthma?
  - What exacerbates asthma?
  - How is asthma controlled?
  - What physical activities can people with asthma do?
  - A question and answer period
  - A short (approximately 5 to 15 minutes per client) one-on-one private direct (face-to-face) encounter with the physician consisting of:
    - A physical exam
    - The gathering, monitoring, and reviewing of laboratory and diagnostic tests
    - Medical decision making that includes an individual treatment plan
Documentation in the client’s medical record must support level of E/M service as per the CMS and CPT manual approved guidelines.

The documentation of the individual treatment plan retained in the client’s medical record must include data collected (physical exam and lab findings), educational services provided, patient participation, and the beginning and ending time of the visit.

Group visits for conditions of diabetes or asthma are limited to a maximum of four per year for any provider.

9.2.56.2.1 Group Clinical Visits for Diabetes

Group clinical visits are benefits of Texas Medicaid for the management of the condition of diabetes when submitted with one of the following diagnosis codes:

<table>
<thead>
<tr>
<th>Diagnosis Codes</th>
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<tbody>
<tr>
<td>E0800</td>
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9.2.56.3 Group Clinical Visits for Asthma

Group clinical visits are benefits of Texas Medicaid for the management of the condition of asthma when submitted with one of the following diagnosis codes:

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<th>Diagnosis Codes</th>
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<td>E1365</td>
</tr>
</tbody>
</table>

9.2.56.3.1 Group Clinical Visits for Pregnancy

Group clinical visits are benefits of Texas Medicaid for the management of the condition of pregnancy when submitted with procedure code 99078 and modifier TH, along with one of the following diagnosis codes:

<table>
<thead>
<tr>
<th>Diagnosis Codes</th>
</tr>
</thead>
<tbody>
<tr>
<td>J440</td>
</tr>
<tr>
<td>J4532</td>
</tr>
<tr>
<td>J45902</td>
</tr>
</tbody>
</table>

Diagnosis Codes

O0900 | O0901 | O0902 | O0903 | O0910 | O0911 | O0912 | O0913 |
| O09211 | O09212 | O09213 | O09219 | O09291 | O09292 | O09293 | O09299 |
Providers are encouraged to provide a comprehensive curriculum or use materials from the Centering Pregnancy Program that will be incorporated into the educational portion of the group clinical visit. Comprehensive curriculums will allow clinical issues to be identified to promote a healthy pregnancy. The education material may include screenings and preparations, health maintenance, counseling, and birth plans:

- **Screenings and preparations** may consist of the following:
  - Expected course of the pregnancy
  - Anticipated outline of the scheduled visits
  - Signs and symptoms, which should be reported to the physician as soon as possible
  - Laboratory services
  - Appropriate use of medications
  - Proper weight monitoring
  - Immunizations (e.g., hepatitis, varicella, or RhoGAM)
  - Complications of pregnancy that may occur (e.g., preeclampsia, diabetes, or edema)

- **Health maintenance** may consist of the following:
  - Hygiene (e.g., hot tubs or baths)
  - Sexual activity
  - Exercise
  - Nutrition and dietary needs

- **Counseling** may consist of the following:
  - Use of seat belts
  - Job activity
  - Air travel
  - Dental care appointments
  - Domestic abuse or violence
  - Tobacco or drug use

- **Birth planning** may consist of the following:
  - What to expect during labor and delivery

### Diagnosis Codes

<table>
<thead>
<tr>
<th>O0930</th>
<th>O0931</th>
<th>O0932</th>
<th>O0933</th>
<th>O0940</th>
<th>O0941</th>
<th>O0942</th>
<th>O0943</th>
</tr>
</thead>
<tbody>
<tr>
<td>O0951</td>
<td>O0952</td>
<td>O0953</td>
<td>O0954</td>
<td>O0955</td>
<td>O0956</td>
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<td>O0961</td>
<td>O0962</td>
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<td>O0971</td>
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<td>O0981</td>
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<td>O0996</td>
<td>O0997</td>
<td>O0998</td>
</tr>
<tr>
<td>Z331</td>
<td>Z3400</td>
<td>Z3401</td>
<td>Z3402</td>
<td>Z3403</td>
<td>Z3404</td>
<td>Z3405</td>
<td>Z3406</td>
</tr>
<tr>
<td>Z3483</td>
<td>Z3490</td>
<td>Z3491</td>
<td>Z3492</td>
<td>Z3493</td>
<td></td>
<td></td>
<td></td>
</tr>
</tbody>
</table>
• Pain control during labor
• Complications during delivery that may occur (e.g., Caesarean section or episiotomy)
• Breast feeding
• Newborn care
• Postpartum adjustments

Group clinical visits for the management of pregnancy are restricted to female clients who are 10 through 55 years of age and are limited to a maximum of 10 visits per 270 days for any provider.

To be considered for reimbursement, procedure code 99078 with modifier TH must be billed for the same date of service by the same provider as E/M procedure code 99211, 99212, 99213, 99214, or 99215 with modifier TH.

9.2.56.3.2 Preventive Care Visits

Adult preventive services must be provided in accordance with the U.S. Preventive Services Task Force (USPSTF) recommendations with grades A or B. USPSTF recommendations, with specific age and frequency guidelines, are listed on the USPSTF website. The following are recommended screens in addition to USPSTF and are covered separately:

- Tuberculosis screening
- Prostate cancer screening; prostate specific antigen (PSA) for men who are 50 through 64 years of age

Preventive care services are comprehensive visits that may include counseling, anticipatory guidance, and risk-factor-reduction interventions. Documentation must indicate the anticipatory guidance rendered.

Preventive health visits for clients who are birth through 20 years of age are available through THSteps medical checkups.

Refer to: Section 5, “THSteps Medical” in the Children’s Services Handbook (Vol. 2, Provider Handbooks).

Subsection 5.3.11.2.3, “Hearing Screening” in the Children’s Services Handbook (Vol. 2, Provider Handbooks) for additional information about hearing screenings.

Adult preventive services (procedure codes 99385, 99386, 99387, 99395, 99396, and 99397) are a benefit of Texas Medicaid for clients who are 21 years of age and older. Adult preventive services are limited to one service per rolling year, any provider, and must be billed with diagnosis code Z0000, Z0001, Z01411, or Z01419.

The following USPSTF recommendations are not reimbursed separately but must be provided, when applicable, as part of the routine preventive exam:

- Counseling to prevent tobacco use and tobacco-caused disease
- Behavioral counseling in primary care to promote a healthy diet
- Behavioral interventions to promote breast feeding
- Screening for obesity in adults (with intensive counseling and interventions)
- Screening and behavioral counseling interventions in primary care to reduce alcohol misuse
- Screening for depression

The USPSTF recommendation of chemoprevention of breast cancer is not a benefit of Texas Medicaid.
Laboratory, immunization, and diagnostic procedures recommended by USPSTF are covered benefits and may be billed separately, as clinically indicated, using the most appropriate diagnosis code that represents the client’s condition. Diagnosis code Z0000 or Z0001 may each be used once per rolling year for each screen if no other diagnosis is appropriate for the service rendered, but no more frequently than recommended by the USPSTF.

Services that exceed USPSTF recommendations are not considered part of a screening and require medical documentation to justify medical necessity of the services performed.

For clients who are 21 years of age and older, breast exams and Pap smears are available through programs related to women’s health, including Texas Medicaid family planning services and the Healthy Texas Women (HTW) program.

Refer to:
Section 2, “Medicaid Title XIX Family Planning Services” in the Gynecological, Obstetrics, and Family Planning Title XIX Services Handbook (Vol. 2, Provider Handbooks).


9.2.56.3.3 Tobacco Use Cessation

Tobacco use cessation counseling is a benefit for male and female clients who are 10 years of age and older and must be submitted with procedure codes 99406 and 99407. Tobacco use cessation services delivered in a group setting will be limited to a maximum of 8 participants per group and must be submitted with modifier HQ.

<table>
<thead>
<tr>
<th>Modifier</th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td>HQ</td>
<td>Group Counseling</td>
</tr>
</tbody>
</table>

Procedure codes 99406 and 99407 may be reimbursed when submitted with the following diagnosis codes:

<table>
<thead>
<tr>
<th>Diagnosis Codes</th>
</tr>
</thead>
<tbody>
<tr>
<td>F17200</td>
</tr>
<tr>
<td>F17218</td>
</tr>
<tr>
<td>F17291</td>
</tr>
</tbody>
</table>

Procedure codes 99406 and 99407 may be billed in any combination by the same or a different provider, whether individual or group counseling, and are limited to eight services per rolling year.

Additional services require documentation of medical necessity to exceed the established limit.

Procedure codes 99406 and 99407 are limited to once per day, same or different procedure code, any provider.

Refer to: Subsection 4.1.17, “Tobacco Use Cessation” in the Gynecological, Obstetrics, and Family Planning Title XIX Services Handbook (Vol. 2, Provider Handbooks) for additional information related to the tobacco use cessation counseling and additional covered diagnoses related to pregnancy.

9.2.56.3.4 Office and Outpatient Consultation Services

A consultation is an E/M service provided at the request of another provider for the evaluation of a specific condition or illness. The consultation must meet the following requirement:

- There must be a request from the referring provider for the evaluation of a particular condition or illness.
• There must be correspondence from the consulting provider back to the referring provider indicating the consulting provider’s medical findings.

During a consultation, the consulting provider may initiate diagnostic and therapeutic services if necessary.

The visit is not considered a consultation if any of the following applies:

• If diagnostic or therapeutic treatment is initiated during a consultation and the patient returns for follow-up care, the follow-up visit is considered an established patient visit, and must be billed as an established patient visit.

• If the purpose of the referral is to transfer care.

The medical records maintained by both the referring and consulting providers must identify the other provider and the reason for consultation.

Providers must use procedure code 99241, 99242, 99243, 99244, or 99245 when billing new or established patient consultations in the office, or in an outpatient or other ambulatory facility.

Office or outpatient consultations are limited to one consultation every six months by the same provider for the same diagnosis. Subsequent office or outpatient consultation visits during this six-month period will be denied.

9.2.56.3.5 Physician Services Provided in the Emergency Department

Providers must use procedure codes 99281, 99282, 99283, 99284, and 99285 when billing emergency department services. If an emergency department visit is billed by the same provider with the same date of service as an office visit, outpatient consultation, inpatient consultation, or subsequent nursing facility service, the emergency department visit may be reimbursed and the other services will be denied.

If an emergency department visit is billed by the same provider with the same date of service as an initial nursing facility service, the initial nursing facility service may be reimbursed and the emergency department visit will be denied.

Emergency department visits are denied when billed with the same date of service as an observation service (procedure code 99217) by the same provider.

Multiple emergency department visits provided by the same provider for the same client on the same day must have the times for each visit documented on the claim form. Also, more than one visit billed with the same date of service can be indicated by adding the appropriate modifier to the claim form. Medical documentation is required to support this service.

Reimbursement for physicians in the emergency department is based on Section 104 of TEFRA. TEFRA requires that Medicaid limit reimbursement for nonemergent and nonurgent physicians’ services furnished in hospital outpatient settings that also are ordinarily furnished in physician offices. The emergency department procedure code that is submitted on the claim is used to determine the appropriate reimbursement for these services. The procedure code billed may include, but is not limited to, E/M, surgical or other procedure, or any other service rendered to the client in the emergency room. The procedure code must accurately reflect the services rendered by the physician in the hospital’s emergency department. The reimbursement for each service is determined by multiplying the base allowable fee by 60 percent.

Refer to: Section 4, “Outpatient Hospital (Medical and Surgical Acute Care Outpatient Facility)” in the Inpatient and Outpatient Hospital Services Handbook (Vol. 2, Provider Handbooks) for information on emergency department services by facilities (room and ancillary).

Subsection 2.2.1.1, “Non-emergent and Non-urgent Evaluation and Management (E/M) Emergency Department Visits” in “Section 2: Texas Medicaid Fee-for-Service Reimbursement” (Vol. 1, General Information) for more information.
9.2.56.3.6 After-Hours Services

Texas Medicaid limits reimbursement for after-hours charges to office-based providers rendering services after routine office hours.

An office-based provider must bill an after-hours charge in addition to a visit charge for providing services after routine office hours. This after-hours charge must be billed when a provider judges it medically necessary to provide after-hours care for a patient with an emergent condition. A provider’s routine office hours are the hours posted at the physician’s office as the usual office hours. Medicaid reimburses office-based physicians an inconvenience or after-hours charge when any of the following situations exist:

- The physician leaves the office or home to see a client in the emergency room.
- The physician leaves the home and returns to the office to see a client after the physician’s routine office hours.
- The physician is interrupted from routine office hours to attend to another client’s emergency outside of the office.

Charges for inconvenience or after-hours services by emergency department-based physicians or emergency department-based groups are not allowed.

After-hours procedure codes are limited to one per day, same provider.

Providers must use one of the following procedure codes to report after-hours services:

<table>
<thead>
<tr>
<th>After-Hours Procedure Codes</th>
</tr>
</thead>
<tbody>
<tr>
<td>99050</td>
</tr>
</tbody>
</table>

9.2.56.3.7 Observation Services

Hospital observation (procedure codes 99217, 99218, 99219, and 99220) are professional services provided for a period of more than 6 hours but fewer than 24 hours regardless of the hour of the initial contact, even if the client remains under physician care past midnight.

Hospital observation services must be submitted using the procedure code 99234, 99235, or 99236.

Observation care discharge day management procedure code 99217 must be billed to report services provided to a client upon discharge from observation status if the discharge is on a date other than the initial date of admission. If an observation care discharge day management visit is billed by the same provider with the same date of service as an established office or outpatient visit, the observation care discharge visit may be reimbursed and the other services will be denied. The following procedure codes are denied if submitted with the same date of service as procedure code 99217:

<table>
<thead>
<tr>
<th>Procedure Codes</th>
</tr>
</thead>
<tbody>
<tr>
<td>99218</td>
</tr>
</tbody>
</table>

If a physician observation visit and a prolonged service are billed by the same provider for the same date of service, only one procedure code may be reimbursed.

If dialysis treatment and a physician observation visit are billed by the same provider (and same specialty other than an internist or nephrologist) with the same date of service, the dialysis treatment may be reimbursed and the physician observation visit will be denied. Subsequent observation care, per day (procedure codes 99224, 99225, and 99226), is also a benefit of Texas Medicaid.

Observation may take place in any patient care area of the hospital or outpatient setting.
9.2.56.4  Other Inpatient and Outpatient Hospital Services

A hospital care visit submitted by the same provider for the same client within three days of a new patient office, home, nursing facility, or skilled nursing facility (SNF) visit, for the same or for a similar diagnosis must be submitted as a subsequent care visit.

Refer to: Subsection 9.2.69.6, “Global Fees” in this handbook for more information about global services.

9.2.56.5  Prolonged Physician Services

Prolonged services involve face-to-face patient contact and may be provided in the office, outpatient hospital, or inpatient hospital settings. The face-to-face patient contact must exceed the time threshold of the following E/M procedure codes submitted for the date of service and be beyond the usual service.

<table>
<thead>
<tr>
<th>Procedure Codes</th>
<th>Limitation</th>
</tr>
</thead>
<tbody>
<tr>
<td>99202 99203 99204 99205 99211 99212 99213 99214 99215 99221</td>
<td>Used in conjunction with the E/M procedure code to report the first hour of prolonged service and are limited to one per day.</td>
</tr>
<tr>
<td>99222 99223 99231 99232 99233 99241 99242 99243 99244 99245</td>
<td>Used to report each additional 30 minutes and are limited to a quantity of 3 units or 1.5 hours per day.</td>
</tr>
</tbody>
</table>

Note: Prolonged services that are less than 30 minutes in duration cannot be reported separately.

Procedure code 99417 should only be used when an office or other outpatient evaluation and management service has been selected using time alone as the basis, and the minimum time required to report the highest level (procedure code 99205 or 99215) has been exceeded by 15 minutes.

Procedure code 99417 is limited to 4 units (1 hour) per day and should not be used to report an additional time increment of less than 15 minutes.

Prolonged services in the inpatient setting involving face-to-face client contact that is beyond the usual service may be reimbursed when provided on the same day as an initial hospital visit (procedure codes 99221, 99222, 99223, 99251, 99252, 99253, 99254, and 99255) or a subsequent hospital visit (99231, 99232, 99233).

Prolonged physician services are denied when billed with critical care or emergency room visits billed with the same date of service, same provider.

Prolonged physician services without a face-to-face contact (procedure codes 99358 and 99359) are not a benefit of Texas Medicaid.

Note: For non-face-to-face prolonged physician services, and for use of the unlisted evaluation and management procedure code for clients who are birth through 20 years of age, refer to subsection 2.5.1.1.6, “Non-Face-to-Face Prolonged Services” in the Children’s Services Handbook (Vol. 2, Provider Handbooks).

Refer to: Subsection 4.2.2, “Psychotherapy” in the Behavioral Health and Case Management Services Handbook (Vol. 2, Provider Handbooks) for more information about prolonged psychotherapy services.

Physician standby services are not a benefit of Texas Medicaid.
9.2.56.5.1 Hospital Admissions, Initial Visits, and Subsequent Visits

Inpatient hospital visits must be submitted using procedure codes 99221, 99222, 99223, 99231, 99232, and 99233.

If a subsequent hospital visit (procedure code 99231, 99232, or 99233) following admission is billed by the same provider with the same date of service as any of the following emergency department visits, office visits, or outpatient consultations, the subsequent hospital visit may be reimbursed and the other visits will be denied:

<table>
<thead>
<tr>
<th>Procedure Code</th>
</tr>
</thead>
<tbody>
<tr>
<td>99281</td>
</tr>
<tr>
<td>99212</td>
</tr>
</tbody>
</table>

Only one initial hospital care visit may be reimbursed to the same provider within a 30-day period for the same diagnosis. Additional initial hospital visits with the same diagnosis within a 30-day period will be denied.

A subsequent hospital visit (procedure code 99231, 99232, or 99233) will be denied when billed on the same day to the same provider as critical care services (procedure codes 99291 and 99292).

E/M services provided in a hospital setting following a major procedure and provided by the same provider or in direct follow-up for postsurgical care are included in the surgeon’s global surgical fee and are denied as included in another procedure.

Referto: Subsection 9.2.44, “Newborn Services” in this handbook for information about newborn services.

9.2.56.5.2 Concurrent Care

Concurrent care exists when services are provided to a patient by more than one physician on the same day during a period of hospitalization in the inpatient hospital setting. Concurrent care is appropriate when the level of care and the documented clinical circumstances require the skills of different specialties to successfully manage the patient in accordance with accepted standards of good medical practice. Concurrent care may be reimbursed to providers of different specialties when the services are for unrelated diagnoses involving different organ systems.

Concurrent care will be denied when billed for providers of the same specialty for the same or related diagnoses. Denied concurrent care may be appealed when accompanied by documentation of medical necessity.

Each appeal submitted for concurrent care must contain the following information:

- Documentation of the medical necessity for the physician’s services (care and treatment)
- Diagnosis and indication of the severity of the client’s condition (acute or critical)
- Role of the physician in the care of the client, including the name of the admitting physician
- Specialty and subspecialty of each physician and any limitations of practice

Claims appealed without clear documentation of medical necessity as described above will be denied.

Important: If the attending physician requests only a consultation, the request must be clearly stated in the orders.

All concurrent care is subject to retrospective review. Documentation of medical necessity for concurrent care must be retained by the physician as required by federal law and must include, but is not limited to, documentation of:

- The orders for concurrent care or valid reasons for the request by the attending physician.
The name of the requesting physician by the physician rendering concurrent care.

9.2.56.5.3 Consultations

Consultations provided to hospital inpatients, residents of nursing facilities, or patients in a partial hospital setting must be billed using procedure codes 99251, 99252, 99253, 99254, and 99255.

One initial inpatient consultation (procedure code 99251, 99252, 99253, 99254, or 99255) is allowed for each hospitalization within a 30-day period. Subsequent consultations billed as initial consultations during this time period will be denied.

Refer to: Subsection 9.2.56.3.4, “Office and Outpatient Consultation Services” in this handbook for additional criteria information.

9.2.56.5.4 Critical Care

Critical care includes the care of critically ill clients that require the constant attention of the physician. The physician must either be at bedside or immediately available to the client. The physician’s full attention must be devoted to the client so that the physician cannot render E/M to any other client during the same period of time. Critical care is usually given in a critical care area, such as the coronary care unit, intensive care unit, respiratory care unit, neonatal intensive care unit, or the emergency department care facility. The following procedure codes are used to bill critical care services:

<table>
<thead>
<tr>
<th>Procedure Code</th>
<th>Limitations</th>
</tr>
</thead>
<tbody>
<tr>
<td>99291</td>
<td>A per day charge for the first 30 to 74 minutes of critical care (time spent by the physician does not have to be continuous on that day).</td>
</tr>
<tr>
<td>99292</td>
<td>A per day charge for each additional 30 minutes beyond the first 74 minutes of critical care for up to 6 units or 3 hours per day.*</td>
</tr>
<tr>
<td>99471</td>
<td>A per day charge for initial inpatient pediatric critical care of the critically ill client who is 29 days through 24 months of age.</td>
</tr>
<tr>
<td>99472</td>
<td>A per day charge for subsequent inpatient pediatric critical care of the critically ill client who is 29 days through 24 months of age.</td>
</tr>
<tr>
<td>99475</td>
<td>A per day charge for initial inpatient pediatric critical care of the critically ill client who is 2 years through 5 years of age.</td>
</tr>
<tr>
<td>99476</td>
<td>A per day charge for subsequent inpatient pediatric critical care of the critically ill client who is 2 years through 5 years of age.</td>
</tr>
</tbody>
</table>

* If the number of units is not stated on the claim, a quantity of one is allowed.

Services for a client who is not critically ill and unstable but who was treated in a critical care unit must be reported using subsequent hospital visit codes or hospital consultation codes.

If the same provider who performed a major surgery must also perform critical care on the same day for the same client, the provider must bill the critical care with documentation that the critical care was unrelated to the specific anatomic injury or general surgical procedure.

Critical care (procedure codes 99291, 99292, 99471, 99472, 99475, and 99476) may be reimbursed only to the provider rendering the critical care service at the time of crisis. Critical care involves high-complexity decision-making to access, manipulate, and support vital system functions. While providers from various specialties may be consulted to render an opinion and assist in the management of a particular portion of the care, only the provider managing the care of the critically ill patient during a life threatening crisis may bill the critical care procedure codes.

Critical care procedure codes 99291 and 99292 are used to report the total duration of time spent by a physician providing critical care services to a critically ill or critically injured client, even if the time spent by the physician on that date is not continuous.
Actual time spent with the individual client must be recorded in the client’s record and reflect the time billed on the claim. The time that can be reported as critical care is the time spent engaged in work directly related to the individual client’s care whether that time was spent at the immediate bedside or elsewhere on the floor or unit.

Time spent under the following circumstances may not be reported as critical care:

- Activities that occur outside of the unit or off the floor
- Activities that do not directly contribute to the treatment of the client
- While performing separately reportable procedures or services

Critical care of less than 30 minutes total duration per day must be reported with the appropriate E/M procedure code.

If critical care that meets the initial 30-minute time requirement is provided to the same client by different physicians, the initial provider’s claim may be reimbursed. The second provider’s claim will be denied but may be appealed. The time spent by each physician cannot overlap; two physicians cannot bill critical care for care delivered at the same time. Supporting medical record documentation that includes the time in which the critical care was rendered must be provided by the second physician. In addition, a statement must be submitted indicating the physician was the only provider managing the care of the critically ill patient during the life threatening crisis.

If the provider’s time exceeds the 74-minute threshold for procedure code 99291, procedure code 99292 may be billed for each additional 30 minutes. Procedure code 99292 must be billed by the same performing provider or by a member of the same performing provider’s group practice and is limited to 6 units per day for any provider.

Inpatient critical care services provided to infants 29 days through 24 months of age are reported with pediatric critical care procedure codes 99471 and 99472. The pediatric critical care procedure codes are reported as long as the infant or young child qualifies for critical care services during the hospital stay through 24 months of age.

Pediatric critical care (procedure codes 99471, 99472, 99475, and 99476) is a per-day charge. Only one physician can bill pediatric critical care per day. If an inpatient or outpatient E/M service is billed by the same provider with the same date of service as pediatric critical care, the E/M service is denied.

Critical care provided to a neonatal, pediatric, or adult client in an outpatient setting (e.g., emergency room), which does not result in admission must be billed using procedure codes 99291 and 99292. Critical care provided to a neonatal or pediatric client in both the outpatient and inpatient settings on the same day must be billed using the appropriate neonatal or pediatric critical care procedure code.

If critical care (procedure code 99291 or 99292) is provided to a patient at a distinctly separate time from another outpatient E/M service by the same provider, both services may be reimbursed with supporting medical record documentation.

Prolonged physician services (procedure codes 99354, 99355, 99356, and 99357) will be denied when billed by the same provider with the same date of service as critical care (procedure code 99291, 99292, 99471, 99472, 99475, or 99476).

Claims may be subject to retrospective review to ensure documentation supports the medical necessity of the service when billing the claim.

Critical care procedure codes 99291 and 99292 will be denied when submitted with the same date of service by the same provider as neonatal intensive care procedure code 99468, 99469, 99478, 99479, or 99480.

9.2.56.5 Hospital Discharge

Hospital discharge must be submitted using procedure code 99238 or 99239.
Hospital discharge management billed by the same provider with the same date of service as the admission will be denied.

Discharge management billed by the same provider with the same date of service as an emergency room visit will be considered for reimbursement and the emergency visit will be denied.

Subsequent hospital visits billed by the same provider with the same date of service as discharge management will be denied.

9.2.56.5.6 Nursing Facility Services

Providers must use the following when billing initial nursing facility assessments, subsequent nursing facility care, and annual nursing facility assessments in a nursing facility:

<table>
<thead>
<tr>
<th>Procedure Codes</th>
</tr>
</thead>
<tbody>
<tr>
<td>99304* 99305* 99306* 99307 99308 99309 99310 99315 99316 99318</td>
</tr>
</tbody>
</table>

* Initial nursing facility assessments include all services related to an admission to the nursing facility.

Comprehensive initial nursing facility assessments performed by the same provider for the same diagnosis are limited to one every six months. The second initial nursing facility assessment within the six-month period will be denied.

Prolonged services in the nursing facility involving direct (face-to-face) patient contact that is beyond the usual service may be reimbursed on the same day as a nursing facility visit (procedure code 99304, 99305, 99306, 99307, 99308, 99309, or 99310).

All E/M services, regardless of setting, are considered part of the initial nursing facility care when performed by the same provider on the same day as the admission.

Subsequent nursing facility care E/M procedure codes 99307, 99308, 99309, and 99310 are limited to one per day regardless of diagnosis.

9.2.56.5.7 Observation

When a patient is admitted to the hospital as an inpatient and is discharged in less than 48 hours, the hospital may request that the physician change the admission order from inpatient status to outpatient observation status. This is an acceptable billing practice under Texas Medicaid when the physician makes the changes to the admitting order from inpatient status to outpatient observation status before the hospital submits the claim for reimbursement.

Refer to: Subsection 9.2.56.3.7, “Observation Services” in this handbook for more information about hospital observation.

9.2.56.6 Domiciliary, Rest Home, or Custodial Care Services

The following procedure codes are used to report E/M in a facility that provides room, board, and other personal assistance services:

<table>
<thead>
<tr>
<th>New Patient Procedure Codes</th>
</tr>
</thead>
<tbody>
<tr>
<td>99324 99325 99326 99327 99328</td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th>Established Patient Procedure Codes</th>
</tr>
</thead>
<tbody>
<tr>
<td>99334 99335 99336 99337</td>
</tr>
</tbody>
</table>

Established patient visits billed on the same date of service as a new patient visit, by the same provider, will be denied as part of another procedure. Established patient visits are limited to one per day regardless of diagnosis.

Established patient visits billed on the same day as a consultation visit, by the same provider, are denied.
9.2.56.7  Home Services

Home services are provided in a private residence. New patient visits will be limited to once every three years. Providers must utilize procedure codes 99341, 99342, 99343, 99344, and 99345 when billing for new patient services provided in the home setting. New patient visits are limited to one every three years. Providers must use procedure codes 99347, 99348, 99349, and 99350 when billing established patient services provided in the home setting.

A subsequent home visit (procedure codes 99347, 99348, 99349, and 99350) billed with the same date of service as a new patient home visit (procedure codes 99344 and 99345) by the same provider will be denied as part of another procedure, regardless of the diagnosis.

Subsequent home E/M codes are limited to one per day, regardless of diagnosis.

9.2.56.8  Referrals

A referral is defined as the transfer of the total or specific care of a patient from one physician to another; a referral does not constitute a consultation. These services must be billed using the appropriate E/M visit code.

When a Texas Medicaid provider refers a Texas Medicaid client to another provider for additional treatment or services, the referring provider must forward notification of the client’s eligibility and his provider identifier. The client must be made aware that the provider he/she is referred to does or does not participate in Texas Medicaid. Some clients not eligible for Medicaid are eligible for family planning through the HHSC Family Planning Program. These clients should be referred to contracted agency providers for family planning services.

9.2.56.8.1  Referral Requirements for Children with Disabilities

All health-care professionals are required by state and federal legislation to refer children who are 35 months of age or younger with developmental delays to early childhood intervention services provided under the authority of the Texas Health and Human Services Commission (HHSC).


9.2.57  Physician Services in a Long Term Care (LTC) Nursing Facility

HHSC requires initial certification and recertification of Medicaid clients in nursing facilities by physicians in accordance with guidelines set forth in federal regulations. Physician visits for certification and recertification are considered medically necessary, and are reimbursable by Medicaid whether performed in the physician’s office or the nursing facility.

Additional information is available on the HHSC website at https://hhs.texas.gov.

9.2.58  Podiatry and Related Services

Podiatry and related services are a benefit of Texas Medicaid.

9.2.58.1  Clubfoot Casting

Procedure code 29450 is limited to clients who are birth through 3 years of age and is payable to a physician in the management of clubfoot when a previous surgery has been performed. The physician may bill the appropriate E/M code with a casting code and be reimbursed for both. Procedure code 29750 is limited to clients who are birth through 3 years of age and is payable to a physician in addition to the initial casting or strapping procedure.

Use modifiers LT (left) and RT (right) with all procedures, as appropriate.
Casting and wedging are benefits if the client has one of the following conditions:

<table>
<thead>
<tr>
<th>Diagnosis Codes</th>
</tr>
</thead>
<tbody>
<tr>
<td>M21541 M21542 M21549 Q6600 Q6601 Q6602 Q6610 Q6611</td>
</tr>
<tr>
<td>Q6612 Q66211 Q66212 Q66219 Q66221 Q66222 Q66229 Q6630</td>
</tr>
<tr>
<td>Q6631 Q6632 Q6640 Q6641 Q6642 Q6651 Q6652 Q666</td>
</tr>
<tr>
<td>Q6670 Q6671 Q6672 Q6681 Q6682 Q6689 Q6690 Q6691</td>
</tr>
<tr>
<td>Q6692</td>
</tr>
</tbody>
</table>

### 9.2.58.2 Flat Foot Treatment

Reimbursement for treatment of deformities of the foot and lower extremity that includes flat foot as a component of the deformity may be considered when the client presents with significant pain in the foot, leg, or knee, resulting in a loss of or decrease in function, along with a secondary condition such as valgus deformity or plantar fasciitis.

Treatment of flat foot (flexible pes planus) that is solely cosmetic in nature is not a benefit of Texas Medicaid.

### 9.2.58.3 Routine Foot Care

Routine foot care must be medically necessary and billed with the following procedure codes. No specific diagnosis restrictions exist. The following procedures are limited to one service every six months per client, regardless of provider specialty: 11055, 11056, 11057, 11719, and G0127.

### 9.2.59 Prostate Surgery

A transurethral resection of the prostate (TURP) is the most common procedure performed to treat benign prostatic hyperplasia (BPH). A TURP may be billed with procedure code 52601, 52630, or 52640. If a provider submits separate charges for any of the TURP procedure codes listed above and procedure code 52351 or 52354, the charges for procedure codes 52351 and 52354 will be denied as part of the TURP procedure.

### 9.2.60 Radiation Therapy

Radiation treatment management may be reimbursed by Texas Medicaid as defined in the Current Procedure Terminology (CPT) manual under the “Radiation Treatment Management” section.

The following radiation therapy services are limited to once per day unless documentation submitted with an appeal supports the need for the service to be provided more frequently:

- Therapeutic radiation treatment planning
- Therapeutic radiology simulation-aided field setting
- Teletherapy
- Brachytherapy isodose calculation
- Treatment devices
- Proton beam delivery/treatment
- Intracavitary radiation source application
- Interstitial radiation source application
- Remote afterloading high intensity brachytherapy
- Radiation treatment delivery
- Localization
• Radioisotope therapy

Laboratory and diagnostic radiological services provided in the office setting may be reimbursed to physicians as a total component. Radiation treatment centers may also be reimbursed for the total component for these services in the outpatient hospital setting. Injectable medications given during the course of therapy in any setting may be reimbursed separately.

Routine follow-up care by the same physician on the day of any therapeutic radiology service will be denied. Medical services within program limitations may be reimbursed on appeal when documentation supports the medical necessity of the visit due to services unrelated to the radiation treatment or radiation treatment complication.

The professional component and the technical component will be denied when billed with the total component. The total component includes the professional and the technical components.

The professional component may be reimbursed for services rendered in the inpatient hospital setting, radiation treatment center setting, or outpatient hospital setting. Physicians billing client services rendered in the office setting or in a facility recognized by Medicaid as a radiation treatment center may be reimbursed for total components.

9.2.60.1 Brachytherapy

9.2.60.1.1 Prior Authorization for Brachytherapy

Prior authorization is not required for brachytherapy.

9.2.60.1.2 Other Limitations on Brachytherapy

Clinical brachytherapy services include admission to the hospital and daily care. Initial and subsequent hospital care will be denied as part of another service when billed with the same date of service as clinical brachytherapy services.

An office visit will be denied as part of another service when billed with the same date of service by the same provider as clinical treatment planning and clinical brachytherapy.

Normal follow-up care by the same physician will be denied as part of another service when billed with the same dates of service as any therapeutic radiology service. Any other E/M office visit will be denied as part of another service when billed with the same date of service by the same provider as the radiation treatment or radiation treatment complication.

Providers may use modifier 25 to indicate that the additional visit was for a separate, distinct service unrelated to the radiation treatment or radiation treatment complication. Documentation that supports the provision of a significant, separately identifiable E/M service must be maintained in the client’s medical record and made available upon request.

Each service provided using procedure codes 77321 and 77470 are limited to once per two calendar months.

Documentation that supports the provision of special procedures must be maintained in the client’s medical record and made available upon request.

9.2.60.2 Stereotactic Radiosurgery

9.2.60.2.1 Prior Authorization for Stereotactic Radiosurgery

The following procedure codes are a benefit of Texas Medicaid with prior authorization and documentation of medical necessity:

<table>
<thead>
<tr>
<th>Procedure Codes</th>
</tr>
</thead>
<tbody>
<tr>
<td>32701</td>
</tr>
<tr>
<td>63621</td>
</tr>
</tbody>
</table>
Prior authorization requests received after the requested start date of service will be denied for dates of service prior to the date the request was received.

Prior authorization requirements for stereotactic radiosurgery and stereotactic body radiation therapy may include, but are not limited to, diagnoses indicating one of the following medical conditions:

- Benign and malignant tumors of the central nervous system
- Vascular malformations
- Soft tissue tumors in chest, abdomen, or pelvis
- Trigeminal neuralgia refractory to medical management

Stereotactic radiosurgery and stereotactic body radiation therapy are considered investigational and not a benefit of Texas Medicaid for all other indications including, but not limited to, epilepsy, chronic pain, and pancreatic adenocarcinoma.

Prior authorization requirements for proton beam (procedure codes 77520, 77522, 77523, 77525, and S8030) and helium ion radiosurgery (procedure code 77423) may include, but are not limited to, diagnoses indicating one of the following medical conditions:

- Melanoma of the uveal tract (iris, choroid, ciliary body)
- Postoperative treatment for chordomas or low-grade chondrosarcomas of the skull or cervical spine
- Prostate cancer
- Pituitary neoplasms
- Other central nervous system tumors located near vital structures

Prior authorization for neutron beam radiosurgery may be considered for malignant neoplasms of the salivary gland.

Prior authorization requirements for procedure code 77399 include, but are not limited to, diagnosis, documentation of medical necessity, a specific description of the procedure to be performed, and an indication that the procedure would not be covered by a more specific procedure code.

Stereotactic radiosurgery and stereotactic body radiation therapy will not be prior authorized for clients with metastatic disease and a projected life span of less than six months or for clients with widespread cerebral or extracranial metastasis that is not responsive to systemic therapy.

### Other Limitations on Stereotactic Radiosurgery

In the following table, the procedure codes in Column A may be reimbursed when at least one corresponding procedure code from Column B has been paid to the same provider for the same date of service:

<table>
<thead>
<tr>
<th>Column A Procedure Code</th>
<th>Column B Procedure Code</th>
</tr>
</thead>
<tbody>
<tr>
<td>77522</td>
<td>77523</td>
</tr>
<tr>
<td>77525</td>
<td>G0339</td>
</tr>
<tr>
<td>G0340</td>
<td>G6002</td>
</tr>
<tr>
<td>S8030</td>
<td></td>
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</tbody>
</table>

Procedure codes 61796 and 63620 must not be billed more than once per course of treatment.
Procedure codes 61797 and 61799 must not be billed more than once per lesion, and may only be billed up to four times for the entire course of treatment, regardless of the number of lesions treated.

Procedure code 63621 may only be billed up to two times for the entire course of treatment, regardless of the number of lesions treated.

Procedure codes 77336 and 77370 may be reimbursed to the ordering physician if consultation is performed with a qualified medical physicist for stereotactic radiosurgery, brachytherapy, or any other method of radiation therapy for which medical necessity is determined.

Procedure code 77336 will be limited to once per week of treatment.

Procedure code 77370 will be limited to once per course of treatment. A special medical physics consultation may be reimbursed when the input and complex analysis of a qualified medical physicist are beyond that of a continuing medical physics consultation and are necessary to address a patient-specific reason or scenario.

9.2.61 Radiology Services

In compliance with HHS regulations, physicians (MDs and DOs), group practices, and clinics may not bill for radiology services provided outside their offices. These services must be billed directly by the facility/provider that performs the service.

This restriction does not affect radiology services performed by physicians or under their supervision in their offices. The radiology equipment must be owned by physicians and be located in their office to allow for billing of TOS 4 (complete procedure) or TOS T with modifier TC to Texas Medicaid. If physicians are members of a clinic that owns and operates radiology facilities, they may bill for these services. However, if physicians practice independently and share space in a medical complex where radiology facilities are located, they may not bill for these services even if they own or share ownership of the facility, unless they supervise and are responsible for the operation of the facilities on a daily basis.

Providers billing for three or more of the same radiology procedures on the same day must indicate the time the procedure was performed to indicate that it is not a duplicate service. The use of modifiers 76 and 77 does not remove the requirement of indicating the times services were rendered. The original claim will be denied but can be appealed with the documentation of procedure times.

When billing for services in an inpatient or outpatient hospital setting, the radiologist may only bill the professional interpretation of procedures (modifier 26). This also applies when providing services to a client who is in an inpatient status even if the client is brought to the radiologist’s office for the service. The hospital is responsible for all facility services (the technical component) even if the service is supplied by another facility/provider.

A separate charge for an X-ray interpretation billed by the attending or consulting physician is not allowed concurrently with that of the radiologist. Interpretations are considered part of the attending or consulting physician’s overall work-up and treatment of the patient.

Providers other than radiologists are sometimes under agreement with facilities to provide interpretations in specific instances. Those specialties may be paid if a radiologist does not bill for the professional component of X-ray procedures.

If duplicate billings are found between radiologists and the other specialties, the radiologist may be paid, and the other provider is denied.

Oral preparations for X-rays are included in the charge for the X-ray procedure when billed by a physician. Separate charges for the oral preparation are denied as part of another procedure on the same day.

Separate charges for injectable radiopharmaceuticals used in the performance of specialized X-ray procedures may be paid. If a procedure code is not indicated, an unlisted code must have a drug name, route of administration, and dosage written on the claim.
9.2.61.1 Diagnosis Requirements
Physicians enrolled and practicing as radiologists are not routinely required to send a diagnosis with their request for payment except when providing the following services:

- Arteriograms
- Venography
- Chest X-rays
- Cardiac blood pool imaging
- Echography

Radiologists are required to identify the referring provider by full name and credentials in Block 17 of the CMS-1500 claim form. Radiology procedures submitted by all other physician specialties must reference a diagnosis with every procedure billed. As with all procedures billed to Texas Medicaid, baseline screening and/or comparison studies are not a benefit.

9.2.61.2 Cardiac Blood Pool Imaging
Cardiac blood pool imaging may be reimbursed with procedure codes 78472, 78473, 78481, 78483, 78494, and 78496. Prior authorization is required for outpatient diagnostic services.

Refer to: Subsection 9.2.25.9, “Myocardial Perfusion Imaging” in this handbook for more information about myocardial perfusion imaging.

Section 3, “Radiological and physiological laboratory services” in the Radiology and Laboratory Services Handbook (Vol. 2, Provider Handbooks) for additional information and authorization requirements.

9.2.61.3 Chest X-Rays
All providers including radiologists billing for chest X-rays must supply a diagnosis code.

Screening, baseline, or rule-out studies do not qualify for reimbursement.

9.2.61.4 Magnetic Resonance Angiography (MRA)
MRA is an effective diagnostic tool used to detect, diagnose, and aid the treatment of heart disorders, stroke, and blood vessel diseases.

Refer to: Section 3, “Radiological and physiological laboratory services” in the Radiology and Laboratory Services Handbook (Vol. 2, Provider Handbooks) for additional information and authorization requirements.

9.2.61.5 Magnetic Resonance Imaging (MRI)
MRIs may be an effective diagnostic tool for detecting defects, diseases, and trauma.

Refer to: Section 3, “Radiological and physiological laboratory services” in the Radiology and Laboratory Services Handbook (Vol. 2, Provider Handbooks) for additional information and authorization requirements.

9.2.61.6 Technetium TC 99M
Procedure codes A9500 (Sestamibi) and A9502 (Tetrofosmin) are limited to three per day when billed by the same provider.

9.2.62 Magnetoencephalography (MEG)
Magnetoencephalography is a benefit of Texas Medicaid when medically necessary for the presurgical evaluation of clients with intractable epilepsy (i.e., refractory or drug-resistant epilepsy), brain tumors, vascular malformations of the brain, or when one or more conventional measures of localizing the seizure focus have failed to provide sufficient information.
MEG is a noninvasive method of measuring magnetic fields in the brain and is used to precisely localize both the essential functional cortex (i.e., eloquent cortex) and abnormal epileptogenic brain activity as part of a presurgical evaluation. The origin of abnormal MEG brain activity can be precisely localized (source localization) and displayed as a map or image.

The term magnetic source imaging (MSI) refers to an imaging technique that combines a MEG scan with an anatomic magnetic resonance imaging (MRI) image of the brain to map or visualize brain activity. MEG may assist in guiding the placement of intracranial Electroencephalography (EEG) and, in some patients, avoid an unnecessary intracranial EEG. In the case of pre-surgical mapping of patients with operable lesions, MEG provides non-invasive localization of eloquent cortices (e.g., motor, sensory, language, auditory, or visual).

Physicians must provide MEG services in a comprehensive level IV epilepsy center or a physiological laboratory. A neurologist, epileptologist, or neurosurgeon must order the MEG test.

MEG is not a stand-alone test. Pre-surgical evaluation with MEG testing must include a comprehensive evaluation by the medical team.

Procedure codes 95965, 95966, and 95967 may be reimbursed for MEG services. Procedure code 95967 is an add-on code and must be submitted with procedure code 95966.

Physicians may be reimbursed for the professional component of MEG services.

**9.2.62.1 Prior Authorization for MEG**

Prior authorization is required for MEG. Prior authorization requests must be submitted using the Special Medical Prior Authorization (SMPA) Request Form. The ordering physician must sign and date the form and submit it to the SMPA department. Requests must include documentation supporting the medical necessity of the study. The ordering physician must maintain all documentation.

Providers must include information about the MEG test facility. This information must be documented on the SMPA form.

Prior authorization requests must include a completed SMPA request form and all of the following documentation:

- Documentation of one of the following conditions: intractable epilepsy, brain tumors, or vascular malformations of the brain
- The statement of medical necessity from the ordering physician, which must support the need for MEG with identified medical conditions as applicable, including:
  - History of treatment methods used
  - Length of treatment and treatment outcomes
  - Date of onset of supporting diagnoses
  - Types of previous diagnostic testing used or considered and documentation that indicates how these tests have failed to provide the necessary information to address the client’s medical needs or when one or more conventional measures of localizing the seizure focus have failed to provide sufficient information

Documentation from the ordering physician outlining how the MEG test will assist in identifying the area to be resected in instances when an MEG test is needed due to a tumor and surgery is the first option.

Documentation that includes the name and number of medications, tried and failed, to control the client’s seizure activity when the MEG request is related to intractable epilepsy.

The date of prior MEG, the results of the previous MEG tests, and supporting medical documentation outlining the medical reasons for the repeat MEG requested if the request is for a repeat MEG.
Providers may submit prior authorization requests electronically, through the provider website, fax, or by standard mail.

The provider may complete and submit the required prior authorization documentation through any approved electronic method. The provider must maintain a copy of the prior authorization request as well as all submitted documentation in the client’s medical record at the performing provider’s place of business, in order to complete the prior authorization process electronically.

The provider may complete and submit the required prior authorization documentation through fax or standard mail and must maintain a copy of the prior authorization request as well as all submitted documentation in the client’s medical record at the performing provider’s place of business, to complete the prior authorization process by paper.

Providers must include correct and complete information, such as documentation of medical necessity for the service(s) requested, in order to avoid unnecessary denials. Providers must maintain documentation of medical necessity in the client’s medical record. The requesting provider may be asked for additional information to clarify or complete a request.

Requests for prior authorization with documentation supporting the medical necessity for the number of studies requested must be received on, or before, the requested date(s) of service.

**Note:** Requests received after the services are performed will be denied for dates of service that occurred before the date the request was received.

### 9.2.62.2 Documentation Requirements

In addition to documentation requirements outlined in the “Prior Authorization for MEG” section, the following requirements apply:

- All MEG services are subject to retrospective review to ensure that the documentation in the client’s medical record supports the medical necessity of the service(s) provided.

- Magnetic Source Imaging procedure code S8035 is not a benefit of Texas Medicaid, but it may be used for informational purposes.

### 9.2.62.3 Noncovered Services

The following MEG services are not benefits of Texas Medicaid:

- MEG when used as a stand-alone test for epilepsy
- MEG used as a first-line diagnostic screening
- MEG when used for evaluation of:
  - Alzheimer’s disease
  - Autism
  - Cognitive and mental disorders
  - Developmental dyslexia
  - Learning disorders
  - Migraines
  - Multiple sclerosis
  - Parkinson’s disease
  - Schizophrenia
  - Stroke rehabilitation
- Traumatic brain injury

Note: This list is not all inclusive.

9.2.63 Reduction Mammaplasties

9.2.63.1 Prior Authorization for Reduction Mammaplasty

Procedure code 19318 is the removal of breast tissue and is a benefit of Texas Medicaid when prior authorized.

For prior authorization of reduction mammaplasty, a completed “Medicaid Certificate of Medical Necessity for Reduction Mammaplasty” form signed and dated by the physician, must be submitted and include at least one of the following criteria:

- Evidence of severe neck and/or back pain with incapacitation from the pain.
- Evidence of ulnar pain or paresthesia from thoracic nerve root compression.
- Submammary dermatological conditions such as intertrigo and acne that are refractory to conventional medication.
- Shoulder grooving with ulceration due to breast size.

In addition to the above criteria, documentation must indicate:

- The minimum weight of tissue expected to be removed from each breast with consideration to height and weight is as follows:

<table>
<thead>
<tr>
<th>Height and Weight Chart</th>
</tr>
</thead>
<tbody>
<tr>
<td>Under 5’</td>
</tr>
<tr>
<td>5’-5’.4”</td>
</tr>
<tr>
<td>5’.4”-5’.7”</td>
</tr>
<tr>
<td>5’.7”- and up</td>
</tr>
</tbody>
</table>

- The client, if 40 years of age or older, has had a mammogram within the past year that was negative for cancer.

The following services are not a benefit of Texas Medicaid:

- Reduction mammaplasty for cosmetic purposes (such as the equalization of breast size)
- Augmentation mammaplasty to increase breast size

The physician is required to maintain the following documentation in the client’s clinical records:

- A complete history and physical
- Pulmonary function studies results
- Past treatments, therapies, and outcomes for pain control and weight reduction

The physician is required to maintain preoperative photographs (frontal and lateral views) in the client’s clinical records and must be made available to Texas Medicaid upon request.

For reimbursement purposes on a bilateral procedure, the full allowed amount will be paid to the surgeon and assistant surgeon for the first breast reduction and one half the allowed amount will be paid for the second reduction. Facilities are paid for one surgical procedure.
When submitting for prior authorization, requests must be sent to TMHP Special Medical Prior Authorization. Sending requests directly to the TMHP Medical Director delays the processing of the request. Providers are to mail prior authorization requests for reduction mammoplasty to the following address:

Texas Medicaid & Healthcare Partnership
Special Medical Prior Authorization
12357-B Riata Trace Parkway, Suite 100
Austin, TX 78727
Fax 1-512-514-4213

9.2.64 Renal Disease

9.2.64.1 Dialysis Patients

Physician reimbursement for supervision of patients on dialysis is based on a monthly capitation payment (MCP) calculated by Medicare. The MCP is a comprehensive payment that covers all physician services associated with the continuing medical management of a maintenance dialysis patient for treatments received in the facility. An original onset date of dialysis treatment must be included on claims for all renal dialysis procedures in all POSs except inpatient hospital. The original onset date must be the same date entered on the 2728 form sent to the Social Security office.

9.2.64.1.1 Physician Supervision of Dialysis Patients

Physician supervision of outpatient ESRD services includes services provided in the course of office visits where any of the following occur:

- The routine monitoring of dialysis.
- The treatment or follow-up of complications of dialysis, including:
  - The evaluation of related diagnostic tests and procedures.
  - Services involved in prescribing therapy for illnesses unrelated to renal disease, if the treatment occurs without increasing the number of physician-client contacts.

Use the following procedure codes when billing for physician supervision of outpatient ESRD dialysis services:

<table>
<thead>
<tr>
<th>Procedure Codes</th>
</tr>
</thead>
<tbody>
<tr>
<td>90951</td>
</tr>
<tr>
<td>90961</td>
</tr>
</tbody>
</table>

The procedure codes must be billed as described below:

- In the circumstances where the client is not on home dialysis and has had a complete assessment visit during the calendar month and ESRD-related services are provided for a full month, procedure codes 90951, 90952, 90953, 90954, 90955, 90956, 90957, 90958, 90959, 90960, 90961, or 90962 must be used, determined by the number of face-to-face visits the physician has had with the client during the month, and the client’s age.
- When a full calendar month of ESRD-related services are reported for clients on home dialysis, procedure codes 90963, 90964, 90965, or 90966 must be used, determined by the client’s age.
- Report procedure codes 90967, 90968, 90969, and 90970 when ESRD related services are provided for less than a full month, per day, under the following conditions:
  - The client is seen for a partial month and is not on home dialysis and received one or more face-to-face visits but did not receive a complete assessment.
  - The client is on home dialysis and received less than a full month of services.
• The client is a transient client.
• The client was hospitalized during a month of services before a complete assessment could be performed.
• Dialysis was stopped due to recovery or death of client.
• The client received a kidney transplant.
• Procedure codes 90967, 90968, 90969, and 90970 are limited to one per day by any provider. When billing procedure code 90967, 90968, 90969, or 90970, the date of service must indicate each day that supervision was provided.
• Procedure codes 90967, 90968, 90969, and 90970 will be denied when billed within the same calendar month by any provider as procedure code 90951, 90952, 90953, 90954, 90955, 90956, 90957, 90958, 90959, 90960, 90961, 90962, 90963, 90964, 90965, or 90966.
• Procedure codes 90951, 90952, 90953, 90954, 90955, 90956, 90957, 90958, 90959, 90960, 90961, 90962, 90963, 90964, 90965, or 90966 are limited to once per calendar month by any provider, and only one service may be reimbursed per calendar month by any provider.

The following services may be provided in conjunction with physician supervision of ESRD dialysis but are considered non-routine and may be billed separately:
• Declotting of shunts when performed by the physician.
• Physician services to inpatient clients. If a client is hospitalized during a calendar month of ESRD related services before a complete assessment is performed, or the client receives one or more face-to-face assessments, but the timing of inpatient admission prevents the client from receiving a complete assessment, the physician must bill procedure code 90967, 90968, 90969, or 90970 for each date of outpatient supervision and bill the appropriate hospital evaluation and management code for individual services provided on the hospitalized days. If a client has a complete assessment during a month in which the client is hospitalized, procedure code 90951, 90952, 90953, 90954, 90955, 90956, 90957, 90958, 90959, 90960, 90961, or 90962 must be reported for the month of supervision, determined by the number of face-to-face physician visits with the client during the month, and the client’s age. The appropriate inpatient evaluation and management codes must be reported for procedures provided during the hospitalization.
• Dialysis at an outpatient facility other than the usual dialysis setting for a patient of a physician who bills the MCP. The physician must bill procedure code 90967, 90968, 90969, or 90970 for each date supervision is provided. The physician may not bill for days that the client dialyzed elsewhere.
• Physician services beyond those that are related to the treatment of the patient’s renal condition that cause the number of physician-patient contacts to increase. Physicians may bill on a fee-for-service basis if they supply documentation on the claim that the illness is not related to the renal condition and that additional visits are required.

Use procedure codes 90935, 90937, 90945, and 90947 for inpatient dialysis services for ESRD or non-ESRD clients when the physician is present during dialysis treatment. The physician must be physically present and involved during the course of the dialysis. These codes are not payable for a cursory visit by the physician; hospital visit codes must be used for a cursory visit.

The hospital procedure codes 90935, 90937, 90945, and 90947 are for complete care of the patient; hospital visits cannot be billed on the same day as these codes. However, if the physician only sees the patient when they are not dialyzing, the physician must bill the appropriate hospital visit code. The inpatient dialysis code must not be submitted for payment.

Only one of procedure code 90935, 90937, 90945, or 90947 may be reimbursed per day, any provider.

Procedure codes 90935, 90937, 90945, and 90947 may also be used for outpatient dialysis services for non-ESRD clients.
Inpatient services provided to hospitalized clients for whom the physician has agreed to bill monthly, may be reimbursed in one of the following three ways:

- The physician may elect to continue monthly billing, in which case she or he may not bill for individual services provided to the hospitalized clients.
- The physician may reduce the monthly bill by 1/30th for each day of hospitalization and charge fees for individual services provided on the hospitalized days.
- The physician may bill for inpatient dialysis services using the inpatient dialysis procedure codes. The physician must be present and involved with the clients during the course of the dialysis.

Clients may receive dialysis at an outpatient facility other than his or her usual dialysis setting, even if their physician bills for monthly dialysis coordination. The physician must reduce the monthly billed amount by 1/30th for each day the client is dialyzed elsewhere.

Physician services beyond those related to the treatment of the client’s renal condition may be reimbursed on a fee-for-service basis. The physician should provide documentation stating the illness is not related to the renal condition and added visits are required.

Payment is made for physician training services in addition to the monthly capitation payment for physician supervision rendered to maintenance facility clients.

### 9.2.64.2 Laboratory Services for Dialysis Patients

Texas Medicaid may reimburse for laboratory services performed for dialysis patients.

Charges for routine laboratory services performed according to established frequencies are included in the facility’s composite rate billed to Texas Medicaid regardless of where the tests were performed. Routine laboratory testing processed by an outside laboratory are billed to the facility and billed by a renal dialysis facility, unless they are inclusive tests.

Nonroutine laboratory services for people dialyzing in a facility and all laboratory work for people on CAPD may be billed separately from the dialysis charge.

**Refer to:** Subsection 6.2.9, "Laboratory and Radiology Services" in the Clinics and Other Outpatient Facility Services Handbook (Vol. 2, Provider Handbooks) for more information on laboratory services.

### 9.2.64.3 Self-Dialysis Patients

Physician reimbursement for supervision of patients on self-dialysis is made after completion of the patient’s training. If the training is not completed, payment is proportionate to the amount of time spent in training. Payment for training may be made in addition to payment under the MCP for physician supervision of an in-facility maintenance dialysis patient. Use procedure codes 90989 and 90993 for dialysis training regardless of the type of training performed. These procedure codes must be billed as specified:

- When complete dialysis training is provided, bill procedure code 90989. Providers are to use modifier AT when using this procedure code. The date of service indicates the date training was completed, and the quantity is 1.
- When dialysis training is not completed, bill procedure code 90993. The date of service must list each day that a session of training was provided and the quantity must indicate the number of training sessions provided.

The amount of reimbursement of subsequent training is determined by prorating the physician’s payment for initial training sessions. The amount of payment for each additional training session does not exceed $20.
9.2.64.3.1 Physician Supervision
All physician services required to create the capacity for self-dialysis must include:

- Direction of and participation in training of dialysis patients.
- Review of family and home status and environment, and counseling and training of family members.
- Review of training progress.

9.2.64.3.2 Initial Training
The following services are included in the physician charge for supervision of a client on self-dialysis:

- Physician services rendered during a dialysis session including those backup dialyses that occur in outpatient facility settings.
- Office visits for the routine evaluation of patient progress, including the interpretation of diagnostic tests and procedures.
- Physician services rendered by the attending physician in the course of an office visit, the primary purpose of which is routine monitoring or the follow-up of complications of dialysis, including services involved in prescribing therapy for illnesses unrelated to renal disease, which may be appropriately treated without increasing the number of contacts beyond those occurring at regular monitoring sessions or visits for treatment of renal complications.
- General support services (for example, arranging for supplies).

9.2.64.3.3 Subsequent Training
No additional payment is made after the initial self-dialysis training course unless subsequent training is required for one of the following reasons:

- A change from the client’s treatment machine to one the client had not been trained to use in the initial training course
- A change in setting
- A change in dialysis partner

The physician must document the reason for additional training sessions on the CMS-1500 paper claim form.

Dialysis equipment and supplies used by the client who dialyzes in the home are not benefits of Texas Medicaid, including the lease or purchase of dialysis machines and disposable supply kits.

9.2.65 Sign Language Interpreting Services
Sign language interpreting services are benefits of Texas Medicaid. Providers must use procedure code T1013 with modifier U1 for the first hour of service, and T1013 with modifier UA for each additional 15 minutes of service. Procedure code T1013 billed with modifier U1 is limited to once per day, same provider, and procedure code T1013 billed with modifier UA is limited to a quantity of 28 per day, same provider.

Sign language interpreting services are available to Medicaid clients who are deaf or hard of hearing or to a parent or guardian of a Medicaid client if the parent or guardian is deaf or hard of hearing.

Physicians in private or group practices with fewer than 15 employees may be reimbursed for this service. The physician will be responsible for arranging and paying for the sign language interpreting services to facilitate the medical services being provided. The physician will then seek reimbursement from Texas Medicaid for providing this service.
Sign language interpreting services must be provided by an interpreter who possesses one of the following certification levels (i.e., levels A through H) issued by either the Office of Deaf and Hard of Hearing Services, Board for Evaluation of Interpreters (BEI) or the National Registry of Interpreters for the Deaf (RID).

Certification Levels:
- BEI Level I/Ii and BEI OC: B (Oral Certificate: Basic)
- BEI Basic and RID NIC (National Interpreter Certificate) Certified
- BEI Level II/IIi, RID CI (Certificate of Interpretation), RID CT (Certificate of Transliteration), RID IC (Interpretation Certificate), and RID TC (Transliteration Certificate)
- BEI Level III/IIIi, BEI OC: C (Oral Certificate: Comprehensive), BEI OC: V (Oral Certificate: Visible), RID CSC (Comprehensive Skills Certificate), RID IC/TC, RID CI/CT, RID RSC (Reverse Skills Certificate), and RID CDI (Certified Deaf Interpreter)
- BEI Advanced and RID NIC Advanced
- BEI IV/IVi, RID MCSC (Master Comprehensive Skills Certificate), and RID SC: L (Specialist Certificate: Legal)
- BEI V/VI
- BEI Master; and RID NIC Master

Interpreting services include the provision of voice-to-sign, sign-to-voice, gestural-to-sign, sign-to-gestural, voice-to-visual, visual-to-voice, sign-to-visual, or visual-to-sign services for communication access provided by a certified interpreter.

The physician requesting interpreting services must maintain documentation verifying the provision of interpreting services. Documentation of the service must be included in the client’s medical record and must include the name of the sign language interpreter and the interpreter’s certification level. Documentation must be made available if requested by HHSC or its designee.

9.2.66 Skin Therapy

Skin therapy is a benefit of Texas Medicaid and may be reimbursed with the following procedure codes:

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Claims for incision and drainage of acne when the diagnosis states there is infection or pustules may be paid.

Procedure codes 96900, 96910, 96912, 96913, 96920, 96921, and 96922 are covered benefits for the following diagnosis codes:

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Intralesional injection(s) may be considered for reimbursement in addition to an office visit.

Procedure codes 11900 and 11901 are covered benefits for intralesional injections for the following diagnosis codes:

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Diagnosis Codes

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<tr>
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<td>T2622S</td>
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<td>T2651XD</td>
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<tr>
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<td>T2682XS</td>
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<td>T2691XD</td>
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<td>T275XXA</td>
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<td>T280XXA</td>
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<tr>
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<td>T282XXA</td>
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<td>T28411A</td>
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<tr>
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<td>T28412D</td>
<td>T28412S</td>
<td>T2849XA</td>
<td>T2849XD</td>
<td>T2849XS</td>
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</tbody>
</table>
Procedure codes 15782, 15783, 15792, 15793, and 17999 require prior authorization. Requests for prior authorization must be submitted by the physician to the Special Medical Prior Authorization (SMPA) department with documentation supporting the medical necessity of the anticipated procedure. This documentation must remain in the client’s medical record and is subject to retrospective review. To avoid unnecessary denials, the physician must provide correct and complete information.

Dermabrasion procedures (procedure codes 15782 and 15783) and chemical peel procedures (procedure codes 15792 and 15793) may be prior authorized with documentation that the client meets all of the following criteria:

- A diagnosis of actinic keratosis with more than three lesions.
- Failed conservative treatment or documentation that conservative treatment is contraindicated.

Prior authorization requests for procedure code 17999 must include the following documentation:

- A clear, concise description of the procedure to be performed.
- Reason for recommending the particular procedure.
- Documentation that a specific procedure code is not available for the procedure requested.
- The client’s diagnosis.
- Medical records indicating prior treatment for the diagnosis and the medical necessity of the requested procedure.
- Place of service the procedure is to be performed.
- Documentation that the procedure is not investigational or experimental.
- The physician’s intended fee for the procedure including a comparable procedure code.

### 9.2.67 Sleep Studies

Sleep study procedure code 95806 is not a benefit of Texas Medicaid.
9.2.67.1 Actigraphy

Actigraphy (procedure code 95803) may be reimbursed in the office or outpatient hospital setting with a limit of one per day, and two per rolling year by any provider. Claims denied for more than two times per year may be appealed with documentation of medical necessity.

Actigraphy can be performed as a stand-alone procedure or as an adjunct to polysomnography or multiple sleep latency test (MSLT).

Actigraphy (procedure code 95803) must be billed with one of the following diagnosis codes:

<table>
<thead>
<tr>
<th>Diagnosis Codes</th>
</tr>
</thead>
<tbody>
<tr>
<td>F5104, F5105, F5113, G2581, G4700, G4701, G4709, G4710, G4711, G4712, G4713, G4714, G4719, G4720, G4721, G4722, G4723, G4724, G4725, G4726, G4727, G4729, G4761</td>
</tr>
</tbody>
</table>

If the primary care physician performs the actigraphy, the technical component must be billed (procedure code 95803 with modifier TC).

Documentation of actigraphy must include a hard-copy printout or electronic file. Interpretation and treatment recommendations must be completed by a sleep specialist. The physician’s professional interpretation and report must include inspection of the entire recording and integration of the information gathered from other professionals’ analysis and observations. Documentation of the interpretation must be maintained by the interpreting physician.

Under the following conditions, actigraphy may be a useful adjunct to a detailed history, examination, and subjective sleep diary for the diagnosis and treatment of insomnia, circadian-rhythm disorders, and excessive sleepiness:

- When demonstration of multiday rest-activity patterns is necessary to diagnose, document severity, and guide the proper treatment.
- When more objective information regarding the day-to-day timing or the amount or patterns of a client’s sleep is necessary for optimal clinical decision-making.
- When the severity of a sleep disturbance reported by the client or caretaker seems inconsistent with clinical impressions or laboratory findings.
- To clarify the effects of, and under some instances, compliance with pharmacologic, behavioral, phototherapeutic, or chronotherapeutic treatment.
- In symptomatic clients for whom an accurate history cannot be obtained and at least one of the following is true:
  - A polysomnographic study has already been conducted.
  - A polysomnographic study is considered unlikely to be of much diagnostic benefit.
  - A polysomnographic study is not yet clearly indicated (because of the absence of accurate historical data).
  - A polysomnographic study is not immediately available.

Actigraphy may be useful in the assessment of specific aspects of the following disorders:

- Insomnia. Assessment of sleep variability, measurement of treatment effects, and detection of sleep phase alterations in insomnia secondary to circadian rhythm disturbance.
- Restless legs syndrome or periodic limb movement disorder. Assessment of treatment effects.
9.2.67.2  **Pneumocardiograms**

Pneumocardiograms (procedure code 95807) are limited to clients who are birth through 12 months of age.

Pneumocardiograms are limited to one per day, and two per rolling year by any provider. Claims denied for more than two times per year may be appealed with documentation of medical necessity.

Procedure code 95807 must be billed with one of the following diagnosis codes:

<table>
<thead>
<tr>
<th>Diagnosis Codes</th>
</tr>
</thead>
<tbody>
<tr>
<td>G4731</td>
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</tbody>
</table>

Documentation of the complete readings associated with the pneumocardiogram and the physician’s interpretation must be maintained in the client’s medical record in a hard-copy printout or electronic file at the facility where the procedure is performed.

The physician’s interpretation and report must include inspection and integration of the information gathered from all physiological systems and other professionals’ analysis and observations.

9.2.67.3  **Polysomnography**

Polysomnography (procedure codes 95782, 95783, 95808, 95810, and 95811) is a benefit of Texas Medicaid.

Polysomnography is distinguished from sleep studies by the inclusion of sleep staging that includes a 1-to 4-lead electroencephalogram (EEG), electro-oculogram (EOG), and a limb or submental electromyogram (EMG).

Additional parameters of sleep that are evaluated in polysomnography include, but are not limited to, the following:

- ECG
- Airflow (by thermistor or intra-nasal pressure monitoring)
- Respiratory effort
- Adequacy of oxygenation by oximetry or transcutaneous monitoring
- Extremity movement or motor activity
- EEG monitoring for sleep staging
- Nocturnal penile tumescence
- Esophageal pH or intraluminal pressure monitoring
- Continuous blood pressure monitoring
- Snoring
- Body positions
- Adequacy of ventilation by end-tidal or transcutaneous CO2 monitoring

For a sleep study to be reported as a polysomnography, sleep must be recorded and staged. Use the following procedure codes to bill for polysomnography studies: 95782, 95783, 95808, 95810, and 95811.
Polysomnography (procedure codes 95782, 95783, 95808, 95810, and 95811) is limited to one per day and two per rolling year by any provider and is allowed for the following diagnosis codes:

<table>
<thead>
<tr>
<th>Diagnosis Codes</th>
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<tbody>
<tr>
<td>E6601 E662 F10182 F10282 F10982 F11182 F11282 F11982</td>
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<td>F13182 F13282 F13982 F14182 F14282 F14982 F15182 F15282</td>
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<tr>
<td>F15982 F19182 F19282 F19982 F5101 F5102 F5103 F5104</td>
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<tr>
<td>F5105 F5109 F5111 F5112 F5113 F5119 F513 F514</td>
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<tr>
<td>F515 F518 F519 G120 G121 G1221 G128 G2581</td>
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<tr>
<td>G373 G4700 G4701 G4710 G4711 G4712 G4713 G4719</td>
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<tr>
<td>G4720 G4721 G4722 G4723 G4724 G4725 G4726 G4727</td>
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<tr>
<td>G4729 G4730 G4731 G4732 G4733 G4734 G4735 G4736</td>
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<tr>
<td>G4737 G4739 G47411 G47419 G47421 G47429 G4750 G4751</td>
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<td>G4752 G4753 G4754 G4759 G4761 G4762 G4763 G4769</td>
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<tr>
<td>G478 G479 G7100 G7101 G7102 G7109 G7120 G7121</td>
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<tr>
<td>G71220 G71228 G7129 G809 G8250 G901 G931 J353</td>
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<tr>
<td>J9610 J9611 J9612 N5201 N5202 N5203 N521 N5235</td>
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<tr>
<td>N5236 N5237 Q040 Q041 Q042 Q078 Q308 Q311</td>
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<tr>
<td>Q775 Q777 Q778 Q779 Q781 Q789 Q870 R0681</td>
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<tr>
<td>R0902</td>
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</table>

Claims denied for more than two times per year may be appealed with documentation of medical necessity.

Documentation of the polysomnography testing must be maintained in the client’s medical record at the sleep facility and include approximately 1,000 pages or the electronically-stored equivalent of data during a single nighttime recording. Each record must be for sleep-wake states and stages, cardiac arrhythmias, respiratory events, motor activity, oxygen desaturations, and behavioral observations.

Documentation must also include the technologist’s analysis and report, the patient’s subjective report, and the influence of intervention applied during the night.

Interpretation and treatment recommendations must be completed by a sleep specialist. The physician’s professional interpretation and report must include inspection of the entire recording, examination of the technologist’s analysis and observations, and integration of the information gathered from all physiological systems. Documentation of the interpretation must be maintained in the sleep facility and by the interpreting physician.

9.2.67.4 Multiple Sleep Latency Test (MSLT)

Multiple sleep latency test (procedure code 95805) is limited to one per day and two per rolling year by any provider, and is restricted to the following diagnosis codes:

<table>
<thead>
<tr>
<th>Diagnosis Codes</th>
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<tbody>
<tr>
<td>E662 F5104 F5105 G2581 G4700 G4701 G4709 G4730</td>
</tr>
<tr>
<td>G47411 G47419 G47421 G47429 G4753 G4761</td>
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</tbody>
</table>
Claims denied for more than two times per year may be appealed with documentation of medical necessity.

Documentation of MSLT must be maintained in the client’s medical record at the sleep facility and include a hard copy or electronic copy of four to five 20-minute recordings of sleep-wake states and stages spaced at two-hour intervals throughout the day, taking approximately seven to nine hours to complete. In addition, documentation must include the physiological recordings typically made during daytime testing. These typically include:

- EEG
- Electro-oculogram (EOG)
- EMG
- EKG
- Audio and video recordings made during the monitored portion of the day

Documentation must also include the technologist’s analysis and report, the client’s subjective report, and the influence of intervention applied during the night.

Interpretation and treatment recommendations must be completed by a sleep specialist. The physician’s interpretation and report must include inspection of the entire recording, examination of the technologist’s analysis and observations, and integration of the information gathered from all physiological systems. Documentation of the interpretation must be maintained in the sleep facility and by the interpreting physician.

MSLT procedure code 95805 must be performed in conjunction with polysomnography procedure code 95782, 95783, 95808, 95810, or 95811. Polysomnography must be performed on the date before MSLT. MSLT that is not performed in conjunction with polysomnography will be denied, but may be considered on appeal with documentation that explains why the polysomnography did not occur.

### 9.2.67.5 Home Sleep Study Test

Home sleep study tests are unattended studies that are performed in the client’s home using a portable monitoring device. The portable monitoring device must meet American Academy of Sleep Medicine (AASM) practice parameters and clinical guidelines.

Home sleep study testing is a benefit of Texas Medicaid only when performed in conjunction with a comprehensive sleep evaluation that has been performed by a physician who is board-certified or board-eligible, as outlined in the AASM guidelines. Documentation of the comprehensive sleep evaluation must be kept in the client’s medical record. The evaluation must indicate probability of moderate to severe obstructive sleep apnea to support medical necessity for home sleep study testing.

Procedure codes G0398, G0399, and G0400 are a benefit for Texas Medicaid clients who are 18 years of age and older with suspected or proven simple, uncomplicated obstructive sleep apnea. Procedure codes G0398, G0399, and G0400 are restricted to diagnosis code G4733.

Home sleep study tests are payable to physicians in the office setting. Procedure codes G0398, G0399, and G0400 are limited to one per day and a combined total of two tests per rolling year, with any provider. If a client needs more than two tests in a rolling year, subsequent tests must be performed in a sleep facility.
9.2.67.6 Sleep Facility Restrictions for Polysomnography and Multiple Sleep Latency Testing

Sleep facilities that perform services for Medicaid clients must be accredited with the AASM or the Joint Commission of Accreditation of Healthcare Organizations (JCAHO). Sleep facilities must maintain documentation with proof that the facility is accredited. Documentation is subject to retrospective review. Sleep facilities that perform services for Texas Medicaid clients must also follow current AASM practice parameters and clinical guidelines.

Physicians who provide supervision in sleep facilities must be board-certified or board-eligible, as outlined in the AASM guidelines.

Sleep facility technicians, technologists, and trainees must demonstrate that they have the skills, competencies, education, and experience that are set forth by their certifying agencies and AASM as necessary for advancement in the profession.

Polysomnographic technologists, technicians, and trainees must meet the following supervision requirements:

- A polysomnographic trainee provides basic polysomnographic testing and associated interventions under the direct supervision of a polysomnographic technician, polysomnographic technologist, or a physician.

  **Note:** Direct supervision means that the supervising licensed/certified professional must be present in the office suite or building and immediately available to furnish assistance and direction throughout the performance of the service. It does not mean that the supervising professional must be present in the room while the service is provided.

- A polysomnographic technologist provides comprehensive evaluation and treatment of sleep disorders under the general supervision of the clinical director (MD or DO).

- A polysomnographic technician provides comprehensive polysomnographic testing and analysis and associated interventions under the general supervision of a polysomnographic technologist or clinical director (MD or DO).

- The supervising physician must be readily available to the performing technologist throughout the duration of the study, but is not required to be in the building.

The sleep facility must have one or more supervising physicians who are responsible for the direct and ongoing oversight of the quality of the testing performed, the proper operation and calibration of equipment used to perform tests, and the qualifications of the nonphysician staff who use the equipment.

Services provided without the required level of supervision are not considered medically appropriate and will be recouped upon retrospective record review.

Claims denied for more than two times per year may be appealed with documentation of medical necessity.

Documentation of MSLT must be maintained in the client’s medical record at the sleep facility and include a hard copy or electronic copy of four to five, 20-minute recordings of sleep-wake states and stages spaced at two-hour intervals throughout the day, taking approximately seven to nine hours to complete. In addition, documentation must include the physiological recordings typically made during daytime testing. These typically include:

- EEG
- Electro-oculogram (EOG)
- EMG
- EKG
• Audio and video recordings made during the monitored portion of the day

Documentation must also include the technologist’s analysis and report, the client’s subjective report, and the influence of intervention applied during the night.

Interpretation and treatment recommendations must be completed by a sleep specialist. The physician’s interpretation and report must include inspection of the entire recording, examination of the technologist’s analysis and observations, and integration of the information gathered from all physiological systems. Documentation of the interpretation must be maintained in the sleep facility and by the interpreting physician.

MSLT procedure code 95805 must be performed in conjunction with polysomnography procedure code 95808, 95810, or 95811. Polysomnography must be performed on the date before MSLT. MSLT that is not performed in conjunction with polysomnography will be denied, but may be considered on appeal with documentation that explains why the polysomnography did not occur.

9.2.68 Speech Therapy (ST) Services

Speech therapy (ST) is a payable benefit to physicians.

Refer to: Section 4, “Therapy Services Overview” in the Physical Therapy, Occupational Therapy, and Speech Therapy Services Handbook (Vol. 2, Provider Handbooks) for information about speech therapy services provided by a physician.

9.2.69 Surgery Billing Guidelines

9.2.69.1 Primary Surgeon

A primary surgeon may be reimbursed for services provided in the inpatient hospital, outpatient hospital setting, and ASC/HASC Center.

A surgeon billing for a surgery and an assistant surgery fee on the same day may be reimbursed if two separate procedures are performed.

Refer to: Subsection 9.2.69.7, “Multiple Surgeries” in this handbook.

9.2.69.2 Anesthesia Administered by Surgeon

If the physician bills for a surgical procedure and anesthesia for the same procedure, the surgery is paid and the anesthesia is denied as part of the surgical procedure. The exception to this policy is an epidural during labor and delivery.

Refer to: Subsection 9.2.7, “Anesthesia” in this handbook.

9.2.69.3 Assistant Surgeon

Assistant surgeons may be reimbursed 16 percent of the TMRM fee for the surgical procedures performed.

Medicaid follows the TEFRA regulations for assistant surgeons in teaching hospitals. TEFRA states that an assistant surgeon will not be paid in a hospital classified by Medicare as a teaching facility with an approved graduate training program in the performing physician’s specialty. Medicaid may consider reimbursement for an assistant surgeon at a teaching hospital classified by Medicare as a teaching facility with approved graduate training program if one of the following situations is present and documented on the claim:

• No qualified resident was available. (Modifier 82 may be used to document this exception.)

• There were exceptional medical circumstances such as an emergency or life-threatening situation requiring immediate attention (modifiers 80 and KX).

• The primary surgeon has a policy of never, without exception, involving a resident in the preoperative, operative, or postoperative care of a patient (modifiers 80 and KX).
• The surgical procedure was complex and required a team of physicians (modifiers 80 and KX).

Use of these modifiers is not required but expedites claims processing. Therefore, it is recommended that these modifiers be used in conjunction with the procedure code rather than a narrative statement when these specific circumstances exist.

All claims for assistant surgeon services must include in Block 32 of the CMS-1500 paper claim form the name, address, and provider identifier of the hospital in which the surgery was performed. If the physician seeks an exception to this TEFRA regulation based on unavailability of a qualified resident, the following certification statement must appear on or attached to the claim form:

“I understand that section 1842(b)(6)(D) of the Social Security Act generally prohibits reasonable charge payment for the services of assistants at surgery in teaching hospitals when qualified residents are available to furnish such services. I certify that the services for which payment is claimed were medically necessary, and that no qualified residents were available to perform the services. I further understand that these services are subject to postpayment review by TMHP.”

Surgical procedures that do not ordinarily require the services of an assistant, as identified by Medicare, are denied when billed as an assistant surgery. One assistant surgeon is reimbursed for surgical procedures when appropriate.

Use modifier AS when the physician assistant is not enrolled as an individual provider and provides assistance at surgery. The claim must include the PA’s name and license number. Only procedures currently allowed for assistant surgeons are payable.

PAs actively enrolled as a Medicaid provider with an assigned provider identifier may bill assistant surgery services on a separate claim form using the PA’s individual provider identifier and modifiers U7 and 80.

9.2.69.4 Bilateral Procedures

When a bilateral procedure is performed and an appropriate bilateral code is not available, a unilateral code must be used. The unilateral code must be billed twice with a quantity of 1 for each code. For all procedures, use modifiers LT (left) and RT (right) as appropriate. For example, bilateral application of short leg cast is billed as follows:

<table>
<thead>
<tr>
<th>Procedure Code</th>
<th>Modifier</th>
</tr>
</thead>
<tbody>
<tr>
<td>29405</td>
<td>LT</td>
</tr>
<tr>
<td>29405</td>
<td>RT</td>
</tr>
</tbody>
</table>

9.2.69.5 Cosurgery

Cosurgery (two surgeons) may be reimbursed when the skills of two surgeons (usually with different skills) are required in the management of a specific surgical procedure. Cosurgery is for a surgery where the two surgeons’ separate contributions to the successful outcome of the procedure are considered to be of equal importance.

Note: No additional reimbursement will be made for an assistant surgeon.

Cosurgeons may be reimbursed for surgical procedure codes that are billed with modifier 62 if the CMS fee schedule indicates that the procedure allows for cosurgeons. Claims will not suspend for manual review of the documentation of medical necessity. Reimbursement will be calculated at 62.5 percent of the amount allowed for the intraoperative portion of the surgical procedure’s fee.

No cosurgery payment is made for claims submitted without modifier 62. In instances where the surgeons do not use modifier 62, the first claim received at TMHP for the service is considered that of the primary surgeon, and the subsequent claim is denied as a previously paid service.
9.2.69.6 Global Fees

Texas Medicaid uses global surgical periods to determine reimbursement for services that are related to surgical procedures. The following services are included in the global surgical period:

- Preoperative care, including history and physical
- Hospital admission work-up
- Anesthesia (when administered and monitored by the primary surgeon)
- Surgical procedure (intraoperative)
- Postoperative follow-up and related services
- Complications following the surgical procedure that do not require return trips to the operating room

Texas Medicaid adheres to a global fee concept for minor and major surgeries and invasive diagnostic procedures. Global surgical periods are defined as follows:

- 0-day Global Period-Reimbursement includes the surgical procedure and all associated services that are provided on the same day.
- 10-day Global Period-Reimbursement includes the surgical procedure, any associated services that are provided on the same day of the surgery, and any associated services that are provided for up to 10 days following the date of the surgical procedure.
- 90-day Global Period-Reimbursement includes the surgical procedure, preoperative services that are provided on the day before the surgical procedure, any associated services that are provided on the same day of the surgery, and any associated services that are provided for up to 90 days following the date of the surgical procedure.

Procedure codes that are designated as “Carrier Discretion” will have their global periods determined by HHSC.

Note: All unlisted surgical procedure codes have a 42 day global period assigned by Texas Medicaid.

The global surgical fee period applies to both emergency and nonemergency surgical procedures. Physicians who are in the same group practice and specialty must bill, and are reimbursed, as if they were a single provider.

Modifiers

For services that are rendered in the preoperative, intraoperative, or postoperative period to be correctly reimbursed, providers must use the appropriate modifiers from the following table. Failure to use the appropriate modifier may result in recoupment.

<table>
<thead>
<tr>
<th>Modifiers Related to Surgical Fees</th>
</tr>
</thead>
<tbody>
<tr>
<td>24</td>
</tr>
<tr>
<td>58</td>
</tr>
</tbody>
</table>

For services that are billed with modifier 54, 55, or 56, medical record documentation must be maintained by both the surgeon and the physician who provides preoperative or postoperative care. Reimbursement for claims associated with modifier 54, 55, or 56 is limited to the same total amount as would have been paid if only one physician provided all of the care, regardless of the number of physicians who actually provide the care.

If a physician provided all of the preoperative, intraoperative, and postoperative care, claims may be considered for reimbursement when they are submitted without a modifier.
Documentation Requirements
For services that are billed with any of the listed modifiers to be considered for reimbursement, providers must maintain documentation in the client’s medical record that supports the medical necessity of the services. Acceptable documentation includes, but is not limited to, progress notes, operative reports, laboratory reports, and hospital records.

On a case-by-case basis, providers may be required to submit additional documentation that supports the medical necessity of services before the claim will be reimbursed.

*Note:* Retrospective review may be performed to ensure that the submitted documentation supports the medical necessity of the surgical procedure and any modifier used to bill the claim.

Preoperative Services
Preoperative physician E/M services (such as office or hospital visits) that are directly related to the planned surgical procedure and provided during the preoperative limitation period will be denied if they are billed by the surgeon or anesthesiologist who was involved in the surgical procedure.

Reimbursement will be considered when the E/M services are performed for distinct reasons that are unrelated to the procedure. E/M services that meet the definition of a significant, separately identifiable service may be billed with modifier 25 if they are provided on the same day by the same provider as the surgical procedure.

Modifier 25 is not used to report an E/M service that results in a decision to perform a surgical procedure. Documentation that supports the provision of a significant, separately identifiable E/M service must be maintained in the client’s medical record and made available to Texas Medicaid upon request. If the decision to perform a minor procedure is made during an E/M visit immediately before the surgical procedure, the E/M visit is considered a routine preoperative service and is not separately billable.

Physicians who provide only preoperative services for surgical procedures with a 10- or 90-day global period may submit claims using the surgical procedure code with the identifying modifier 56. Reimbursement will be limited to a percentage of the fee for the surgical procedure.

E/M services that are provided during the preoperative period (one day before or the same day) of a major surgical procedure (90-day global period) and result in the initial decision to perform the surgical procedure may be considered for reimbursement when billed with modifier 57. The client’s medical record must clearly indicate when the initial decision to perform the procedure was made.

Intraoperative Services
Physicians who perform a surgical procedure with a 10- or 90-day global period but do not render postoperative services must bill the surgical procedure code with modifier 54. Modifier 54 indicates that the surgeon provided the surgical care only. Documentation in the medical record must support the transfer of care and must indicate that an agreement has been made with another physician to provide the postoperative management.

Postoperative services
Postoperative services that are directly related to the surgical procedure are included in the global surgical fee and are not reimbursed separately. Postoperative services include, but are not limited to, all of the following:

- Postoperative follow-up visits (any place of service)
- Postoperative pain management
- Miscellaneous services, including:
  - Dressing changes
  - Local incision care
• Platelet gel
• Removal of operative packs
• Removal of cutaneous sutures, staples, lines, wires, drains, casts, or splints
• Replacement of vascular access lines
• Insertion, irrigation, and removal of urinary catheters, routine peripheral intravenous lines, nasogastric tubes, and rectal tubes
• Changes or removal of tracheostomy tubes

Note: Removal of postoperative dressings or anesthetic devices is not eligible for separate reimbursement as the removal is considered part of the allowance for the primary surgical procedure.

If the surgeon provides the surgery and only the postoperative care for a procedure that has a 10- or 90-day global period, the surgeon must include the following details on the claim form:

• The surgical procedure, date of the surgery, and modifier 54, which indicates that he or she was the surgeon.
• The surgical procedure, date of service, and modifier 55 to denote the postoperative care.

Note: Providers must not submit a claim for the postoperative care until after the client has been seen during a face-to-face follow-up visit.

When a transfer of care occurs for postoperative care for procedures that have a 10- or 90-day global period, the following conditions apply:

• When transfer of care occurs immediately after surgery, the surgeon or other provider assuming in-hospital postoperative care must bill subsequent care procedure code 99231, 99232, or 99233.
• When the transfer of care occurs after hospital discharge, the surgeon or other provider who provides postdischarge care must bill the appropriate surgical code with modifier 55. Reimbursement will be limited to a percentage of the allowable fee for the surgical procedure.

Documentation in the medical record must include all of the following:

• A copy of the written transfer agreement.
• The dates the care was assumed and relinquished.
• The claim must indicate in the comments field of the claim form the dates on which care was assumed and relinquished, and the units field must reflect the total number of postoperative care days provided. Claims that are submitted on the CMS-1500 paper claim form must include the date of surgery in Block 14 and the dates on which care was assumed and relinquished in Block 19.

Staged or related surgical procedures or services that are performed during the postoperative period may be reimbursed when they are billed with modifier 58. A postoperative period will be assigned to the subsequent procedure. Documentation must indicate that the subsequent procedure or service was not the result of a complication and any of the following:

• It was planned at the time of the initial surgical procedure.
• It is more extensive than the initial surgical procedure.
• It is for therapy following an invasive diagnostic surgical procedure.

Note: Modifier 58 does not apply to procedure codes that are already defined as staged or sessioned services in the Current Procedural Terminology (CPT) Manual (e.g., 65855 or 66821).
Hospital visits by the surgeon during the same hospitalization as the surgery are considered to be related to the surgery and, as a result, not separately billable; however, separate payment for such visits can be allowed if any of the following conditions apply:

- Immunotherapy management is provided by the transplant surgeon. Immunosuppressant therapy following transplant surgery is covered separately from other postoperative services, so postoperative immunosuppressant therapy is not part of the global fee allowance for the transplant surgery. This coverage applies regardless of the setting.

- Critical care is provided by the surgeon for a burn or trauma patient.

- The hospital visit is for a diagnosis that is unrelated to the original surgery.

E/M services that are provided by the same provider for reasons that are unrelated to the operative surgical procedure may be considered for reimbursement if they are billed with modifier 24. The submitted documentation must substantiate the reasons for providing E/M services.

- Modifier 24 may be billed with modifier 25 if a significant, separately identifiable E/M service that was performed on the day of a procedure falls within the postoperative period of another unrelated procedure.

- Modifier 24 may be billed with modifier 57 if an E/M service that was performed within the postoperative period of another unrelated procedure results in the decision to perform major surgery.

Return Trips to the Operating Room

Return trips to the operating room for a repeat surgical procedure on the same part of the body may be considered for reimbursement when billed with modifiers 76 and 77. Billing with modifier 76 or 77 initiates the beginning of a new global period. Medical record documentation must support the need for a repeat procedure.

All surgical procedure codes with a predefined limitation (e.g., once per lifetime, one every 5 years) must not be submitted with modifier 76 or 77.

For modifiers 76 and 77, the repeated procedure must be the same as the initial surgical procedure. The repeat procedure must be billed with the appropriate modifier. The reason for the repeat surgical procedure should be entered in the narrative field on the claim form.

Return trips to the operating room for surgical procedures that are related to the initial surgery (i.e., complications) may be considered for reimbursement when they are billed with modifier 78 by the same provider.

- When a surgical procedure has a 0-day global period, the full value of the surgical procedure will be reimbursed; when the procedure has a 10- or 90-day global period only the intraoperative portion will be reimbursed.

- When an unlisted procedure is billed because no code exists to describe the treatment for the complications, reimbursement is a maximum of 50 percent of the value of the intraoperative services that were originally performed.

Reimbursement for the postoperative period of the first surgical procedure includes follow-up services from both surgical procedures, and no additional postoperative reimbursement is allotted. The global period will be based on the first surgical procedure.

Billing with modifier 78 does not begin a new global period.

Surgical procedures that are performed by the same provider during the postoperative period may be considered for reimbursement when they are billed with modifier 79 for any of the following:

- When the same procedure is performed with a different diagnosis.
• When the same procedure is performed on the left and right side of the body in different operative sessions and that procedure is billed with the RT or LT modifier.

• When a different procedure is performed with the same diagnosis.

• When a different procedure is performed with a different diagnosis.

Billing with modifier 79 initiates a new global surgical period.

9.2.69.7 Multiple Surgeries

Medicaid payment for multiple surgeries is based on the following guidelines:

• When two surgical procedures are performed on the same day at the same operative session, the primary procedure (such as the higher paying procedure) is paid at the full TMRM allowance. Secondary procedures performed on the same day are paid at half of the TMRM allowance when medically justified.

• Surgical procedures performed at different operative sessions on the same day are paid at the full TMRM allowance for each primary procedure at each session.

• Vaginal deliveries followed by tubal ligations are considered different operative sessions and are paid at full allowance for each primary procedure at a different session (i.e., both vaginal delivery and tubal ligation are paid at full allowance).

• Procedure code 58611 performed in conjunction with a Cesarean section is reimbursed at full allowance in cases where the allowance already represents half of the primary procedure.

• When a surgical procedure and a biopsy on the same organ or structure is done on the same day, the charges will be reviewed and reimbursement will be made only for the service with the higher of the allowed amounts.

9.2.69.8 Office Procedures

CMS has identified certain surgical procedures that are more appropriately performed in the office setting rather than as outpatient hospital, ASC/HASC procedures. The following list of surgical procedure codes should be billed in POS 1 (physician’s office). The medical necessity and/or special circumstances that dictate that these surgical procedures be performed in a POS other than the office must be documented on the claim. These surgical procedures are evaluated on a retrospective basis that may cause recoupment and/or adjustment of the original claim payment. This list is not all inclusive.

<table>
<thead>
<tr>
<th>Procedure Codes</th>
<th>Excision benign lesions</th>
<th>Excision malignant lesions</th>
<th>Manipulation (urethral)</th>
</tr>
</thead>
<tbody>
<tr>
<td>11400</td>
<td>11600</td>
<td>53600</td>
<td></td>
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<tr>
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<td>11601</td>
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<tr>
<td>11442</td>
<td>11641</td>
<td></td>
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</tbody>
</table>
9.2.69.9 Orthopedic Hardware

Reimbursement for the orthopedic hardware (e.g., buried wire, pin, screw, metal band, nail, rod, or plate) is part of the surgeon’s global fee or the facility’s payment group. The hardware is not reimbursed separately to either the surgeon or the facility.

The removal of orthopedic hardware is not payable to the same provider who inserted it, if removed within the global operative care period of the original insertion.

Services for removal of orthopedic hardware may be reimbursed separately after the global post operative care period.

9.2.69.10 Second Opinions

Texas Medicaid benefits include payment to physicians when eligible clients request second opinions about specific problems. The claim must be coded with the appropriate office or hospital visit codes, and the notation “Client Initiated Second Opinion” should be identified in Block 24D of the CMS-1500 paper claim form.

Refer to: Subsection 9.2.56.3.4, “Office and Outpatient Consultation Services” in this handbook.

9.2.69.11 Supplies, Trays, and Drugs

Payment to physicians for supplies is not allowed under Texas Medicaid. All supplies, including anesthetizing agents, inhalants, surgical trays, or dressings are included in the surgical payment on the day of surgery when the surgery is performed in the office or home setting.

Reimbursement for office visits includes overhead for supplies. If any of these items are submitted separately, they are denied as included in the surgical fee. If the supplies are submitted with a place of service (POS) other than the office, these supplies are denied as services that must be billed by the hospital, or as services that are included in nursing facility charges.

Silver nitrate applicators, used to treat granulated tissue around gastrostomy tubes and tracheostomies, are considered part of the office/hospital visit. Silver nitrate applicators are not a benefit for home use.

9.2.70 Telemedicine Services

Telemedicine services are a benefit of Texas Medicaid.

Refer to: The Telecommunication Services Handbook (Vol. 2, Provider Handbooks) for information about telemedicine services.

9.2.71 Therapeutic Apheresis

The following conditions must be met for therapeutic apheresis:

- To perform the medical services, including all nonphysician services, and to respond to medical emergencies at all times during client care, direct supervision by a physician is required.
Each client must be under the care of a physician.

Procedure codes 36511, 36512, 36513, 36514, and 36516 are limited to the following diagnosis codes:

<table>
<thead>
<tr>
<th>Diagnosis Codes</th>
</tr>
</thead>
<tbody>
<tr>
<td>C880 C882 C883 C888 C9000 C9002 C9010 C9011</td>
</tr>
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<td>C9012 C9020 C9021 C9022 C9030 C9031 C9032 C9100</td>
</tr>
<tr>
<td>C9101 C9102 C9110 C9111 C9112 C9130 C9131 C9132</td>
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<tr>
<td>C9140 C9141 C9142 C9150 C9151 C9152 C9160 C9161</td>
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<tr>
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<td>C9242 C9250 C9251 C9252 C9260 C9261 C9262 C9290</td>
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<tr>
<td>C9291 C9292 C92A0 C92A1 C92A2 C92Z0 C92Z1 C92Z2</td>
</tr>
<tr>
<td>C9300 C9301 C9302 C9310 C9311 C9312 C9330 C9331</td>
</tr>
<tr>
<td>C9332 C9391 C9392 C93Z0 C93Z1 C93Z2 C9400 C9401</td>
</tr>
<tr>
<td>C9402 C9420 C9421 C9422 C9430 C9431 C9432 C9440</td>
</tr>
<tr>
<td>C9441 C9442 C9480 C9481 C9482 C9500 C9501 C9502</td>
</tr>
<tr>
<td>C9510 C9511 C9512 C9590 C9591 C9592 D45 D472</td>
</tr>
<tr>
<td>D473 D474 D47Z2 D5700 D5701 D5702 D5703 D5709</td>
</tr>
<tr>
<td>D571 D5720 D57211 D57212 D57213 D57218 D57219 D57412</td>
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<tr>
<td>D57413 D57418 D5742 D57431 D57432 D57433 D57438 D57439</td>
</tr>
<tr>
<td>D5744 D57451 D57452 D57453 D57458 D57459 D5780 D57811</td>
</tr>
<tr>
<td>D57812 D57813 D57818 D57819 D588 D589 D590 D5910</td>
</tr>
<tr>
<td>D5911 D5912 D5913 D5919 D592 D593 D594 D599</td>
</tr>
<tr>
<td>D6182 D65 D682 D68311 D6851 D6852 D6859 D6861</td>
</tr>
<tr>
<td>D6862 D6869 D688 D690 D691 D692 D693 D6941</td>
</tr>
<tr>
<td>D6942 D6949 D696 D698 D699 D72828 D732 D740</td>
</tr>
<tr>
<td>D748 D749 D750 D751 D7589 D759 D761 D762</td>
</tr>
<tr>
<td>D763 D77 D890 D892 D8940 D8941 D8942 D8943</td>
</tr>
<tr>
<td>D8949 E0842 E0942 E1042 E1142 E7800 E7801 E7841</td>
</tr>
<tr>
<td>E7849 G603 G610 G6181 G6182 G6189 G620 G621</td>
</tr>
<tr>
<td>G622 G6281 G6282 G63 G64 G650 G7000 G7001</td>
</tr>
<tr>
<td>G731 I00 I010 I012 I018 I019 I773 I776</td>
</tr>
<tr>
<td>I7789 K716 K7200 K7201 K7581 K759 K760 K762</td>
</tr>
<tr>
<td>K767 K7689 K77 K8041 K8043 K8045 K8047 K8061</td>
</tr>
<tr>
<td>K8063 K8065 K8081 L100 L101 L102 L103 L104</td>
</tr>
<tr>
<td>L105 L1081 L1089 L109 L900 L940 L941 L943</td>
</tr>
<tr>
<td>M05011 M05021 M05022 M05023 M05031 M05032 M05041 M05042</td>
</tr>
<tr>
<td>M05051 M05052 M05061 M05062 M05071 M05072 M0509 M05411</td>
</tr>
<tr>
<td>M05412 M05421 M05422 M05431 M05432 M05441 M05442 M05451</td>
</tr>
<tr>
<td>M05452 M05461 M05462 M05471 M05472 M0549 M05611 M05612</td>
</tr>
<tr>
<td>M05621 M05622 M05631 M05632 M05641 M05642 M05651 M05652</td>
</tr>
</tbody>
</table>
Procedure code 36516 may be considered for reimbursement when billed for the low-density lipoprotein (LDL) apheresis (such as Liposorber LA 15) or the protein A immunoadsorption (such as Prosorba) columns.

The protein A immunoadsorption column is indicated for use in either of the following cases:

- Clients who have a platelet count of less than 100,000 mm3.
- Adult clients who have signs and symptoms of moderate to severe rheumatoid arthritis with long-standing disease who have failed, or are intolerant to, DMARDs.

The LDL apheresis column is indicated for use in clients who have severe familial hypercholesterolemia whose cholesterol levels remain elevated despite a strict diet and ineffective or untolerated maximum drug therapy. Coverage is considered for the following high-risk population, for whom diet has been ineffective and maximum drug therapy has either been ineffective or not tolerated:

- Functional hypercholesterolemia homozygotes with LDL-C > 500 mg/dL.
- Functional hypercholesterolemia heterozygotes with LDL-C > 300 mg/dL.
- Functional hypercholesterolemia heterozygotes with LDL-C > 200 mg/dL and documented coronary heart disease.

Baseline LDL-C levels are to be obtained after the client has had, at a minimum, a six-month trial on an American Heart Association (AHA) Step II diet or equivalent and maximum tolerated combination drug therapy designed to reduce LDL-C. Baseline lipid levels are to be obtained during a two- to four-week period and should be within 10 percent of each other, indicating a stable condition.

Therapeutic apheresis using the LDL apheresis column may be reimbursed for diagnosis code E780.

Apheresis is denied for all other diagnosis codes. Other diagnosis codes can be reviewed by the TMHP Medical Director or designee on appeal with documentation of medical necessity.
Laboratory work before and during the apheresis procedure is covered when apheresis is performed in the outpatient setting (POS 5). Laboratory work billed in conjunction with apheresis performed in the inpatient setting (POS 3) is included in the DRG reimbursement and is not paid separately.

9.2.72 Therapeutic Phlebotomy

Therapeutic phlebotomy is a treatment whereby a prescribed amount of blood is withdrawn for medical reasons. Conditions that cause an elevation of the red blood cell volume or disorders that cause the body to accumulate too much iron may be treated by therapeutic phlebotomy.

Therapeutic phlebotomy is a benefit of Texas Medicaid and may be billed using procedure code 99195. This procedure code should be used only for the therapeutic form of phlebotomy and not for diagnostic reasons.

Reimbursement of therapeutic phlebotomy is limited to the following diagnosis codes:

<table>
<thead>
<tr>
<th>Diagnosis Codes</th>
</tr>
</thead>
<tbody>
<tr>
<td>D45</td>
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<tr>
<td>E8029</td>
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</table>

Therapeutic phlebotomy will be automatically denied for all other diagnosis codes.

9.2.73 Therapeutic Radiopharmaceuticals

Therapeutic radiopharmaceuticals, when used for therapeutic treatment, are a benefit of Texas Medicaid.

The following procedure codes may be submitted for therapeutic radiopharmaceuticals:

<table>
<thead>
<tr>
<th>Procedure Codes</th>
</tr>
</thead>
<tbody>
<tr>
<td>79403</td>
</tr>
</tbody>
</table>

9.2.73.1 Prior Authorization for Therapeutic Radiopharmaceuticals

Prior authorization is required for ibritumomab tiuxetan procedure codes A9542 and A9543.

Only one ibritumomab tiuxetan (procedure codes A9542 and A9543) may be prior authorized and reimbursed once per lifetime, any provider with one of the following diagnosis codes:

<table>
<thead>
<tr>
<th>Diagnosis Codes</th>
</tr>
</thead>
<tbody>
<tr>
<td>C8259</td>
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<tr>
<td>C8599</td>
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</tbody>
</table>

Ibritumomab tiuxetan may be prior authorized when all of the following criteria are met:

- Client has a diagnosis of either a low-grade follicular or transformed B-cell non-Hodgkin’s lymphoma.
- Client has failed, relapsed, or become refractory to conventional chemotherapy and the following is documented:
  - Marrow involvement is less than 26 percent.
  - Platelet count is 100,000 cell/mm3 or greater.
  - Neutrophil count is 1,500 cell/mm3 or greater.
- Client has failed a trial of rituximab.

Prior authorization must be submitted through Special Medical Prior Authorization department.
Prior authorization is required for lutetium lu 177 dotatate (Lutathera) procedure code A9513. Lutetium lu 177 dotatate (Lutathera) procedure code A9513 will be considered with documentation that meets all of the following criteria:

- The client has a diagnosis of somatostatin receptor-positive gastroenteropancreatic neuroendocrine tumors (GEP-NETs) (one of the following diagnosis codes must be submitted on the prior authorization request: C7A00, C7A010, C7A011, C7A012, C7A019, C7A020, C7A021, C7A022, C7A023, C7A024, C7A025, C7A026, C7A029, C7A092, C7A094, C7A095, C7A096).
- The client is 18 years of age or older.
- The client is not pregnant or breastfeeding.
- The official pathology report documents a GEP-NET with Ki67 index less than 20 percent.
- The disease is metastatic, or locally advanced and unresectable as indicated by one of the following:
  - Positive somatostatin receptor scintigraphy with correlative magnetic resonance imaging (MRI)
  - Computed tomography (CT) imaging of metastatic measurable disease
  - 68-Ga-Dotate positron emission tomography (PET) scan positive for metastatic disease
- The client experienced disease progression while on a long-acting somatostatin analog (e.g. octreotide, lanreotide).
- The client has not had prior treatment with Peptide Receptor Radionuclide Therapy (PRRT), and has not had prior external radiation therapy to more than 25 percent of the bone marrow.
- The documentation includes an oncologist’s or nuclear medicine specialist’s complete written order and prescription for Lutetium lu 177 dotatate intravenous infusion.
- A treatment plan that includes all of the following documentation:
  - Lutetium lu 177 dotatate 7.4 GBq (200 mCi) every 60 days for a total of 4 doses that is administered in a facility under the control of a physician who is licensed and authorized to administer radiopharmaceuticals
  - The recommended use of premedication and concomitant medications of somatostatin analogs, antiemetics , and specialized amino acid solution
  - The restrictions and usage of long- and short-acting octreotide agents before, during, and after lutetium lu 177 dotatate intravenous infusions
  - Details of withholding the treatments for contraindicated circumstances including, but not limited to:
    - Thrombocytopenia
    - Anemia neutropenia
    - Renal toxicity
    - Hepatotoxicity
    - Other non-hematologic toxicities

Prior authorization requests must be submitted to the TMHP Prior Authorization Department by mail, fax, or the electronic portal. Prescribing or ordering providers, dispensing providers, clients’ responsible adults, and clients may sign prior authorization forms and supporting documentation using electronic or wet signatures.
Refer to: Subsection 5.5.1.2, “Document Requirements and Retention” in “Section 5: Fee-for-Service Prior Authorizations” (Vol. 1, General Information) for additional information about electronic signatures.

An SMPA Request Form must be completed, signed, and dated by the prescribing provider. The SMPA form will not be accepted after 90 days from the date of the prescribing provider’s signature.

The completed SMPA Request Form must be maintained by the prescribing provider in the client’s medical record and is subject to retrospective review.

Section C of the SMPA Request Form under Statement of Medical Necessity must contain the following:

- Documentation of the client’s dosage
- The administration schedule
- The number of injections to be administered during the prior authorization period
- The requested units/millicuries per injection
- The dosage calculation

To facilitate the determination of medical necessity and avoid unnecessary denials, the prescribing provider must submit correct and complete information, including documentation of medical necessity for the equipment or supplies requested, the procedure codes, and the numerical quantities for services requested. The provider must maintain documentation of medical necessity in the client’s medical record.

Prior authorization must be requested through the Special Medical Prior Authorization (SMPA) department with appropriate documentation.

Requests can be mailed or faxed to:

Texas Medicaid & Healthcare Partnership
Special Medical Prior Authorization
12357-B Riata Trace parkway
Austin, TX 78727
Fax: 1-512-514-4213

Requests for prior authorization can also be submitted online through the TMHP website at www.tmhp.com.

9.2.73.1.1 Reimbursement Limitations for Ibritumomab tiuxetan and Lutetium lu 177 dotate

Ibritumomab tiuxetan is indicated for the treatment of clients that have failed rituximab and have CD20 antigen-expressing relapsed or refractory, low grade, follicular, or transformed non-Hodgkin’s lymphoma or refractory non-Hodgkin’s lymphoma.

Ibritumomab tiuxetan may only be considered once per lifetime, any provider, and only one of the agents.

Lutetium Lu 177 dotate (Lutathera) intravenous injection (procedure code A9513) is indicated for the treatment of adult clients who are 18 years of age or older with a diagnosis of somastatin receptor-positive gastroenteropancreatic neuroendocrine tumors (GEP-NETs), including foregut, midgut, and hindgut neuroendocrine tumors. For all other indications, Lutetium Lu 177 dotate (Lutathera) injection for intravenous use is not proven to be medically effective and is considered experimental.

Lutetium Lu 177 dotate (Lutathera) must be administered under the control of an oncologist or a nuclear medicine specialist who is licensed and authorized to administer radiopharmaceuticals and must be administered in an outpatient setting.

Lutetium Lu 177 dotate (Lutathera) procedure code A9513 is limited to one service every 60 days for a total of four services per lifetime, any provider.
9.2.73.2 Other Limitations on Therapeutic Radiopharmaceuticals

Strontium-89 chloride (procedure code A9600) may be reimbursed when submitted with diagnosis code C7951 or C7952.

Strontium-89 chloride is limited to a total of 10 mci intravenously injected every 90 days, any provider.

Sodium phosphate P-32, therapeutic (procedure code A9563) may be reimbursed when submitted with the following diagnosis codes:

<table>
<thead>
<tr>
<th>Diagnosis Codes</th>
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</thead>
<tbody>
<tr>
<td>C7951</td>
</tr>
<tr>
<td>C9512</td>
</tr>
</tbody>
</table>

Chromic phosphate P-32 suspension (procedure code A9564) may be reimbursed when submitted with diagnosis codes C782 and C786.

An appropriate modifier may be used when billing for services more than once per day, same provider.

Iodine i-131 iobenguane procedure code A9590 is a benefit for clients who are 12 years of age and older.

Iodine i-131 iobenguane is a radiopharmaceutical indicated for the treatment of adult and pediatric clients who are 12 years of age and older with iobenguane scan positive, unresectable, locally advanced or metastatic pheochromocytoma or paraganglioma who require systemic anticancer therapy. Iodine i-131 iobenguane should be handled with appropriate safety measures to minimize radiation exposure and should be administered by or under the control of physicians who are licensed and authorized to administer radiopharmaceuticals.

Procedure code A9590 is limited to the following diagnosis codes:

<table>
<thead>
<tr>
<th>Diagnosis Codes</th>
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<tr>
<td>C7410</td>
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</table>

9.2.74 Urethral Dilation

If urethral dilation (procedure code 53600, 53601, 53605, 53620, 53621, 53660, 53661, or 53665) is billed on the same date of service by the same provider as procedure code 52000, the charges will be combined and processed as procedure code 52281.

Urethral dilation will be denied when billed on the same date of service by the same provider as any other cystoscopy.

9.2.75 Ventilation Assist and Management for the Inpatient

Use the following procedure codes and guidelines for reimbursement of ventilation assist and management: 94002 and 94003. Procedure codes 94002 and 94003 may be reimbursed only when the client is in observation or inpatient status. Respiratory care billed in any other POS will be denied.

Use the ventilation assist and management subsequent code (procedure code 94003) when respiratory support must be established for a patient in the postoperative period in the hospital (POS 3). Subsequent days of ventilation assistance are payable when documentation indicates a respiratory problem.

When the use of a ventilator is required as part of a major surgery, initial ventilation assist and management will be denied. It should be billed as ventilation assist and management subsequent procedure code 94003.
Procedure codes 94002 and 94003 apply only to hospital care for critically ill patients. They do not apply to routine recovery room ventilation services. Separate support service charges billed on the same day as ventilatory support are denied (for example, arterial or venous punctures; interpretations of arterial blood gases; or pulmonary function tests and management of the hemodynamic functions of the patient).

Use ventilation assist and management and initiation of pressure or volume preset ventilators for assisted or controlled breathing–first day (procedure code 94002) when respiratory support must be established for a patient. It is a one-time charge per hospitalization that may be paid when the claim documents that a respiratory problem exists (for example, respiratory distress, asphyxia). After the first day, use subsequent days (procedure code 94003).

### 9.2.76 Wearable Cardiac Defibrillator (WCD)

A WCD (procedure codes 93292, 93745, and K0606) are a benefit of Texas Medicaid.

The rental of a WCD (procedure code K0606) is limited to once per month and must be submitted with modifier RR.

Modifier 25 may be used to identify a significant separately identifiable evaluation and management service performed (for example, different diagnosis) on the same day as the initial set up of a WCD by the same provider for the same client. Documentation that supports the provision of a significant, separately identifiable E/M service must be maintained in the client’s medical record and made available to Texas Medicaid upon request.

Procedure code 93292 will be denied as part of procedure code 93745 when submitted on the same date of service by any provider.

Procedure codes 93000, 93005, 93010, 93040, 93041, and 93042 will be denied as part of procedure code 93745 when submitted on the same date of service by any provider.

### 9.2.76.1 Prior Authorization for WCD

Prior authorization is required for the rental of WCD (procedure code K0606).

The WCD may be prior authorized for clients at high-risk of sudden cardiac arrest who meets one of the following criteria:

- Has completed electrophysiologic studies to determine the type of arrhythmia present and confirm that a wearable cardiac defibrillator is the best course of treatment.
- Is contraindicated for an implantable cardiac defibrillator (ICD) at the current time, such as with a systemic infection.
- Is waiting for ICD implantation.
- Is waiting for ICD implantation and is undergoing treatment for a systemic infection.
- Has had an ICD explantation due to pocket infection.
- Is waiting for heart transplantation.
- Has self-limiting arrhythmias from iatrogenic (drug loading with potentially pro-arrhythmic medications) or other causes.
- Has a familial or inherited condition with a high risk of life-threatening ventricular tachyarhythmias, such as long QT syndrome or hypertrophic cardiomyopathy.
- Has had either documented prior myocardial infarction or dilated cardiomyopathy and a measured left ventricular ejection fraction (LVEF) less than or equal to 35 percent.
- Has received a documented diagnosis of any one of the following conditions:
• Clinically inducible hemodynamically significant ventricular tachycardia (HSVT) or ventricular fibrillation (VF), where drug treatment has been ineffective, or the side effects of the medication used to treat the arrhythmia are intolerable.

• Inducible VT or VF despite endocardial ablation or surgical excision when drug therapy has failed.

• VF or syncopal ventricular tachycardia.

• Specific ST-T wave changes, borderline CPK-MB isoenzymes, and dangerous ventricular arrhythmias are exhibited in a postmyocardial infarction patient.

• VT caused by ischemic heart disease not associated with an acute myocardial infarction, and where drug therapy or surgical therapy has failed.

• Recurrent syncope of undetermined etiology in a patient with HSVT or VF induced by EPS in whom no effective or tolerated drug is available or appropriate. Symptoms must be linked to HSVT or VF.

• Recurrent syncope of undetermined etiology with positive EPS studies where ventricular arrhythmia is documented as the cause.

• Palliative treatment for VT or VF in clients awaiting heart transplant.

The WCD is contraindicated in clients with an active ICD and should not be used in clients who meet the following criteria:

• Have a vision or hearing problem that may interfere with the perception of alarms or messages from the WCD.

• Is taking medications that would interfere with responding to the alarms or message from the WCD by depressing buttons.

• Is unwilling or unable to wear the device continuously, except when bathing or showering.

• Is pregnant or breastfeeding.

• Is of childbearing age and is not attempting to prevent pregnancy.

The WCD is considered investigational and not medically necessary for all other indications, including but not limited to, the following:

• Clients with drug-refractory class IV congestive heart failure who is not candidates for heart transplantation.

• Clients who have a history of psychiatric disorders that interfere with the necessary care and follow-up.

• Clients in whom a reversible triggering factor for VT/VF can be definitely identified, such as ventricular tachyarrhythmias in evolving acute myocardial infarction or electrolyte abnormalities.

• Clients with terminal illnesses.

A completed Home Health Services (Title XIX) DME/Medical Supplies Physician Order Form (Title XIX Form) prescribing the DME and/or medical supplies must be signed and dated by the ordering physician familiar with the client prior to requesting authorization.

• The completed Title XIX Form must be maintained by the DME provider.

• The ordering physician must maintain the completed, originally signed and dated Title XIX Form in the client’s medical record.

• The completed Title XIX Form must include the procedure codes and quantities requested for the services.
To complete the prior authorization process the provider must submit the completed Title XIX Form by fax to the Home Health Unit at 1-512-514-4209 or in writing to the following address:

Texas Medicaid & Healthcare Partnership  
Home Health Services  
PO Box 202977  
Austin, TX 78720-2977

When a WCD is not covered as a home health service, it may be considered for reimbursement through the CCP for clients who are 20 years of age and younger. All of the following criteria must be met for CCP reimbursement for a WCD:

- The client is eligible for CCP benefits.
- The documentation submitted with the request supports the determination of medical necessity based on the criteria listed in the policy.
- Federal financial participation is available.
- The client’s cardiac status would be compromised without the requested equipment.
- The requested equipment is safe in the home setting.

**Note:** For clients who are 21 years of age or older, requests for a WCD that does not meet the criteria through Title XIX Home Health Services may be considered under the Texas Medicaid Home Health—Durable Medical Equipment (DME) Exceptional Circumstances process.

Rental of an automatic external defibrillator, with integrated electrocardiogram analysis, garment type (procedure code K0606) may be prior authorized (initially for up to three months) with documentation supporting the medical necessity and appropriateness of the device.

The provider may be reimbursed only for the length of time the device is used even though the authorization for the rental may be for a longer period of time.

The rental of the device includes the monitor, electrode belt (four sensors or electrodes and three treatment pads), garment, two rechargeable batteries, a battery charger and modem.

The purchase of a replacement battery (procedure code K0607), the purchase of a garment (procedure code K0608), and electrodes (procedure code K0609) will be considered part of the rental.

Prior authorization extensions for WCDs beyond the initial three-month rental may be considered by the medical director when documentation supports continued medical necessity for the device.

Providers must submit new documentation to support continued medical necessity for an extension of the rental to be considered.

To avoid unnecessary denials, the physician must provide correct and complete information, including documentation for medical necessity of the device. The physician must maintain documentation of medical necessity in the client’s medical record. The requesting provider may be asked for additional information to clarify or complete a request for the WCD.

Retrospective review may be performed to ensure documentation supports the medical necessity of the service when billing the claim.

**9.2.77 Wound Care Management**

Wound care management includes the care of acute and chronic wounds, which include, but are not limited to, open ulcers (venous pressure or diabetic ulcers), fistulas, or erosion of skin related to cancer. Acute and chronic wounds are defined as the following:

- Acute wounds: Wounds taking less than 30 days for complete healing
- Chronic wounds: Wounds taking more than 30 days for complete healing
Wound care includes the following:

- Optimization of nutritional status
- Debridement by any means to remove devitalized tissue
- Maintenance of a clean, moist bed of granulation tissue
- Necessary treatment to resolve any infection that may be present

For clients with an ulcer, wound care may include the following:

- Frequent repositioning of a client who has a pressure ulcer
- Off-loading pressure and good glucose control for a client who has a diabetic ulcer
- Establishment of adequate circulation for a client who has an arterial ulcer
- Use of a compression system for clients who have a venous ulcer

Wound care management includes first- and second-line therapies. First-line wound care is used for acute wounds. If the wound does not improve with first-line treatment, adjunctive second-line therapy may be used. Measurable signs of improved healing include the following:

- A decrease in wound size, either in surface area or volume
- A decrease in amount of exudate
- A decrease in amount of necrotic tissue

Wound care must be performed by a licensed health professional who is qualified to safely and effectively provide the medically necessary care. Providers are expected to exercise their clinical judgment to render the most appropriate care in accordance with their scope of practice as designated by their regulatory and governing boards.

The following services are not a benefit of Texas Medicaid:

- Infrared therapy
- Ultraviolet therapy
- Topical hyperbaric oxygen therapy
- Low-energy ultrasound wound cleanser (MIST therapy)
- Services that are submitted as debridement but do not include the removal of devitalized tissue. Examples include removal of non-tissue integrated fibrin exudates, crusts, biofilms, or other materials from a wound, without the removal of tissue.
- Electrical stimulation and electromagnetic therapy

### 9.2.77.1 First-Line Wound Care Therapy

First-line wound care therapy includes the following:

- Cleansing, antibiotics, and pressure off-loading
- Compression
- Debridement
- Dressing
- Whirlpool for burns

#### 9.2.77.1.1 Cleansing, Antibiotics, and Pressure Off-loading

Wound cleansing helps to create an optimal healing environment and decreases the potential for infection by loosening and removing cellular debris and residual topical agents from previous dressings.
Wound cleansing agents may include normal saline, commercial wound cleansers, providone iodine, hydrogen peroxide, or sodium hydrochlorite. Cleansing solutions and methods vary based on effectiveness and individual client needs.

Systemic or topical antibiotics may be used to prevent or treat wound infections and to aid in the healing of wounds.

Pressure off-loading devices, such as pillows, boots, mattresses, and protectors, may also be used as part of first-line wound care therapy to prevent or relieve pressure on the wound.

Procedure code 29445 may be reimbursed for pressure off-loading performed as part of wound care management.

9.2.77.1.2  **Compression**

Compression performed as a part of wound care management is a benefit and may be reimbursed when billed with procedure code 29580 or 29581.

9.2.77.1.3  **Debridement**

Wound debridement includes the pre-debridement wound assessment, the debridement, and the post-procedure instructions provided to the client on the date of service.

Selective debridement consists of the following:
- Conservative sharp debridement
- High-pressure lavage to selected areas

Non-selective debridement consists of the following:
- Autolytic debridement
- Blunt debridement
- Enzymatic debridement
- Hydrotherapy and wound immersion ross
- Mechanical debridement

The following procedure codes may be reimbursed for wound debridement:

<table>
<thead>
<tr>
<th>Procedure Codes</th>
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<tbody>
<tr>
<td>11000</td>
</tr>
<tr>
<td>97602</td>
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</tbody>
</table>

Professional services for selective wound debridement (procedure codes 97597 and 97598) may be reimbursed to a licensed physical therapist, when it is determined to be within the provider’s scope of practice, and the service is prescribed by a Medicaid-enrolled supervising physician or qualified non-physician provider.

The following procedure codes may be reimbursed for debridement of partial-thickness burns:

<table>
<thead>
<tr>
<th>Procedure Codes</th>
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</thead>
<tbody>
<tr>
<td>16020</td>
</tr>
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</table>

Prior authorization is not required for debridement of partial-thickness burns (procedure codes 16020, 16025, or 16030).

Prior authorization is required for wound debridement procedure codes 11042, 11043, and 11044. A request for prior authorization must be submitted to TMHP with the Special Medical Prior Authorization (SMPA) Request Form before the procedure is performed. Providers must retain a copy of the
signed and dated form in the client’s medical record at the provider’s place of business. The requesting provider may be asked for additional information to clarify or complete a request for the equipment/supply requested.

Requests for prior authorization for wound debridement procedure codes 11042, 11043, and 11044 must include the following documentation:

- Location of the wound
- Characteristics of the wound, which include all of the following:
  - Dimensions (diameter and depth)
  - Drainage (amount and type)
  - Related signs and symptoms (swelling, pain, inflammation)
  - Presence of necrotic tissue/slough

When submitting an initial prior authorization request, the treating provider (registered nurse, physician, physical therapist) must submit a signed and dated wound care treatment plan or letter of medical necessity that includes all the following documentation:

- Planned interventions for the problem identified
- Treatment goals
- Expected outcomes

The treatment plan or letter of medical necessity is considered current when it is signed and dated within 30 calendar days prior to or on the date the procedure is performed. Otherwise, a new treatment plan must be submitted.

Retroactive authorization requests for wound debridement performed on an urgent or emergent basis (procedure code 11042, 11043, or 11044) will be denied if not submitted within 14 calendar days, beginning the day after the procedure is performed.

For retroactive authorization to be considered, the treatment plan or letter of medical necessity must be signed and dated within 14 calendar days beginning the day after the procedure is performed. If the treatment plan is not signed and dated within the 14-calendar-day period, the request will be denied.

For procedure codes 11043 and 11044, at least one of the following conditions must be present and documented:

- Stage III or IV wounds
- Venous or arterial insufficiency ulcers
- Dehisced wounds or wounds with exposed hardware or bone
- Neuropathic ulcers
- Complications of surgically created or traumatic wound where accelerated granulation therapy is necessary but cannot be achieved by other available topical wound treatment

Wound debridement procedure codes 11042, 11043, and 11044 are not appropriate and will not be approved for the following:

- Washing bacteria or fungal debris from the feet
- Paring or cutting of corns or calluses
- Incision and drainage of an abscess
- Trimming or debridement of nails, or avulsion of nail plates
• Acne surgery
• Destruction of warts
• Burn debridement

Prior authorization requests must be submitted by the provider within 30 calendar days prior to, or on the date the procedure is performed. If the prior authorization request is not submitted within 30 calendar days prior to, or on the date the procedure is performed, the request will be denied.

The physician's signature on the Special Medical Prior Authorization (SMPA) Request Form is considered current when signed and dated within 30 calendar days prior to, or on the date the procedure is performed. If the physician's signature is not signed and dated within the 30-calendar-day period prior to or on the date the procedure is performed, the request will be denied.

Prior authorization requests for procedure codes 11042, 11043, and 11044 will be considered for 7 calendar days, beginning on the requested procedure date.

Retroactive authorization is required for wound debridement procedure codes 11042, 11043, and 11044 that are performed on an urgent or emergent basis. The provider must submit a request for retroactive authorization within 14 calendar days, beginning the day after the procedure is performed.

For retroactive authorization requests, the physician's signature on the Special Medical Prior Authorization (SMPA) Request Form is considered current when signed and dated within 14 calendar days, beginning the day after the procedure is performed. Requests with the physician's signature not signed and dated within the 14-calendar-day period will be denied.

Prior authorization requests for subsequent debridement will be considered on a case-by-case basis with documentation of medical necessity. These requests will be reviewed by the Medical Director.

The wound debridement procedure code submitted on the prior authorization or retroactive authorization request must reflect the level of debrided tissue, e.g., partial-thickness skin, full-thickness skin, subcutaneous tissue, muscle, and/or bone, and not the extent, depth, or grade of the ulcer or wound.

9.2.77.1.4 Dressings and Metabolically Active Skin Equivalents

Wound dressings may include wet and dry dressings.

All dressings applied to the wound during a wound debridement procedure are considered part of the service for wound debridement.

9.2.77.1.5 Whirlpool for Burns

Whirlpool may be a benefit when used as first-line wound care therapy for the treatment of burn wounds.

9.2.77.2 Second-Line Wound Care Therapy

Second-line wound care therapy is limited to chronic Stage III or IV wounds and may be covered only after first-line therapy has been tried for at least 30 days without measurable signs of improved healing. First-line wound care therapy may continue as appropriate, with the addition of second line wound care measures as indicated by the client's medical condition.

Second-line wound care therapy includes the following:

• Whirlpool
• Irrigation, including pulsatile jet irrigation
• Application of metabolically active skin equivalents/skin substitutes
9.2.77.2.1 Pulsatile-Jet Irrigation

Pulsatile-jet irrigation is a benefit for the treatment of Stage III or IV wounds when other forms of treatment have failed. Removal of devitalized tissue using pulsatile-jet irrigation may be reimbursed when claims are submitted for procedure code 97597 or 97598.

9.2.77.2.2 Skin Substitutes and Surgical Wound Preparation

The application of skin substitutes is a benefit for the treatment of chronic Stage 3 or 4 wounds that have failed to respond to standard wound care treatment after 30 days. A failed response is defined as a wound that has increased in size or depth, or has not changed in baseline size or depth, and shows no measurable signs of healing improvements after 30 days of appropriate wound-care measures.

Use of the appropriate specific skin substitute product(s) for the episode of each documented wound is expected. Compliance with the Food and Drug Administration (FDA) assessments and submitted guidelines for the specific skin substitute product(s) used is expected. Skin substitute products not used within the scope of the FDA's intended use and indications are considered experimental and/or investigational. All wound care services require documentation of the wound, and a comprehensive treatment plan is required to be maintained in the client's medical record.

The following procedure codes may be reimbursed for the application of skin substitute grafts:

<table>
<thead>
<tr>
<th>Procedure Codes</th>
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<tr>
<td>15271</td>
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Approved skin substitute products used in wound care services that are provided in an office-based setting will be considered for separate reimbursement when submitted with an appropriate application procedure code from the table above.

The approved skin substitute product(s) must have a published average sales price, must be FDA cleared/approved or be designated as 361 HCT/P exempt, and should be used in accordance with each product’s individualized labeling and application guidelines. The approved list of skin substitute products are reviewed and updated biannually. Providers should refer to the Center for Medicare & Medicaid Services (CMS) Medicare Part B Drug Average Sales Price web page at www.cms.gov for updates to the list of approved skin substitute products.

All skin substitute products used in wound care services that are provided in a facility setting are considered part of the application services and are not separately reimbursed.

Procedure code C9250 is considered part of the application services and will not be separately reimbursed, regardless of the setting.

Surgical Wound Preparation

Appropriate surgical wound preparation may be expected at least once at the initiation of care, prior to placement of the skin substitute graft. Repeated use of surgical preparation services in conjunction with skin substitute application codes will be considered not reasonable or necessary and will not be reimbursed.

Procedure codes 15002, 15003, 15004, 15005, 15040, and 15050 may be reimbursed for surgical wound preparation.

*Note:* Procedure code 15005 is not a benefit for ambulatory surgical center providers.

Limitations

The treatment of any chronic skin wound will typically last no more than 12 weeks.
Skin substitute applications and grafts are limited to 10 per episode of care in a 12-week period, per rolling year beginning on the first day of the first skin substitute application. If more than one specific product is used or a product change occurs during the 12-week period of care, the expectation remains that the cumulative number of applications will not exceed 10.

More than 10 skin substitute applications in a 12-week period will be considered on a case-by-case basis with documentation of medical necessity. These requests will be reviewed by the Medical Director.

**Re-Treatment of Healed, Stage 3, or Stage 4 Chronic Wounds**

Retreating healed skin wounds showing greater than 75 percent in size reduction and smaller than 0.5 square cm is not considered medically reasonable or necessary and will not be reimbursed.

Retreating a venous stasis ulcer or diabetic neuropathic foot ulcer with any skin substitute product(s) within one year of previous treatment is considered treatment failure. This unsuccessful treatment does not meet reasonable and necessary criteria for re-treatment and will not be reimbursed.

Unsuccessful treatment is defined as an increase in size or depth of an ulcer, or no change in baseline size or depth and no sign of improvement or indication that improvement is likely (such as granulation, epithelialization, or progress towards closing) for a period of 4 weeks past the start of therapy.

**Contraindications**

Skin substitute grafts are contraindicated for the following:

- Clients with known hypersensitivity to any component of the specific skin substitute graft (e.g., allergy to avian, bovine, porcine, or equine products).
- Skin substitute grafts will not be considered reasonable and necessary for clients with inadequate control of underlying conditions or exacerbating factors, such as the following:
  - Clients with uncontrolled diabetes
  - Clients with active infection
  - Clients with active Charcot arthropathy of the ulcer extremity
  - Clients with vasculitis
  - Clients who continue smoking tobacco and have not received smoking cessation guidance from their physician

**9.2.77.3 Documentation Requirements**

For all wound care management services, documentation that supports the medical necessity of the service must be maintained in the client’s medical records, which includes but is not limited to the following information:

- Accurate diagnostic information that pertains to the underlying diagnosis and condition as well as any other medical diagnoses and conditions, which include the client’s overall health status.
- Appropriate medical history related to the current wound, including the following:
- Wound location
  - Wound measurements, which includes length, width, and depth, any tunneling and/or undermining
  - Wound color, drainage (type and amount), and odor, if present
  - The prescribed wound care regimen, which includes frequency, duration, and supplies needed
  - Treatment for infection, if present
  - All previous wound care therapy regimens, if appropriate
• The client’s use of a pressure reducing support surface, mattress, and/or cushion, when appropriate

Documentation maintained in the client’s medical record must support the level of debridement service provided.

Fewer than five surgical debridements that involve removal of muscle or bone are typically required for management of most wounds. Documentation that is maintained in the client’s medical record must support the number of debride ments involving muscle or bone that are performed.

9.2.77.3.1 Skin Substitutes

Documentation maintained in the client’s medical record must support the need for skin substitute applications and the product used.

Documentation for all wound care treatments involving the application of skin substitute products must include, but is not limited to, the following:

• Wound treatments are accompanied by the appropriate adjunctive measures, and identify the specific adjunctive therapies being provided to the client as part of the wound treatment regimen.

• Clients who use tobacco must satisfy one of the following documentation requirements:
  • The client will have ceased smoking or have refrained from systemic tobacco intake for at least 4 weeks prior to beginning skin substitute applications and during the conservative wound care.
  • Smoking history, cessation counseling on the effects of smoking on surgical outcomes, treatment for smoking cessation (if applicable), and the outcome of counseling must be recorded in the client's medical record.

• Adequate circulation/oxygenation to support tissue growth/wound healing must be present as evidenced by physical examination (e.g., Ankle-Brachial Index [ABI] of no less than 0.60, toe pressure greater than 30 millimeters of mercury [mmHg]).

• The wound has a skin deficit at least 1.0 square centimeter in size.

• For diabetic foot ulcers, the client’s medical record reflects a diagnosis of Type 1 or Type 2 diabetes.

• Partial or full thickness ulcers must have a clean granular base without tendon and or muscle involvement, bone exposure, or sinus tracts.

• Documentation of the wound’s response to the treatment is required at least every 30 days for each treatment episode. The documentation requirements must include measurements of the initial wound, measurements at the completion of appropriate wound care every 30 days, and measurements immediately prior to placement and with each subsequent placement of the skin substitute.

9.2.77.4 Exclusions

The following services are not a benefit of Texas Medicaid:

• Separately billed, repeated use of a skin substitute product after 12 weeks for a single wound or episode

• Skin substitute grafting for partial thickness loss with the retention of epithelial appendages is not covered, as epithelium will repopulate the deficit from the appendages, negating the benefit of over grafting

9.3 Doctor of Dentistry Practicing as a Limited Physician

This section outlines the guidelines for the Doctor of Dentistry practicing as a limited physician. The THSteps dental program is not addressed in these guidelines.
Services by a dentist (DDS or DMD) are covered by Texas Medicaid in accordance with the Omnibus Budget Reconciliation Act (OBRA) of 1987 §4103 and Title 2 Texas Human Resources Code §32.054, if the services are furnished within the dentist’s scope of practice as defined by Texas state law and would be covered under Texas Medicaid when provided by a licensed physician (MD or DO).

Dentist (DDS or DMD) who want to participate as a dentist-physician in Texas Medicaid must be separately enrolled as a Doctor of Dentistry practicing as a limited physician even if they are enrolled in the THSteps Dental Program.

Dual licensure (MD, DO, and DDS) is not required for a dentist to enroll as a limited physician. Medicare enrollment is required for a dentist to enroll as a limited physician.

9.3.1 Prior Authorization for General Dental Services Due to Life-Threatening Medical Condition

Reimbursement for general dental services by any provider, irrespective of the medical or dental qualifications of the provider, is not a Medicaid benefit for Medicaid clients who are 21 years of age and older (who do not reside in an ICF-IID facility).

The TMHP Medical Director or designee may allow an exception for a dental condition causally related to a life-threatening medical condition. Mandatory prior authorization is required and the dental diagnoses must be secondary to a life-threatening medical condition.

Examples of dental procedures that may be authorized for a general dentist who is enrolled as a limited physician are:

- Extractions.
- Alveolectomies (in limited situations).
- Incision and drainage.
- Curettement.

Examples of dental procedures that may be authorized for an oral and maxillofacial surgeon who is enrolled as a limited physician are:

- Extractions.
- Alveolectomies (in limited situations).
- Incision and drainage.
- Curettement maxillofacial surgeries to correct defects caused by accident or trauma.
- Surgical corrections of craniofacial dysostosis.

Note: Therapeutic procedures such as restorations, dentures, and bridges are not a benefit of the program and will not be authorized.

9.3.1.1 Guidelines for Requesting Mandatory Prior Authorization

The limited physician dentist must request the mandatory prior authorization, and the request must include:

- A treatment plan that clearly outlines the dental condition as related to the life-threatening medical condition.
- Narrative describing the current medical problem, client status, and medical need for requested services.
- The client name and Medicaid number.
- The limited physician dentist’s provider identifier.
• The name and address of the facility.
• CPT procedure codes.
• The history and physical.
• The limited physician dentist’s signature.

Note: The "limited physician" dentist who will perform the procedure(s) must submit the request for prior authorization.

All supporting documentation must be included with the request for authorization. Providers are to send requests and documentation to the following address:

Texas Medicaid & Healthcare Partnership
Special Medical Prior Authorization
12357-B Riata Trace Parkway, Suite 100
Austin, TX 78727
Fax: 1-512-514-4213

9.3.2 Benefits and Limitations
Dental procedure codes and their corresponding CPT procedures may not be billed on the same date of service by any provider.

Cosmetic procedures are not a benefit of Texas Medicaid. Certain procedure codes, including, but not limited to, the procedure codes in the following table, may be considered cosmetic and are not a benefit except when the procedure is performed as a result of trauma or injury for the purpose of:

• Reconstructing tissues/body structures.
• Repairing damaged tissues.

<table>
<thead>
<tr>
<th>Procedure Codes</th>
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<tbody>
<tr>
<td>11950 11951 11952 11954 11970 15780 15781 15786 15787 15788</td>
</tr>
<tr>
<td>15789 15838 15876 21089 21497 41820 41821 41828 61501 Q3031</td>
</tr>
</tbody>
</table>

9.3.2.1 Additional Payable Procedure Codes
The following procedure codes are a benefit when prior authorized and the dentist is qualified and licensed to perform the procedures:

<table>
<thead>
<tr>
<th>Procedure Codes</th>
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</thead>
<tbody>
<tr>
<td>Surgery</td>
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<tr>
<td>10004 10005 10006 10007 10008 10009 10010 10011 10012 10021</td>
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<tr>
<td>Procedure Codes</td>
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<tr>
<td>20920 20922 20955 20956 20957 20962 20969 20970 20971 20973</td>
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<td>21082 21083 21085 21087 21088 21089 21100 21110 21116 21120</td>
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<td>21337 21338 21339 21340 21343 21344 21345 21346 21347 21348</td>
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<td>21355 21356 21360 21365 21366 21366 21367 21368 21369 21395</td>
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</tr>
<tr>
<td>41113 41114 41115 41116 41120 41130 41135 41140 41145 41150</td>
</tr>
</tbody>
</table>
9.3.2.2 Immune Globulin by a Doctor of Dentistry as a Limited Physician

A Doctor of Dentistry Practicing as a Limited Physician may be reimbursed for immune globulin injection procedure code J1571 when billed with one of the following diagnosis codes:

<table>
<thead>
<tr>
<th>Diagnosis Codes</th>
</tr>
</thead>
<tbody>
<tr>
<td>D611 D612 D613 D619 D804 D805 D838 D8982</td>
</tr>
</tbody>
</table>

9.3.2.3 Radiographs by a Doctor of Dentistry Practicing as a Limited Physician

When a Doctor of Dentistry Practicing as a Limited Physician uses appropriate radiograph equipment to produce required radiographs, the following procedure codes are eligible for reimbursement:

<table>
<thead>
<tr>
<th>Procedure Codes</th>
</tr>
</thead>
<tbody>
<tr>
<td>70100 70110 70120 70130 70140 70150 70160 70190 70200 70250</td>
</tr>
<tr>
<td>70260 70300 70310 70320 70328 70332 70336 70350 70355 70370</td>
</tr>
<tr>
<td>70371 70380 70390 73100 70450 70460 70470 70480 70481 70482</td>
</tr>
</tbody>
</table>
9.3.2.4 Dental Anesthesia by a Doctor of Dentistry Practicing as a Limited Physician

A Doctor of Dentistry Practicing as a Limited Physician who is licensed by the Texas State Board of Dental Examiners (TSBDE) practicing in Texas, who has obtained an Anesthesia Permit from the TSBDE in accordance with Title 22 TAC §§110.1 through 110.15, may be reimbursed for anesthesia services on clients having dental/oral and maxillofacial surgical procedures in the dental office or hospital in accordance with all applicable rules for physician administration and supervision of anesthesia services.

Dentists providing sedation/anesthesia services must have the appropriate permit from TSBDE for the level of sedation/anesthesia provided.

The following anesthesia services are payable to dentists as physician services:

<table>
<thead>
<tr>
<th>Procedure Codes</th>
<th>70486</th>
<th>70487</th>
<th>70488</th>
<th>70490</th>
<th>70491</th>
<th>70492</th>
</tr>
</thead>
</table>

9.4 Documentation Requirements

All services require documentation to support the medical necessity of the service rendered, including physician services. Physician services are subject to retrospective review and recoupment if documentation does not support the service billed.

9.5 Claims Filing and Reimbursement

9.5.1 Claims Information

Claims for physician and doctor services must be submitted to TMHP in an approved electronic format or on the CMS-1500 paper claim form. Providers may purchase CMS-1500 paper claim forms from the vendor of their choice. TMHP does not supply them.

When completing a CMS-1500 paper claim form, all required information must be included on the claim, as information is not keyed from attachments. Superbills and itemized statements are not accepted as claim supplements.

Physicians who submit a claim using the physician’s own provider identifier for services provided by an NP, CNS, PA, or CNM must submit one of the following modifiers on each claim detail if the physician does not make a decision regarding the client’s care or treatment on the same date of service as the billable medical visit:

- SA – Services were provided by an NP or CNS
- U7 – Services were provided by a physician assistant
- SB – Services were provided by a CNM

Refer to: “Section 3: TMHP Electronic Data Interchange (EDI)” (Vol. 1, General Information) for information on electronic claims submissions.

“Section 6: Claims Filing” (Vol. 1, General Information) for general information about claims filing.

Subsection 6.5, “CMS-1500 Paper Claim Filing Instructions” in “Section 6: Claims Filing” (Vol. 1, General Information) for instructions on completing paper claims. Blocks that are not referenced are not required for processing by TMHP and may be left blank.
9.5.2 National Drug Codes (NDC)

Refer to: Subsection 6.3.4, “National Drug Code (NDC)” in “Section 6: Claims Filing” (Vol. 1, General Information).

9.5.3 Reimbursement

Texas Medicaid rates for physicians and other practitioners are calculated in accordance with TAC §355.8085. Providers can refer to the online fee lookup (OFL) or the applicable fee schedule on the TMHP website at www.tmhp.com.

Physicians may be reimbursed 92 percent of the established reimbursement rate for services provided by an NP, CNS, PA, or CNM if the physician does not make a decision regarding the client’s care or treatment on the same date of service as the billable medical visit. The 92 percent reimbursement rate will not apply to laboratory services, X-ray services, and injections provided by an NP, CNS, PA, or CNM.

Texas Medicaid implemented mandated rate reductions for certain services. The OFL and static fee schedules include a column titled “Adjusted Fee” to display the individual fees with all mandated percentage reductions applied. Additional information about rate changes is available on the TMHP website at www.tmhp.com/resources/rate-and-code-updates.

Section 104 of the Tax Equity and Fiscal Responsibility Act (TEFRA) of 1982 requires that Medicare/Medicaid limit reimbursement for those physician services furnished in outpatient hospital settings (e.g., clinics and emergency situations) that are ordinarily furnished in physician offices. Reimbursement for these services will be 60 percent of the Texas Medicaid rate for the service furnished in the physician’s office. The following table identifies the services applicable to the 60-percent limitation when furnished in outpatient hospital settings:

<table>
<thead>
<tr>
<th>Procedure Codes</th>
</tr>
</thead>
<tbody>
<tr>
<td>99202</td>
</tr>
<tr>
<td>99282</td>
</tr>
</tbody>
</table>

These procedures are designated with note code “1” in the current physician fee schedule, which is available at www.tmhp.com. The following list shows the services excluded from the 60-percent limitation:

- Services furnished in rural health clinics (RHCs).
- Surgical services that are covered ambulatory surgical center (ASC)/hospital-based ambulatory surgical center (HASC) services.
- Anesthesiology and radiology services.
- Prenatal services when billed with modifier TH and the appropriate E/M procedure code to the highest level of specificity.
- Emergency services provided in a hospital emergency room after the sudden onset of a medical condition manifesting itself by acute symptoms of sufficient severity (including severe pain), such that the absence of immediate medical attention could reasonably be expected to result in one of the following:
  - Serious jeopardy to the client’s health.
  - Serious impairment to bodily functions.
  - Serious dysfunction of any bodily organ or part.
Because of TEFRA, Texas Medicaid reimbursement for a payable nonemergency office service that is performed in the outpatient department of a hospital is limited to 60 percent of Texas Medicaid rate for that service. If the condition qualifies as an emergency or if the client is critically ill or critically injured, the 60 percent professional service reimbursement limit does not apply.

**Refer to:** Subsection 2.2, “Fee-for-Service Reimbursement Methodology” in “Section 2: Texas Medicaid Fee-for-Service Reimbursement” (Vol. 1, General Information) for more information about reimbursement.

Subsection 2.2.1.1, “Non-emergent and Non-urgent Evaluation and Management (E/M) Emergency Department Visits” in “Section 2: Texas Medicaid Fee-for-Service Reimbursement” (Vol. 1, General Information) for more information about conditions that are excluded from the 60-percent limitation.

Subsection 9.2.7, “Anesthesia” in this handbook for information on anesthesia services that are reimbursed according to relative value units (RVUs).

### 9.5.3.1 Affordable Care Act of 2010 (ACA) Rate Increase for Primary Care Services

To qualify for the Affordable Care Act of 2010 (ACA) rate increase for primary care services, a physician must have a specialty designated of general internal medicine, family practice, or pediatrics and must attest to one of the following:

- **The provider has a certification recognized by the American Board of Allergy and Immunology (ABAI), American Board of Medical Specialties (ABMS), American Board of Physician Specialties (ABPS), or American Osteopathic Association (AOA) and meets the requirements as required by federal and state regulation to receive the increased payment.**

- **The provider does not have a certification recognized by the ABAI, ABMS, ABPS, or AOA, but at least 60 percent of the provider’s Medicaid billings for the previous calendar year (or for the previous calendar month if the provider has been enrolled in Medicaid for less than one year) were for the evaluation and management (E/M) and vaccine administration procedure codes as published in the final federal and state regulations and the provider meets the requirement to receive payment.**

**Note:** New providers with no history of Medicaid billings can attest that 60 percent of their Medicaid billing will be for primary care services.

Providers can attest using the Texas Medicaid Attestation for ACA Primary Care Services Rate Increases form. ABAI-certified allergists must indicate “ABAI-allergy” in the “List subspecialties” field of the attestation form.

**Important:** By signing the form, providers attest that they qualify for the rate increase, and that the increase will be applied to paid claims for primary care services on or after the effective date. Payment of the rate increase may be subject to retrospective review and recoupment if it is determined at a later time that the provider did not qualify for the ACA primary care services rate increase. Federal regulations require states to conduct an annual audit of provider attestations.

Non-physician practitioners who are under the supervision of a provider who has self-attested, are not required to submit a separate provider attestation form. Increased payment may be available to the supervising physician when the following conditions are met:

- **The non-physician practitioner renders services under the personal supervision of a provider who has self-attested to meeting the requirements.**

- **Services are billed under the qualifying provider’s provider identification number.**
10 Physician Assistant

10.1 Enrollment

To enroll in Texas Medicaid, a PA must be licensed and recognized as a PA by the Texas Physician Assistant Board. Texas Medicaid accepts a signed letter of certification from the Texas Physician Assistant Board as acceptable documentation of appropriate licensure and certification for enrollment. The PA must identify their supervising physician in the appropriate field of the enrollment application.

Providers cannot be enrolled if their license is due to expire within 30 days.

Enrollment as an individual provider is optional. PAs currently treating clients and billing under the supervising physician’s provider identifier may continue this billing arrangement.

All PA services must be delivered according to protocols developed jointly within the scope of practice and state law governing PAs.

All providers of laboratory services must comply with the rules and regulations of CLIA. Providers not complying with CLIA are not reimbursed for laboratory services.

PAs may enroll as providers of THSteps medical checkups.

Referato: Subsection 1.1, “Provider Enrollment” in “Section 1: Provider Enrollment and Responsibilities” (Vol. 1, General Information).


Subsection 5.2, “Enrollment” in the Children’s Services Handbook (Vol. 2, Provider Handbooks) for more information about enrolling as a THSteps provider.

PAs may be included as primary care providers in the provider network for Medicaid and CHIP programs (both fee-for-service and managed care), regardless of whether the physician supervising the PA is enrolled in Medicaid or in the provider network.

10.2 Services, Benefits, Limitations, and Prior Authorization

Services performed by PAs are covered if the services meet the following criteria:

- Are within the scope of practice for PAs, as defined by Texas state law
- Are consistent with rules and regulations promulgated by the Texas Medical Board or other appropriate state licensing authority
- Are covered by Texas Medicaid when provided by a licensed physician (MD or DO)
- Are reasonable and medically necessary as determined by HHSC or its designee

Services provided to Medicaid clients must be documented in the client’s medical record to include the following:

- Services provided
- Date of service
- Pertinent information about the client’s condition supporting the need for service
- The individual practitioner of the service

PAs who are employed or remunerated by a physician, hospital, facility, or other provider must not bill Texas Medicaid for their services if the billing results in duplicate payment for the same services.
Physicians who submit a claim using the physician’s own provider identifier for services provided by a PA must submit modifier U7 on each claim detail if the physician does not make a decision regarding the client’s care or treatment on the same date of service as the billable medical visit.

Laboratory (including pregnancy tests) and radiology services provided during pregnancy must be billed separately from antepartum care visits and claims must be received within 95 days from the date of service.

**Note:** Payment to providers for supplies is not a benefit of Texas Medicaid. Costs of supplies are included in the reimbursement for office visits.

**Refer to:**
- Section 2, “Medicaid Title XIX Family Planning Services” in the *Gynecological, Obstetrics, and Family Planning Title XIX Services Handbook* (Vol. 2, Provider Handbooks).
- Section 9, “Physician” in this handbook.
- Section 5, “THSteps Medical” in the *Children’s Services Handbook* (Vol. 2, Provider Handbooks).

### 10.2.1 Prior Authorization

Services performed by a PA are subject to the same prior authorization guidelines as services performed by other provider types.

### 10.3 Documentation Requirements

All services require documentation to support the medical necessity of the service rendered, including PA services. PA services are subject to retrospective review and recoupment if documentation does not support the service billed.

### 10.4 Claims Filing and Reimbursement

#### 10.4.1 Claims Information

Claims for PA services must include modifier U7 on the claim details to indicate that the client was treated by a PA.

PA services must be submitted to TMHP in an approved electronic format or on the CMS-1500 claim form. Providers may purchase CMS-1500 claim forms from the vendor of their choice. TMHP does not supply the forms.

When completing a CMS-1500 claim form, all required information must be included on the claim, as information is not keyed from attachments. Superbills, or itemized statements, are not accepted as claim supplements.

**Refer to:**
- “Section 3: TMHP Electronic Data Interchange (EDI)” *(Vol. 1, General Information)* for information on electronic claims submissions.
- “Section 6: Claims Filing” *(Vol. 1, General Information)* for general information about claims filing.
- Subsection 6.5, “CMS-1500 Paper Claim Filing Instructions” in “Section 6: Claims Filing” *(Vol. 1, General Information)* for instructions on completing paper claims. Blocks that are not referenced are not required for processing by TMHP and may be left blank.
10.4.2 Reimbursement

According to 1 TAC §355.8093, the Medicaid rate for PAs is 92 percent of the rate paid to a physician (MD or DO) for the same professional service and 100 percent of the rate paid to physicians for laboratory services, X-ray services, and injections.

**Note:** PA providers who are enrolled in Texas Medicaid as THSteps providers also receive 92 percent of the rate paid to a physician for THSteps services when a claim is submitted with their THSteps provider identifier as the billing provider.

PAs who bill Medicaid directly for services they perform must use their individual provider identifier. If the services were performed by the PA but billed by a physician or physician group, the billing provider is the physician or physician group. Physicians may be reimbursed 92 percent of the established reimbursement rate for services provided by a PA if the physician does not make a decision regarding the client’s care or treatment on the same date of service as the billable medical visit. This 92 percent reimbursement rate does not apply to laboratory services, X-ray services, or injections provided by a PA.

Providers can refer to the online fee lookup (OFL) or the applicable fee schedule on the TMHP website at [www.tmhp.com](http://www.tmhp.com). To request a hard copy, call the TMHP Contact Center at 1-800-925-9126.

Texas Medicaid implemented mandated rate reductions for certain services. The OFL and static fee schedules include a column titled “Adjusted Fee” to display the individual fees with all mandated percentage reductions applied. Additional information about rate changes is available on the TMHP website at [www.tmhp.com/resources/rate-and-code-updates](http://www.tmhp.com/resources/rate-and-code-updates).

**Refer to:** Subsection 1.1, "Provider Enrollment” in ”Section 1: Provider Enrollment and Responsibilities” (Vol. 1, General Information).”

“Section 3: TMHP Electronic Data Interchange (EDI)” (Vol. 1, General Information) for information on how to obtain electronic fee schedules from the TMHP website.

11 Claims Resources

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<th>Resource</th>
<th>Location</th>
</tr>
</thead>
<tbody>
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<td>Acronym Dictionary</td>
<td>“Appendix C: Acronym Dictionary” (Vol. 1, General Information)</td>
</tr>
<tr>
<td>Automated Inquiry System (AIS)</td>
<td>Subsection A.10, “TMHP Telephone and Fax Communication” in “Appendix A: State, Federal, and TMHP Contact Information” (Vol. 1, General Information)</td>
</tr>
<tr>
<td>CMS-1500 Paper Claim Filing Instructions</td>
<td>Subsection 6.5, “CMS-1500 Paper Claim Filing Instructions” in ”Section 6: Claims Filing” (Vol. 1, General Information)</td>
</tr>
<tr>
<td>Family Planning Claim Form Examples</td>
<td>Section 10, “Claim Form Examples” in the Gynecological, Obstetrics, and Family Planning Title XIX Services Handbook (Vol. 2, Provider Handbooks)</td>
</tr>
<tr>
<td>State, federal, and TMHP contact information</td>
<td>“Appendix A: State, Federal, and TMHP Contact Information” (Vol. 1, General Information)</td>
</tr>
<tr>
<td>TMHP electronic claims submission information</td>
<td>Subsection 6.2, “TMHP Electronic Claims Submission” in “Section 6: Claims Filing” (Vol. 1, General Information)</td>
</tr>
<tr>
<td>TMHP Electronic Data Interchange (EDI)</td>
<td>“Section 3: TMHP Electronic Data Interchange (EDI)” (Vol. 1, General Information)</td>
</tr>
</tbody>
</table>
12 Contact TMHP

The TMHP Contact Center at 1-800-925-9126 is available Monday through Friday from 7 a.m. to 7 p.m., Central Time.

13 Forms

The following linked forms can also be found on the Forms page of the Provider section of the TMHP website at www.tmhp.com:

<table>
<thead>
<tr>
<th>Resource</th>
<th>Location</th>
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14 Claim Form Examples

The following linked claim form examples can also be found on the Claim Form Examples page of the Provider section of the TMHP website at www.tmhp.com:

<table>
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<th>Claim Form Examples</th>
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</tr>
<tr>
<td>Certified Nurse-Midwife (CNM)</td>
</tr>
<tr>
<td>Certified Registered Nurse Anesthetist (CRNA)</td>
</tr>
<tr>
<td>Chiropractic Services</td>
</tr>
<tr>
<td>Dental (Doctor of Dentistry)</td>
</tr>
<tr>
<td>Dialysis Training</td>
</tr>
<tr>
<td>Claim Form Examples</td>
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<tr>
<td>Genetics</td>
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<td>Radiation Therapy</td>
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<td>Surgery</td>
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1 General Information

The Medical Transportation Program (MTP), under the direction of the Texas Health and Human Services Commission (HHSC), arranges transportation and travel-related services for eligible Medicaid, Children with Special Health Care Needs (CSHCN) Services Program, and Transportation for Indigent Cancer Patients (TICP) clients who have no other means of transportation. MTP is responsible for the prior authorization of all MTP services.

MTP provides for the following general services:

- Mass transit (intercity and intracity): Passes or tickets for client transport within a city and from city to city. Air travel is also an allowable service.
- Demand response transportation: Common carriers such as taxi, wheelchair van, and other transportation according to contractual requirements.
- Mileage reimbursement for enrolled individual transportation provider (ITP): The enrolled ITP can be the responsible party, family member, friend, neighbor, or client.
- Meals: Contracted vendors (e.g., hospital cafeteria).
- Lodging: Contracted hotels and motels.
- Advanced funds: Financial services contractor.
- Attendant: Responsible party, parent/guardian, etc., who accompanies the client to a health-care service.

Under the contract between Texas Medicaid & Healthcare Partnership (TMHP) and MTP, TMHP is responsible for enrollment of providers and processing of MTP provider claims.

MTP contracts with various provider types to arrange transportation and travel-related services for eligible MTP clients and their attendants.

There are three MTP provider types that enroll directly with TMHP:

- ITPs
- Lodging providers
- Meal providers

All other transportation providers arrange enrollment through MTP (e.g., transportation service area providers, client services providers).

1.1 Contacting MTP

If health-care providers have MTP-eligible clients who express difficulty accessing health-care services, advise the clients or their advocates to call the statewide MTP toll-free number at 1-877-633-8747 to request transportation services. MTP clients in the Houston/Beaumont area can call 1-855-687-4786 to request services. Clients in the Dallas/Ft. Worth area can call 1-855-687-3255 to request services. For transportation services within the county where the client lives, clients or their advocates must call the MTP office at least 2 business days before the scheduled appointment. For clients who need to travel beyond the county where they live, clients or their advocates must call the MTP office at least 5 business days before the scheduled appointment.

The client must provide the following information to the intake operator at the time of the call:

- Client name, address, and, if available, the telephone number
- Medicaid, TICP or CSHCN Services Program client identification number (if applicable) or Social Security number, and date of birth
• Name, address, and telephone number of health-care provider and/or referring health-care provider
• Purpose and date of trip and time of appointment
• Affirmation that other means of transportation are unavailable
• Special needs, including wheelchair lift or attendant(s)
• Medical necessity verified by the Health Care Provider’s Statement of Need, if applicable
• Affirmation that advance funds are needed in order for the recipient to access health-care services

Note: Clients must reimburse the department for any advance funds, and any portion thereof, that are not used for the specific prior authorized service.

2 Individual Transportation Provider (ITP)

ITPs are individuals who volunteer to use their personal vehicle to drive themselves, a friend, or a family member safely to the doctor, dentist, or drug store.

2.1 Enrollment for ITPs

ITPs must follow all rules for enrollment that other providers follow when enrolling with TMHP.

To initiate the enrollment process, the MTP client must contact MTP to request a ride from an individual who is a potential ITP. This request is the first step in the enrollment process for the ITP.

After the client’s call, MTP sends the potential ITP’s information to TMHP, and TMHP mails the potential ITP an enrollment package. The ITP must fill out the Individual Transportation Provider Enrollment Application and mail it to TMHP with all requested documentation.

The provider must identify the MTP clients they will be transporting and whether they are related to the client. The application packet also includes an Electronic Funds Transfer (EFT) Agreement form that authorizes TMHP to deposit payments directly into a bank account, which results in faster payments.

After the ITP application has been processed, the ITP will receive a letter from TMHP that includes the Atypical Provider Identifier (API) and the Texas Provider Identifier (TPI) to be used when the ITP submits claim forms for mileage reimbursement.

2.2 Prior Authorization for ITPs

Once an ITP is enrolled with TMHP and a client calls MTP to request a ride, MTP will mail a preprinted ITP Service Record (Form H3017) to the MTP client. The H3017 is the form the provider must mail to TMHP to be reimbursed for the ride.

Important: Only claims that are authorized by MTP will be considered for payment. All claims must be prior authorized to be paid.

Refer to: Subsection 3, “Prior Authorization” in this handbook.

2.3 Claims Filing for ITPs

To file a claim, the ITP must complete the H3017 form that was sent to the Medicaid client and mail it to TMHP at the following address:

Texas Medicaid & Healthcare Partnership
Claims
PO Box 200555
Austin, TX 78720-0555
The H3017 includes the following transportation details:

- Date of the ride
- Number of miles authorized
- MTP Authorization Number
- MTP client’s name
- ITP’s name

The H3017 claim form must be signed by the doctor, dentist, or drug store representative that rendered services to the MTP client. This signature stands as proof that the ride authorized by MTP was taken. The ITP must also sign the claim form and include the API and TPI that was assigned to them by TMHP. If any of this required information is missing, the claim will be denied.

The provider must mail the completed claim form to TMHP after the client’s authorized ride, but no later than 95 days from the date of the ride. Any claims received by TMHP more than 95 days after the date of the ride will be denied.

An ITP may not charge an MTP client a fee for completing claim forms. TMHP also cannot be charged for the filing of claim forms.

3 Prior Authorization

All MTP services must be prior authorized by MTP, which issues all prior authorizations for transportation services. The eligible MTP client must contact MTP to obtain an authorization. Claims that are submitted without proper prior authorization will be denied.

3.1 Retention of Prior Authorization Documents

MTP prior authorization documents relating to Medicaid services or benefits provided to clients who are 20 years of age and younger must not be destroyed until the provider receives notice from HHSC. Examples of such documents include but are not limited to:

- Correspondence with HHSC/MTP;
- Invoices
- Receipts
- Contacts with clients who are class members

3.2 Definition of Prior Authorization Documents

The term “prior authorization document” is broad and includes, but is not limited to, the following:

- Paper records
- Electronic files in any format
- Database entries
- The original and any drafts or non-identical copies of any document
- Exhibits or attachments to documents
- Handwritten documents
- Emails
- Drawings, graphs, charts
• Electronic or videotape recordings
• Computer disks
• Other forms of computer memory storage

3.3 Copies of Prior Authorization Documents
Providers are not required to retain multiple exact copies of a document. For example:
• An exact electronic copy (e.g., scanned computer image, microfiche) may be retained instead of a paper copy.
• If the last in a chain of emails is retained, it is not necessary to retain each of the individual emails included in the chain, as long as the email that is retained reflects all of the earlier emails.

However, a document containing any substantive editorial comment, margin notes, underlining, etc., is not an exact copy and becomes a new original that must be retained.

3.4 Storage of Prior Authorization Document Storage
Relevant information and documents should be stored in a way that is protected from unintentional disclosure or destruction.

4 Claims Filing
This section contains instructions for completion of Medicaid-required claim forms. When filing a claim, providers should review the instructions carefully and complete all requested information. A correctly completed claim form is processed faster.

Texas Medicaid cannot make payments to clients, so the provider who performs the service must file an assigned claim. Federal regulations prohibit providers from charging clients a fee for completing or filing Medicaid claim forms. Providers are not allowed to charge TMHP for filing claims. The cost of claims filing is part of the usual and customary rate for doing business. Providers cannot bill Texas Medicaid or Medicaid clients for missed appointments or failure to keep an appointment. Only claims for services rendered are considered for payment.

Medicaid providers are also required to complete and sign authorized medical transportation forms (e.g., Form 3103, Individual Driver Registrant (IDR) Service Record, or Form 3111, Verification of Travel to Healthcare Services by Mass Transit) or provide an equivalent (e.g., provider statement on official letterhead) to attest that services were provided to a client on a specific date. The client presents these forms to the provider.

Providers are not allowed to bill clients or Texas Medicaid for completing these forms.

Medicaid claims are subject to the following procedures:
• TMHP verifies all required information is present.
• Claims filed under the same provider identifier and program and ready for disposition at the end of each week are paid to the provider with an explanation of each payment or denial. The explanation is called the Remittance and Status (R&S) Report, which may be received as a downloadable portable document format (PDF) version or on paper. A Health Insurance Portability and Accountability Act (HIPAA)-compliant 835 transaction file is also available for those providers who wish to import claim dispositions into a financial system.
An R&S Report is generated for providers that have weekly claim or financial activity with or without payment. The report identifies pending, paid, denied, and adjusted claims. If no claim activity or outstanding account receivables exist during the time period, an R&S Report is not generated for the week.

Providers can participate in the most efficient and effective method of submitting claims to TMHP by submitting claims through the TMHP Electronic Data Interchange (EDI) claims processing system using TexMedConnect or a third party vendor. Claims must contain the provider’s complete name, address, and provider identifier to avoid unnecessary delays in processing and payment.

### 4.1 Claims Filing Deadlines

All claims for services rendered to eligible MTP clients are subject to a filing deadline from the DOS of:

- 95 days for in-state providers
- 365 days for out-of-state providers

Claims submitted by newly-enrolled MTP providers must be received within 95 days of the date the atypical provider identifier (API) is issued, and within 365 days of the date of service (DOS). Providers with a pending application should submit any claims that are nearing the 365-day deadline from the DOS. TMHP will reject all claims until an API is issued. MTP providers can use the TMHP rejection report or Return to Provider (RTP) letters as proof of meeting the 365-day deadline and submit an appeal.

### 4.2 Auditing of Claims

Reimbursement may be recouped when the medical record does not document that the level of service provided accurately matches the level of service claimed. Furthermore, the level of service provided and documented must be medically necessary based on the clinical situation and needs of the patient.

HHSC and TMHP routinely perform retrospective reviews of all providers. HHSC ultimately is responsible for Texas Medicaid utilization review activities. This review includes comparing services billed to the client’s clinical record. The following requirements are general requirements for all providers. Any mandatory requirement not present in the client’s medical record subjects the associated services to recoupment.

### 4.3 Important Codes for All MTP Providers

MTP providers must use the following codes when submitting claims:

- Benefit Code = MTP
- Provider Type = MT
- Diagnosis Code = Z753
- Place of Service = 09 for paper claims, 99 for TexMedConnect claims
- Type of Service = 9

The following table shows additional codes that TMHP recommends for filing MTP claims. The codes are based on transportation provider type:

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<tr>
<th>MTP Provider Description</th>
<th>Provider Specialty</th>
<th>Taxonomy Code</th>
<th>Recommended Procedure Code</th>
<th>Modifier Codes</th>
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<tr>
<td>Client Financial Services (CFS)</td>
<td>T1</td>
<td>347E00000X</td>
<td>A0170</td>
<td></td>
</tr>
<tr>
<td>Individual Transportation Provider (ITP)</td>
<td>T4</td>
<td>347C00000X</td>
<td>S0215</td>
<td></td>
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</tbody>
</table>
4.4 Delegation of Signature Authority

A provider that delegate signatory authority to a member of the office staff or to a billing service remains responsible for the accuracy of all information on a claim submitted for payment. A provider’s employees or a billing service and its employees are equally responsible for any false billings in which they participated or directed.

If the claim is prepared by a billing service or printed by data processing equipment, it is permissible to print “Signature on File” in place of the provider’s signature. When claims are prepared by a billing service, the billing service must obtain and keep a letter on file that is signed by the provider authorizing claim submission.

4.5 Electronic Claims

4.5.1 TMHP Electronic Data Interchange (EDI)

Providers are encouraged to submit claims using electronic methods. Providers can participate in the most efficient and effective method of submitting requests to TMHP by submitting through the TMHP EDI Gateway. TMHP uses the HIPAA-compliant American National Standards Institute (ANSI) ASC X12 4010A1 file format through secure socket layer (SSL) and virtual private networking (VPN) connections for maximum security. Providers can access TMHP’s electronic services through the TMHP website at www.tmhp.com, TexMedConnect, vendor software, and billing agents. Providers may also submit claims using paper forms. Version 2001 0805 3 MTP Claim Filing.

4.5.2 TexMedConnect

TexMedConnect is a free, web-based, claims submission application provided by TMHP. Technical support and training for TexMedConnect are also available free from TMHP. Providers can submit claims, eligibility requests, claim status inquiries, appeals, and download ER&S Reports (in either PDF or ANSI 835 formats) using TexMedConnect. TexMedConnect can interactively submit individual claims that are processed in seconds. To use TexMedConnect, providers must have:

- An internet service provider (ISP)
- Microsoft Internet Explorer®

<table>
<thead>
<tr>
<th>MTP Provider Description</th>
<th>Provider Specialty</th>
<th>Taxonomy Code</th>
<th>Recommended Procedure Code</th>
<th>Modifier Codes</th>
</tr>
</thead>
<tbody>
<tr>
<td>Lodging</td>
<td>T6</td>
<td>177F00000X</td>
<td>A0180</td>
<td>U1</td>
</tr>
<tr>
<td>Meals</td>
<td>T8</td>
<td>174200000X</td>
<td>A0190</td>
<td>U1, U2, U3</td>
</tr>
<tr>
<td>Transportation Service Area Provider (TSAP)</td>
<td>T0</td>
<td>343800000X</td>
<td>A0100</td>
<td>U1, U2, U3, U4</td>
</tr>
</tbody>
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A broadband connection is recommended but not required. Providers that use TexMedConnect can find the online instruction manual on the home page and on the EDI page of the TMHP website at www.tmhp.com.

4.5.3 Vendor Software

Providers that do not use TexMedConnect may use vendor software to create, submit, and retrieve data files. Providers can use software from any vendor listed on the EDI Submitter List, which is located on the EDI Vendor Testing web page of the TMHP website at www.tmhp.com. There are hundreds of software vendors that have a wide assortment of services and that have been approved to submit electronic files to TMHP. Providers that plan to access TMHP’s electronic services with vendor software should contact the vendor for details on software requirements. TMHP does not make vendor recommendations or provide any assistance for vendor software. Not all vendor software offers the same features or levels of support. Providers are encouraged to research their software thoroughly to make certain that it meets their needs and that it has completed testing with TMHP.

Providers must setup their software or billing agent services to access the TMHP EDI Gateway. Providers who use billing agents or software vendors should contact those organizations for information on installation, settings, maintenance, and their processes and procedures for exchanging electronic data.

Providers that download the ANSI 835 file through TexMedConnect and providers that use vendor software must request a submitter ID. A submitter ID is necessary for vendor software to access TMHP’s electronic services. It serves as an electronic mailbox for the provider and TMHP to exchange data files. To order a submitter ID, providers must call the EDI Help Desk at 1-888-863-3638. Providers that use a billing agent do not need a submitter ID.

Providers may receive an ER&S Report by completing the Electronic Remittance and Status (ER&S) Agreement and submitting it to the EDI Help Desk after setting up access to the TMHP EDI Gateway. 4 Version 2001 0805 MTP Claim Filing.

4.5.4 Third Party Vendor Implementation

TMHP requires all software vendors and billing agents to complete EDI testing before access to the production server is allowed. Vendors that wish to begin testing may either call the EDI Help Desk at 1-888-863-3638 or visit the EDIFECs testing site at edittesting.tmhp.com and use the TMHP Support link. An EDIFECs account will be created for the vendor to begin testing EDI formats once they have enrolled for testing. After the successful completion of EDIFECs testing and the submission of a Trading Partner Agreement, vendors must then complete end-to-end testing on the TMHP test server. Software vendors and billing agents must be partnered with at least one Texas provider before a test submitter ID can be issued. When end-to-end testing has been completed, the software vendor or billing agent will be added to the EDI Submitter List. Providers and billing agents may then order production submitter IDs for use with the vendor’s software. Companion guides and vendor specifications are available in the EDI section of the TMHP website at www.tmhp.com.

4.6 Paper Claims

MTP providers can also file claims using the CMS-1500 paper claim form. Providers obtain copies of the CMS-1500 paper claim form from a vendor of their choice; TMHP does not supply them.
Providers must submit paper claims to TMHP at the following address:

Texas Medicaid & Healthcare Partnership
Claims
PO Box 200555
Austin, TX 78720-0555

4.6.1 Tips on Expediting Paper Claims

Use the following guidelines to enhance the accuracy and timeliness of paper claims processing.

4.6.1.1 General requirements

- Use original claim forms. Don’t use copies of claim forms.
- Detach claims at perforated lines before mailing.
- Use 10 x 13 inch envelopes to mail claims. Don’t fold claim forms, appeals, or correspondence.
- Don’t use labels, stickers, or stamps on the claim form.
- Don’t send duplicate copies of information.
- Use 8 ½ x 11 inch paper. Don’t use paper smaller or larger than 8 ½ x 11 inches.
- Don’t mail claims with correspondence for other departments.

4.6.1.2 Data Fields

- Print claim data within defined boxes on the claim form.
- Use black ink, but not a black marker. Don’t use red ink or highlighters.
- Use all capital letters.
- Print using 10-pitch (12-point) Courier font, 10 point. Don’t use fonts smaller or larger than 12 points. Don’t use proportional fonts, such as Arial or Times Roman.
- Use a laser printer for best results. Don’t use a dot matrix printer, if possible.
- Don’t use dashes or slashes in date fields.

4.6.1.3 Attachments

- Use paper clips on claims or appeals if they include attachments. Don’t use glue, tape, or staples.
- Place the claim form on top when sending new claims, followed by any medical records or other attachments.
- Number the pages when sending when sending attachments or multiple claims for the same client (e.g., 1 of 2, 2 of 2).
- Don’t total the billed amount on each claim form when submitting multi-page claims for the same client.

Note: It is strongly recommended that providers who submit paper claims keep a copy of the documentation they send.

- All paper claims must be submitted with a TPI and NPI.
- Modifiers describe and qualify the services provided by Texas Medicaid. A modifier is placed after the five-digit procedure code.
4.6.1.4 Attachments to Claims

To expedite claims processing, providers must supply all information on the claim form itself and limit attachments to those required by TMHP or necessary to supply information to properly adjudicate the claim.

4.6.2 CMS-1500 Instruction Table

The table below describes what information must be entered in each of the block numbers of the CMS-1500 claim form. Providers obtain copies of the CMS-1500 paper claim form from a vendor of their choice; TMHP does not supply them.

Block numbers not referenced in the table may be left blank. They are not required for TMHP to process MTP claims.

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<td><strong>Description</strong></td>
</tr>
<tr>
<td>1a</td>
<td>Insured’s ID No. (for program checked above, include all letters)</td>
</tr>
<tr>
<td>2</td>
<td>Patient’s name</td>
</tr>
<tr>
<td>21</td>
<td>Diagnosis or nature of illness or injury</td>
</tr>
<tr>
<td>23</td>
<td>Prior authorization number</td>
</tr>
<tr>
<td>24a</td>
<td>Date(s) of service</td>
</tr>
<tr>
<td>24b</td>
<td>Place of service</td>
</tr>
<tr>
<td>24d</td>
<td>Fully describe procedures, medical services, or supplies furnished for each date given</td>
</tr>
<tr>
<td>24e</td>
<td>Diagnosis pointer</td>
</tr>
<tr>
<td>24f</td>
<td>Charges</td>
</tr>
<tr>
<td>24g</td>
<td>Days or units</td>
</tr>
<tr>
<td>27</td>
<td>Accept assignment</td>
</tr>
<tr>
<td>28</td>
<td>Total charge</td>
</tr>
<tr>
<td>31</td>
<td>Signature of physician or supplier</td>
</tr>
<tr>
<td>33</td>
<td>Billing provider info &amp; PH #</td>
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5 Claim Form Examples

The following linked claim form examples can also be found on the Claim Form Examples page of the Provider section of the TMHP website at www.tmhp.com:

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OUTPATIENT DRUG SERVICES HANDBOOK

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1 General Information

The information in this handbook provides information about Texas Medicaid’s benefits, policies, and procedures applicable to clinician-administered drugs.

Important: All providers are required to read and comply with “Section 1: Provider Enrollment and Responsibilities” (Vol. 1, General Information). In addition to required compliance with all requirements specific to Texas Medicaid, it is a violation of Texas Medicaid rules when a provider fails to provide healthcare services or items to Medicaid clients in accordance with accepted medical community standards and standards that govern occupations, as explained in Title 1 Texas Administrative Code (TAC) §371.1659. Accordingly, in addition to being subject to sanctions for failure to comply with the requirements that are specific to Texas Medicaid, providers may also be subject to Texas Medicaid sanctions for failure, at all times, to deliver healthcare items and services to Medicaid clients in full accordance with all applicable licensure and certification requirements including, without limitation, those related to documentation and record maintenance.

Refer to: “Section 1: Provider Enrollment and Responsibilities” (Vol. 1, General Information).

1.1 About the Vendor Drug Program

The Texas Vendor Drug Program (VDP) provides statewide access to prescription drugs as authorized by a prescribing provider for clients enrolled in:

- Medicaid (fee-for-service and managed care).
- Children’s Health Insurance Program (CHIP).
- Children with Special Health Care Needs (CSHCN) Services Program.
- Healthy Texas Women (HTW) Program.
- Kidney Health Care (KHC) Program.

VDP manages the Medicaid and CHIP drug formularies and Medicaid Preferred Drug List (PDL).

Note: Pharmacy services for clients in Medicaid managed care are administered by a client’s managed care organization (MCO).

Refer to: The Medicaid Managed Care Handbook (Vol. 2, Provider Handbooks) for additional information about managed care prescription drug and pharmacy benefits.

Clinician-administered drugs or biologicals, also known as physician-administered drugs, are injectable medications given in an office or outpatient clinic setting when oral medications are not appropriate and may be reimbursable as a medical benefit through Texas Medicaid and CHIP. Newly-released Healthcare Common Procedure Coding System (HCPCS) codes for CADs are reviewed throughout the year. If a CAD is determined to be an appropriate benefit for Medicaid, then the HCPCS code is presented at a rate hearing as part of the process to become a benefit. CADs may become a benefit prior to issuance of a HCPCS code. In this instance, the CAD may be listed on the Texas NDC-to-HCPCS Crosswalk with an unclassified HCPCS code. VDP manages the CAD benefit and crosswalk. The Texas NDC-to-HCPCS Crosswalk identifies relationships between National Drug Codes (NDC) and payable Healthcare Common Procedure Coding System (HCPCS) codes in Medicaid and CHIP. The crosswalk assists with billing and coding and is a reference for converting HCPCS billing units to valid NDC unit calculations. The crosswalk is published quarterly and based on revisions to the CMS list of rebate-eligible drugs and new drugs and biologicals added to First Databank.
1.2 Pharmacy Enrollment

VDP enrolls any eligible, in-state pharmacy licensed as Class A or C by the Texas State Board of Pharmacy.

Any out-of-state pharmacies or pharmacies that hold any other class of pharmacy license are considered for inclusion in the program on a case-by-case basis. Consideration is relative to the benefits made available to the client eligible for pharmacy benefits. Enrollment is not granted unless additional benefits are established.

Pharmacy providers must be enrolled with VDP prior to providing outpatient prescription services and prior to participating in any Medicaid managed care network. To participate in the Medicaid or CHIP managed care networks the pharmacy must contact the health plan.

Pharmacy providers enrolled with VDP should refer to the VDP Pharmacy Provider Procedure Manual for policies and procedures pertaining to fee-for-service outpatient pharmacy claims, including drug benefit guidance, pharmacy prior authorization, coordination of benefits, drug pricing, and reimbursement.

Refer to: The VDP Pharmacy Provider Procedure Manual on the VDP website.

1.3 Program Contact Information

<table>
<thead>
<tr>
<th>Vendor Drug Program</th>
<th>Telephone Number</th>
</tr>
</thead>
<tbody>
<tr>
<td>Pharmacy Benefits Access: for questions about outpatient drug and billing (the 800 number is for pharmacy use only and can be used to reach any area within VDP).</td>
<td>1-800-435-4165</td>
</tr>
<tr>
<td>Program Management</td>
<td>1-512-707-6108</td>
</tr>
<tr>
<td>Program Policy</td>
<td>1-512-707-6108</td>
</tr>
<tr>
<td>Drug formulary (Texas listing of national drug codes)</td>
<td>1-512-462-6390</td>
</tr>
<tr>
<td>Texas Pharmacy Prior Authorization Center Hotline</td>
<td>1-877-728-3927</td>
</tr>
<tr>
<td>Texas Pharmacy Third Party Call Center</td>
<td>1-866-389-5594</td>
</tr>
</tbody>
</table>

2 Enrollment

Refer to: Subsection 1.1, “Provider Enrollment” in “Section 1: Provider Enrollment and Responsibilities” (Vol. 1, General Information) for more information about procedures for enrolling as a Medicaid provider.

Subsection 2.2, “Provider Enrollment and Responsibilities” in the Medicaid Managed Care Handbook (Vol. 2, Provider Handbooks).

3 Services, Benefits, Limitations, and Prior Authorization

Clinician-administered drugs or biologicals (CADs), also known as physician-administered drugs, are injectable medications given in an office or outpatient clinic setting when oral medications are not appropriate and may be reimbursable as a medical benefit through Texas Medicaid and CHIP.

Newly released HCPCS codes for CADs and biologicals are reviewed by Texas Medicaid throughout the year. If the CADs are determined to be appropriate benefits for Medicaid, then the HCPCS codes are presented at a rate hearing as part of the process to become a benefit. An application to initiate this
review process is not necessary. HHSC’s review of any new CAD does not guarantee that the new CAD will become a benefit. If a manufacturer is interested in having a CAD included on the Texas Medicaid Vendor Drug Program (VDP) formulary list it is necessary to contact VDP for an application.

If a HCPCS code that already is a benefit of Texas Medicaid has a new NDC that needs to be added to the Texas NDC-to-HCPCs crosswalk, contact the Texas Medicaid Vendor Drug Program. A new NDC for a currently payable HCPCS code generally does not require a new rate hearing.

Refer to: Subsection 9, “Pharmacy Benefit” in this handbook for more information.

The Medicaid Managed Care Handbook (Vol. 2, Provider Handbooks) for information about the managed care prescription drug and pharmacy benefits.

3.1 Prior Authorization Requests

Prior authorization requests for CADs must be submitted to the TMHP Prior Authorization Department by mail, fax, or the electronic portal.

To facilitate determination of medical necessity and avoid unnecessary denials, the prescribing provider must submit correct and complete information, including documentation for medical necessity for the equipment or supplies requested, procedure codes, and numerical quantities for services requested. The provider must maintain documentation of medical necessity in the client’s medical record.

The requesting provider may be asked for additional information to clarify or complete a request.

To complete the prior authorization process by paper, the provider must fax or mail the completed prior authorization request form to the Special Medical Prior Authorization (SMPA) unit.

To complete the prior authorization process electronically, the provider must complete the prior authorization requirements through any approved electronic methods and retain a copy of the signed and dated SMPA Request form in the client’s medical record.

A SMPA Request Form must be completed, signed, and dated by the prescribing provider. The SMPA form will not be accepted beyond 90 days from the date of the prescribing provider’s signature.

The completed SMPA Request Form must be maintained by the prescribing provider in the client’s medical record and is subject to retrospective review.

Documentation of the client’s dosage, administration schedule, number of injections to be administered during the prior authorization period, requested units per injection, and the dosage calculation must be submitted in Section C of the Special Medical Prior Authorization (SMPA) Request Form under Statement of Medical Necessity.

3.2 Electronic Signatures in Prior Authorizations

Prior authorization requests may be submitted to the TMHP Prior Authorization Department by mail, fax, or the electronic portal. Prescribing or ordering providers, dispensing providers, clients’ responsible adults, and clients may sign prior authorization forms and supporting documentation using electronic or wet signatures.

Refer to: Subsection 5.5.1.2, “Document Requirements and Retention” in “Section 5: Fee-for-Service Prior Authorizations” (Vol. 1, General Information) for additional information about electronic signatures.
4 Reimbursement

Clinician-administered drugs, vaccines, and biologicals are reimbursed under Texas Medicaid in accordance with 1 TAC rule §355.8085. Reimbursement for clinician-administered drugs, vaccines, and biologicals are based on the lesser of the billed amount, a percentage of the Medicare rate, or one of the following methodologies:

- If the drug or biological is considered a new drug or biological (that is, approved for marketing by the Food and Drug Administration within 12 months of implementation as a benefit of Texas Medicaid), it may be reimbursed at an amount equal to 89.5 percent of average wholesale price (AWP).
- If the drug or biological does not meet the definition of a new drug or biological, it may be reimbursed at an amount equal to 85 percent of AWP.
- Vaccines may be reimbursed at an amount equal to 89.5 percent of AWP.
- Infusion drugs furnished through an item of implanted durable medical equipment may be reimbursed at an amount equal to 89.5 percent of AWP.
- Drugs, other than vaccines and infusion drugs, may be reimbursed at an amount equal to 106 percent of the average sales price (ASP).

HHSC may use other data sources to determine Medicaid fees for physician-administered drugs, vaccines, and biologicals when HHSC determines that the above methodologies are unreasonable or insufficient.

Texas Medicaid reimburses providers using several different reimbursement methodologies, including fee schedules, reasonable cost with interim rates, hospital reimbursement methodology, provider-specific encounter rates, reasonable charge payment methodology, and manual pricing. Each Texas Medicaid service describes the appropriate reimbursement for each service area.

Note: If a client is covered by a Medicaid managed care organizations (MCO) or dental plan, providers must contact the client’s MCO or dental plan for reimbursement information. The MCOs and dental plans are not required to follow the Texas Medicaid fee schedules, so there may be some differences in reimbursement based on decisions made by the individual health and dental plans.

When services or products do not have an established reimbursement amount, the detail or claim is manually reviewed to determine an appropriate reimbursement.

Texas Medicaid (FFS and MCO) providers can bill and receive reimbursement for the unused portion of weight-based or variable dosing CADs that are only manufactured in single-dose vials. A multi-dose vial is a vial of liquid medication that is intended for parenteral administration (injection or infusion), contains more than one dose of medication, and may be used for more than one patient preparation or administration. Multi-dose vials are excluded from reimbursement under Texas Medicaid for any unused or discarded portions.

Claims will only be considered for reimbursement if an HHSC review has determined that the medication has a weight-based, variable dosing schedule or that it requires dosing adjustments for pharmacokinetic or pharmacodynamic considerations. The administration of the medication for the recommended dosing must result in a patient dose portion plus a discarded portion of a drug vial. The provider may be eligible for reimbursement up to the amount of drug or biological in accordance with the drug label.

Claims submitted for the unused portion and discarded portion of weight-based or variable dosing clinician-administered drugs (CADs) manufactured only in single-dose vials must include the modifier JW for consideration of reimbursement.
This only applies to medical claims for weight-based or variable dosing CADs billed with Healthcare Common Procedure Coding System (HCPCS) procedure codes, manufactured only in single-dose vials, and provided in a professional or outpatient setting.

Providers must use the modifier JW to identify the unused portion of the vial contents and the discarded amount of the drug or biological. Medicaid and CHIP providers must bill the JW on a separate line.

**Example:** A single use vial labeled to contain 100 units of a drug has 95 units administered to the client and 5 units discarded. The 95-unit dose is billed on one line, while the discarded 5 units are billed on another line by using the JW modifier. Both line items are reviewed for reimbursement. The provider must record the discarded amounts of drugs and biologicals in the client’s medical record.

**Refer to:** Subsection 8.1, “JW Modifier Claims Filing Instructions” in this handbook for further instructions about the JW modifier.

### 5 Injectable Medications as a Pharmacy Benefit

Some injectable drugs or biologicals are available by prescription and are reimbursable as a pharmacy benefit through the Vendor Drug Program (VDP) under Texas Medicaid.

**Refer to:** Subsection 9, “Pharmacy Benefit” in this handbook for more information.

Oral medications that are given in the hospital or physician’s office are considered part of the hospital or office visit and cannot be reimbursed separately. Take-home and self-administered drugs may be a pharmacy benefit when they are provided to eligible Texas Medicaid fee-for-service clients through VDP with a valid prescription.

Providers may utilize the “white-bagging” delivery method, in which the treating provider submits prescriptions to pharmacies and the prescription is shipped or mailed to the provider’s office.

**Refer to:** Subsection 10.5.1, “Pharmacy Delivery Method for Clinician-Administered Drugs” in this handbook for additional information on the “white-bagging” delivery method.

Providers must use oral medication in preference to injectable medication in the office and outpatient hospital. If an oral medication cannot be used, the KX modifier must be submitted on the claim. The following situations are acceptable reasons for the use of administering an injectable medication instead of administering an oral medication.

<table>
<thead>
<tr>
<th>Claim Form</th>
<th>Reason for Injection</th>
</tr>
</thead>
<tbody>
<tr>
<td>Modifier KX</td>
<td>• No acceptable oral equivalent is available.</td>
</tr>
<tr>
<td></td>
<td>• Injectable medication is the standard treatment of choice.</td>
</tr>
<tr>
<td></td>
<td>• The oral route is contraindicated.</td>
</tr>
<tr>
<td></td>
<td>• The client has a temperature over 102 degrees Fahrenheit (documented on the claim and in the medical record) and a high blood level of antibiotic is needed quickly.</td>
</tr>
<tr>
<td></td>
<td>• The client has demonstrated noncompliance with orally prescribed medication (must be documented on the claim and in the medical record).</td>
</tr>
<tr>
<td></td>
<td>• Previously attempted oral medication regimens have proven ineffective (must be supported by documentation in the medical record).</td>
</tr>
<tr>
<td></td>
<td>• Situation is emergent.</td>
</tr>
</tbody>
</table>
The claim and the client’s medical record must include documentation of medical necessity to support the need for the service. Retrospective review may be performed to ensure that the documentation supports the medical necessity of the service and any modifier used when billing the claim.

6 National Drug Code (NDC)

The NDC is an 11-digit number on the package or container from which the medication is administered. Some packages may display less than 11 digits. In those cases, leading zeros can be assumed and are required for billing. For example, 5678-0123-01 becomes 05678-0123-01. In another example, 78513-677-2 becomes 78513-0677-02.

Note: The NDCs in the examples show hyphens between the segments for easier visualization. NDCs submitted on claims should not include hyphens or spaces between the segments.

Refer to: Subsection 6.3.4, “National Drug Code (NDC)” in “Section 6: Claims Filing” (Vol. 1, General Information) for more information on NDC requirements as well as drug rebates.

6.1 Calculating Billable HCPCS and NDC Units

All drug claims must include HCPCS billing units as well as NDC billing units. HCPCS billing units are calculated by dividing the amount administered by the units found in the procedure code description. The calculated HCPCS billing unit is also needed to determine the correct NDC billing unit. NDC billing units are calculated by multiplying the HCPCS billing unit by the conversion factor. The conversion factor is calculated by dividing the HCPCS unit (found in the code description) by the NDC unit (found on the box or packaging). See calculation examples in the following sections. Conversion factors are already computed and included on the Texas NDC-to-HCPCS Crosswalk. The NDC billing unit also requires a unit of measurement. For example, if the NDC is for a liquid medication the submitted units must be in milliliters (ML). If the NDC is for a powder form then the submitted units are Unit (UN). Other allowable NDC units are GR for gram, F2 for international unit, and ME for milligram. For all claims, the HCPCS and NDC billing units are required, along with the specific NDC and HCPCS procedure code. Claims submitted with incorrect unit calculations may cause delayed or incorrect payment.

6.1.1 Single-Dose Vials Calculation Examples

Below are three examples of how to calculate the HCPCS and NDC billing units using single-dose vials.

1) A patient receives 4 mg Zofran IV in the physician’s office. The NDC of the product used is 00173-0442-02 (Zofran 2 mg/ml in solution form). There are 2 milliliters per vial. The provider should bill J2405 for ondansetron hydrochloride with 4 HCPCS units and the NDC units submitted should be 2 ML.

2) A patient receives 8mg of Avastin IV in the physician’s office. The NDC of the product used is 50242-060-01 (Avastin 25mg/ml). The provider should bill J9035 for bevacizumab with 0.8 HCPCS unit. The NDC unit is 0.32 ML.

3) A patient receives 1 gm Rocephin IM in the physician’s office. The NDC of the product used is 00004-1963-02 (Rocephin 500 mg vial in a powder form that is reconstituted prior to the injection). The provider should bill J0696 for ceftriaxone sodium with 4 HCPCS units. The NDC units are 2 UN because this NDC is in powder form.

<table>
<thead>
<tr>
<th></th>
<th>Zofran</th>
<th>Avastin</th>
<th>Rocephin</th>
</tr>
</thead>
<tbody>
<tr>
<td>Dose Administered to Patient</td>
<td>4 mg</td>
<td>8mg</td>
<td>1gm = 1000mg</td>
</tr>
<tr>
<td>HCPCS Code and Unit found in description</td>
<td>J2405 Per 1 mg</td>
<td>J9035 Per 10 mg</td>
<td>J0696 Per 250 mg</td>
</tr>
</tbody>
</table>
6.1.2 Multi-Dose Vials Calculation Examples

Below is an example of calculating the correct billing units for a drug administered from a multi-dose vial. Calculations for multi-dose vials differ from those for single-dose vials.

A patient receives 8 mg Dexamethasone in the physician’s office. A 20 mg multi-dose vial is used. The NDC of the product used is 63323-0165-05 (Dexamethasone 20 MG/5 ml Vial). The provider should bill J1100 for dexamethasone with 8 HCPCS units and the NDC units submitted should be 2 ML. There are 12 mg (3 ml) remaining in the vial.

<table>
<thead>
<tr>
<th>Multi-Dose Calculation Examples for Dexamethasone</th>
<th>Zofran</th>
<th>Avastin</th>
<th>Rocephin</th>
</tr>
</thead>
<tbody>
<tr>
<td>Dose Administered to Patient</td>
<td>8 mg</td>
<td></td>
<td></td>
</tr>
<tr>
<td>HCPCS Code and Given Unit</td>
<td>J1100 Per 8 mg</td>
<td></td>
<td></td>
</tr>
<tr>
<td>HCPCS CODE BILLING UNIT(s) = Dose divided by units found in HCPCS code</td>
<td>8mg/1mg = 8</td>
<td></td>
<td></td>
</tr>
<tr>
<td>NDC Information on Vial/Box</td>
<td>20mg/5ml = 4mg/1ml</td>
<td></td>
<td></td>
</tr>
<tr>
<td>NDC BILLING UNIT(s) = Dose divided by NDC unit from vial/box</td>
<td>8/4 = 2</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Quantity Information Required on Claim (HCPCS and NDC)</td>
<td>8 and 2 ML</td>
<td>0.8 and 0.32 ML</td>
<td>4 and 2 UN</td>
</tr>
</tbody>
</table>

6.1.3 Single and Multi-Use Vials

A single-dose (or single-use) vial of medication intended for administration through injection or infusion contains a single dose of medication. A multi-dose (or multi-use) vial of medication intended for administration through injection or infusion contains more than one dose of medication.

Many drugs have recommended doses that are based on factors such as height, weight, and initial tolerance for the drug. It is important to clearly document how the dosage is calculated so those who review the patient health record can verify the dosage amount when reviewing the claim.

Other resources on clinician-administered drugs may be found online by visiting the TMHP, CDC and CMS websites.
6.1.4 Nonspecific, Unlisted, or Miscellaneous Procedure Codes

Drugs or biologicals that do not have a unique CPT or HCPCS procedure code must be billed using a nonspecific, unlisted, unclassified, or miscellaneous procedure code. All claims for nonspecific, unlisted, unclassified, or miscellaneous procedure codes are processed manually and must be submitted on paper with accompanying documentation. The billing provider must include the following required documentation:

- The name and NDC number of the drug administered.
- The quantity of the drug administered, the amount discarded (if applicable for reimbursement), and the units of measurement.
- A brief description of the recipient’s condition(s) that supports the medical need for the drug.
- One of the following pricing information sources:
  - The manufacturer’s average wholesale price (AWP)
  - A copy of the invoice for the drug

The claim and attached information will suspend for manual review to determine whether the drug is clinically appropriate based on the information provided and to price the claim using the information provided. Miscellaneous drug or biological procedure codes are reimbursed a percentage of the average wholesale price (AWP). HHSC reserves the option to use other data sources to determine Texas Medicaid fees for drugs when AWP calculations are determined to be unreasonable or insufficient.

The claim will be denied when:

- The information is not sufficient to determine medical necessity.
- The pricing information is insufficient for pricing the claim.
- There is a more appropriate billing procedure code for the drug or biological.
- The NDC and HCPCs (if applicable) codes are missing.

Providers are responsible for administering drugs based on the U.S. Food and Drug Administration (FDA)-approved guidelines. In the absence of FDA indications, a drug needs to meet the following criteria:

- The drug is recognized by the American Medical Association Drug Evaluations (AMA-DE), American Hospital Formulary Service Drug Information, the U.S. Pharmacopoeia Dispensing Information, Volume I, or two articles from major peer-reviewed journals that have validated and uncontested data supporting the proposed use for the specific medical condition as safe and effective.
- It is medically necessary to treat the specific medical condition, including life-threatening conditions or chronic and seriously debilitating conditions.
- The off-label use of the drug is not investigational or experimental.

Retrospective review may be performed to ensure documentation supports the medical necessity of the service.

Some injectable medications require prior authorization, which is a condition for reimbursement; it is not a guarantee of payment. To avoid unnecessary denials, the physician must provide correct and complete information, including documentation for medical necessity for the service requested. The
physician must maintain documentation of medical necessity in the client’s medical record. Providers may fax or mail prior authorization requests, including all required documentation, to the TMHP Special Medical Prior Authorization Department at:

Texas Medicaid & Healthcare Partnership
Special Medical Prior Authorization Department
12357-B Riata Trace Parkway, Suite 100
Austin, TX 78727
Fax: 1-512-514-4213

The following injections in the table below are benefits of Texas Medicaid but are subject to the indicated limitations. Those with an asterisk have more information and can be found listed after the table.

7 Outpatient Drugs—Benefits and Limitations

The clinician administered drugs identified throughout this handbook are benefits of Texas Medicaid, but are subject to the indicated limitations.

Clinician administered drugs may be considered for reimbursement in the home, office, or outpatient hospital settings. Certain procedure codes may be restricted for place of service and provider types. Providers should verify the restrictions for specific procedure codes of injectable or infused medications prior to rendering services.

7.1 Abatacept (Orencia)

Abatacept is a synthetic protein produced by recombinant deoxyribonucleic acid (DNA) technology that is used for treating rheumatoid arthritis. Abatacept slows the damage to bones and cartilage and relieves the symptoms and signs of arthritis. Abatacept (procedure code J0129) is a benefit of Texas Medicaid for clients who have moderately to severely active rheumatoid arthritis. These clients may also have an inadequate response to one or more non-biological, disease modifying antirheumatic drugs (DMARDs).

7.1.1 Prior Authorization for Abatacept (Orencia)

Prior authorization may be given for an initial six months for eight doses. Prior authorization for an initial request for abatacept injections will be considered when all of the following criteria are met:

- Dates of treatment
- The number of anticipated doses
- The dosage to be administered
- Diagnosis of adult RA or juvenile idiopathic arthritis (JIA)

Note: A diagnosis of adult RA must conform to the American College of Rheumatology (ACR) RA classification that requires the following:

- Presence of synovitis in at least one joint
- Absence of an alternative diagnosis to explain the synovitis
- A combined score of at least six out of ten on the level of involved joints, abnormality, and symptom duration from the individual scores in four domains:
  - The number and sites of involved joints
  - Serologic abnormality
  - Elevated acute-phase response
• Symptom duration

Prior authorization for an initial request for abatacept injections may be granted for six months for eight doses. Prior authorization will be considered when the client has an inadequate response after 12 weeks to a nonbiological DMARD such as methotrexate or sulfasalazine or one or more biological (injectable) DMARDs, such as adalimumab, etanercept, or tumor necrosis factor (TNF) antagonists. The inadequate response must be indicated by all of the following commonly used prognostic factors:

• Visual Analogue scale (VAS) (4 or greater on a pain scale from 0-10)
• Global Arthritis Score (GAS) (3 or greater with remission defined as less than 3)
• Health Assessment Questionnaire Disability Index (HAQDI) score (greater than 1)
• Evidence of radiographic erosions
• Elevated erythrocyte sedimentation rate (greater than 20 millimeters/hour)
• Elevated C-reactive protein level (greater than zero milligrams/deciliter)
• Elevated rheumatoid factor (RF) level (greater than 60 units/millimeter or a titer greater than 1:80 titer)
• Elevated anti-cyclic citrullinated peptide (anti-CCP) antibody level (20 units/millimeter or greater)

Prior authorization for subsequent dosing may be given for a maximum of six doses when documentation supports medical necessity for continued treatment with abatacept. Prior authorization for a subsequent request must include all of the following:

• Documentation from the physician stating that there has been at least a 20-percent improvement as defined by the ACR
• The number of anticipated doses
• The dosage to be administered

The documentation of medical necessity must be maintained by the requesting provider in the client's medical record and is subject to retrospective review.

7.2 Adalimumab

Procedure code J0135 is a benefit when billed with one of the following diagnosis codes:

<table>
<thead>
<tr>
<th>Diagnosis Codes</th>
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<tbody>
<tr>
<td>K5000 K50011 K50012 K50013 K50014 K50018 K5010 K50111</td>
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<td>M05011 M05012 M05019 M05021 M05022 M05029 M05031 M05032</td>
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7.3 Ado-trastuzumab entansine (Kadcyla)

Ado-trastuzumab entansine (Kadcyla), procedure code J9354, is a benefit of Texas Medicaid for clients of any age when all of the following indications are present:

- Individuals have a diagnosis of HER2 positive metastatic breast cancer
- Individuals have previously received trastuzumab and a taxane separately or in combination, and have either:
  - Received prior therapy for metastatic disease
  - Experienced disease reoccurrence during or within six months of completing adjuvant therapy

Documentation must be maintained by the treating physician in the client’s medical record to support administration of Ado-trastuzumab entansine (Kadcyla). Prior authorization is not required for ado-trastuzumab entansine (Kadcyla).
At initiation of treatment, documentation must include all of the following:

- Evidence of HER2 positive breast cancer as evidenced by immunochemistry (IHC) test or fluorescent in situ hybridization (FISH) test
- Evidence of metastatic breast cancer
- Evidence demonstrating prior treatment for this diagnosis with trastuzumab and a taxane oncology agent separately or in combination
- Evidence demonstrating receipt of prior therapy for this diagnosis or recurrent disease, including the previous treatment protocol, within six months of completing adjuvant therapy.

### 7.4 Alglucosidase Alfa (Myozyme)

Alpha-glucosidase, a recombinant human enzyme alpha-glucosidase (rhGAA), is an essential enzyme for normal muscle development and function. Alglucosidase alfa may be a benefit of Texas Medicaid for clients of any age who are diagnosed with glycogen storage disease Type II (GSD Type II, also known as Pompe disease), using procedure codes J0220 and J0221. The most appropriate diagnosis code must be indicated on the prior authorization request and on the claim.

Prior authorization is required for alglucosidase alfa and documentation must include all of the following:

- A request for alglucosidase alfa.
- Laboratory evidence of acid alpha-glucosidase (GAA) deficiency, (i.e., below the laboratory-defined cut-off value as determined by the laboratory performing the GAA enzyme activity assay). Tissues used for determination of GAA deficiency may include blood, muscle, or skin fibroblasts.

The physician must maintain supporting documentation in the client’s medical record.

### 7.5 Amifostine

Amifostine is a benefit of Texas Medicaid for the reduction of the cumulative renal toxicity associated with administration of cisplatin in clients who have advanced ovarian cancer or non-small cell lung cancer with documentation of a creatinine clearance of 50 or less and where no other chemotherapeutic agent can be used.

Amifostine may also be used to reduce the incidence of moderate-to-severe xerostomia in clients undergoing postoperative radiation treatment for head and neck cancers where the radiation port includes a substantial portion of the parotid glands.

Amifostine may be reimbursed for the following indications:

- Bone marrow toxicity
- Cisplatin- and cyclophosphamide-induced (prophylaxis)
- Advanced solid tumors
- Head and neck carcinoma
- Malignant lymphoma
- Non-small cell lung cancer
- Myelodysplastic syndromes
- Nephrotoxicity
- Advanced ovarian carcinoma
- Melanoma
• Advanced solid tumors of non-germ cell origin
• Neurotoxicity
• Reduction in the incidence of mucositis in clients receiving radiation therapy, or radiation combined with chemotherapy
• Reduction in the incidence of xerostomia associated with postoperative radiation treatment of head and neck cancer, where the radiation port includes a substantial portion of the parotid glands

Providers must use procedure code J0207 with one of the following diagnosis codes:

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</table>
7.6 Antibiotics and Steroids
Injectable antibiotic or steroid medications may be considered for reimbursement even if the same oral medications are appropriate and available. Injected antibiotics or steroid medications, when used in place of oral medications, require the use of the modifier KX.

Physicians billing for injectable antibiotic and steroid medications must indicate the appropriate modifiers with the appropriate injection code and quantity:

<table>
<thead>
<tr>
<th>Modifier</th>
<th>Use</th>
</tr>
</thead>
<tbody>
<tr>
<td>AT</td>
<td>For acute conditions*</td>
</tr>
<tr>
<td>KX</td>
<td>To indicate any of the following:</td>
</tr>
<tr>
<td></td>
<td>• Oral route contraindicated or an acceptable oral equivalent is not available.</td>
</tr>
<tr>
<td></td>
<td>• Injectable medication is the accepted treatment of choice. Oral medication regimen has proven ineffective or is not applicable.</td>
</tr>
<tr>
<td></td>
<td>• The patient has a temperature over 102 degrees and a high level of antibiotic is needed immediately.</td>
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<tr>
<td></td>
<td>• Injection is medically necessary into joints, bursae, tendon sheaths, or trigger points to treat an acute condition or the acute flare-up of a chronic condition.</td>
</tr>
</tbody>
</table>

If a steroid medication is injected into joints, bursae, tendon sheaths, or trigger points, modifier AT must be used to indicate an acute condition. When performed for a chronic condition, these procedures are denied.

7.7 Antisense Oligonucleotides (eteplirsen, golodirsen, and nusinersen)
Antisense oligonucleotides, eteplirsen (Exondys 51) (procedure code J1428), golodirsen (Vyondys 53) (procedure code J1429), nusinersen (Spinraza) (procedure code J2326), or viltolarsen (Viltepso) (procedure code C9071) may be benefits of Texas Medicaid with prior authorization.

An antisense oligonucleotide is a synthetic single stranded nucleic acid that binds to RNA and thereby alters or reduces expression of the target RNA. This may result in an improvement in physical function.

7.7.1 Prior Authorization Requirements
Prior authorization requests for procedure codes J1428, J1429, J2326, and C9071 must be submitted by the prescribing provider to the Special Medical Prior Authorization (SMPA) department at TMHP using the Special Medical Prior Authorization (SMPA) Request Form.

Prior authorization is not required for physician services associated with the administration of eteplirsen, golodirsen, nusinersen, or viltolarsen. Physician services include the procedural costs and the associated supplies for the administration of the medication.

For situations in which procedure code J1428, J1429, J2326, or C9071 are being dispensed by a pharmacy via white bagging, the prescribing provider must provide the dispensing durable medical equipment (DME) pharmacy the authorization approval number.

The dispensing DME pharmacy may not request prior authorization.

The DME pharmacy provider billing for nusinersen (Spinraza) (procedure code J2326) will be responsible for coordinating with the rendering provider to obtain the prior authorization request approval number.
The requesting provider (physician or hospital) may coordinate with the DME Pharmacy provider for the initial or recertification prior authorization request for the specific oligonucleotide. DME Pharmacy providers may assist in providing necessary information such as their National Provider Identifier (NPI) number, fax number, and business address to the requesting provider. However, the Special Medical Prior Authorization (SMPA) form must be signed and dated and submitted by the Medicaid-enrolled requesting provider, not the DME Pharmacy provider.

The dispensing pharmacy must submit the authorization approval number when billing for the drug. Reimbursement for dispensing of the drug by the pharmacy may not occur unless an approved prior authorization for the specific oligonucleotide is in place.

**Note:** For additional information on white bag delivery, providers may refer to Subsection 10.5.1, “Pharmacy Delivery Method for Clinician-Administered Drugs” in this handbook.

A neurologist’s consultation must be dated no more than six months prior to an initial request and no more than one rolling year prior to a recertification or extension request. The consultation must include the neurologist’s name, credentials, contact information, and a recommendation for treatment with the specific antisense oligonucleotide.

Documentation of the client’s dosage, administration schedule, the number of injections to be administered during the prior authorization period, the requested units per injection, and the dosage calculation must be submitted in Section C of the SMPA Request Form under Statement of Medical Necessity. When the FDA approves dosing guidelines that require a weight based calculation, the client’s current weight must be included.

**Refer to:** Subsection 3.1, “Prior Authorization Requests” in this handbook for additional prior authorization information.

### 7.7.1.1 Initial Requests (for all Antisense Oligonucleotides)

Initial prior authorization requests for all antisense oligonucleotides will be considered by Medical Director Review for a six month period. The initial request must include documentation supporting medical necessity for the requested antisense oligonucleotide in addition to the SMPA request form completed, signed, and dated by the prescribing provider.

Documentation supporting medical necessity for an initial prior authorization for all requested antisense oligonucleotides must include the following information:

- A diagnosis specific to the requested antisense oligonucleotide
- Genetic testing specific to the requested antisense oligonucleotide
- Client age specific to the requested antisense oligonucleotide
- Documentation of baseline physical function. Testing tools used to measure physical function must be age appropriate for the client being tested.
- A neurologist’s consultation dated no more than six months prior to the initially requested authorization start date. The consultation must include the neurologist’s name, credentials, contact information, and a recommendation for treatment with the requested antisense oligonucleotide.
- Documentation of the requested antisense oligonucleotide dosage and administration schedule, including the number of injections to be administered during the prior authorization period, the requested units per injection, and the dosage calculation. This information must be submitted in Section C of the SMPA request form under Statement of Medical Necessity.

Each antisense oligonucleotide has specific clinical indications and unique documentation requirements.
The initial request for eteplirsen (Exondys 51) must include the following documentation to support medical necessity for eteplirsen:

- Genetic testing must confirm that the client’s DMD gene is amenable to exon 51 skipping.
- Client age
- Current client weight, including the date the weight was obtained; the weight must be dated no more than 30 days before the request date.

Testing tools that can be used to demonstrate physical function include, but are not limited to:

- Brooke Upper Extremity Scale
- Baseline 6-minute walk test (6MWT)
- Pediatric Evaluation of Disability Inventory

Exondys 51 should not be used concomitantly with other exon skipping therapies for DMD.

The initial request for golodirsen (Vyondys 53) must include the following documentation to support medical necessity for golodirsen:

- Genetic testing must confirm that the client’s Duchenne muscular dystrophy (DMD) gene is amenable to exon 53 skipping.
- Baseline renal function test (i.e. Glomerulus Filtration Rate, GFR) with therapy initiation and continuation.
- Current client weight, including the date the weight was obtained; the weight must be dated no more than 30 days before the request date.
- Baseline function testing documented in patient chart or electronic health record.

Testing tools that can be used to demonstrate physical function include, but are not limited to:

- Brooke Upper Extremity Scale
- Baseline 6MWT (6-minute walk test)
- Pediatric Evaluation of Disability Inventory

Vyondys 53 should not be used concomitantly with other exon skipping therapies for DMD.

The initial request for nusinersen (Spinraza) must include the following documentation to support medical necessity for nusinersen:

- Genetic testing must confirm biallelic pathogenic variants in the client’s survival motor neuron 1 (SMN1) gene
- Client age
- Baseline pulmonary status, including any requirements for invasive or non-invasive ventilation

Testing tools that can be used to demonstrate physical function include, but are not limited to:

- The Hammersmith Infant Neurological Exam (HINE).
- The Hammersmith Functional Motor Scale Expanded (HFMSE).
- The Upper Limb Module (UML).
- Baseline 6MWT.
- Children’s Hospital of Philadelphia Infant Test of Neuromuscular Disorders (CHOP-INTEND).

The initial request for viltolarsen (Viltepso) must include the following documentation:
• Genetic testing must confirm that the client’s Duchenne muscular dystrophy (DMD) gene is amenable to exon 53 skipping.

• Current client weight, including the date the weight was obtained; the weight must be dated no more than 30 days before the request date.

• Baseline renal function test (i.e. Glomerulus Filtration Rate) and urine protein-to-creatinine ratio should be measured before starting treatment.

Testing tools that can be used to demonstrate physical function include, but are not limited to:

• Brooke Upper Extremity Scale
• Baseline 6MWT (6-minute walk test)
• Pediatric Evaluation of Disability Inventory
• Viltepso should not be used concomitantly with other exon skipping therapies for DMD.

7.7.1.2 Recertification/Extension Requests (for all Antisense Oligonucleotides)

Recertification/extension prior authorization requests for antisense oligonucleotides will be considered by Medical Director Review for additional six month periods. The recertification/extension request must include documentation supporting the ongoing medical necessity for the requested antisense oligonucleotide in addition to a new SMPA request form completed, signed, and dated by the prescribing provider.

A complete recertification/extension request must be received no earlier than 30 days before the current authorization period expires. Requests for recertification/extension of prior authorization received after the current prior authorization expires will be denied for dates of service that occurred before the date the request is received.

Documentation supporting a recertification/extension prior authorization for all requested antisense oligonucleotide must include the following:

• A diagnosis specific to the requested antisense oligonucleotide

• Client age

• Current documentation of physical function

• Testing tools used to measure physical function must be age appropriate for the client being tested. Providers must use the same testing instrument as used in the initial evaluation. If re-use of the initial testing instrument is not appropriate, for example, due to change in client status or restricted age range of the testing tool, the provider must explain the reason for the change.

• The physical function testing tool results must include one of the following:
  • An increase in physical function from baseline has been observed
  • Baseline physical function has been maintained

• A neurology consultation dated no more than one rolling year of the recertification/extension date that includes the name, credentials, and contact information for the consulting neurologist recommending ongoing treatment with the requested antisense oligonucleotide

• Statement from prescribing clinician that the client has been compliant with the treatment

• Documentation of the requested antisense oligonucleotide dosage and administration schedule, including the number of injections to be administered during the prior authorization period, the requested units per injection, and the dosage calculation must be submitted in Section C of the Special Medical Prior Authorization (SMPA) Request Form under Statement of Medical Necessity.
The medical necessity documentation for eteplirsen (Exondys 51) recertification/extension requests must include the client’s current weight the date on which the weight was obtained. The weight must be dated no more than 30 days before the request date.

The medical necessity documentation for golodirsen (Vyondys 53) and viltolarsen (Viltepso) recertification/extension requests must include the client’s continual renal function test while on therapy and current client weight, including the date the weight was obtained. The weight must be dated no more than 30 days before the request date.

The medical necessity documentation for nusinersen (Spinraza) recertification/extension requests must include the client’s pulmonary status, including any requirements for invasive or non-invasive ventilation. Any changes in pulmonary status that have occurred since the previous prior authorization request must be addressed.

### 7.7.1.3 Exclusions

Eteplirsen (Exondys 51™), Golodirsen (Vyondys 53™), nusinersen (Spinraza™), and viltolarsen (Viltepso™) should not be continued on clients who experience decreasing physical function while on the medication.

Eteplirsen (Exondys 51™) and Golodirsen (Vyondys 53™) should not be used concomitantly or with other exon skipping therapies for DMD.

Nusinersen (Spinraza™) is not a continuing benefit for clients with decreasing pulmonary function while on the medication.

### 7.8 Aripiprazole Lauroxil, (Aristada Initio)

Aripiprazole lauroxil (procedure codes J1943 and J1944) are benefits of Texas Medicaid for clients who are 18 years of age or older.

### 7.9 Azacitidine (Vidaza)

Procedure code J9025 is a benefit when billed with one of the following diagnosis codes:

| Diagnosis Codes | C9202 | C9210 | C9212 | C9220 | C9222 | C9232 | C9242 | C9252 | C9262 | C9290 | C9292 | C92A2 | C92Z2 | C9310 | C9312 | C9330 | C9332 | C9502 | C9510 | C9512 | C9592 | D460 | D461 | D46B | D46C | D46Z | D640 | D641 | D642 | D643 |
|-----------------|-------|-------|-------|-------|-------|-------|-------|-------|-------|-------|-------|-------|-------|-------|-------|-------|-------|-------|-------|-------|-------|-------|-------|-------|-------|-------|-------|-------|-------|-------|-------|-------|-------|

### 7.10 Botulinum Toxin Type A and Type B

OnabotulinumtoxinA (Botox brand of botulinum toxin type A), abobotulinumtoxinA (Dysport brand of botulinum toxin type A), incobotulinumtoxin A (Xeomin brand of botulinum toxin type A), and rimabotulinumtoxinB (Myobloc brand of botulinum toxin type B) are benefits of Texas Medicaid.

Botulinum toxins are potent neuromuscular blocking agents that are useful in treating various focal muscle spastic disorders and excessive muscle contractions, such as dystonias, spasms, and twitches. They produce a presynaptic neuromuscular blockade by preventing the release of acetylcholine from the nerve endings. Since the resulting chemical denervation of muscle produces local paresis or paralysis, selected muscles can be treated. Two of the seven naturally occurring serotypes of botulinum toxin have been approved by the FDA for human use in the United States-type A and type B.
Due to the unique manufacturing process of each toxin, botulinum toxins are chemically, clinically, and pharmacologically distinct; as a consequence, these products are not interchangeable. The units of biological activity of one botulinum toxin product cannot be compared to, nor converted into, units of any other botulinum toxin product. The established drug names of the botulinum products emphasize the differing dose-to-potency ratios of these products.

Procedure code J0585 is a benefit when billed with one of the following diagnosis codes:

<table>
<thead>
<tr>
<th>Diagnosis Codes</th>
</tr>
</thead>
<tbody>
<tr>
<td>G114 G2401 G241 G243 G244 G245 G248 G250</td>
</tr>
<tr>
<td>G251 G252 G253 G35 G360 G370 G371 G372</td>
</tr>
<tr>
<td>G374 G375 G378 G379 G43701 G43709 G43711 G43719</td>
</tr>
<tr>
<td>G800 G801 G802 G803 G804 G808 G809 G8110</td>
</tr>
<tr>
<td>G8111 G8112 G8113 G8114 G8220 G8221 G8222 G8250</td>
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<tr>
<td>G8251 G8252 G8253 G8254 G830 G8310 G8311 G8312</td>
</tr>
<tr>
<td>G8313 G8314 G8320 G8321 G8322 G8323 G8324 G8330</td>
</tr>
<tr>
<td>G8331 G8332 G8333 G8334 G834 H4901 H4902 H4903</td>
</tr>
<tr>
<td>H4911 H4912 H4913 H4921 H4922 H4923 H4931 H4932</td>
</tr>
<tr>
<td>H4933 H4941 H4942 H4943 H499 H5000 H50011 H50012</td>
</tr>
<tr>
<td>H50021 H50022 H50031 H50032 H50041 H50042 H5005 H5006</td>
</tr>
<tr>
<td>H5007 H5008 H5010 H5011 H50112 H50121 H50122 H50131</td>
</tr>
<tr>
<td>H50132 H50141 H50142 H5015 H5016 H5017 H5018 H5021</td>
</tr>
<tr>
<td>H5016 H5017 H5018 H5021 H5022 H5030 H50311 H50312</td>
</tr>
<tr>
<td>H5032 H50331 H50332 H5034 H5040 H50411 H50412 H5042</td>
</tr>
<tr>
<td>H5043 H5050 H5051 H5052 H5053 H5054 H5055 H5060</td>
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<td>H50611 H50612 H5069 H50811 H50812 H5089 H510 H5111</td>
</tr>
<tr>
<td>H5112 H5121 H5122 H5123 H518 H519 I6903 I69032</td>
</tr>
<tr>
<td>I69033 I69034 I69041 I69042 I69043 I69044 I69051 I69052</td>
</tr>
<tr>
<td>I69053 I69054 I69061 I69062 I69063 I69064 I69065 I69098</td>
</tr>
<tr>
<td>I69131 I69132 I69133 I69134 I69141 I69142 I69143 I69144</td>
</tr>
<tr>
<td>I69151 I69152 I69153 I69154 I69161 I69162 I69163 I69164</td>
</tr>
<tr>
<td>I69165 I69198 I69231 I69232 I69233 I69234 I69241 I69242</td>
</tr>
<tr>
<td>I69243 I69244 I69251 I69252 I69253 I69254 I69261 I69262</td>
</tr>
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<td>I69263 I69264 I69265 I69298 I69331 I69332 I69333 I69334</td>
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</tr>
<tr>
<td>I69361 I69362 I69363 I69364 I69365 I69398 I69831 I69832</td>
</tr>
<tr>
<td>I69833 I69834 I69841 I69842 I69843 I69844 I69851 I69852</td>
</tr>
<tr>
<td>I69853 I69854 I69861 I69862 I69863 I69864 I69865 I69898</td>
</tr>
<tr>
<td>J385 K117 K220 K600 K601 K602 M436 M62838</td>
</tr>
<tr>
<td>M722 N318 N3281 N3644 R490 R498</td>
</tr>
</tbody>
</table>

Procedure code J0586 is a benefit when billed with one of the following diagnosis codes:

<table>
<thead>
<tr>
<th>Diagnosis Codes for J0586</th>
</tr>
</thead>
<tbody>
<tr>
<td>G114 G241 G243 G244 G245 G248 G35 G360</td>
</tr>
</tbody>
</table>
Procedure code J0587 is a benefit when billed with diagnosis code G243 or K117.

Procedure code J0588 is a benefit when billed with one of the following diagnosis codes:

<table>
<thead>
<tr>
<th>Diagnosis Codes for J0588</th>
</tr>
</thead>
<tbody>
<tr>
<td>G243 G245 G800 G801 G802 G830 G8110 G8111</td>
</tr>
<tr>
<td>G8112 G8113 G8114 G8253 G8254 G8320 G8321 G8322</td>
</tr>
<tr>
<td>G8323 G8324 I69059 I69259 I69359 I69859 I69959 I69051</td>
</tr>
<tr>
<td>I69052 I69151 I69152 I69251 I69252 I69351 I69352 I69851</td>
</tr>
<tr>
<td>I69852 I69951 I69952 I69053 I69054 I69153 I69154 I69253</td>
</tr>
<tr>
<td>I69254 I69353 I69354 I69853 I69854 I69953 I69954 I69039</td>
</tr>
<tr>
<td>I69139 I69239 I69339 I69839 I69939 I69031 I69032 I69131</td>
</tr>
<tr>
<td>I69932 I69033 I69034 I69133 I69134 I69233 I69234 I69333</td>
</tr>
<tr>
<td>I69334 I69833 I69834 I69933 I69934 J385 M436 M62838</td>
</tr>
</tbody>
</table>

Procedure codes J0588, J0586, and J0587 are denied when billed on the same date of service by any provider as procedure code J0585. Procedure codes J0588 and J0587 are denied when billed on the same date of service by any provider as procedure code J0586. Procedure code J0587 is denied when billed on the same date of service by any provider as procedure code J0588.

IncobotulinumtoxinA, procedure code J0588, is FDA-approved for the treatment of adults with blepharospasm previously treated with onabotulinumtoxinA (J0585).

Physicians, hospitals, and other providers and suppliers should care for and administer drugs to patients in such a way that they can use drugs or biologicals most efficiently, in a clinically appropriate manner. Texas Medicaid encourages scheduling patients to make the most efficient use of the drugs administered. Safe handling guidelines per manufacturer must be observed (e.g., shelf life, cold chain requirements). The smallest size vial to cover the dose is encouraged to be used.
Claims for botulinum toxin type A and B must indicate the number of units used. If the number of units is not specified, the claim will be paid a quantity of one. Claims that exceed the following quantity limitations, per day, may be considered on appeal with documentation of medical necessity:

<table>
<thead>
<tr>
<th>Procedure Codes</th>
<th>Quantity Limitations of Medication</th>
<th>Billing Units</th>
</tr>
</thead>
<tbody>
<tr>
<td>J0585</td>
<td>400 units</td>
<td>One billing unit is equal to 1 unit of medication. <strong>Example:</strong> A provider that administers 400 units of medication would submit a claim for a quantity of 400.</td>
</tr>
<tr>
<td>J0586</td>
<td>1,500 units</td>
<td>One billing unit is equal to 5 units of medication. <strong>Example:</strong> A provider that administers 1,500 units of medication would submit a claim for a quantity of 300.</td>
</tr>
<tr>
<td>J0587</td>
<td>10,000 units</td>
<td>One billing unit is equal to 100 units of medication. <strong>Example:</strong> A provider that administers 10,000 units of medication would submit a claim for a quantity of 100.</td>
</tr>
<tr>
<td>J0588</td>
<td>400 units</td>
<td>One billing unit is equal to 1 unit of medication. <strong>Example:</strong> A provider that administers 400 units of medication would submit a claim for a quantity of 400.</td>
</tr>
</tbody>
</table>

Procedures performed in conjunction with botulinum toxin injections are subject to guidelines set forth in the policies specific for those procedures. Any supplies billed by the provider for the administration of botulinum toxin type A or B are not separately payable.

Botulinum toxins administered more frequently than every 12 weeks must include documentation of medical necessity justifying why the medication was given at an interval sooner than 12 weeks.

Documentation in the client’s medical record must include the following elements:

- Support for the medical necessity of the botulinum toxin injection:
- A covered diagnosis
- Dosage and frequency of the injections
- Support of the clinical effectiveness of the injections
- Specific site(s) injected

All documentation is subject to retrospective review.

### 7.11 Brexanolone (Zulresso)

Brexanolone (Zulresso) (procedure code J1632) is a benefit for female clients who are 18 years of age and older, and is indicated for the treatment of postpartum depression in adults. Brexanolone (Zulresso) must be prescribed by, or in consultation with, a psychiatrist or obstetrics/gynecologist.

Prior authorization is required for brexanolone (Zulresso) (procedure code J1632).

Brexanolone (Zulresso) is not a benefit for clients with active psychosis or history of bipolar disorder or schizophrenia.
7.11.1 Risk Evaluation and Mitigation Strategy Program
Brexanolone (Zulresso) is only available through a restricted program under a Risk Evaluation and Mitigation Strategy (REMS) due to the risk of excessive sedation or sudden loss of consciousness.

Health-care facilities must enroll in the brexanolone (Zulresso) REMS program.

Prior to brexanolone (Zulresso) treatment, clients must also be enrolled in the brexanolone (Zulresso) REMS program. Certified facilities must ensure that brexanolone (Zulresso) is only administered to clients who are enrolled in the brexanolone (Zulresso) REMS program.

Pharmacies must be certified in the brexanolone (Zulresso) REMS program and must only dispense to health-care facilities certified to administer brexanolone (Zulresso).

7.11.2 Prior Authorization Requirements
Prior authorization requests for procedure code J1632 must be submitted with a Special Medical Prior Authorization Request Form, and may be approved for one continuous 60-hour intravenous infusion per pregnancy or postpartum period.

For treatment of postpartum depression with brexanolone (Zulresso) therapy, all of the following criteria must be met:

- The client has a diagnosis of postpartum depression (diagnosis code F530) with a HAM-D total score of at least 20, or as scored by an alternative comparable rating scale that measures depressive symptoms.
- The onset of the major depressive episode is within the third trimester and no later than the first four weeks postpartum.
- The client is six months or less postpartum at screening.
- The client does not have active psychosis or history of bipolar disorder or schizophrenia.
- The client has not received treatment with brexanolone (Zulresso) for the current postpartum depressive episode.
- The client must have continuous pulse oximetry monitoring during the infusion period due to risk of serious harm, and must be accompanied when interacting with their children as the drug can cause loss of consciousness.
- A health-care provider must be available on site for continuous monitoring of the client for the duration of the infusion.

Refer to: Subsection 3.1, “Prior Authorization Requests” in this handbook for additional prior authorization information.

7.12 Burosumab-Twza (Crysvita)
Burosumab-Twza (Crysvita) (procedure code J0584) is a benefit of Texas Medicaid with prior authorization. Burosumab-Twza (Crysvita) may be approved for a duration of every 12 months per prior authorization request.

Burosumab-twza (Crysvita) is a fibroblast growth factor 23 (FGF23) blocking antibody, indicated to treat the following:

- X-linked hypophosphatemia (XLH, a rare, inherited form of rickets) in adult and pediatric clients who are 6 months of age and older
- FGF23-related hypophosphatemia in tumor-induced osteomalacia associated with phosphaturic mesenchymal tumors that cannot be localized or is not amenable by surgical excision in adult and pediatric clients who are two years of age and older
The initial therapy for X-linked hypophosphatemia (XLH) must meet the following criteria:

- The client is 6 months of age or older.
- The client has a diagnosis of X-linked hypophosphatemia (XLH) (diagnosis code E8330 or E8331) that is supported by one of the following:
  - Confirmed phosphate regulating gene with homology to endopeptidases located on the X chromosome (PHEX-gene) mutation
  - Serum fibroblast growth factor -23 (FGF23) level >30 pg/ml
- The prescriber discontinues any oral phosphate or active vitamin D analog supplementation at least one week prior to starting burosumab-twza (Crysvita) therapy.
- The prescriber agrees to measure serum phosphate throughout therapy and withhold medication when serum phosphorus is above 5 mg/dl.

The initial therapy for FGF23-related hypophosphatemia in tumor-induced osteomalacia must meet the following criteria:

- The client is two years of age or older.
- The client has a diagnosis of FGF23-related hypophosphatemia produced by an underlying tumor that cannot be localized or is not amenable to surgical excision.
- The prescriber discontinues any oral phosphate or vitamin D analog supplement at least two weeks prior to starting burosumab-twza (Crysvita) therapy.
- The prescriber agrees to measure serum phosphate throughout therapy.

For renewal or continuation therapy, the following criteria must be met:

- The client has previously received treatment with burosumab-twza (Crysvita).
- Documentation from physician confirming one of the following:
  - The client has achieved normal level of serum phosphate.
  - The client has demonstrated a positive clinical response to burosumab-twza (Crysvita) (e.g., enhanced height velocity, improvement in askeletal deformity, reduction of fractures, and reduction of generalized bone pain).
- The physician continues to monitor serum phosphate level.

Burosumab-twza (Crysvita) must be prescribed by a nephrologist or endocrinologist, or be in consultation with a nephrologist or endocrinologist.

Burosumab-twza (Crysvita) is not a benefit for the following:

- Clients who currently use oral phosphates and active vitamin D analogs.
- Clients whose serum phosphorus is within or above the normal range for client’s age.
- Clients with severe renal impairment or end stage renal disease.

Refer to: Subsection 3.1, “Prior Authorization Requests” in this handbook for additional prior authorization information.

7.13 Calaspargase Pegol-Mkn1

Calaspargase pegol-mkn1 (procedure code J9118) is a benefit of Texas Medicaid for clients who are birth through 21 years of age.

Procedure code J9118 is limited to diagnosis codes C9100, C9101, C9102.
7.14 Cemiplimab-rwlc

Cemiplimab-rwlc (procedure code J9119) is a benefit for clients who are 18 years of age and older and is limited to the following diagnosis codes:

<table>
<thead>
<tr>
<th>Diagnosis Codes</th>
</tr>
</thead>
<tbody>
<tr>
<td>C4402 C44121 C441221 C44122 C441291 C441292 C44221 C44222</td>
</tr>
<tr>
<td>C44229 C44320 C44321 C44329 C4442 C44520 C44521 C44529</td>
</tr>
<tr>
<td>C44621 C44622 C44629 C44721 C44722 C44729 C4482 C4492</td>
</tr>
</tbody>
</table>

7.15 Chelating Agents

Chelating agent procedure codes J0470, J0600, and J0895 are benefits of Texas Medicaid when billed with an appropriate diagnosis code.

7.15.1 Dimercaprol

Procedure code J0470 is a benefit when billed with one of the following diagnosis codes:

<table>
<thead>
<tr>
<th>Diagnosis Codes</th>
</tr>
</thead>
<tbody>
<tr>
<td>T560X1A T560X1D T560X1S T560X2A T560X2D T560X2S T560X3A T560X3D</td>
</tr>
<tr>
<td>T560X3S T560X4A T560X4D T560X4S T561X1A T561X1D T561X1S T561X2A</td>
</tr>
<tr>
<td>T561X2D T561X2S T561X3A T561X3D T561X3S T561X4A T561X4D T561X4S</td>
</tr>
<tr>
<td>T564X1A T564X1D T564X1S T564X2A T564X2D T564X2S T564X3A T564X3D</td>
</tr>
<tr>
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</tr>
<tr>
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</tr>
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</tr>
<tr>
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</tr>
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</tr>
<tr>
<td>T570X3S T570X4A T570X4D T570X4S</td>
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</tbody>
</table>

7.15.2 Edetate calcium disodium

Procedure code J0600 is a benefit when billed with one of the following diagnosis codes:

<table>
<thead>
<tr>
<th>Diagnosis Codes</th>
</tr>
</thead>
<tbody>
<tr>
<td>T560X1A T560X1D T560X1S T560X2A T560X2D T560X2S T560X3A T560X3D</td>
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<tr>
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</tr>
<tr>
<td>T564X2D T564X2S T564X3A T564X3D T564X3S T564X4A T564X4D T564X4S</td>
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<tr>
<td>T565X1A T565X1D T565X1S T565X2A T565X2D T565X2S T565X3A T565X3D</td>
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<tr>
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</tr>
<tr>
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</tr>
<tr>
<td>T56892D T56892S T56893A T56893D T56893S T56894A T56894D T56894S</td>
</tr>
</tbody>
</table>
7.15.3 Deferoxamine mesylate (Desferal)

Procedure code J0895 must be billed with one of the following diagnosis codes:

<table>
<thead>
<tr>
<th>Diagnosis Codes</th>
</tr>
</thead>
<tbody>
<tr>
<td>T5691XA T5691XD T5691XS T5692XA T5692XD T5692XS T5693XA T5693XD</td>
</tr>
<tr>
<td>T5693XS T5694XA T5694XD T5694XS</td>
</tr>
</tbody>
</table>

7.16 Chimeric Antigen Receptor (CAR) T-Cell Therapy

Axicabtagene ciloleucel (Yescarta) (procedure code Q2041), Brexucabtagene autoleucel (Tecartus) (procedure code C9073), and Tisagenlecleucel (Kymriah) (procedure code Q2042) are benefits of Texas Medicaid with prior authorization and must be prescribed by an oncologist or in consultation with an oncologist.

Procedure codes C9073, Q2041, and Q2042 are limited to once per lifetime, any provider.

Axicabtagene ciloleucel (Yescarta) and tisagenlecleucel (Kymriah) infusions must take place at a certified healthcare facility. Certified healthcare facilities must enroll with the Risk Evaluation and Mitigation Strategies (REMS) and comply with its requirements for each drug administered within this section.

Certified healthcare facilities must ensure that providers that prescribe, dispense, or administer axicabtagene ciloleucel (Yescarta) and tisagenlecleucel (Kymriah) receive training for the management of cytokine release syndrome (CRS) and neurological toxicities.
It is recommended that severe or life-threatening CRS be treated with tocilizumab. Facilities must have on-site at least 2 doses of tocilizumab per patient for administration within 2 hours of infusion if needed for treatment of CRS.

Providers and facilities must ensure that the client:
- Receives the recommended pre-medications before treatment.
- Is closely monitored for toxicity post infusion.
- Is instructed to remain within proximity of the certified healthcare facility for at least 4 weeks post-infusion.

### 7.16.1 Prior Authorization Criteria for Axicabtagene Ciloleucel (Yescarta)
Prior authorization approval of axicabtagene ciloleucel (Yescarta) (procedure code Q2041) infusion therapy will be considered when all of the following criteria are met:
- The client must have a histologically confirmed diagnosis of one of the following types of aggressive non-Hodgkin’s lymphoma (diagnosis codes C8330, C8331, C8332, C8333, C8334, C8335, C8336, C8337, C8338, and C8339):
  - Diffuse large B-cell lymphoma, not otherwise specified
  - High-grade B-cell lymphoma
  - Primary mediastinal large B-cell lymphoma
  - Transformed follicular lymphoma
- The client is 18 years of age or older.
- The client must have relapsed or refractory disease, defined as progression after two or more lines of systemic therapy (which may or may not include therapy supported by autologous stem cell transplant).
- The client must have received adequate prior therapy including, at a minimum, all of the following:
  - An anthracycline-containing chemotherapy regimen
  - For CD20+ disease, anti-CD20 monoclonal antibody
  - For clients with transformed follicular lymphoma, prior chemotherapy for follicular lymphoma with chemotherapy refractory disease after transformation to DLBCL
- There must be documentation of all of the following clinical findings:
  - Client has an Eastern Cooperative Oncology Group performance status of 0 or 1.
  - Client does not have primary central nervous system lymphoma/disease.
  - Client does not have an active infection or inflammatory disorder.
  - Client has not received prior CD-19 directed CAR-T therapy.

### 7.16.2 Prior Authorization for Brexucabtagene autoleucel (Tecartus)
Prior authorization approval for Brexucabtagene autoleucel (Tecartus) (procedure code C9073) infusion therapy will be considered when all of the following criteria are met:
- The client has histologically confirmed diagnosis of relapse or refractory mantle cell lymphoma (diagnosis codes C8310, C8311, C8312, C8313, C8314, C8315, C8316, C8317, C8318 and C8319).
- The client is 18 years of age or older.
- The client has received adequate therapy and has had at least one of the following systemic treatment for MCL prior to brexucabtagene autoleucel therapy:
• Anthracycline or bendamustine-containing chemotherapy
• Anti-CD 20 monoclonal antibody therapy
• Bruton Tyrosine Kinase inhibitors (e.g. ibrutinib or acalabrutinib)
• The client does not have primary central nervous system lymphoma/disease.
• The client does not have an active infection or inflammatory disorder.
• The client has not received prior CD-19 directed CAR-T therapy.

7.16.3 Prior Authorization Criteria for Tisagenlecleucel (Kymriah)

Prior authorization approval of tisagenlecleucel (Kymriah) (procedure code Q2042) infusion for the treatment of clients with refractory or second relapse B-cell precursor acute lymphoblastic leukemia will be considered when all of the following criteria are met:

• The client has a confirmed diagnosis of B-cell acute lymphoblastic leukemia (diagnosis codes C9100, C9101, and C9102).
• The client is 25 years of age or younger.
• The client has a confirmed CD-19 tumor expression.
• The client does not have an active infection or inflammatory disorder.
• The Eastern Cooperative Oncology Group performance status is between 0 to 3.
• The client has not received prior CAR-T therapy.

Prior authorization approval of tisagenlecleucel (Kymriah) infusion for the treatment of clients with relapsed or refractory diffuse large B-cell lymphoma will be considered when all of the following criteria are met:

• The client has a confirmed diagnosis of relapsed or refractory large B-cell lymphoma (diagnosis codes C8330, C8331, C8332, C8333, C8334, C8335, C8336, C8337, C8338, and C8339):
  • Diffuse large B-cell lymphoma, not otherwise specified
  • High grade B-cell lymphoma
  • Diffuse large B-cell lymphoma arising from follicular lymphoma
• The client is 18 years of age or older.
• The client must have relapsed or refractory disease as progression after two or more lines of systemic therapy (which may or may not include therapy supported by autologous stem cell transplant).
• The client must have received adequate prior therapy including, at a minimum, the following:
  • An anthracycline-containing chemotherapy regimen
  • For CD20+ disease, an anti-CD20 monoclonal antibody
  • For clients with transformed follicular lymphoma, prior chemotherapy refractory disease after transformation to DLBCL
• The client has an Eastern Cooperative Oncology Group performance status of 0 or 1
• The client does not have primary central nervous system lymphoma
• The client does not have an active infection or inflammatory disorder
• The client has not received prior CD-19 directed CAR-T therapy

Refer to: Subsection 3.1, “Prior Authorization Requests” in this handbook for additional prior authorization information.
7.16.4 Exclusions
Axicabtagene ciloleucel (Yescarta), Brexucabtagene autoleucel (Tecartus), and Tisagenlecleucel (Kymriah) are not benefits for clients who have any of the following:
- An active infection
- An inflammatory disorder
- Primary central nervous system lymphoma

7.17 Clofarabine
Clofarabine is used for the treatment of relapsed or refractory acute lymphoblastic leukemia. Clofarabine is administered by IV infusion once daily for five days and is repeated every two to six weeks, as needed.

7.17.1 Prior Authorization for Clofarabine
Prior authorization is required for treatment with clofarabine (procedure code J9027) and may be granted for a maximum of six weeks.

Clofarabine may be prior authorized for the treatment of relapsed or refractory acute lymphoblastic leukemia. The following criteria apply to requests for prior authorization:
- The number of anticipated injections needed as well as the dosage per injection must be submitted with the request for prior authorization.
- Prior authorization must be obtained before services are rendered whenever possible. If authorization cannot be obtained prior to the rendering of the service, the authorization request must be submitted within three business days from the date the treatment is initiated.

Prior authorization requests may be considered with documentation of both of the following:
- A diagnosis of refractory or relapsed acute lymphoblastic leukemia
- A history of at least two prior failed chemotherapy regimens

The prior authorization number must be included on the claim along with the number of units, based on the dosage given. Failure to place the prior authorization number on the claim or to obtain prior authorization within the allotted timeframe will result in denied claims.

Refer to: Subsection 3.1, “Prior Authorization Requests” in this handbook for additional prior authorization information.

7.18 Colony Stimulating Factors (Filgrastim, Pegfilgrastim, and Sargramostim)
Colony stimulating factors (CSFs) are growth factors (glycoproteins) that support survival, clonal expansion and differentiation of blood forming cells and are a benefit of Texas Medicaid. CSFs reduce the likelihood of neutropenic complications due to chemotherapy and bone marrow transplant.

Filgrastim (procedure codes J1442, J1447, and Q5101) and pegfilgrastim (procedure code J2505) are granulocyte colony stimulating factors (G-CSFs). The biosimilars to filgastrim, nivestym (procedure code Q5110), pegfilgrastim-bmez (Ziextenzo) (procedure code Q5120), and pegfilgrastim (Fulphila) (procedure code Q5108) are G-CSFs. Sargramostim (procedure code J2820) is a granulocyte-macrophage colony stimulating factor (GM-CSF). GM-CSF and G-CSF stimulate neutrophil production after autologous bone marrow transplant and significantly reduce the duration and impact of neutropenia.

To submit claims for reimbursement of colony stimulating factors, providers must submit the most appropriate procedure code with the number of units administered.

Procedure code J2505 will be denied if billed on the same date of service as procedure code J1442.
Procedure code 96377 must be billed with procedure code J2505 on the same day by the same provider.

One of the following diagnosis codes must be billed with the appropriate procedure code:

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7.19 Crizanlizumab-tmca (Adakveo)

Crizanlizumab-tmca (Adakveo) (procedure code J0791) is a benefit of Texas Medicaid for clients who are 16 years of age and older.

Crizanlizumab-tmca (Adakveo) is indicated for clients with sickle cell disease to reduce the frequency of vaso-occlusive crises (VOCs) and must be prescribed by, or in consultation with, a hematologist or a sickle cell disease specialist.

7.19.1 Prior Authorization

Prior authorization is required for crizanlizumab-tmca (Adakveo) and may be approved for a duration of 12 months.

Initial therapy requests for Crizanlizumab-tmca (Adakveo) may be approved for a 12-month duration if all of the following criteria are met:

- The client must be 16 years of age or older.
- The client has a diagnosis of sickle cell disease of any genotype.
- The client has experienced two or more vaso-occlusive events in the past 12 months.
- Clients will not receive crizanlizumab-tmca (Adakveo) therapy concomitantly with voxelotor (Oxbryta).

For renewal or continuation therapy requests, the client must meet all of the following requirements:

- The client continues to meet the following initial approval criteria:
  - The client must be 16 years of age or older.
  - The client has a diagnosis of sickle cell disease of any genotype.
  - The client is not receiving crizanlizumab-tmca (Adakveo) therapy concomitantly with voxelotor (Oxbryta).
  - The client experienced positive clinical response to therapy as demonstrated by reduced frequency of vaso-occlusive crisis.
  - The client has previously received treatment with Crizanlizumab-tmca (Adakveo) without complications.
Refer to: Subsection 3.1, “Prior Authorization Requests” in this handbook for additional prior authorization information.

7.20 Denileukin diftitox (Ontak)
Denileukin diftitox (Ontak) (procedure code J9160) is a benefit for clients who have advanced or recurrent cutaneous T-cell lymphoma with the CD25 component of IL-2 and failure of at least one type of traditional therapy. Documentation of diagnosis and treatment must be submitted with the claim.

7.21 Dimethyl sulfoxide
Dimethyl sulfoxide (procedure code J1212) is a benefit of Texas Medicaid and is limited to diagnosis codes N3010 and N3011.

7.22 Eculizumab
Eculizumab (procedure code J1300) is a benefit of Texas Medicaid and is limited to the following diagnosis codes:

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7.23 Edaravone (Radicava)
Procedure code J1301 is a benefit of Texas Medicaid for clients who are 18 years of age and older with prior authorization. Client must have a diagnosis of amyotrophic lateral sclerosis (ALS).

7.24 Emapalumab-lzsg (Gamifant)
Emapalumab-lzsg (Gamifant) is an interferon gamma (IFNy) blocking antibody that is indicated for the treatment of adult and pediatric (newborn and older) clients with primary Hemophagocytic Lymphohistiocytosis (HLH) with refractory, recurrent or progressive disease or intolerance with conventional HLH therapy.

Emapalumab-lzsg (Gamifant) (procedure code J9210) is a benefit of Texas Medicaid for clients (newborn and older) with prior authorization.

Emapalumab-lzsg (Gamifant) must be prescribed by, or in consultation with, a hematologist, oncologist or a specialist in hemophagocytic lymphohistiocytosis disorder.

Prescriber must administer dexamethasone concomitantly with emapalumab-lzsg (Gamifant) therapy.

Emapalumab-lzsg (Gamifant) is administered as part of the induction/maintenance phase of hematopoietic stem cell transplant (HSCT) and therapy will be discontinued once client is at the initiation phase for HCST and no longer requires emapalumab-lzsg (Gamifant) therapy for HLH.

7.24.1 Prior Authorization Requirements
Prior authorization is required for emapalumab-lzsg (Gamifant) (procedure code J9210).

Prior authorization for initial therapy will be approved for a duration of six months for clients who meet the following criteria:

- Client has a documented diagnosis of primary HLH (diagnosis code D761) based on the following:
  - Genetic mutation of the gene known to cause primary HLH (e.g., PRF1, UNC13D, STX11, or STXB2) or a family history consistent with primary HLH or
  - Confirmation of least 5 of the following criteria:
• Fever ≥101.3 °F
• Splenomegaly
• Cytopenia defined by at least 2 of the following: Hemoglobin <9 g/dl; OR platelet count <100 x 10^9/L; OR neutrophils <1 x 10^9/L; OR fasting triglycerides >265 mg/dl OR fibrinogen ≤1.5 g/L
• Hemophagocytosis in the liver, bone marrow, spleen or lymph node
• Low or absent natural killer (NK) cell activity
• Serum ferritin concentration ≥ 500 mg/L
• High plasma concentration of soluble CD25 (i.e., soluble interleukin-2 receptor) >2,400 U/mL

• Client has refractory, recurrent or progressive disease or intolerance with conventional HLH therapy (e.g., etoposide, cyclosporine A, or anti-thymocyte globulin).
• Client has been tested/screened for latent tuberculosis infection prior to initiation of emapalumab-lzsg (Gamifant) therapy.
• Client has not undergone hematopoietic stem cell transplant and is candidate for HSCT once emapalumab-lzsg (Gamifant) therapy has been discontinued.

For renewal or continuation therapy with emapalumab-lzsg (Gamifant), the client must meet all of the following requirements:
• Client continues to meet the initial approval criteria.
• Client continues to require emapalumab-lzsg (Gamifant) as HLH treatment pending the initiation of HSCT.

Refer to: Subsection 3.1, “Prior Authorization Requests” in this handbook for additional prior authorization information.

7.25 Enfortumab Vedotin-ejfv (Padcev)
Enfortumab Vedotin-ejfv (procedure code J9177) is a benefit of Texas Medicaid for clients who are 18 years of age or older.

7.26 Eravacycline (Xerava)
Eravacycline (procedure code J0122) is a benefit of Texas Medicaid for clients who are 18 years of age or older.

7.27 Esketamine (Spravato)
Esketamine (Spravato) is a benefit of Texas Medicaid for clients who are 18 years of age or older with prior authorization. Providers must submit claims for esketamine (Spravato) with procedure code S0013.

Esketamine (Spravato) nasal spray is an N-methyl-D-aspartate (NMDA) receptor antagonist that is indicated in conjunction with an oral antidepressant in adult clients for the treatment of the following:
• Treatment-resistant depression (TRD)
• Depressive symptoms in clients with major depressive disorder (MDD) with acute suicidal ideation or behavior

Esketamine (Spravato) must be prescribed by, or in consultation with a psychiatrist.
Esketamine (Spravato) is only available through a restricted program under a Risk Evaluation and Mitigation Strategy (REMS).

Providers/health-care settings must be certified in the Spravato REMS program to administer Spravato to clients enrolled in the REMS program. Administration of the drug must take place in a health-care facility under the direct observation of a health-care provider.

Pharmacies must be certified in the Spravato REMS program to dispense to health-care settings who are authorized to administer the drug.

Clients must be enrolled in the Spravato REMS program and comply with the ongoing requirements for Spravato treatment.

Esketamine (Spravato) is not a benefit for clients who have aneurysmal vascular disease, arteriovenous malformation or history of intracerebral hemorrhage.

### 7.27.1 Prior Authorization

Prior authorization is required for esketamine (Spravato) (procedure code S0013) and may be approved for a duration of six months.

Initial therapy for esketamine (Spravato) nasal spray may be approved when all of the following criteria is met:

- The client is 18 years of age or older.
- Client has either of the following:
  - A treatment-resistance depression that has been confirmed as an inadequate response/failure to previous antidepressant treatment.
  - A major depressive disorder with acute suicidal ideation or behavior, and the prescriber’s evaluation shows that the client:
    - Has suicidal ideation with intent.
    - Needs acute psychiatric hospitalization due to an imminent risk of suicide.
- The client must receive esketamine (Spravato) nasal spray concomitantly with an oral antidepressant agent (esketamine [Spravato] should not be used as monotherapy).
- Esketamine (Spravato) must be administered under the direct observation of a health-care provider and the client must be monitored for at least 2 hours after each treatment.
- Prior to starting esketamine (Spravato) treatment, there must be an attestation of baseline scoring of clinical assessment of MDD.
- The client must not have contraindications to esketamine (Spravato), such as aneurysmal vascular disease, arteriovenous malformation, or intracerebral hemorrhage.

For renewal or continuation therapy, the client must meet all of the following requirements:

- The client continues to meet the initial prior authorization approval criteria.
- The client demonstrates positive clinical response to esketamine (Spravato) therapy by an improvement from baseline assessment.
- The client has previously received treatment esketamine (Spravato) without complications.

**Refer to:** Subsection 3.1, “Prior Authorization Requests” in this handbook for additional prior authorization information.
7.28 Fam-trastuzumab Deruxtecan-nxki
Fam-trastuzumab Deruxtecan-nxki (procedure code J9358) is a benefit of Texas Medicaid for clients who are 18 years of age or older.

7.29 Fluocinolone Acetonide (Retisert)
Procedure code J7311 is a benefit of Texas Medicaid for clients of all ages but is only considered for reimbursement with a posterior uveitis diagnosis of more than six months in duration and only when the condition has been unresponsive to oral or systemic medication treatment. Prior authorization is required.

7.30 Fremanezumab-vfrm
Fremanezumab-vfrm (procedure code J3031) is a benefit of Texas Medicaid for clients who are 18 years of age or older.

7.31 Galsulfase
Galsulfase (procedure code J1458) is a benefit of Texas Medicaid and is limited to the following diagnosis codes:

<table>
<thead>
<tr>
<th>Diagnosis Codes</th>
</tr>
</thead>
<tbody>
<tr>
<td>E7601</td>
</tr>
<tr>
<td>E7629</td>
</tr>
</tbody>
</table>

7.32 Granisetron Hydrochloride
Granisetron hydrochloride is a benefit of Texas Medicaid and is limited to diagnosis codes Z5189, Z510, Z5111, and Z5112.

7.33 Hematopoietic Injections
Hematopoietic agents erythropoietin alfa, epoetin alfa (EPO), methoxy polyethylene glycol-epoetin beta (Mircera), and darbepoetin alfa are benefits of Texas Medicaid and reimbursed using procedure codes J0881, J0882, J0885, J0887, and J0888 with an appropriate diagnosis code.

Providers must maintain medical records in their offices that document regular monitoring of hemoglobin or hematocrit levels and explain the rationale for the dosing of epoetin alfa, methoxy polyethylene glycol-epoetin beta, and darbepoetin alfa. These records are subject to retrospective review to determine appropriate utilization and reimbursement for this service.

When billing procedure code J0882 providers must submit the client’s most recent dated hemoglobin or hematocrit levels in the comments section of the claim form.

EPO, Mircera, and darbepoetin alfa injections are limited to specific diagnosis codes as indicated in this section.

Refer to: Subsection 6.2.9.4, “Hematopoietic Injections” in the Clinics and Other Outpatient Facility Services Handbook (Vol. 2, Provider Handbooks) for information about outpatient facility criteria.
7.33.1 Darbepoetin Alfa

Darbepoetin alfa (procedure codes J0881 and J0882) is an erythropoiesis-stimulating protein closely related to erythropoietin. Darbepoetin stimulates erythropoiesis by the same mechanism as EPO. Darbepoetin alfa has approximately a three-fold longer half-life than EPO, resulting in a sustained erythropoietic effect and less frequent dosing. Darbepoetin alfa is indicated for:

- Treatment of anemia associated with chronic renal failure (CRF), including clients on dialysis and clients not on dialysis.
- Treatment of anemia in clients who have non-myeloid malignancies where anemia is due to the effect of chemotherapy.

Procedure code J0881 must be billed with one of the following diagnosis codes:

<table>
<thead>
<tr>
<th>Diagnosis Codes</th>
</tr>
</thead>
</table>

Procedure code J0882 must be billed with one of the following diagnosis codes:

<table>
<thead>
<tr>
<th>Diagnosis Codes</th>
</tr>
</thead>
<tbody>
<tr>
<td>D631 N181 N182 N1830 N1831 N1832 N184 N185 N186 N189 N19</td>
</tr>
</tbody>
</table>

Darbepoetin alfa injections are limited to once weekly (Sunday through Saturday).

7.33.2 Epoetin Alfa (EPO)

EPO (procedure code J0885) is a glycoprotein that stimulates the formation of red blood cells and the production of the precursor red blood cells of the bone marrow. EPO is indicated for:

- Anemia associated with chronic renal failure (CRF), including clients on dialysis (end-stage renal disease or ESRD) and clients not on dialysis.
- Anemia related to therapy with zidovudine (AZT) in HIV-infected clients.
- Anemia due to the effects of concomitantly administered chemotherapy in clients who have non-myeloid malignancies.
- Anemia of prematurity.
- Clients scheduled to undergo elective noncardiac, nonvascular surgery to decrease need for allogenic blood transfusion.

Procedure code J0885 must be billed with one of the following diagnosis codes:

<table>
<thead>
<tr>
<th>Diagnosis Codes</th>
</tr>
</thead>
</table>
EPO may be considered for reimbursement when the dose is titrated consistent with prevailing, evidence-based clinical guidelines, as published by the National Kidney Foundation Kidney Disease Outcomes Quality Initiative, including appropriate monitoring of the rise and fall of the hemoglobin or hematocrit levels.

EPO is limited to three injections per calendar week (Sunday through Saturday).

### 7.33.3 Methoxy Polyethylene Glycol-Epoetin Beta (Mircera)

Methoxy polyethylene glycol-epoetin beta (Mircera) is indicated for the treatment of anemia associated with chronic kidney disease (CKD) and may be administered subcutaneously or intravenously. Mircera is not indicated for use in the treatment of anemia due to cancer chemotherapy or as a substitute for red blood cell transfusions in clients who require immediate correction of anemia. Mircera is indicated for the following:

- Treatment of anemia associated with CKD in adult clients on hemodialysis and adult clients not on hemodialysis.
- Treatment of anemia associated with CKD in pediatric clients who are 5 through 17 years of age. These clients must be on hemodialysis and converting from another erythropoiesis-stimulating agent (ESA) after their hemoglobin level was stabilized with an ESA.

**Note:** In pediatric clients, Mircera is administered by intravenous injection only.

Methoxy polyethylene glycol-epoetin beta (Mircera) (procedure code J0887) is limited to diagnosis codes D631 and N186.

Methoxy polyethylene glycol-epoetin beta (Mircera) (procedure code J0888) is limited to the following diagnosis codes:

<table>
<thead>
<tr>
<th>Diagnosis Codes</th>
</tr>
</thead>
<tbody>
<tr>
<td>D631</td>
</tr>
<tr>
<td>N1830</td>
</tr>
</tbody>
</table>

Methoxy polyethylene glycol-epoetin beta (Mircera) may be considered for reimbursement when the dose is titrated consistent with prevailing, evidence-based, clinical guidelines as published by the National Kidney Foundation Kidney Disease Outcomes Quality Initiative, including appropriate monitoring of the rise and fall of the hemoglobin or hematocrit levels.

Methoxy polyethylene glycol-epoetin beta (Mircera) is limited to one injection every 2 calendar weeks, any provider (Sunday through Saturday).

### 7.34 Hydroxyprogesterone Caproate

**Refer to:** The Gynecological, Obstetrics, and Family Planning Title XIX Services Handbook (Vol. 2, Provider Handbooks) for information about procedure codes J1726 and J1729.

### 7.35 Ibalizumab-uiyk (Trogarzo)

Ibalizumab-uiyk (Trogarzo) (procedure code J1746) is a benefit of Texas Medicaid for clients who are 18 years of age and older with prior authorization. Ibalizumab-uiyk (Trogarzo) may be approved for a duration of every 12 months per prior authorization request.

For initial therapy, all of the following criteria must be met:

- The client is 18 years of age or older.
- The client has a documented diagnosis of multi-drug resistant human immunodeficiency virus (diagnosis code B20) from the provider and meets the following criteria:
• Has received antiretroviral treatment for at least 6 months and is failing or has recently failed therapy
• Has documented resistance, measured by resistance testing, to at least one antiretroviral medication from each of the following 3 classes of ARV:
  • Nucleoside reverse transcriptase inhibitors (NRTI)
  • Non-Nucleoside reverse transcriptase inhibitors (NNRTI)
  • Protease inhibitor (PI)
• The client has documented RNA viral load greater than 1,000 copies/mL.

Providers must use ibalizumab-uiyk (Trogarzo) concomitantly with another antiretroviral medication to which the client’s virus is susceptible.

For renewal or continuation of therapy, all of the following criteria must be met:
• The client has previously received treatment with ibalizumab-uiyk (Trogarzo).
• Documentation from the physician confirming that the client has achieved a clinical viral response defined as one of the following:
  • Decrease in viral load
  • Sustained viral load reduction
• The physician continues ibalizumab-uiyk (Trogarzo) therapy with another antiretroviral.

Ibalizumab-uiyk (Trogarzo) must be prescribed by a physician, in consultation with an infectious disease physician or a physician who specializes in the treatment of HIV infection.

Trogarzo is not a benefit for clients who fail to demonstrate heavily treated multi-drug resistance.

Refer to: Subsection 3.1, “Prior Authorization Requests” in this handbook for additional prior authorization information.

### 7.36 Ibutilide fumarate

Ibutilide fumarate (procedure code J1742) is a benefit of Texas Medicaid and is limited to diagnosis codes I480, I481, I482, I483, and I484.

### 7.37 Idursulfase (Elaprase)

Idursulfase (procedure code J1743) is a benefit of Texas Medicaid and is limited to the following diagnosis codes:

<table>
<thead>
<tr>
<th>Diagnosis Codes</th>
</tr>
</thead>
<tbody>
<tr>
<td>E7601</td>
</tr>
<tr>
<td>E7629</td>
</tr>
</tbody>
</table>
7.38 * Immune Globulin

Immune globulins may be indicated for treatment of certain immune disorders and states of immunodeficiency. The following immune globulin procedure codes are benefits of Texas Medicaid:

<table>
<thead>
<tr>
<th>Procedure Codes</th>
</tr>
</thead>
<tbody>
<tr>
<td>90284 90291 C9072 J0850 J1459 J1460 J1555 J1556 J1557 J1559</td>
</tr>
<tr>
<td>J1561 J1566 J1569 J1572 J1575 J1599 J1670 J2788 J2791</td>
</tr>
<tr>
<td>J2792 J7504 J7511</td>
</tr>
</tbody>
</table>

Note: Procedure codes 90291 and J0850 may only be reimbursed when billed with diagnosis code Z940, Z941, Z942, Z943, Z944, or Z9483.

7.39 Immunosuppressive Drugs

Immunosuppressive drugs weaken or modulate the activity of the immune system and are most often used in organ transplantation to prevent rejection or to treat autoimmune diseases such as rheumatoid arthritis.

The following procedure codes are benefits of Texas Medicaid:

<table>
<thead>
<tr>
<th>Procedure Codes</th>
</tr>
</thead>
<tbody>
<tr>
<td>J0202 J0257 J0480 J0485 J0490 J0717 J1595 J1602 J7501 J7516 J7525</td>
</tr>
</tbody>
</table>

The following procedure codes may be indicated for, but are not limited to, treatment of the following conditions:

<table>
<thead>
<tr>
<th>Procedure Code</th>
<th>Conditions</th>
</tr>
</thead>
<tbody>
<tr>
<td>J0202</td>
<td>Multiple sclerosis (MS): For treatment of relapsing forms of MS and should be reserved for clients who have had an inadequate response to two or more drugs indicated for the treatment of MS.</td>
</tr>
<tr>
<td>J0257</td>
<td>Alpha-1 proteinase inhibitor deficiency: For the treatment of clients who have a deficiency of the alpha-1 proteinase inhibitor enzyme (also known as alpha-1 antitrypsin deficiency) in the treatment of emphysema.</td>
</tr>
<tr>
<td>J0480</td>
<td>Organ rejection: For the prophylaxis of acute organ rejection in patients receiving renal transplantation when used as part of an immunosuppressive regimen that includes cyclosporine and corticosteroids.</td>
</tr>
<tr>
<td>J0485</td>
<td>Organ rejection: For the prophylaxis of organ rejection in adults receiving a kidney transplant, to be used in combination with basiliximab injection, mycophenolate mofetil, and corticosteroids.</td>
</tr>
<tr>
<td>J0490</td>
<td>Systemic lupus erythematosus (SLE): For use in clients with moderate to severe SLE when other forms of treatment have failed to control moderate to severe symptoms</td>
</tr>
<tr>
<td>J0717</td>
<td>Psoriatic arthritis, Ulcerative colitis, Ankylosing spondylitis, Crohn’s disease</td>
</tr>
<tr>
<td>J1595</td>
<td>Multiple sclerosis (MS): For the reduction of the frequency of relapses in clients with relapsing remitting MS, including clients who have experienced a first clinical episode and have magnetic resonance imaging (MRI) features consistent with MS.</td>
</tr>
</tbody>
</table>
### Procedure Code | Conditions
---|---
J1602 | Psoriatic arthritis, Rheumatoid arthritis, Ankylosing spondylitis
J7501 | Renal homotransplantations: Adjunct for the prevention of rejection in renal homotransplantation. Rheumatoid arthritis: Azathioprine is indicated only in adult patients meeting the criteria for classic or definite rheumatoid arthritis as specified by the American Rheumatism Association.
J7516 | Allogeneic transplants: For prophylaxis of organ rejection in kidney, liver, and heart allogeneic transplants.
J7525 | Organ rejection prophylaxis: For the prophylaxis of organ rejection in clients receiving allogeneic liver, kidney, or heart transplants.

**Note:** Oral, self-administered immunosuppressive drugs may be reimbursed for Medicaid fee-for-service clients through the Medicaid Vendor Drug Program (VDP).

Retrospective review may be performed to ensure documentation supports the medical necessity of the service. Authorization is not required for immunosuppressive drugs.

### 7.40 Inebilizumab-cdon (Uplizna)
Inebilizumab-cdon (Uplizna) (procedure code J1823) is a benefit of Texas Medicaid for clients who are 18 years of age or older with prior authorization and must be prescribed by or in consultation with a neurologist.

Inebilizumab-cdon (Uplizna) is indicated for the treatment of adult clients with neuromyelitis optica spectrum disorder (NMOD/NMOSD) who are anti-aquaporin-4 antibody positive.

#### 7.40.1 Prior Authorization Criteria
Prior authorization for initial therapy for inebilizumab-cdon (Uplizna) may be approved for a 12-month duration if all of the following criteria are met:

- Client must be 18 years of age or older.
- Client has a diagnosis of neuromyelitis optica spectrum disorder (G360).
- Client is anti-aquaporin 4 (AQP4) antibody seropositive.
- Client has been screened for hepatitis B virus (HBV), quantitative serum immunoglobulins, and tuberculosis (TB) prior to initiating treatment.
- Client has at least one attack requiring rescue therapy in the last year OR two attacks requiring rescue therapy in the last 2 years.
- Client is not receiving Inebilizumab-cdon (Uplizna) concomitantly with the following therapies:
  - Anti-CD20 monoclonal antibody treatments
  - Complement inhibitors (e.g. Eculizumab, Ravulizumab)
  - Immunosuppressant drugs (e.g. Cyclosporine, Methotrexate)
  - Satralizumab
For renewal or continuation of therapy, the client must meet all of the following requirements:

- Client continues to meet the following initial approval criteria.
- Client experienced positive clinical response to therapy as demonstrated by decreased attacks or disease stabilization.
- Client has previously received Inebilizumab-cdon (Uplizna) treatment without complications.

### 7.41 Infliximab (Remicade), Inflectra*, Renflexis*

Procedure code J1745, Q5103, and Q5104 are benefits of Texas Medicaid, and may not be reimbursed for the same date of service by any provider. Procedure codes J1745, Q5103, and Q5104 are limited to the following diagnosis codes:

<table>
<thead>
<tr>
<th>Diagnosis Codes</th>
</tr>
</thead>
<tbody>
<tr>
<td>K5000</td>
</tr>
<tr>
<td>K50112</td>
</tr>
<tr>
<td>K50814</td>
</tr>
<tr>
<td>K50919</td>
</tr>
<tr>
<td>K51211</td>
</tr>
<tr>
<td>K51313</td>
</tr>
<tr>
<td>K51518</td>
</tr>
<tr>
<td>K51911</td>
</tr>
<tr>
<td>L401</td>
</tr>
<tr>
<td>L4054</td>
</tr>
<tr>
<td>M05032</td>
</tr>
<tr>
<td>M05072</td>
</tr>
<tr>
<td>M05441</td>
</tr>
<tr>
<td>M0549</td>
</tr>
<tr>
<td>M05542</td>
</tr>
<tr>
<td>M05611</td>
</tr>
<tr>
<td>M05651</td>
</tr>
<tr>
<td>M05712</td>
</tr>
<tr>
<td>M05752</td>
</tr>
<tr>
<td>M05811</td>
</tr>
<tr>
<td>M05851</td>
</tr>
<tr>
<td>M06012</td>
</tr>
<tr>
<td>M06052</td>
</tr>
<tr>
<td>M06812</td>
</tr>
<tr>
<td>M06841</td>
</tr>
<tr>
<td>M06869</td>
</tr>
<tr>
<td>M08012</td>
</tr>
<tr>
<td>M08052</td>
</tr>
<tr>
<td>M08821</td>
</tr>
<tr>
<td>M08861</td>
</tr>
</tbody>
</table>
7.42 Inotuzumab ozogamicin (Besponsa)

Inotuzumab ozogamicin (Besponsa) (procedure code J9229) is a benefit of Texas Medicaid for clients who are 18 years of age and older with prior authorization, and must be prescribed by an oncologist or be in consultation with an oncologist.

Inotuzumab ozogamicin (Besponsa) is a CD22-directed antibody-drug conjugate (ADC) that has 3 components:
- The antibody inotuzumab
- N-acetyl-gamma-calicheamicin dimethylhydrazide (a cytotoxic agent)
- An acid cleavable linker

Besponsa is indicated for the treatment of relapsed or refractory precursor B-cell acute lymphoblastic leukemia (ALL).

7.42.1 Prior Authorization Requirements for Inotuzumab ozogamicin (Besponsa)

Prior authorization approval for Besponsa intravenous injection (procedure code J9229) will be considered when all of the following criteria are met:
- Client is 18 years of age or older
- Client has a confirmed diagnosis of precursor B-cell acute lymphoblastic leukemia (ALL) that is refractory or in relapse
- The prescriber must agree to monitor the client for signs and symptoms of hepatic veno-occlusive disease (VOD) for the duration of Besponsa therapy

Refer to: Subsection 3.1, “Prior Authorization Requests” in this handbook for additional prior authorization information.

7.42.2 Documentation Requirements

In addition to documentation requirements outlined above all services are subject to retrospective review to ensure that the documentation in the client’s medical record supports the medical necessity of the service(s) provided.

7.42.3 Exclusions

Besponsa is not a benefit for patients who have hepatic veno-occlusive disease.

7.43 * Interferon

Interferons are a family of naturally-occurring proteins that are produced by cells of the immune system. Three classes of interferons have been identified: alfa, beta, and gamma. Each class has different effects, though their activities overlap. Together, the interferons direct the immune system’s attack on viruses, bacteria, tumors, and other foreign substances that may invade the body. Once interferons have detected and attacked a foreign substance, they alter it by slowing, blocking, or changing its growth or function.
The following interferon procedure codes are benefits of Texas Medicaid:

<table>
<thead>
<tr>
<th>Revised Procedure Codes</th>
</tr>
</thead>
<tbody>
<tr>
<td>J1826</td>
</tr>
</tbody>
</table>

[Revised] Interferon Alfa-2B (procedure code J9214) may be indicated for, but is not limited to, treatment of the conditions listed below:
- Acute leukemias
- AIDS-related Kaposi sarcoma Basal- and squamous-cell cancer Behcet syndrome
- Bladder tumors (local use for superficial tumors) Carcinoid tumor
- Chronic granulocytic leukemia Chronic hepatitis B virus Chronic hepatitis C virus Chronic myelogenous leukemia Condylomata acuminata Cutaneous T-cell lymphoma Cytomegalovirus
- Essential thrombocytopenia Essential thrombocytosis Follicular lymphoma Hairy cell leukemia Herpes simplex Hodgkin’s disease
- Hypereosinophilic syndrome Melanoma Multiple myeloma Mycosis fungoides
- Non-Hodgkin’s lymphoma Ovarian and cervical carcinoma Papilloma viruses Polycythemia vera
- Renal cell carcinoma Rhino viruses Varicella zoster

[Revised] Interferon Gamma-1B (procedure code J9216) may be indicated for, but is not limited to, treatment of the following:
- Chronic granulomatous disease
- Malignant osteoporosis

[Revised] Interferon Beta-1A (procedure codes J1826, Q3027, and Q3028), and Interferon Beta-1B (procedure code J1830) may be indicated for, but are not limited to, treatment of relapsing forms of multiple sclerosis.

Note: Pegylated interferons are self-administered weekly and are available through Texas Medicaid Vendor Drug Program for Medicaid fee-for-service clients.

### 7.44 Iron Injections

Iron is a hematinic, essential to the synthesis of hemoglobin to maintain oxygen transport and to the function and formation of other physiologically important heme and non-heme compounds.

Ferric carboxymaltose (procedure code J1439) may be indicated for, but is not limited to, treatment of iron deficiency anemia for adult clients with:
- Intolerance or unsatisfactory response to oral iron.
- Non-dialysis-dependent chronic kidney disease.

Iron Dextran injection (procedure code J1750) may be indicated for, but is not limited to treatment of iron deficiency anemia when oral administration is unsatisfactory or impossible.

Iron Sucrose injection (procedure code J1756) may be indicated for, but is not limited to treatment of iron deficiency anemia for the following conditions:
- Non-dialysis-dependent chronic kidney disease (NDD-CKD) for clients who are receiving erythropoietin.
- NDD-CKD for clients who are not receiving erythropoietin.
• Hemodialysis-dependent chronic kidney disease (HDD-CKD) for clients who are receiving erythropoietin.

• Peritoneal dialysis-dependent chronic kidney disease (PDD-CKD) clients who are receiving erythropoietin.

Sodium Ferric Gluconate Complex injection (procedure code J2916) may be indicated for, but is not limited to treatment of Iron deficiency anemia in clients who are six years of age or older who are undergoing long term hemodialysis treatments and who are receiving supplemental epoetin therapy.

Ferumoxytol injection (procedure codes Q0138 and Q0139) may be indicated for, but is not limited to treatment of Iron deficiency anemia in adults who have chronic kidney disease (CKD).

Note: Report procedure code Q0138 for non-end stage renal disease (ESRD) and Q0139 for ESRD injections.

Authorization is not required for iron injections. Retrospective review may be performed to ensure documentation supports the medical necessity for the service being billed.

### 7.45 Joint Injections and Trigger Point Injections

The following procedure codes must be used to submit claims for injections into joints:

<table>
<thead>
<tr>
<th>Procedure Codes for Joint Injections</th>
</tr>
</thead>
<tbody>
<tr>
<td>20600</td>
</tr>
</tbody>
</table>

The following procedure codes must be used to submit claims for trigger point injections:

<table>
<thead>
<tr>
<th>Procedure Codes for Trigger Point Injections</th>
</tr>
</thead>
<tbody>
<tr>
<td>20526</td>
</tr>
</tbody>
</table>

These procedures are valid only in the treatment of acute problems. Procedures billed for reimbursement with chronic diagnosis codes are denied. The provider must use the AT modifier to indicate an acute condition.

<table>
<thead>
<tr>
<th>Modifier</th>
<th>Use</th>
</tr>
</thead>
<tbody>
<tr>
<td>AT</td>
<td>For acute conditions</td>
</tr>
</tbody>
</table>

The cost of the injection does not include the drugs used. The drug can be reimbursed separately. Multiple joint injections may be reimbursed when billed with the same date of service if the claim indicates the specific site of each injection. The first injection or aspiration is reimbursed at the full profile allowance and any subsequent injections are reimbursed at half allowance.

### 7.46 Lactated Ringer’s

Lactated Ringer’s (procedure code J7121) is a benefit of Texas Medicaid for clients who are birth through 20 years of age.

### 7.47 Lanadelumab-flyo

Lanadelumab-flyo (procedure code J0593) is a benefit of Texas Medicaid for clients who are 12 years of age or older.
7.48  **Leuprolide Acetate (Lupron Depot)**

Procedure codes J9217, J1950, J9218, or J9219 may be reimbursed for leuprolide acetate injections with the following limitations:

<table>
<thead>
<tr>
<th>Procedure Code</th>
<th>Limitation(s)</th>
</tr>
</thead>
<tbody>
<tr>
<td>J1950</td>
<td>Reimbursed once per month</td>
</tr>
<tr>
<td>J9219</td>
<td>Reimbursed once per year</td>
</tr>
</tbody>
</table>

Procedure code J9217 may be reimbursed in monthly, three-month, four-month, and six-month doses as follows:

<table>
<thead>
<tr>
<th>Frequency</th>
<th>Dosage</th>
<th>Limitations</th>
</tr>
</thead>
<tbody>
<tr>
<td>Monthly</td>
<td>7.5 mg</td>
<td>Billed with a quantity of 1</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Reimbursed once per month</td>
</tr>
<tr>
<td>3-month</td>
<td>22.5 mg</td>
<td>Billed with a quantity of 3</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Reimbursed once every three months</td>
</tr>
<tr>
<td>4-month</td>
<td>30 mg</td>
<td>Billed with a quantity of 4</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Reimbursed once every 4 months</td>
</tr>
<tr>
<td>6-month</td>
<td>45 mg</td>
<td>Billed with a quantity of 6</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Reimbursed once every 6 months</td>
</tr>
</tbody>
</table>

The total dosage allowed within a 6-month period is 45 mg.

7.49  **Medroxyprogesterone Acetate (Depo Provera)**

Refer to: **The Gynecological, Obstetrics, and Family Planning Title XIX Services Handbook (Vol. 2, Provider Handbooks)** for information about procedure code J1050.

7.50  **Melphalan**

Procedure code J9245 is a benefit when billed with one of the following diagnosis codes:

<table>
<thead>
<tr>
<th>Diagnosis Codes</th>
</tr>
</thead>
<tbody>
<tr>
<td>C50011</td>
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<tr>
<td>C50012</td>
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<tr>
<td>C50019</td>
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<td>C9000</td>
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<td>C9001</td>
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</tbody>
</table>

7.51  **Luspatercept-aamt (Reblozyl)**

Luspatercept-aamt (Reblozyl) (procedure code J0896) is a benefit of Texas Medicaid with prior authorization.

Luspatercept-aamt (Reblozyl) is restricted to clients who are 18 years of age or older, and may be approved for treatment of the following:

- Anemia in adult clients with beta thalassemia requiring red blood cell (RBC) transfusions
• Anemia failing an erythropoiesis stimulating agent and requiring two or more red blood cell units over eight weeks in adult clients with low to intermediate-risk myelodysplastic syndrome with ring sideroblasts (MDS-RS) or with myelodysplastic/myeloproliferative neoplasm with ring sideroblasts and thrombocytosis (MDS/MPN-RS-T)

Luspatercept-aamt (Reblozyl) must be prescribed by, or in consultation with, a hematologist.

7.51.1 Prior Authorization for Luspatercept-aamt (Reblozyl)

Prior authorization requests for procedure code J0896 must be submitted with a Special Medical Prior Authorization (SMPA) Request Form.

For initial prior authorization requests for Luspatercept-aamt (Reblozyl), the client must be 18 years of age or older and meet the following criteria:

• Client who has anemia with beta thalassemia requiring regular RBC transfusions:
  • The client must have a diagnosis of beta thalassemia.
  • The client requires regular RBC transfusions of six or more units within the previous 24 weeks and has had no transfusion-free period for 35 days or longer during the review period.

• Client who has anemia failing an erythropoiesis stimulating agent:
  • The client must have a diagnosis of myelodysplastic syndrome classified as low to intermediate risk disease.
  • The client must require RBC transfusions of two or more units over a period of eight weeks.
  • The client must be ineligible or must have failed prior erythropoietin stimulating agent treatment.

For renewal or continuation of therapy, the client must meet the initial age and diagnosis criteria, in addition to the following requirements:

• Client who has anemia with beta thalassemia requiring regular RBC transfusions:
  • The client had a positive response/hematological improvement demonstrated by a reduction in RBC transfusion as indicated by the prescribing physician.
  • The client previously received treatment with luspatercept-aamt (Reblozyl) without complications.

• Client who has anemia failing an erythropoiesis stimulating agent:
  • The client had a positive response demonstrated by RBC transfusion independence during any consecutive eight-week period or a decrease in transfusion requirement as indicated by the prescribing physician.
  • The client previously received treatment with luspatercept-aamt (Reblozyl) without complications.

7.52 Mepsevii (Vestronidase alfa-vjbk)

Vestronidase alfa-vjbk (Mepsevii) (procedure code J3397) is a benefit of Texas Medicaid for pediatric and adult clients with prior authorization. Vestronidase alfa-vjbk (Mepsevii) may be approved for a duration of every 12 months per prior authorization request.

For initial therapy, the following criteria must be met:

• Documentation of clinical signs and symptoms of Mucopolysaccharidosis VII (MPS VII) (e.g., skeletal deformities; enlarged liver, spleen, or both; airway obstruction or pulmonary problems; joint limitations; etc.)
• Diagnosis of Mucopolysaccharidosis VII (MPS VII, Sly syndrome) (diagnosis code E7629 or
diagnosis code E763) supported by elevated urine glycosaminoglycans excretion at a minimum of
3-fold over the mean normal for age at screening and either or the following:
  • Beta-glucuronidase enzyme deficiency in peripheral blood based on leukocytes or cultured
    fibroblasts
  • Mutation in the glucuronidase beta (GUSB) gene, confirmed by molecular genetic testing

For renewal or continuation of therapy, the following criteria must be met:
• Client has previously received treatment with vestronidase alfa-vjbk without an adverse reaction.
• Documentation from physician confirms client has experienced an improvement in clinical
  response compared to pretreatment baseline (e.g., stability in skeletal deformities; reduction in liver
  volume; reduction in spleen volume, or both; stable or improved pulmonary function; improved
  endurance; and functional capacity, etc.).

Refer to: Subsection 3.1, “Prior Authorization Requests” in this handbook for additional prior
authorization information.

7.53  Mogamulizumab-kp (Poteligeo)
Mogamulizumab-kp (Poteligeo) procedure code J9204 is a benefit of Texas Medicaid. Mogamulizumab-kp (Poteligeo) is a CCR4-directed monoclonal antibody indicated for the treatment of an adult client with relapsed or refractory mycosis fungoides or Sézary syndrome after at least one prior systemic therapy.

Mogamulizumab-kp (Poteligeo) requires prior authorization, and must be prescribed by, or in consultation with, an oncologist or hematologist. Mogamulizumab-kp (Poteligeo) (procedure code J9204) may be approved for a duration of every 12 months.

7.53.1  Prior Authorization Criteria
Prior authorization for initial therapy using mogamulizumab-kp (Poteligeo) infusion will be considered when all of the following criteria are met:
• The client is 18 years of age or older
• The client has relapsed or refractory disease
• The client has received at least one prior systemic therapy
• The client has a histologically confirmed diagnosis of Mycosis fungoides or Sézary syndrome

One of the following diagnosis codes must be submitted with procedure code J9204 if the diagnosis is mycosis fungoides:

<table>
<thead>
<tr>
<th>Diagnosis Codes for Mycosis Fungoides</th>
</tr>
</thead>
<tbody>
<tr>
<td>C8400</td>
</tr>
<tr>
<td>C8408</td>
</tr>
</tbody>
</table>

One of the following diagnosis codes must be submitted with procedure code J9204 if the diagnosis is sézary syndrome:

<table>
<thead>
<tr>
<th>Diagnosis for Sézary Syndrome</th>
</tr>
</thead>
<tbody>
<tr>
<td>C8410</td>
</tr>
<tr>
<td>C8418</td>
</tr>
</tbody>
</table>
For renewal or continuation of therapy, the client must meet all the following requirements:

- The client demonstrates partial or complete response to treatment or stabilization of disease, shown by a decrease in spread or size of the tumor
- The absence of unacceptable drug toxicity, such as dermatological toxicity, severe infection, infusion reactions (Stevens-Johnson Syndrome or toxic epidermal necrolysis), and life-threatening autoimmune complications

Refer to: Subsection 3.1, “Prior Authorization Requests” in this handbook for additional prior authorization information.

7.54 Monoclonal Antibodies

7.54.1 Omalizumab

Omalizumab (procedure code J2357) is an injectable drug that is FDA-approved for the treatment of clients who are 6 years of age and older with moderate to severe asthma (as defined by the National Heart, Lung, and Blood Institute’s Guidelines for the Diagnosis and management of Asthma). Omalizumab is also FDA-approved for the treatment of clients who are 12 years of age or older and have chronic idiopathic urticaria (CIU) who remain symptomatic despite H1 antihistamine treatment.

Omalizumab may be a benefit of Texas Medicaid when medically necessary with prior authorization. Clients who are younger than the FDA approved age will be considered on a case-by-case basis by the TMHP medical director.

Providers may not bill for an office visit if the only reason for the visit is an omalizumab injection.

7.54.2 Benralizumab

Benralizumab (procedure code J0517) is a benefit of Texas Medicaid for clients who are 12 years of age and older with prior authorization.

Benralizumab is an injectable drug that is FDA-approved and indicated for the treatment of clients who are 12 years of age and older and have severe asthma with an eosinophilic phenotype.

Treatment of benralizumab may not be used concurrently with omalizumab or any other interleukin-5 antagonist.

Procedure codes J0517, J2182, J2357, and J2786 may not be billed in any combination for the same date of service by any provider.

Providers may not bill for an office visit if the only reason for the visit is a benralizumab injection.

7.54.3 Mepolizumab

Mepolizumab (procedure code J2182) is a benefit of Texas Medicaid when medically necessary with prior authorization.

Mepolizumab is an injectable drug that is approved by the FDA and indicated for the following treatments:

- Clients who are 6 years of age or older and have severe asthma with an eosinophilic phenotype.
- Adult clients who are 18 years of age or older with eosinophilic granulomatosis with polyangiitis
- Adult and pediatric clients who are 12 years of age or older with hypereosinophilic symptoms (HES) for 6 months or longer without identifiable non-hematologic secondary cause

Clients who are younger than the FDA-approved age will be considered on a case-by-case basis by the TMHP medical director.

Treatment with mepolizumab may not occur concurrently with omalizumab or any other interleukin-5 antagonist.
Providers may not bill for an office visit if the only reason for the visit is a mepolizumab injection.

### 7.54.4 Reslizumab

Reslizumab (procedure code J2786) is a benefit of Texas Medicaid when medically necessary with prior authorization.

Reslizumab is an injectable drug that is FDA-approved and indicated for the treatment of clients who are 18 years of age and older and have severe asthma (as defined by the National Heart, Lung, and Blood Institute's Guidelines for the Diagnosis and Management of Asthma) with an eosinophilic phenotype. Clients who are younger than the FDA-approved age will be considered on a case-by-case basis by the TMHP medical director.

Treatment of reslizumab may not be used concurrently with omalizumab or any other interleukin-5 antagonist.

Procedure codes J2182, J2786 and J2357 may not be billed in any combination for the same date of services by any provider.

Providers may not bill for an office visit if the only reason for the visit is a reslizumab injection.

### 7.54.5 Prior Authorization for Omalizumab, Benralizumab, Mepolizumab, and Reslizumab

When requesting prior authorization, the exact dosage must be included with the request using omalizumab (procedure code J2357), benralizumab (J0517), mepolizumab (procedure code J2182), or reslizumab (procedure code J2786). Prior authorization for omalizumab will be considered for clients who are 6 years of age or older with moderate to severe asthma and for clients who are 12 years of age or older with CIU. Prior authorization for reslizumab may be approved for clients who are 18 years of age or older with severe asthma. Prior authorization for benralizumab will be considered for clients who are 12 years of age and older with severe asthma with eosinophilic phenotype.

Prior authorization for mepolizumab will be considered for the following:

- Clients who are 6 years of age or older and have severe asthma with an eosinophilic phenotype (as defined by the National Heart, Lung, and Blood Institute's Guidelines for the Diagnosis and Management of Asthma)

- Adult clients with eosinophilic granulomatosis with polyangiitis

- Clients who are 12 years of age or older with hypereosinophilic symptoms (HES) for 6 months or longer without identifiable non-hematologic secondary cause

Prior authorization approvals for omalizumab, benralizumab, mepolizumab, or reslizumab are for intervals of six months at a time. Clients must be compliant with their omalizumab, benralizumab, mepolizumab, or reslizumab regimen in order to qualify for additional prior authorizations. The provider must submit a statement documenting compliance with the requests for each renewal.

**Referato:** Subsection 3.1, “Prior Authorization Requests” in this handbook for additional prior authorization information.

### 7.54.6 Prior Authorization Criteria for Chronic Idiopathic Urticaria

Prior authorization for omalizumab will be considered for clients who are 12 years of age or older with CIU. Documentation supporting medical necessity for treatment of CIU with omalizumab must be submitted with the request and include all of the following:

- Documented failure of, or contraindication to, antihistamine and leukotriene inhibitor therapies.

- Evidence of an evaluation that excludes other medical diagnoses associated with chronic urticaria.
7.54.7 Prior Authorization Criteria for Asthma — Moderate to Severe (Omalizumab) and Severe (Benralizumab, Mepolizumab, and Reslizumab)

Requests for prior authorization must be submitted by the treating physician to the Special Medical Prior Authorization (SMPA) department by mail or approved electronic method using the SMPA request form.

Documentation supporting medical necessity for treatment of asthma with omalizumab, benralizumab, mepolizumab, or reslizumab must be submitted with the request and must indicate the following:

- Symptoms are inadequately controlled with use of one of the following combination therapies:
  - 12 months of high-dose inhaled corticosteroid (ICS) given in combination with a minimum of 3 months of controller medication (either a long-acting beta2-agonist [LABA], leukotriene receptor antagonist [LTRA], or theophylline), unless the individual is intolerant of, or has a medical contraindication to these agents; or
  - 6 months of ICS with daily oral glucocorticoids given in combination with a minimum of 3 months of controller medication (a LABA, LTRA, or theophylline), unless the individual is intolerant of, or has a medical contraindication to these agents.

  **Note:** Exceptions to the criteria above will be considered on a case-by-case basis, which will require a letter from the prescribing provider stating the medical necessity for omalizumab, benralizumab, mepolizumab, or reslizumab, the client’s asthma severity level, and the duration of current and past therapies and lack of asthma control. Consideration for these exceptions will be reviewed by the TMHP medical director.

- Pulmonary function tests must have been performed within a three-month period and be documented for all clients.

  **Note:** Exceptions may be considered with documentation of medical reasons explaining why pulmonary function tests cannot be performed.

- Client is not currently smoking.

7.54.7.1 Eosinophilic Granulomatosis with Polyangiitis

Prior authorization for mepolizumab will be considered for adult clients who are 18 years and older with eosinophilic granulomatosis with polyangiitis (EGPA).

Documentation supporting medical necessity for treatment of EGPA with mepolizumab must be submitted with the request and meet all of the following:

- Diagnosis of EGPA
- Medical history of asthma
- Presence of at least 2 of the following EGPA characteristics below:
  - Histopathological findings of eosinophilic vascularitis, perivascularitis eosinophilic infiltration or eosinophil-rich granulomatous inflammation
  - Neuropathy
  - Pulmonary infiltrates, non-fixed; Sino-nasal abnormality
  - Cardiomyopathy
  - Glomerulonephritis
  - Alveolar hemorrhage
  - Palpable purpura
  - Anti-neutrophils cytoplasmic antibody
• Refractory disease or has had a history of EGPA relapse within the past 2 years from the requested date of service.
• Attestation from prescriber that client is on a stable dose of corticosteroids.

7.54.7.2 Hypereosinophilic Syndrome (HES)
Prior authorization for mepolizumab will be considered for clients who are 12 years of age and older with hypereosinophilic syndrome for 6 months or longer without non-hematologic secondary cause.

Documents supporting medical necessity for treatment of HES in clients who are 12 years of age or older with mepolizumab must be submitted with the request and meet all of the following:
• Diagnosis of HES for 6 months or longer without any non-hematologic secondary cause
• History of 2 or more HES flares (flare defined as worsening clinical symptoms or blood eosinophil counts requiring an increase in prior therapy) within the past 12 months prior to the initiation of mepolizumab therapy
• Prescriber’s attestation that client has been on a stable dose of HES therapy which includes, but not limited to corticosteroids, immunosuppressive and cytotoxic therapy

7.54.7.3 Mepolizumab
The following additional documentation for treatment with mepolizumab must also be submitted:
• One of the following blood eosinophil counts in the absence of other potential causes of eosinophilia, including hypereosinophilic syndromes, neoplastic disease, and known or suspected parasitic infection:
  • Greater than or equal to 150 cells/microliter at initiation of therapy; or
  • Greater than or equal to 300 cells/microliter within 12 months prior to initiation of therapy

Note: 1 microliter (ul) is equal to 1 cubic millimeter (mm3)

7.54.7.4 Omalizumab
The following additional documentation for treatment with omalizumab also must be submitted:
• Positive skin test or RAST to a perennial (not seasonal) aeroallergen within the past 36 months
• Total IgE level greater than 30 IU/ml but less than 1300 IU/ml within the past 12 months

7.54.7.5 Benralizumab
The following additional documentation for treatment with benralizumab must also be submitted with the initial prior authorization request:
• Documented diagnosis of severe eosinophilic asthma
• Blood eosinophil count greater than or equal to 150 cells/microliter before the initiation of therapy, in the absence of other potential causes of eosinophilia including hypereosinophilic syndromes, neoplastic disease, and known or suspected parasitic infection

Note: 1 microliter (ul) is equal to 1 cubic millimeter (mm3)

7.54.7.6 Reslizumab
The following additional documentation for treatment with reslizumab must also be submitted:
• Has an eosinophilic phenotype as determined by blood eosinophils of 400 cells/microliter or higher prior to initiation of therapy (within 3-4 weeks of dosing).

Note: 1 microliter (ul) is equal to 1 cubic millimeter (mm3).
• When requesting prior authorization, the exact dosage must be included with the request.
7.54.8 Requirements for Continuation of Therapy

For continuation of therapy with omalizumab, benralizumab, mepolizumab, or reslizumab after 6 continuous months, the requesting provider must submit the following documentation of the client’s compliance and satisfactory clinical response to omalizumab, benralizumab, mepolizumab, or reslizumab:

• Documentation of clinical improvement must include one or more of the following:
  • Decreased utilization of rescue medications; or
  • Increase in predicted FEV1 (forced expiratory volume) from pretreatment baseline; or
  • Reduction in reported asthma-related symptoms, as evidenced by decreases in frequency or magnitude of one or more of the following symptoms:
    • Asthma attacks
    • Chest tightness or heaviness
    • Coughing or clearing throat
    • Difficulty taking deep breath or difficulty breathing out
    • Shortness of breath
    • Sleep disturbance, night wakening, or symptoms upon awakening
    • Tiredness
    • Wheezing/heavy breathing/fighting for air, and
  • Member has not exhibited symptoms of hypersensitivity or anaphylaxis (bronchospasm, hypotension, syncope, urticaria, and/or angioedema) after administration of omalizumab, benralizumab, mepolizumab, or reslizumab.

After lapses in treatment of 3 months or greater, prior authorization requests submitted with documentation will be reviewed by the TMHP medical director.

Requests for clients who do not meet the above criteria will be reviewed for medical necessity by the TMHP medical director.

7.55 Moxetumomab Pasudotox-tdfk (Lumoxiti)

Moxetumomab pasudotox-tdfk (Lumoxiti) (procedure code J9313) is a benefit of Texas Medicaid when indicated for the treatment of relapsed or refractory hairy cell leukemia in adults who have received at least two prior systemic therapies, including treatment with a purine nucleoside analog (PNA).

Moxetumomab pasudotox-tdfk (Lumoxiti) must be prescribed by, or in consultation with, an oncologist or hematologist.

Moxetumomab pasudotox-tdfk is not a benefit for clients who have renal impairment with CrCl < 29 ml/min.

7.55.1 Prior Authorization Requirements

Prior authorization is required for moxetumomab pasudotox-tdfk (Lumoxiti) (procedure code J9313). Prior authorization may be granted for a duration of 6 months (6 cycles).

Prior authorization approval for moxetumomab pasudotox-tdfk (Lumoxiti) infusion will be considered if all the following criteria are met:

• The client is 18 years of age or older.
• The client has a confirmed diagnosis of hairy cell leukemia (diagnosis codes C9140 and C9142).
• The client has relapsed or refractory disease.
• The client has received at least two prior systemic therapies, including treatment with a PNA.
• The client does not have severe renal impairment, defined as creatinine clearance (CrCl) or 29 ml/min or less.

Refer to: Subsection 3.1, “Prior Authorization Requests” in this handbook for additional prior authorization information.

7.56 Natalizumab

Procedure code J2323 is a benefit when billed with one of the following diagnosis codes:

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<thead>
<tr>
<th>Diagnosis Codes</th>
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<td>G35</td>
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<tr>
<td>K50813</td>
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<tr>
<td>K50918</td>
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</tbody>
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7.57 Onasemnogene abeparvovec-xioi (Zolgensma)

Onasemnogene abeparvovec-xioi (Zolgensma) is a benefit of Texas Medicaid, with prior authorization. Claims for onasemnogene abeparvovec-xioi (Zolgensma) must be submitted with unlisted procedure code J3399.

Onasemnogene abeparvovec-xioi (Zolgensma) (procedure code J3399) is limited to one treatment per lifetime, per client.

Onasemnogene abeparvovec-xioi (Zolgensma) is a one-time infusion therapy indicated for the treatment of a client who meets all of the following requirements:
• Has spinal muscular atrophy (SMA)
• Has biallelic mutations in the survival motor neuron 1 (SMN1) gene
• Is 24 months of age or younger

Onasemnogene abeparvovec-xioi (Zolgensma) is not a benefit for clients with a tracheostomy or invasive ventilator support.

Onasemnogene abeparvovec-xioi (Zolgensma) must be prescribed by, or in consultation with, a board-certified neurologist or pediatric neurologist who is familiar with the diagnosis and management of spinal muscular atrophy. The consultation must include the neurologist’s name, credentials, and contact information.

7.57.1 Prior Authorization Requirements

To be considered for the approval of a one-time intravenous infusion, prior authorization requests for onasemnogene abeparvovec-xioi (Zolgensma) (unlisted procedure code J3399) must be submitted with a Special Medical Prior Authorization (SMPA) Request Form.

All of the following must be included in Section C (located under Statement of Medical Necessity) of the Special Medical Prior Authorization (SMPA) Request Form:
• Documentation of the client’s dosage
• Administration schedule
• Number of injections to be administered during the prior authorization period
• Requested units per injection
• Dosage calculation

The Special Medical Prior Authorization (SMPA) Request Form must be completed, signed, and dated by the prescribing provider. The completed form must be maintained by the prescribing provider in the client’s medical record and is subject to retrospective review. The form will not be accepted beyond 90 days from the date of the prescribing provider’s signature.

7.57.2 * Documentation Requirements

The prior authorization request for an onasemnogene abeparvovec-xioi (Zolgensma) single-dose intravenous infusion must include documentation of all of the following:

• Client is 24 months of age or younger.
• Medical record supports any of the following mutation or deletion of genes in chromosome 5q:
  • Homozygous gene deletion of the SMN1 gene (e.g., absence of SMN1 gene)
  • Homozygous mutation of the SMN1 gene (e.g., biallelic mutation of exon 7)
  • Compound heterozygous mutation in the SMN1 gene (e.g., deletion of SMN1 exon 7 [allele 1] and mutation of SMN1 [allele 2])
• Confirmed diagnosis of Type I SMA (diagnosis code G120) based on gene mutation analysis with biallelic SMN1 mutation (deletion or point mutation) and 3 or less copies of SMN2.
• [Revised] Evaluation of motor skill and function must be documented using a standardized test. However, it is not a prerequisite of therapy and should not delay treatment. Standardized testing tools that may be used to evaluate motor skill/function include, but are not limited to:
  • [Revised] Children’s Hospital of Philadelphia Infant Test of Neuromuscular Disorder (CHOP-INTEND) score
  • [Revised] Bayley scale of infant and toddler development screening test
  • [Revised] WHO Multicenter Growth Reference Study (WHO MGRS)
• Baseline documentation supports an AAV9 antibody titer of 1:50 or lower, as determined by enzyme-linked immunosorbent assay (ELISA) binding immunoassay.
• Physician attestation supports that the client has not received prior onasemnogene abeparvovec-xioi (Zolgensma) therapy.

[Revised] If nusinersen (Spinraza) (procedure code J2326) or risdiplam (Evrysdi) have been previously prescribed, the prescriber must provide documentation of one of the following before switching to onasemnogene abeparvovec-xioi (Zolgensma) therapy:

• [Revised] Evidence of clinical deterioration (e.g., decreased physical function and motor skill/function test scores) while on nusinersen (Spinraza) or risdiplam (Evrysdi) therapy
• [Revised] Prescriber’s attestation that nusinersen (Spinraza) or risdiplam (Evrysdi) therapy has been discontinued

7.58 Panhematin

Panhematin (procedure code J1640) is a benefit of Texas Medicaid and is limited to diagnosis code E8021.

7.59 Patisiran (Onpattro)

Patisiran (Onpattro) is a benefit of Texas Medicaid with prior authorization for the treatment of the polyneuropathy of hereditary transthyretin-mediated (hATTR) amyloidosis in adults.
Prior authorization requests for patisiran (Onpattro) (procedure code J0222) must be submitted with a Special Medical Prior Authorization (SMPA) Request Form and may be approved for 12 months per prior authorization request.

For initial therapy, all of the following criteria must be met:

- The client is 18 years of age or older.
- The client has a diagnosis of hATTR amyloidosis (diagnosis code E851), supported by the following:
  - Transthyretin (TTR) mutation proven by genetic testing
  - Clinical signs and symptoms of the disease (e.g., peripheral/autonomic neuropathy, motor disability)
- The client will not receive patisiran (Onpattro) therapy in combination with other polyneuropathy hATTR amyloidosis therapies (e.g., inotersen or tafamidis meglumine).
- The client has not had a liver transplant.

For renewal or continuation of therapy, all of the following criteria must be met:

- The client has previously received treatment with patisiran (Onpattro) without an adverse reaction.
- The client has a positive clinical response to patisiran (Onpattro) (e.g., improved neurologic impairment, improved motor function, slowing of disease progression).

Documentation of the client’s dosage, administration schedule, number of injections to be administered during the prior authorization period, requested units per injection, and the dosage calculation must be submitted in Section C of the Special Medical Prior Authorization (SMPA) Request Form under Statement of Medical Necessity.

Refer to: Subsection 3.1, “Prior Authorization Requests” in this handbook for additional prior authorization information.

### 7.60 Plazomicin

Plazomicin (procedure code J0291) is a benefit of Texas Medicaid for clients who are 18 years of age or older.

### 7.61 Porfimer (Photofrin)

Porfimer (procedure code J9600) is a benefit of Texas Medicaid and is limited to the following diagnosis codes:

<table>
<thead>
<tr>
<th>Diagnosis Codes</th>
</tr>
</thead>
<tbody>
<tr>
<td>C153</td>
</tr>
</tbody>
</table>

### 7.62 Ravulizumab-cwvz (Ultomiris)

Ravulizumab-cwvz (procedure code J1303) is a benefit of Texas Medicaid and is limited to diagnosis codes D588, D593, D594, D595, and D598.

Procedure code J1303 is restricted to clients who are 18 years of age or older when billed with diagnosis code D595.

### 7.63 Risperidone (Perseris)

Risperidone (procedure code J2798) is a benefit of Texas Medicaid for clients who are 18 years of age or older.
7.64 Rituximab-Abbs, (Truxima)
Rituximab-abbs (Truxima) (procedure code Q5115) is a benefit of Texas Medicaid for clients who are 18 years of age or older.

7.65 Romosozumab
Romosozumab (procedure code J3111) is a benefit of Texas Medicaid for clients who are 18 years of age or older.

7.66 Sumatriptan succinate (Imitrex)
Procedure code J3030 is a benefit when billed with one of the following diagnosis codes:

7.67 Tagraxofusp-erzs (Elzonris)
Tagraxofusp-erzs (Elzonris) (procedure code J9269) is a CD 123-directed cytotoxin indicated for the treatment of blastic plasmacytoid dendritic cell neoplasm (BPDCN) in adults and pediatric clients who are 2 years of age and older.

Tagraxofusp-erzs (Elzonris) (procedure code J9269) is a benefit of Texas Medicaid with prior authorization.

Tagraxofusp-erzs (Elzonris) must be prescribed by, or in consultation with, a hematologist or oncologist.

7.67.1 Prior Authorization Requirements
Prior authorization is required for tagraxofusp-erzs (Elzonris) (procedure code J9269) and may be approved for a duration of every 12 months.

Prior authorization approval for tagraxofusp-erzs (Elzonris) infusion will be considered once all of the following criteria are met for initial therapy:

- Client has a diagnosis of blastic plasmacytoid dendritic cell neoplasm (diagnosis code C864) excluding, acute promyelocytic leukemia (APL, FAB, M3)
- Client is 2 years of age or older
- Client has a CD-123 positive/expressing disease
- Client has adequate serum albumin level and baseline organ function, including cardiac, renal, and hepatic function prior to each course of therapy:
  - Baseline serum albumin level of 3.2 g/dl or greater
  - Left ventricular ejection fraction (LVEF) of 40% or greater
  - Serum creatinine (Scr) of 1.5 mg/dl or less
  - Bilirubin level of 1.5 mg/dl or less
- For renewal or continuation therapy, the client must meet all of the following criteria:
• Client continues to meet initial approval criteria
• Client has previously received treatment with tagraxofusp-erzs (Elzonris) with absence of drug toxicity (i.e. capillary leak syndrome, severe hepatotoxicity, and nephrotoxicity)
• Client has a positive clinical response demonstrated by disease stabilization

Refer to: Subsection 3.1, “Prior Authorization Requests” in this handbook for additional prior authorization information.

7.68 Teprotumumab-trbw (Tepezza)
Teprotumumab-trbw (Tepezza) (procedure code J3241) is a benefit of Texas Medicaid and prior authorization is required. Teprotumumab-trbw (Tepezza) is indicated for the treatment of thyroid eye disease (TED) and must be prescribed by, or in consultation with, an ophthalmologist or endocrinologist.

7.68.1 Prior Authorization Requirements
Prior authorization requests for procedure code J3241 must be submitted with a Special Medical Prior Authorization (SMPA) Request Form.

The client must meet all of the following requirements for approval of Teprotumumab-trbw (Tepezza):

• The client has a diagnosis of Graves’ disease associated with active TED.
• The client has active TED with a clinical activity score (CAS) of 4 or greater.
• The client is euthyroid, or the client has either mild hypothyroidism or mild hyperthyroidism.
• The client has no history of prior surgical intervention for TED and does not plan to have surgical treatment while on Teprotumumab-trbw (Tepezza).
• The client may not exceed the course of eight total infusions per lifetime.

7.68.2 Exclusions
Teprotumumab-trbw (Tepezza) should not be used in pregnancy as it may potentially lead to fetal loss. Females of reproductive potential should use effective contraception prior to initiation, during treatment with Teprotumumab-trbw (Tepezza) and for 6 months after the last dose of Teprotumumab-trbw (Tepezza).

7.69 Thyrotropin alpha for injection (Thyrogen)
Thyrotropin alpha for injection (Thyrogen) (procedure code J3240) is a benefit of Texas Medicaid and is limited to the following diagnosis codes:

<table>
<thead>
<tr>
<th>Diagnosis Codes</th>
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<tbody>
<tr>
<td>C323</td>
</tr>
<tr>
<td>D449</td>
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<tr>
<td>E049</td>
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7.70 Tildrakizumab (Ilumya)
Tildrakizumab (procedure code J3245) is a benefit of Texas Medicaid for clients who are 18 years of age or older, and is limited to diagnosis code L400.
7.71 **Trastuzumab**
Procedure code J9355 is a benefit of Texas Medicaid. Reimbursement for this drug is considered when it is used as a single agent for the treatment of clients who have metastatic breast cancer whose tumors overexpress the Her-2 protein and who have received one or more chemotherapy regimens for their metastatic disease. Trastuzumab may also be reimbursed when:

- Used in combination with paclitaxel for the treatment of clients who have metastatic breast cancer whose tumors overexpress the Her-2 protein and who have not received chemotherapy for their metastatic disease.
- Used as part of a treatment regimen containing doxorubicin, cyclophosphamide, and paclitaxel for the adjuvant treatment of clients who have Her-2-overexpressing, node-positive breast cancer.

Trastuzumab is a benefit for clients whose tumors have Her-2 protein overexpression.

When billing for the test used to determine whether a client overexpresses the Her-2 protein, use procedure code 83950. Diagnosis of overexpression of the Her-2 protein must be made before Texas Medicaid will consider reimbursement for trastuzumab. This test may be reimbursed only once in a client’s lifetime to the same provider. An additional test by the same provider requires documentation to support the medical necessity.

7.72 **Triamcinolone Acetonide**
Procedure code J3304 is a benefit of Texas Medicaid and is restricted to the following diagnosis codes:

<table>
<thead>
<tr>
<th>Diagnosis Codes</th>
</tr>
</thead>
<tbody>
<tr>
<td>M170 M1711 M1712 M172 M1731 M1732 M174 M175</td>
</tr>
</tbody>
</table>

7.73 **Valrubicin sterile solution for intravesical instillation (Valstar)**
Procedure code J9357 valrubicin sterile solution for intravesical instillation (Valstar), is a benefit for clients with the diagnosis of bladder cancer in situ who have been treated unsuccessfully with BCG therapy and have an unacceptable morbidity or mortality risk if immediate cystectomy should be performed. Documentation of diagnosis and treatment must be submitted with the claim.

7.74 **Vitamin B12 (Cyanocobalamin) Injections**
Vitamin B12 injections are a benefit of Texas Medicaid. Vitamin B12 injections should only be considered for clients with conditions that are refractory to, or have a contraindication to, oral therapy.

Vitamin B12 injections may be considered for the following indications:

- Dementia secondary to vitamin B12 deficiency
- Resection of the small intestine
- Schilling test (vitamin B12 absorption test)

Procedure code J3420 must be used when billing for Vitamin B12 (cyanocobalamin) injections. Vitamin B12 (cyanocobalamin) injections are limited to the following diagnosis codes:

<table>
<thead>
<tr>
<th>Diagnosis Codes</th>
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</thead>
<tbody>
<tr>
<td>B700 D510 D511 D512 D513 D518 D520 D521</td>
</tr>
<tr>
<td>D528 D529 D531 D649 E538 E710 E71110 E71111</td>
</tr>
<tr>
<td>E71118 E71120 E71121 E71128 E7119 E712 E7210 E7211</td>
</tr>
<tr>
<td>E7212 E7219 E723 E7251 E7259 E7281 E7289 G621</td>
</tr>
<tr>
<td>G63 H4611 H4612 H4613 H463 K900 K901 K902</td>
</tr>
</tbody>
</table>
Claims that are denied for indications or other diagnosis codes may be considered on appeal with documentation of medical necessity. For the list of diagnosis codes above, documentation in the medical record must include rationale as to why the client was unable to be treated with oral therapy.

### 7.75 Voretigene neparvovec-rzyl (Luxturna)

Voretigene neparvovec-rzyl (Luxturna) is an adeno-associated virus vector-based gene therapy indicated for the treatment of patients with confirmed biallelic RPE65 mutation-associated retinal dystrophy, who have viable retinal cells in each eye as determined by the treating physician.

Luxturna (procedure code J3398) is a benefit of Texas Medicaid for clients who are 1 year through 65 years of age with prior authorization.

Voretigene neparvovec-rzyl (Luxturna) must be prescribed and administered by a retinal surgeon at an ocular gene therapy treatment center with experience performing intraocular surgery.

#### 7.75.1 Prior Authorization Requirements

Prior authorization is not required for the physician services associated with administration of Luxturna. Physician services include the procedural costs and the associated supplies for administration of the medication.

Prior authorization is required for voretigene neparvovec-rzyl (Luxturna) (procedure code J3398).

For situations in which voretigene neparvovec-rzyl (Luxturna) is being dispensed by a pharmacy via white bagging, the prescribing provider must provide the dispensing DME pharmacy the authorization approval number. The dispensing DME pharmacy may not request prior authorization.

The DME Pharmacy provider billing for voretigene neparvovec-rzyl (Luxturna) will be responsible for coordinating with the rendering provider to obtain the prior authorization request approval number.

The requesting provider (physician or hospital) may coordinate with the DME Pharmacy provider for the initial request for voretigene neparvovec-rzyl (Luxturna). DME Pharmacy providers may assist in providing necessary information, such as their NPI number, fax number, and business address, to the requesting provider. However, the Special Medical Prior Authorization (SMPA) form must be signed, dated, and submitted by the Medicaid-enrolled requesting provider, not the DME Pharmacy provider.

The dispensing pharmacy must submit the authorization approval number when billing for the drug. Reimbursement for dispensing of the drug by the pharmacy may not occur unless an approved prior authorization for voretigene neparvovec-rzyl (Luxturna) is in place.

**Refer to:** Subsection 10.5.1, “Pharmacy Delivery Method for Clinician-Administered Drugs” in this handbook for additional information on the “white bagging” delivery method.

Subsection 3.1, “Prior Authorization Requests” in this handbook for additional prior authorization information.

Prior authorization request for Luxturna injections will be considered when all of the following criteria are met:

- Client is 1 year of age through 65 years of age
- A documented diagnosis of a confirmed biallelic RPE65 mutation-associated retinal dystrophy (e.g., Leber’s congenital amaurosis subtype 2, retinitis pigmentosa, or early onset severe retinal dystrophy)

### Diagnosis Codes

<table>
<thead>
<tr>
<th>K903</th>
<th>K9041</th>
<th>K9049</th>
<th>K9089</th>
<th>K909</th>
<th>K911</th>
<th>K912</th>
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Claims that are denied for indications or other diagnosis codes may be considered on appeal with documentation of medical necessity. For the list of diagnosis codes above, documentation in the medical record must include rationale as to why the client was unable to be treated with oral therapy.
• Genetic testing documenting biallelic mutations of the RPE65 gene
• Systemic corticosteroids equivalent to prednisone 1 mg/kg/day are administered for a total of 7 days, starting 3 days before administration of voretigene neparvovec-rzyl to each eye and followed by a tapering dose
• Client has viable retinal cells in each eye as determined by the treating physician and assessed in the previous 6 months. Verification of viable retinal cells must be documented and evident by one of the following:
  • An area of retina within the posterior pole of greater than 100 µm thickness shown on optimal coherence tomography (OCT)
  • Greater than or equal to 3 disc areas of retina without atrophy or pigmentary degeneration within the posterior pole
    • Remaining visual field within 30 degrees of fixation as measured by a III4e isopter or equivalent
  • Prescribed and administered by retinal surgeon at an ocular gene therapy treatment center with experience performing intraocular surgery
• Patient has not previously received RPE65 gene therapy in intended eye
• Injection of the second eye must be administered at least 6 days after the first eye
• Have not had intraocular surgery within 6 months in either eye indicated for treatment

Benefit not to exceed more than 1 injection per eye per lifetime.

Authorization is valid for a period of 6 months from approval.

All services outlined in this section are subject to retrospective review to ensure that the documentation in the client’s medical record supports the medical necessity of the services provided.

7.75.2 Exclusions
Luxturna is not a benefit for patients who have previously received RPE65 gene therapy and who do not have viable retinal cells in each eye as determined by the treating physician.

8 Claims Filing Information

Claims for clinician-administered drugs must be submitted to TMHP in an approved electronic format or on the CMS-1500 paper claim form. Providers may purchase CMS-1500 paper claim forms from the vendor of their choice. TMHP does not supply them.

When completing a CMS-1500 paper claim form, all required information must be included on the claim, as information is not keyed from attachments. Superbills and itemized statements are not accepted as claim supplements.

Referred: “Section 3: TMHP Electronic Data Interchange (EDI)” (Vol. 1, General Information) for information on electronic claims submissions.

“Section 6: Claims Filing” (Vol. 1, General Information) for general information about claims filing.

Subsection 6.5, “CMS-1500 Paper Claim Filing Instructions” in “Section 6: Claims Filing” (Vol. 1, General Information) for instructions on completing paper claims. Blocks that are not referenced are not required for processing by TMHP and may be left blank.

8.1 JW Modifier Claims Filing Instructions
Providers must not use the JW modifier for medications manufactured in a multi-dose vial format.
Providers must choose the most appropriate vial size(s) required to prepare a dose to minimize the discarded portion of the vial payable.

Claims considered for reimbursement must not exceed the package size of the vial used for preparation of the dose. Providers must not bill for vial contents overfill.

Providers must not use the JW modifier when the actual dose of the drug or biological administered is less than the billing unit.

**Example:** One billing unit for a drug is equal to 10mg of the drug in a single use vial. A 7mg dose is administered to a client while 3mg of the remaining drug is discarded. The 7mg dose is billed using one billing unit that represents 10mg on a single line item. The single line item of 1 unit would process for payment of the total 10mg of drug administered and discarded. Billing another unit on a separate line item with the JW modifier for the discarded 3mg of drug is not permitted because it would result in overpayment. Therefore, when the billing unit is equal to or greater than the total actual dose and the amount discarded, the use of the JW modifier is not permitted.

Reimbursement for JW modifier claims is only available for drugs covered in an outpatient setting.

Inpatient and diagnostic radiopharmaceuticals claims are not eligible for reimbursement, and may not include the JW modifier.

Coverage is for “buy and bill” providers only. Specialty pharmacies billing through the medical benefit must not submit claims with a JW modifier because they are unaware of how much the provider administered or discarded.

Federally Qualified Health Center (FQHC) and Rural Healthcare Clinic RHC) are not eligible to bill with the JW modifier because these providers do not bill for the coverage of drugs or biologicals separately.

Critical Access Hospitals (CAH) are eligible to bill with the JW modifier because these providers bill for the coverage of drugs or biologicals separately.

The JW modifier is not allowed for medications prepared in an institutional setting via “batch processing or bulk productions” methods. An example of a batch processing method is when a hospital or repackaging facility produces multiple non-patient specific doses of medications in advance of anticipated use. These preparations are labeled and distributed with client specific information only when orders are received. Because these doses may be recycled for other client use, they are not eligible.

Providers may utilize automatic systems to calculate dose and discard amounts. However, providers must continue to document the exact usage/discard accurately.

Providers must enter the dose administered (used portion) line item detail of the CAD and also enter the dose discarded (unused portion) line item detail of the CAD on the same claim. The dose discarded (unused portion) line item detail must include the JW modifier to be considered for reimbursement.

When billing for reimbursement of wastage on an outpatient claim, the HCPCS and Current Procedural Terminology (CPT) code should always be provided along with the revenue code.

### 9 Pharmacy Benefit

The Texas Drug Code Index, or formulary (or list of available drugs), includes non-legend (over-the-counter) drugs. Additionally, certain supplies and select vitamin and mineral products are also available as a pharmacy benefit. Some drugs are subject to one or both types of prior authorization, clinical and non-preferred. VDP does not reimburse claims for nutritional products (enteral or parenteral), medical supplies, or equipment other than a limited set of home health supplies.
The PDL is arranged by drug therapeutic class, and contains a subset of many, but not all, drugs that are on the Medicaid formulary. Most drugs are identified as preferred or non-preferred. Drugs listed on the PDL as preferred or not listed at all are available without prior authorization unless there is a clinical prior authorization associated with that drug. For more information about prior authorization, refer to Subsection 12, “Pharmacy Prior Authorization” in this handbook.

9.1 Formulary Search

The VDP Formulary Search is an online tool available to health-care providers to help clients get access to medications.

Users can search by either brand or generic name of the drug or product, the 11-digit national drug code (NDC), the PDL drug class, or HCPCS description (for products).

Detailed filters allow searches as follows:

- **By program:**
  - Medicaid
  - CHIP
  - CSHCN Services Program
  - HTW Program
  - HTW Plus Program
  - KHC Program (e.g., CHIP, CSHCN, HTW, and KHC)

- **By prior authorization status:**
  - PDL prior authorization
  - Clinical prior authorization

- **By drug types:**
  - Family planning
  - Diabetic supplies
  - Injectable drugs
  - LARCs
  - Over-the-counter (OTC)
  - Drugs requiring 90% utilization before refilling (refer to section 10.4)

- **By product types:**
  - Flu vaccines
  - Mosquito repellents
  - COVID-19 vaccines

**Refer to:** The Formulary Search on the VDP website for more information.

The Texas Medicaid formulary and Preferred Drug List are available on the Epocrates drug information system. Epocrates (epocrates.com) is the publisher of mobile device software applications designed to provide information about drugs to doctors and other health care professionals.
Once registered, providers can utilize Epocrates Rx, the free drug reference, and search for brand, generic, or over-the-counter medicines. Providers can select the “Texas Medicaid” formulary option, allowing searches by drug name to find products identified as preferred or non-preferred or are subject to clinical prior authorization criteria. In addition to listing a drug’s preferred status, Epocrates includes drug monographs, dosing information, and warnings.

Epocrates does not mirror the HHSC designations differentiated by dosage form. In these situations, the designation includes an explanatory message.

9.2 Vitamin and Mineral Products
Pharmacies enrolled with VDP can dispense vitamin and mineral products to clients who are 20 years of age and younger and enrolled in traditional Medicaid. These products are also available to clients enrolled in Medicaid managed care, if the dispensing pharmacy is contracted with the client’s health plan.

To expedite pharmacy claim processing for vitamin and mineral products, prescribing providers are encouraged to include the diagnosis on the prescription.

The list of products that can be dispensed at a pharmacy and information about the provision of these products to clients enrolled in fee-for-service can be found on the VDP Product Search website at https://www.txvendordrug.com/formulary/formulary-search. Users can search by product NDC or name or the HCPCS description.

For clients enrolled in Medicaid managed care or CHIP, claims are submitted to the clients’ health plans. Pharmacy staff must work with the health plan’s pharmacy benefit manager to determine the billing requirements, reimbursement rates, and coverage limitations for these products.

9.3 Home Health Supplies
Pharmacies enrolled with VDP can dispense a limited set of home health supplies that are commonly found in a pharmacy to clients enrolled in traditional Medicaid. These supplies are also available to clients enrolled in Medicaid managed care, provided the dispensing pharmacy is contracted by the client’s health plan.

The list of supplies that can be dispensed at a pharmacy and information about the provisions of these supplies for clients enrolled in traditional Medicaid can be found on the VDP Product Search website at https://www.txvendordrug.com/formulary/formulary-search. Users can search by product NDC or name or the HCPCS description.

Providers should contact the appropriate health plan or pharmacy benefit manager for more information about providing these supplies to clients enrolled in managed care.

9.4 Long-Acting Reversible Contraception Products
Long-acting reversible contraception (LARC) products are available to clients through either a Medicaid or HTW pharmacy or medical benefit.

The list of LARC drugs dispensed at a pharmacy can be found on the VDP Drug Search website at https://www.txvendordrug.com/formulary/formulary-search. Users can select the “LARC” check-box for a list of all drugs.

Providers can obtain LARC products with no upfront cost by submitting a completed and signed prescription request form to certain specialty pharmacies. The specialty pharmacy will dispense the LARC product by shipping it to the practice address in care of the client and bill Medicaid or HTW for the product. Providers can only bill for product administration at the time of service. LARC products obtained by providers from specialty pharmacies must be returned if unused and unopened.
9.4.1 Product Billing
The specialty pharmacy bills Medicaid or HTW for LARC products when obtained from a specialty pharmacy. Providers will continue to bill Medicaid or HTW for insertion of the LARC product. Providers may only bill for the LARC product if it was obtained through the buy-and-bill process.

For clients enrolled in traditional Medicaid (fee-for-service (FFS)) or HTW, providers bill HHSC for the insertion of the LARC product using procedure code 58300.

For clients enrolled in Medicaid managed care, providers bill the patient’s managed care organization for the insertion of the LARC product. Contact the patient’s MCO for specific billing instructions.

9.4.2 Specialty Pharmacy Participation
Providers do not need to enroll with specialty pharmacies to obtain LARC products. Any provider currently enrolled with Medicaid or HTW may prescribe and obtain a LARC product and bill Medicaid or HTW for insertion of the LARC product. The participating pharmacies ship statewide.

Prescribing providers should identify whether the patient is enrolled in traditional Medicaid or managed care. For clients enrolled in managed care, the provider’s office should coordinate with the managed care organization (MCO) to determine which pharmacy should receive the prescription because the MCO may be contracted with a single specialty pharmacy. The name, phone number, and national provider identifier (NPI) is provided for each specialty pharmacy is listed in the “Manufacturer Information” section.

9.4.3 Product Returns and Abandoned Units
Manufacturers offer abandoned unit return programs that allow a provider to return an abandoned LARC product. An “abandoned unit” is an unused and unopened product that was shipped by a participating specialty pharmacy with a prescription label that includes the name of the patient. In order to be returnable, the LARC product should be in its original packaging.

9.4.4 Manufacturer Information

9.4.4.1 Bayer (Kyleena, Mirena, and Skyla)

9.4.4.1.1 Specialty Pharmacy Participation
Products are available from CVS CarePlus Specialty Pharmacy or Walgreens Specialty Pharmacy, both shipping statewide.

Walgreens Specialty Pharmacy
Frisco, TX
1-800-424-9002
NPI 1851463087

CVS Caremark Specialty Pharmacy
Fort Worth, TX
817-336-7281
NPI 1366551848

9.4.4.1.2 Obtaining Products
Providers use Bayer’s Specialty Pharmacy Prescription Request Form. This form is available on the VDP website at txvendordrug.com/formulary/formulary/long-acting-reversible-contraception-products.

Complete the Bayer’s Specialty Pharmacy Prescription Request Form as follows:

- Enter the patient and prescriber information, including the patient’s pharmacy drug benefit and medical insurance information and copies of the patient’s pharmacy benefit and medical insurance cards.
- Complete the prescribing information and keep a copy for future use.
• Identify the drug to be administered in the prescription section, including the appropriate diagnosis code, and sign the prescription. Advanced Practice Registered Nurses, Physician Assistants, and Nurse Practitioners should identify who their collaborative agreement is with to write prescriptions, if necessary.

• Have the patient read and sign the “Patient Authorization” section of the form.

• Fax the form, including the “Patient Authorization” section, to the pharmacy.

• The pharmacy will call the patient to confirm the patient’s intent to receive an intrauterine device (IUD). This is done to limit potentially abandoned IUD units. The pharmacy will not mail the IUD to the provider until confirmation from the patient is received.

9.4.4.1.3 Returning Products

Providers may return abandoned units. The box must be sealed and have been abandoned for at least 60 days (2 months) from the dispensing date but no more than 210 days (7 months) past the fill date. Only LARC products obtained through a specialty pharmacy can be returned through this program.

To return abandoned units complete the Bayer Abandoned Unit Program Return Form. This form is available on the VDP website at txvendordrug.com/formulary/formulary/long-acting-reversible-contraception-products.

The Bayer Abandoned Unit Program Return Form may be submitted by mail or fax as follows:

• Fax the form to the dispensing specialty pharmacy for verification.

• Wait for an authorization number and return mailing label, and then

• Confirm the specialty pharmacy identification number matches the ID number listed on the return authorization form.

• Package the unit in one of the cardboard boxes the drug was initially shipped in or a large envelope.

• Mail the unit to the specialty pharmacy.

9.4.4.2 Merck (Nexplanon)

9.4.4.2.1 Specialty Pharmacy Participation

Nexplanon is available from CVS CarePlus Specialty Pharmacy or Accredo Specialty Pharmacy, both of which ship statewide.

CVS Caremark Specialty Pharmacy
Fort Worth, TX
817-336-7281
NPI 1366551848

Accredo Specialty Pharmacy
Irving, TX
972-929-6800
NPI 1073569034

9.4.4.2.2 Obtaining Products

Providers use Merck’s Nexplanon Direct Service Request Form. Request the form from Merck Customer Support Center for Nexplanon (CSCN) at 1-844-639-4321.

Complete the Merck’s Nexplanon Direct Service Request Form as follows:

• At the top of the first page of the form, check the “Prescription Order” box and select a specialty pharmacy.
• Complete the patient and prescriber information including the patient’s pharmacy drug benefit and medical insurance information. Include copies of the patient’s insurance card and prescription drug card.

• Have the patient read and sign the “Patient Authorization” section.

• Ensure the physician signs both the Dispense as Written and prescriber signature lines, and the appropriate diagnosis code is selected.

• Fax the completed form to the CSCN at 1-844-232-2618.

• The CSCN will forward the prescription to the specialty pharmacy you selected after confirming the benefits of the patient.

9.4.4.2.3 Returning Products

Providers may return an abandoned unit. The Nexplanon box must be sealed and been abandoned for at least 120 days (4 months) from date of dispense but no more than 180 days (6 months) past the fill date.

Complete the Merck Abandoned Unit Program Return Form. This form is available on the VDP website at txvendordrug.com/formulary/formulary/long-acting-reversible-contraception-products.

The Merck Abandoned Unit Program Return Form may be submitted by mail or fax as follows:

• Fax the form to the specialty pharmacy for verification.

• Wait to receive the return identification number from the specialty pharmacy and return mailing label and instructions, which will be provided by TeleRx, Merck’s third-party processor.

• Confirm the specialty pharmacy return identification number matches the ID number listed in the return mailing label provided by TeleRx.

• Package the unit in the box in which the Nexplanon was originally shipped or other appropriately sized shipping box/envelope.

• Mail the unit along with the Merck Abandoned Unit Program for Nexplanon Return Form to Pharma Returns. A prepaid shipping label and address will be provided by TeleRx.

Only LARC products obtained through a specialty pharmacy can be returned through this program.

9.4.4.3 Cooper Surgical (Paragard)

9.4.4.3.1 Specialty Pharmacy Participation

Paragard is available from Biologics by McKesson Specialty Pharmacy, or City Drugs Specialty Pharmacy, both of which ships statewide:

Biologics by McKesson Specialty Pharmacy
Cary, NC 27513
1-888-275-8596
NPI 1487640314

City Drugs
New York, NY 10028
Phone: 1-855-988-4500
NPI 1972880623

9.4.4.3.2 Obtaining Products

To obtain products providers should use the Cooper Surgical Patient Authorization Form and Patient Referral Form:

• Patients should complete the Patient Authorization Form and return to the provider.
• Providers should ensure the box next to “PARAGARD T 380A Qty: 1” is checked on the Teva Patient Authorization Form and Patient Referral Form.

• Forms may be returned to the Specialty Pharmacy by fax to 1-855-215-5315 for Biologics by McKesson or fax to 1-212-988-4501 for City Drugs. Upon receipt of your completed forms, Cooper Surgical will send you written confirmation by fax.

9.4.4.3.3 Returning Products
The original Paragard box must be sealed and have been abandoned at least 90 days since it was shipped.

The following will occur ninety days following shipment:

• Paragard will follow-up with your office to confirm the product was placed into the intended Medicaid client.

• If the product was not placed, the Paragard Specialist will obtain your email address to send you the return shipping label.

• Place the original unused and unopened Paragard unit and original packaging with affixed prescription label into a shipping box. You can reuse the original shipping box.

• Print the return shipping label and ship the unused and unopened Paragard unit back to Paragard as soon as possible.

• For additional questions regarding Cooper Surgical Abandoned Unit Return program contact Paragard at 1-877-727-2427.

9.4.4.3.4 Loss of Client Eligibility
If the client was eligible for Medicaid or HTW on the date of service when the LARC product was prescribed and ordered, but the client loses eligibility before the LARC product is inserted, the provider is not required to return the LARC product. If the client is no longer eligible for Medicaid or HTW, the provider may insert the LARC device, but reimbursement for all care and services provided must be resolved between the provider and the client.

If a provider accepts a client as a private pay client, the provider must advise the client she is accepted as a private pay client at the time the service is provided and is responsible for paying for all services received. In this situation, HHSC strongly encourages the provider to ensure the client signs written notification so there is no question how the client was accepted.

9.4.4.3.5 Questions
Contact the specialty pharmacy for questions related to obtaining LARC products. Further questions may be directed to the client’s health plan or the TMHP provider help line for FFS Medicaid or HTW clients at 1-800-925-9126.

9.4.4.4 Makena

9.4.4.4.1 Pharmacy Benefit
Makena (hydroxyprogesterone caproate injection) requires clinical prior authorization for clients enrolled in traditional Medicaid. Providers should complete the Makena Prior Authorization Request (HHS Form 1345) and submit to the Texas Prior Authorization Call Center.

Health Plans may require prior authorization for Makena. Providers should refer to the appropriate health plan for specific requirements and forms.

9.4.4.4.2 Medical Benefit
Makena and the compounded version of 17P are available as a Medicaid medical benefit. For additional information about the medical benefit, providers can visit the TMHP website at www.tmhp.com or call the TMHP Contact Center at 1-800-925-9126.
9.5  Palivizumab (Synagis)

Palivizumab is available to physicians for administration to clients in Medicaid and the CSHCN Services Program through VDP. The enables physicians to have palivizumab shipped directly to their office from a network pharmacy, and not purchase the drug.

Physicians who obtain palivizumab through VDP may not submit claims to TMHP for the drug. The administering provider may submit a claim to TMHP for an injection administration fee and any medically necessary office-based evaluation and management service provided at time of injection.

9.5.1  Schedule and Forms

Refer: The Synagis page on the VDP website for more information about the current season, including forms and schedule.

Prior authorization request forms are reviewed annually. Providers must use the most current version of the Medicaid Synagis Prior Authorization Request (HHS Form 1033) to submit prior authorization requests. Forms received outside the RSV season schedule will not be processed.

Note: Palivizumab is also available for clients enrolled in the Children with Special Health Care Needs (CSHCN) Services Program. Providers can refer to the CSHCN Services Program Provider Manual for details.

10  Prescribing Information

The federal Patient Protection and Affordable Care Act and the Code of Federal Regulations Title 42 §455.410(b) require all physicians or other professionals who order, refer, or prescribe drugs, supplies and services for a recipient of traditional Medicaid, CHIP, CSHCN, and HTW Program to be enrolled as participating Medicaid providers.

Refer: “Section 1: Provider Enrollment and Responsibilities” (Vol. 1, General Information) for more information.

10.1  Tamper-Resistant Prescription Pads

Providers are required by federal law (Public Law 110-28) to use a tamper-resistant prescription pad when writing a prescription for any drug for Medicaid clients. Pharmacies are required to ensure that all written Medicaid prescriptions submitted for payment to the VDP were written on a compliant tamper resistant pad.

The Centers for Medicare & Medicaid Services (CMS) has stated that special copy-resistant paper is not a requirement for electronic medical records (EMRs) or e-prescribing-generated prescriptions and prescriptions that are faxed directly to the pharmacy. These prescriptions may be printed on plain paper and will be fully compliant if they contain at least one feature from each of the following three categories:

- Prevents unauthorized copying of completed or blank prescription forms
- Prevents erasure or modification of information written on the prescription form
- Prevents the use of counterfeit prescription forms

Two features that can be incorporated into computer-generated prescriptions printed on plain paper to prevent passing a copied prescription as an original prescription are as follows:

- Use a very small font that is readable when viewed at 5x magnification or greater and illegible when copied.
- Use a “void” pantograph accompanied by a reverse “Rx,” which causes a word such as “Void” to appear when the prescription is photocopied.
10.2 Prescription Refills and Expirations

Medicaid prescriptions for non-controlled substances are valid for one year from the date written and up to 11 refills if authorized by prescriber.

Medicaid prescriptions for controlled substances in Schedules III, IV, and V are valid for six months from the date written and up to five refills if authorized by prescriber provider. Controlled substance prescriptions written by advanced practice registered nurses and physicians assistants are valid for 90 days.

Medicaid prescriptions for Schedule II drugs cannot be refilled and must be dispensed within 21 days of the date on which the prescription was written.

Prescriptions for Schedule II drugs may be written as multiples of three for a total of a 90 day supply subject to federal and state law.

Refer to: The VDP Pharmacy Provider Procedure Manual on the VDP website.

10.3 Prescription Monitoring of Controlled Substances

The Texas Prescription Monitoring Program (PMP) collects and monitors prescription data for all Schedule II, III, IV and V controlled substances dispensed by a pharmacy in Texas or to a Texas resident from a pharmacy located in another state. The PMP also provides a venue for monitoring patient prescription history for practitioners and the ordering of Schedule II Texas Official Prescription Forms.

Pharmacies that dispense Schedule II, III, IV, and V drugs are required to report the information directly to the Texas State Board of Pharmacy’s contracted vendor. Prescription data is reported by the prescriber’s federal Drug Enforcement Administration (DEA) number. Prescribers and pharmacies are required by statute to have a valid, active DEA numbers in order to possess, administer, prescribe or dispense controlled substances.

Refer to: The Texas Prescription Monitoring Program page of TSBP website.

10.4 Requirements for Early Refills

A refill is considered too soon, or early, if the client has not used at least 75 percent of the previous fill of the medication.

For clients enrolled in traditional Medicaid or the CSHCN Services Program, a refill for certain controlled substances is considered too soon if the client has not used at least 90 percent of the previous fill of the medication.

Note: Some drugs, such as attention deficit hyperactivity disorder drugs and certain seizure medications, are excluded from this change.

To identify drugs that require 90 percent utilization, refer to the VDP Drug Search and select the “90% Utilization” filter. The returned results will include only those drugs that meet this requirement.

Refer to: The Formulary Search on the VDP website for more information.

Justifications for early refills include, but are not limited to, the following:

- A verifiable dosage increase
- An anticipated prolonged absence from the state
If a client requests an early refill of a drug, the pharmacy must contact VDP to request an override of the early refill restriction. Prescribing providers may be asked to verify the reason for the early refill by the dispensing pharmacy or VDP staff.

**Note:** Providers who are members of Medicaid managed care plans should contact the appropriate health plan or pharmacy benefits manager for specific requirements and processes related to dispensing early refills.

### 10.5 Clinician-Administered Drugs

All Texas Medicaid providers must submit a rebate-eligible NDC for professional or outpatient claims submitted to TMHP with a clinician-administered drug procedure code.

The NDC is an 11-digit number on the package or container from which the medication is administered. Providers must enter identifier N4 before the NDC code. The NDC unit and the NDC unit of measure must be entered on all professional or outpatient claims that are submitted to TMHP and Medicaid managed care plans.

Clinician-administered drugs that do not have a rebate-eligible NDC will not be reimbursed by Texas Medicaid.

**Refer to:** Subsection 6.3.4, “National Drug Code (NDC)” in “Section 6: Claims Filing” (Vol. 1, General Information) for additional information on claim filing using NDC.

#### 10.5.1 Pharmacy Delivery Method for Clinician-Administered Drugs

Providers administering clinician-administered drugs in an outpatient setting for clients enrolled in Medicaid (both traditional and managed care) can send a prescription to a pharmacy and wait for the drug to be shipped or mailed to their office. This delivery method is called “white-bagging.”

Providers should use the following steps for this delivery method:

1) The treating provider identifies that the client is enrolled in Medicaid.
2) The treating provider or treating provider’s agent sends a prescription to a Texas Medicaid-enrolled pharmacy and obtains any necessary prior authorizations.
3) If any prior authorization is approved, the dispensing pharmacy fills the prescription and overnight ships an individual dose of the medication, in the client’s name, directly to the treating provider.
4) The treating provider administers the medication in the office setting. The provider bills for an administration fee and any medically necessary service provided at time of administration. The provider should not bill Medicaid for the drug.

The pharmacy contacts the provider each month, prior to dispensing any refills, to ensure that the client received all previously dispensed medication. Auto-refills are not allowed.

These medications cannot be used on any other client and cannot be returned to the pharmacy for credit.

**Exception:** Unused long-acting reversible contraceptives may be returned in certain circumstances.

**Note:** Physicians who use this delivery method will not have to buy the clinician-administered drug, therefore, the physician is allowed to administer the drug and should only bill for the administration of the drug.

### 10.6 Opioid Limitations

For many people, substance use disorder starts after initially receiving opioid prescriptions for an episode of acute pain. To encourage the appropriate use of opioids and reduce the over-prescribing of opioids, Texas Medicaid has implemented the requirements in this section.
10.6.1 Affected Clients
The requirements in this section do not apply to clients who are:

- Receiving hospice care or palliative care
- Being treated for cancer
- Residing in a long-term care facility
- Residing in a facility in which residents receive opioid substitution therapy for the treatment of opioid use disorder (OUD).

The requirements also do not apply to other clients that HHSC elects to exempt based on an objective, confirmable physical pathology known to cause severe chronic pain that is not ameliorated by other therapies and for which opioid treatment is appropriate (e.g., sickle cell disease). If diagnoses are not available in the medical data, prescribers can request exemptions on a case-by-case basis through the pharmacy prior authorization process.

10.6.2 Morphine Milligram Equivalents
Morphine milligram equivalents (MME) per day is used to compare the potency of one opioid to another. The clinical decision for the MME per day recommendations varies depending on the person’s opioid use. Additionally, the Centers for Disease Control and Prevention (CDC) recommends starting opioid treatment with an immediate-release/short-acting formulation at the lowest effective dose instead of an extended-release/long-acting formulation.

A client is considered “opioid naive” if the client has taken opioids for a duration that is less than or equal to seven days in the last 60 days. For clients who are opioid naive, providers must submit a one-time prior authorization request for:

- An opioid prescription that exceeds a ten-day supply.
- A prescription for a long-acting opioid formulation.
- A claim or combination of claims in which the total daily dose of opioids exceeds 90.

The one-time requirement for prior authorization does not apply to subsequent claims because the member will no longer be opioid naive. The duration of the prior authorization is equal to the days’ supply of the claim.

For clients who are not opioid naïve, prior authorization is required for opioid prescriptions if the total daily dose of opioids exceeds 90 MME. For those patients who may require a tapering plan, providers would determine the development and management of a patient specific course of therapy to help manage withdrawal symptoms. A prescriber may request a tapering plan through the pharmacy prior authorization process on a case-by-case basis. Prior authorization approvals last for six-months.

10.6.3 Days’ Supply Limit
Opioid prescriptions for the treatment of acute pain are rarely required for more than ten days. To reduce the risk of addiction and the diversion of unused opioids, opioid prescriptions for clients who are opioid naïve are limited to a maximum ten-day supply without prior authorization.

10.6.4 Fee-For-Service Three Prescription Limit
Prescriptions for opioids to treat acute pain for clients who are 21 years of age and older are exempt from the three-prescription-per-month limit for members in fee-for-service.
11 Patient Information

11.1 Medication Synchronization

11.1.1 Overview
Medication Synchronization establishes processes for early refills in order to align the filling or refilling of multiple medications for a client with chronic illnesses.

The Texas Insurance Code §1369(j) allows a client enrolled in Medicaid, their prescribing physician, or the dispensing pharmacist to initiate the medication synchronization request. This process allows for clients to pick up all their medication on a single day each month versus requiring clients to make multiple pharmacy visits to obtain different prescription medications with different refill dates.

11.1.2 Eligible Medications
A drug is eligible for medication synchronization if it meets the following conditions:

- It is listed on the Medicaid, CHIP, KHC or CSHCN formulary.
- It is used for treatment and management of chronic illnesses.
- It is a formulation or dosage form that can be effectively dispensed in a medication synchronization protocol.
- It must meet all prior authorization criteria applicable to the medication on the date the synchronization request is made. This includes clinical prior authorizations, non-preferred prior authorizations, and drug utilization review edits.
- The original prescription must have refills.

**Exception:** The prescription could be new but the drug is categorized with the same Generic Code Number (GCN) class, and if the pharmacy uses the override code, the claim will pay. Having available refills is not required.

A claim cannot be synchronized if it is a Schedule II controlled substance or a Schedule III controlled substance containing hydrocodone.

11.1.3 Chronic Illness
Medications eligible for synchronization must be used to treat chronic illnesses. A chronic illness is defined as an illness or physical condition that is:

- Reasonably expected to continue for an uninterrupted period of at least three months, and
- Controlled, but not cured by medical treatment. This includes drugs used to treat mental health conditions and substance use.

11.1.4 Traditional Medicaid Claims Processing
A synchronized claim will count as one of the three prescriptions Medicaid will pay if a client is limited.

11.1.5 Medicaid Managed Care and CHIP Claims Processing
Each health plan has an HHSC-approved process for medication synchronization for clients eligible for Medicaid or CHIP. In CHIP, cost sharing or co-payment amounts will be prorated. Dispensing fees will not be prorated.

Pharmacy staff should contact the client’s health plan for medication synchronization requirements using the contact information on the Pharmacy MCO Assistance Chart.
11.2 Medicaid Drug Benefits

Clients enrolled in traditional Medicaid are limited to three prescriptions per month with the following exceptions:

- Clients enrolled in waiver programs such as Community Living Assistance (CLASS) and Community-Based Alternatives (CBA)
- Texas Health Steps (THSteps)-eligible clients (clients who are 20 years of age and younger)
- Clients in skilled nursing facilities

The following categories of drugs do not count against the three prescription per month limit:

- Family planning drugs and supplies
- Smoking cessation drugs
- Insulin syringes

*Note:* Prescriptions for family planning drugs and limited home health supplies are not subject to the three-prescription limit.

Though TMHP reimburses family planning agencies and physicians for family planning drugs and supplies, the following family planning drugs and supplies are also available through the VDP and are not subject to the three-prescription limit:

- Oral contraceptives
- Long-acting injectable contraceptives
- Vaginal ring
- Hormone patch
- Certain drugs used to treat sexually transmitted diseases (STDs)

11.3 Cost Avoidance Coordination of Benefits

Cost avoidance coordination of benefits for pharmacy claims ensures compliance with CMS regulations. Under federal rules, Medicaid agencies must be the payer of last resort. The cost avoidance model checks for other known insurance at point of sale, preventing Medicaid from paying a claim until the pharmacy attempts to obtain payment from the client’s primary third party insurance.

*Refer to:* The VDP Pharmacy Provider Procedure Manual on the VDP website.

11.4 Medicaid Children's Services Comprehensive Care Program

Medically-necessary drugs and supplies that are not covered by the VDP may be available to children and adolescents (birth through 20 years of age) through the Medicaid Comprehensive Care Program (CCP). Drugs and supplies not covered could include, as examples, some over-the-counter drugs, nutritional products, diapers, and disposable or expendable medical supplies.

The Prior Authorization fax number is 1-512-514-4212.

*Refer to:* Subsection 2.7.1.1, “Pharmacies (CCP)” in the Children's Services Handbook (Vol. 2, Provider Handbooks) for more information about pharmacy enrollment in CCP.

11.5 Pharmacy Lock-In

Clients enrolled in traditional Medicaid can be “locked-in” to a specific primary care pharmacy. Those clients will have “Lock-in” identified on the face of their Your Texas Benefits Medicaid card. Clients who are not “locked-in” to a specific pharmacy may obtain their drugs or supplies from any enrolled Medicaid pharmacy.
11.6 Free Delivery of Medicaid Prescriptions

Many Medicaid pharmacies offer free delivery of prescriptions to clients enrolled in Medicaid. To find out which pharmacies offer delivery services:

- Refer clients enrolled in traditional Medicaid to the VDP Pharmacy Search. Click the “Delivers” indicator on the search. The returned results will include only those pharmacies that provide a delivery service. These VDP-enrolled pharmacies have certified their delivery services meet the minimum conditions for the payment of the delivery fee. These certified delivery pharmacies are reimbursed a delivery fee that is included in the medication dispensing formula.

- Refer clients enrolled in Medicaid managed care to their respective health plan. Each health plan develops its own participating pharmacy network for the delivery service.

Deliveries are made to client’s home and not institutions, such as nursing homes. Delivery service is not applicable for mail-order prescriptions and not is available for over-the-counter drugs.

12 Pharmacy Prior Authorization

Some Medicaid drugs are subject to one or both types of prior authorization, clinical and non-preferred.

12.1 Clinical Prior Authorization

Clinical prior authorizations utilize evidence-based clinical criteria and nationally recognized peer-reviewed information. These prior authorizations may apply to an individual drug or a drug class on the formulary, including some preferred and non-preferred drugs. There are certain clinical prior authorizations that all health plans are required to perform. Usage of all other clinical prior authorizations will vary between health plans and at the discretion of each health plan.

Refer to: The Clinical Prior Authorization Assistance Chart on the VDP website. It identifies the prior authorization each health plans uses and how those authorizations relate to the authorizations used for traditional Medicaid claim processing. The chart is updated quarterly.

Subsection 9.5, “Palivizumab (Synagis)” in this section for information about Synagis prior authorizations.

12.2 Non-preferred Prior Authorization

The PDL is arranged by drug therapeutic class and contains a subset of many, but not all, drugs that are on the Medicaid formulary. Drugs are identified as preferred or non-preferred on the PDL. Drugs listed on the PDL as preferred, or those not listed at all, are available without PDL prior authorization. Drugs identified as non-preferred on the PDL require a PDL prior authorization.

Refer to: Medicaid health plans are required to adhere to the Texas Medicaid Preferred Drug List. Note: "CHIP does not have a PDL.

Refer to: The PDL Prior Authorization Criteria Guide that explains the criteria that are used to evaluate the PDL prior authorization requests.
12.3 Obtaining Prior Authorization

Prior authorization for clients enrolled in traditional Medicaid is requested through the Texas Prior Authorization Call Center.

The Texas Prior Authorization Call Center accepts prior authorization requests by phone at 1-877-PA-TEXAS (1-877-728-3927) (Monday through Friday, between 7:30 a.m. and 6:30 p.m., central) or online through PAXpress. Online submissions are only available for non-preferred prior authorization requests.

Refer to:
- The Account Registration Instructions on the PAXpress website.

Note: Pharmacists cannot obtain prior authorization for medications. If the client arrives at the pharmacy without prior authorization for a non-preferred drug and/or a drug requiring clinical prior authorization, the pharmacist will alert the provider’s office and ask the provider to get prior authorization.

12.4 72-Hour Emergency Supply

Federal and Texas law allows for a 72-hour emergency supply of a prescribed drug to be provided when a medication is needed without delay and prior authorization is not available. This rule applies to non-preferred drugs on the PDL and any drug that is affected by a clinical prior authorization.

Drugs not on the PDL may also be subject to clinical prior authorization.

Refer to: The Texas Pharmacy Provider Procedure Manual on the VDP website.

12.5 Retrospective Drug Utilization Reviews

Retrospective DUR provides for the ongoing periodic examination of claims data and other records to identify patterns of fraud, abuse, gross overuse, or inappropriate or medically unnecessary care among prescribing providers, pharmacists, and people associated with specific drugs or groups of drugs.

The retrospective review also allows for active and ongoing educational outreach in the form of letters or face-to-face discussions to educate prescribing providers on common drug therapy problems with the aim of improving prescribing or dispensing practices.

The Texas Drug Utilization Review Board reviews and recommends interventions for traditional Medicaid claims. A fixed-number of interventions are performed each calendar year. MCOs are required to create and perform interventions and education of their population.
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1 General Information

This handbook contains information about Texas Medicaid fee-for-service benefits. For information about managed care benefits, refer to the Medicaid Managed Care Handbook.

Managed care carve-out services are administered as fee-for-service benefits. A list of all carve-out services is available in Section 8, “Carve-Out Services” in the Medicaid Managed Care Handbook (Vol. 2, Provider Handbooks).

The information in this handbook is intended for therapy services for clients of all ages. Therapy services include occupational therapy (OT), physical therapy (PT), and speech therapy (ST). The handbook provides information about Texas Medicaid’s benefits, policies, and procedures applicable to these therapies.

Important: All providers are required to read and comply with Section 1: Provider Enrollment and Responsibilities. In addition to required compliance with all requirements specific to Texas Medicaid, it is a violation of Texas Medicaid rules when a provider fails to provide healthcare services or items to Medicaid clients in accordance with accepted medical community standards and standards that govern occupations, as explained in Title 1 Texas Administrative Code (TAC) §371.1659. Accordingly, in addition to being subject to sanctions for failure to comply with the requirements that are specific to Texas Medicaid, providers can also be subject to Texas Medicaid sanctions for failure, at all times, to deliver health-care items and services to Medicaid clients in full accordance with all applicable licensure and certification requirements including, without limitation, those related to documentation and record maintenance.

Refer to: “Section 1: Provider Enrollment and Responsibilities” (Vol. 1, General Information) for more information about enrollment procedures.

All providers are required to report suspected child abuse and neglect, as outlined in subsection 1.7, “Provider Responsibilities” in “Section 1: Provider Enrollment and Responsibilities” (Vol. 1, General Information).

2 Enrollment

Refer to: Subsection 1.7.17, “Physical, Occupational, and Speech Therapy Providers” in “Section 1: Provider Enrollment and Responsibilities” (Vol. 1, General Information) for enrollment information.

3 Managed Care Organization (MCO) Clients Who Transition to Medicaid Fee-For-Service (FFS)

When client’s transition from an MCO to FFS, providers can request previously approved authorizations be transferred from the MCO to FFS for Comprehensive Care Program (CCP) services, OT, PT, and ST.

3.1 Submission Guidelines

The following submission time frames apply for providers that request to transfer previously approved MCO authorizations for PT, OT, and ST services:

- TMHP will consider the reimbursement of claims for services that were rendered on or after the MCO’s disenrollment date only when the provider submits a request to TMHP to transfer the previously approved authorization for PT, OT, and ST services.
• The request to TMHP must be received on or before the end date of the previously approved MCO authorization. Any requests submitted after the MCO’s authorization end date will have to meet the regular submission guidelines for the specific service type.

3.2 Documentation Requirements
All of the requests to transfer the authorizations from the MCO to FFS must include:

- A copy of the previously approved authorization letter.
- All of the documentation that was sent in the original authorization request, including any physician orders that were used to determine the start of care. TMHP will accept the physician orders as the required documentation for the requested services.
- The completed CCP prior authorization form, Special Medical Prior Authorization (SMPA) form, Home Health Plan of Care, or Texas Medicaid Physical, Occupational, or Speech Therapy (PT, OT, ST) Prior Authorization Form whichever is applicable for the requested service. The form must include the dates of service and quantities that are being requested from TMHP, and they must match the dates of service and quantities that were approved in the original MCO authorization. It is not necessary to obtain signatures or dates on the forms if they are submitted to TMHP for the purpose of transferring an authorization from an MCO to FFS Medicaid.

Note: Authorizations for services transferred from an MCO to FFS Medicaid are subject to retrospective review.

TMHP will verify the client’s eligibility, the dates of service, and the quantities requested. TMHP will process reimbursement claims as follows:

- Claims for services that were rendered before the date on which the transfer request was received will be denied as a late submission, and the provider will be notified of their administrative appeal rights through the Health and Human Services Commission (HHSC).
- Claims for services that were rendered on or after the date of receipt use the required information from the transferred authorization and will be processed as if the request was received in a timely manner.

If a request to transfer an MCO authorization is submitted after the end date of the MCO authorization or the provider does not have an authorization letter from the MCO, TMHP will process the request to transfer the authorization based on established TMHP authorization submission guidelines for PT, OT, and ST services.

All new requests for rendered services must meet the documentation requirements.

3.3 New Services and Extension of Services
For new services that occur after the client’s MCO disenrollment change date, the provider is responsible for submitting all TMHP required paperwork and meeting all established submission guidelines for prior authorization.

Requests for the extension of services that occur after the MCO disenrollment change date must include all of the paperwork that is required by TMHP and meet all established submission guidelines for prior authorization.

3.4 Loss of Eligibility
If an MCO disenrolled a client and the client also loses Medicaid eligibility, providers must anticipate, if and when Medicaid eligibility is restored, that the client will initially be considered a Medicaid FFS client and will have a retroactive eligibility period.
All requests for services that require prior authorization and that occur during the client’s retroactive eligibility period, must be submitted to TMHP following the process that is outlined in subsection 5.1.1, “Prior Authorization Requests for Clients with Retroactive Eligibility” in “Section 5: Fee-for-Service Prior Authorizations” (Vol. 1, General Information).

If a client is retroactively disenrolled by an MCO, all of the services that are rendered by the provider during this retroactive disenrollment period (specifically from the date on which the client was eligible for FFS to the date of the client’s MCO eligibility change) will be denied by TMHP, and the provider will be notified of their administrative appeal rights.

TMHP may consider services for the MCO transition beginning on the date of the client’s MCO eligibility change date and going forward. TMHP uses the MCO transition process for the submission of paperwork and the processing of provider requests.

4 Therapy Services Overview

Physical, occupational and speech therapy services must be medically necessary to the treatment of the individual’s chronic or acute need. A diagnosis alone is not sufficient documentation to support the medical necessity of therapy. To be considered medically necessary, all of the following conditions must be met:

- The services requested must be considered under the accepted standards of practice to be a specific and effective treatment for the patient’s condition.
- The services requested must be of a level of complexity or the patient’s condition must be such that the services required can only be effectively performed by or under the supervision of a licensed occupational therapist, physical therapist, or speech-language pathologist, and requires the skills and judgment of the licensed therapist to perform education and training.
- Functional goals refer to a series of behaviors or skills that allow the client to achieve an outcome relevant to his/her health, safety, or independence within the context of everyday environments. Functional goals must be specific to the client, objectively measurable within a specified time frame, attainable in relation to the client’s prognosis or developmental delay, relevant to client and family, and based on a medical need.
- For clients who are 20 years of age and younger, the following conditions must be met:
  - The goals of the requested services to be provided are directed at improving, adapting, restoring, or maintaining functions which have been lost or impaired due to a recent illness, injury, loss of body part or congenital abnormality or as a result of developmental delay or the presence of a chronic medical condition.
  - Testing must establish a client with developmental delays meets the medical necessity criteria as defined in subsection 5.3, “Developmental Delay Criteria” in this handbook for chronic therapy services.
  - Evidence of care coordination with the prescribed pediatric extended care center (PPECC) provider, when the client receives therapy services in a PPECC setting.
- For clients who are 21 years of age and older, the following conditions must be met:
  - The goals of the requested services to be provided are directed at improving, adapting or restoring functions which have been lost or impaired due to a recent illness, injury, loss of body part and restore client’s function to within normal activities of daily living (ADL).
  - There must be reasonable expectation that therapy will result in a meaningful or practical improvement in the client’s ability to function within a reasonable and predictable time period.
Medical necessity criteria for therapy services provided in the home must be based on the supporting documentation of the medical need and the appropriateness of the equipment, service, or supply prescribed by the prescribing provider for the treatment of the individual.

The therapy service must be related to the client’s medical condition, rather than primarily for the convenience of the client or provider.

Frequency must always be commensurate with the client’s medical and skilled therapy needs, level of disability (for clients who are 20 years of age and younger), and standards of practice; it is not for the convenience of the client or the responsible caregivers.

The following apply:

- Treatment plans and plans of care developed must include not only the initial frequency (high, moderate or low) but the expected changes of frequency throughout the duration period requested based on the client’s anticipated therapy treatment needs.

- An example of a tapered down frequency request initiated with a high frequency is: 3 times a week for 2 weeks, 2 times a week for 2 weeks, 1 time a week for 2 weeks, 1 time every other week).

Refer to: Subsection 4.5, “Frequency and Duration Criteria for PT, OT, and ST Services” in this handbook for the frequency prior authorization criteria.

Therapy services are limited to one evaluation, re-evaluation or treatment up to the limits outlined in this handbook for each therapy discipline per date of service.

### 4.1 Physical Therapy

The practice of physical therapy includes:

- Measurement or testing of the function of the musculoskeletal, or neurological, system.

- Rehabilitative treatment concerned with restoring function or preventing disability caused by illness, injury, or birth defect.

- Treatment, consultative, educational, or advisory services to reduce the incidence or severity of disability or pain to enable, train, or retrain a person to perform the independent skills and activities of daily living.

Texas Medicaid limits physical therapy to the skilled treatment of clients who have acute or acute exacerbation of chronic disorders or chronic medical condition of the musculoskeletal and neuromuscular systems. Physical therapy may be provided by a physician or physical therapist within their licensed scope of practice.

### 4.2 Occupational Therapy

The practice of occupational therapy includes:

- Evaluation and treatment of a person whose ability to perform the tasks of living is threatened or impaired by developmental deficits, sensory impairment, physical injury, or illness.

- Using therapeutic goal-directed activities to:
  - Evaluate, prevent, or correct physical dysfunction.
  - Maximize function in a person’s life.
  - Applying therapeutic goal-directed activities in treating patients on an individual basis, in groups, or through social systems, by means of direct or monitored treatment or consultation.

Texas Medicaid limits occupational therapy to the skilled treatment of clients whose ability to function in life roles is impaired. Occupational therapy may be provided by a physician or occupational therapist within their licensed scope of practice.
Occupational therapy uses purposeful activities to obtain or regain skills needed for activities of daily living (ADL) and/or functional skills needed for daily life lost through acute medical condition, acute exacerbation of a medical condition, or chronic medical condition related to injury, disease, or other medical causes. ADLs are basic self-care tasks such as feeding, bathing, dressing, toileting, grooming, and mobility.

### 4.3 Speech Therapy

Speech therapy is a benefit of Texas Medicaid for the treatment of chronic (for clients who are 20 years of age and younger), acute, or acute exacerbations of pathological or traumatic conditions of the head or neck, which affect speech production, speech communication and oral motor, feeding and swallowing disorders. Speech therapy may be provided by a physician or speech language pathologist within their licensed scope of practice.

Speech-language pathologists treat speech sound and motor speech disorders, stuttering, voice disorders, aphasia and other language impairments, cognitive disorders, social communication disorders and swallowing (dysphagia) deficits.

Speech therapy is designed to ameliorate, restore, or rehabilitate speech language communication and swallowing disorders that have been lost or damaged as a result of a chronic, acute or acute exacerbation of a medical condition due to a recent injury, disease or other medical conditions, or congenital anomalies or injuries.

#### 4.3.1 Types of Communication Disorders

There are three types of communication disorders:

- **Language Disorders**—Impaired comprehension and/or use of spoken, written and/or other symbol systems. This disorder may involve the following components: forms of language (phonology, morphology, and syntax), content and meaning of language (pragmatics) and/or the perception/processing of language. Language disorders may involve one, all or a combination of the above components.

- **Speech Production Disorders**—Impairment of the articulation of speech sounds, voice and/or fluency. Speech Production Disorders may involve one, all or a combination of these components of the speech production system. An articulation disorder may manifest as an individual sound deficiency, i.e., traditional articulation disorder, incomplete or deviant use of the phonological system, i.e., phonological disorder, or poor coordination of the oral-motor mechanism for purposes of speech production, i.e., verbal and/or apraxia, dysarthria.

- **Oral Motor/Swallowing/Feeding Disorders**—Impairment of the muscles, structures and/or functions of the mouth (physiological or sensory-based) involved with the entire act of deglutition from placement and manipulation of food in the mouth through the oral and pharyngeal phases of the swallow. These disorders may or may not result in deficits to speech production.

### 4.4 Co-Treatment

Co-treatment is defined as two different therapy disciplines performing therapy on the same client at the same time by a licensed therapist as defined in this handbook for each therapy discipline, and rendered in accordance with the Executive Council of Physical Therapy and Occupational Therapy Examiners and State Board of Examiners for Speech-Language Pathology and Audiology.

Co-treatment may be a benefit when it is medically necessary for the client to receive therapy from two different therapy disciplines at the same time. The therapy performed requires the expertise of two different disciplines (i.e., licensed physical therapist, licensed occupational therapist, or licensed speech-language pathologist), to perform the therapy safely and effectively to reach the client’s goals as determined by the approved plan of care, signed and dated by the client’s prescribing provider.
When performing co-treatment, a primary therapist must be designated by the two performing therapists. Only the primary performing therapist may bill for the therapy services rendered. The secondary therapist will not be reimbursed for assisting a designated primary performing therapist.

The following co-treatment documentation requirements must be maintained in the client’s medical records as follows:

- Medical necessity for the individual therapy services must be justified before performing co-treatment.
- Documentation supports co-treatment goals and how co-treatment will help the therapist achieve the therapist’s goals for the client, for each therapy discipline.
- An explanation of why the client requires and will receive multi-disciplinary team care, defined as at least two therapy disciplines (physical, occupational, or speech therapy) during the same therapy session.

Retrospective review may be performed to ensure documentation supports that the medical necessity of the co-treatment performed and that the billing was appropriate for the services provided by the designated primary-performing therapist.

### 4.4.1 Group Therapy

Group therapy consists of simultaneous treatment to two or more clients who may or may not be doing the same activities. If the therapist is dividing attention among the clients, providing only brief, intermittent personal contact, or giving the same instructions to two or more clients at the same time, the treatment is recognized as group therapy. The physician or therapist involved in group therapy services must be in constant attendance, but one-on-one client contact is not required.

The following requirements must be met in order to meet the Texas Medicaid criteria for group therapy:

- Prescribing provider’s prescription for group therapy.
- Performance by or under the general supervision of a qualified licensed therapist as defined by licensure requirements.
- The licensed therapist involved in group therapy services must be in constant attendance (in the same room) and active in the therapy.
- Each client participating in the group must have an individualized treatment plan for group treatment, including interventions and short- and long-term goals and measurable outcomes.

Texas Medicaid does not limit the number of clients who can participate in a group therapy session. Providers are subject to certification and licensure board standards regarding group therapy.

#### 4.4.1.1 Group Therapy Documentation Requirements

The following documentation must be maintained in the client’s medical record:

- Prescribing provider’s prescription for group therapy.
- Individualized treatment plan that includes frequency and duration of the prescribed group therapy and individualized treatment goals.
- Name and signature of licensed therapist providing supervision over the group therapy session.
- Specific treatment techniques utilized during the group therapy session and how the techniques will restore function.
- Start and stop times for each session.
- Group therapy setting or location.
- Number of clients in the group.
The client’s medical record must be made available upon request.

4.5 Frequency and Duration Criteria for PT, OT, and ST Services
Frequency must always be commensurate with the client’s medical and skilled therapy needs, level of disability and standards of practice; it is not for the convenience of the client or the responsible adult.

Exceptions to therapy limitations may be covered if the medically necessary criteria are met for the following:

- Presentation of new acute condition
- Therapist intervention is critical to the realistic rehabilitative/restorative goal, provided documentation proving medical necessity is received.

When therapy is initiated, the therapist must provide education and training of the client and responsible caregivers, by developing and instructing them in a home treatment program to promote effective carryover of the therapy program and management of safety issues.

Providers may request high, moderate, or low frequencies on the Texas Medicaid Physical, Occupational or Speech Therapy (PT, OT, ST) Prior Authorization Form by indicating 3, 2, or 1 time per week respectively. Providers may request low or maintenance level by requesting 1, 2, or 3 times per month. Additional documentation is required when requesting a frequency of 3 times a week or more.

Note: The reference to “maintenance” in the above statement is applicable to clients who are 20 years of age and younger.

4.5.1 High Frequency
High frequency (3 times per week) can only be considered for a limited duration (approximately 4 weeks or less) or as otherwise requested by the prescribing provider with documentation of medical need to achieve an identified new skill or recover function lost due to surgery, illness, trauma, acute medical condition, or acute exacerbation of a medical condition, with well-defined specific, achievable goals within the intensive period requested.

Therapy provided three times a week may be considered for 2 or more of these exceptional situations:

- The client has a medical condition that is rapidly changing.
- The client has a potential for rapid progress (e.g., excellent prognosis for skill acquisition) or rapid decline or loss of functional skill (e.g., serious illness, recent surgery).
- The client’s therapy plan and home program require frequent modification by the licensed therapist.

On a case-by-case basis, a high frequency requested for a short-term period (4 weeks or less) which does not meet the above criteria may be considered with all of the following documentation:

- Letter of medical need from the prescribing provider documenting the client’s rehabilitation potential for achieving the goals identified,
- Therapy summary documenting all of the following:
  - Purpose of the high frequency requested (e.g., close to achieving a milestone)
  - Identification of the functional skill which will be achieved with high frequency therapy
  - Specific measurable goals related to the high frequency requested and the expected date the goal will be achieved.

A higher frequency (4 or more times per week) may be considered on a case-by-case basis with clinical documentation supporting why 3 times a week will not meet the client’s medical needs.
4.5.2 Moderate Frequency
Therapy provided two times a week may be considered when documentation shows one or more of the following:

- The client is making very good functional progress toward goals.
- The client is in a critical period to gain new skills or restore function or is at risk of regression.
- The licensed therapist needs to adjust the client’s therapy plan and home program weekly or more often than weekly based on the client’s progress and medical needs.
- The client has complex needs requiring ongoing education of the responsible adult.

4.5.3 Low Frequency
Therapy provided one time per week or every other week may be considered when the documentation shows one or more of the following:

- The client is making progress toward the client’s goals, but the progress has slowed, or documentation shows the client is at risk of deterioration due to the client’s development or medical condition.
- The licensed therapist is required to adjust the client’s therapy plan and home program weekly to every other week based on the client’s progress.
- Every other week therapy is supported for clients whose medical condition is stable, they are making progress, and it is anticipated the client will not regress with every other week therapy.

Note: As the client’s medical need for therapy decreases, it is expected that the therapy frequency will decrease as well.

4.5.4 Maintenance Level/Prevent Deterioration
For clients who are 20 years of age and younger only, this frequency level (e.g., every other week, monthly, every 3 months) is used when the therapy plan changes very slowly, the home program is at a level that may be managed by the client or the responsible adult, or the therapy plan requires infrequent updates by the skilled therapist. A maintenance level or preventive level of therapy services may be considered when a client requires skilled therapy for ongoing periodic assessments and consultations and the client meets one of the following criteria:

- Progress has slowed or stopped, but documentation supports that ongoing skilled therapy is required to maintain the progress made or prevent deterioration.
- The submitted documentation shows that the client may be making limited progress toward goals or that goal attainment is extremely slow.
- Factors are identified that inhibit the client’s ability to achieve established goals (e.g., the client cannot participate in therapy sessions due to behavior issues or issues with anxiety).
- Documentation shows the client and the responsible adult have a continuing need for education, a periodic adjustment of the home program, or regular modification of equipment to meet the client’s needs.

4.5.5 Requesting Therapy Services
Providers may request physical, occupational, or speech therapy services frequency by week for one or more visits per week, or by month for 1, 2, or 3 visits per month.

- A week includes the day of the week on which the prior authorization period begins and continues for seven days. For example, if the prior authorization starts on a Thursday, the prior authorization week runs Thursday through Wednesday.
• The number of therapy services authorized for a week or month must be contained in that prior authorization period.

• Services billed, in excess of those authorized are subject to recoupment.

Missed visits may be made up within the authorization period as long as total number of visits or units authorized does not exceed the amount authorized. Provider should document reason for visits outside of the weekly or monthly frequency in the client’s medical record.

### 4.6 Criteria for Discontinuation of Therapy

Discontinuation of therapy may be considered in one or more of the following situations:

• Client no longer demonstrates functional impairment or has achieved goals set forth in the treatment plan or plan of care.

• Client has returned to baseline function.

• Client can continue therapy with a home treatment program and deficits no longer require a skilled therapy intervention and, for clients who are 20 years of age and younger only, maintain status.

• Client has adapted to impairment with assistive equipment or devices.

• Client is able to perform ADLs with minimal to no assistance from caregiver.

• Client has achieved maximum functional benefit from therapy in progress or will no longer benefit from additional therapy.

• Client is unable to participate in the treatment plan or plan of care due to medical, psychological, or social complications; and responsible adult has had instruction on the home treatment program and the skills of a therapist are not needed to provide or supervise the service.

• Testing shows client no longer has a developmental delay.

• Plateau in response to therapy/lack of progress towards therapy goals. Indication for therapeutic pause in treatments or, for those under age 21, transition to chronic status and maintenance therapy.

• Non-compliance due to poor attendance and with client or responsible adult, non-compliance with therapy and home treatment program.

### 4.7 Exclusions (Non-covered Services)

The following services are not a benefit of Texas Medicaid:

• Speech therapy provided in the home to adult clients who are 21 years of age and older

• Therapy services that are provided after the client has reached the maximum level of improvement or is now functioning within normal limits

• Massage therapy that is the sole therapy or is not part of a therapeutic plan of care to address an acute condition

• Separate reimbursement for VitalStim therapy for dysphagia. VitalStim must be a component of a comprehensive feeding treatment plan to be considered a benefit.

• Repetitive therapy services that are designed to maintain function once the maximum level of improvement has been reached, which no longer require the skills of a therapist to provide or oversee

• Therapy services related to activities for the general good and welfare of clients who are not considered medically necessary because they do not require the skills of a therapist, such as:
  • General exercises to promote overall fitness and flexibility or improve athletic performance
- Activities to provide diversion or general motivation
- Supervised exercise for weight loss
- Treatment solely for the instruction of other agency or professional personnel in the client’s physical, occupational or speech therapy program
- Emotional support, adjustment to extended hospitalization and/or disability, and behavioral readjustment
- Therapy prescribed primarily as an adjunct to psychotherapy
- Treatments not supported by medically peer-reviewed literature, including but not limited to investigational treatments such as sensory integration, vestibular rehabilitation for the treatment of attention deficit hyperactivity disorder, anodyne therapy, craniosacral therapy, interactive metronome therapy, cranial electro stimulation, low-energy neuro-feedback, and the Wilbarger brushing protocol.
- Therapy not expected to result in practical functional improvements in the client’s level of functioning
- Treatments that do not require the skills of a licensed therapist to perform in the absence of complicating factors (i.e., massage, general range of motion exercises, repetitive gait, activities and exercises that can be practiced by the client on their own or with a responsible adult’s assistance)
- The therapy requested is for general conditioning or fitness, or for educational, recreational or work-related activities that do not require the skills of a therapist
- Equipment and supplies used during therapy visits are not reimbursed separately; they are considered part of the therapy services provided
- Therapy services provided by a licensed therapist who is the client’s responsible adult (e.g., biological, adoptive, or foster parents, guardians, court-appointed managing conservators, other family members by birth or marriage)

Auxiliary personnel (aide, orderly, student, or technician) may participate in physical therapy, occupational therapy, or speech therapy sessions when they are appropriately supervised according to each therapy discipline’s scope of practice and provider licensure requirements. Providers may not bill Texas Medicaid for therapy services provided solely by auxiliary personnel

Auxiliary personnel, a licensed therapy assistant, and a licensed speech-language pathology intern (Clinical Fellow) are not eligible to enroll as therapist providers in Texas Medicaid.

5 Children’s Therapy Services Clients birth through 20 years of age

5.1 Services, Benefits, and Limitations

This section addresses acute and chronic physical therapy (PT), occupational therapy (OT), and speech therapy (ST) services for clients who are 20 years of age or younger. This section does not address freestanding inpatient rehabilitation services.

Unless otherwise specified, “days” refers to calendar days.

PT, OT, and ST are benefits of Texas Medicaid in Comprehensive Outpatient Rehabilitation Facilities (CORFs) and Outpatient Rehabilitation Facilities (ORFs) for clients who are 20 years of age or younger.

Note: CORF and ORF services provided at schools, homes, daycare facilities, or any other non-Medicare approved ORF or CORF facility is not a covered Comprehensive Care Program (CCP) benefit.
Services provided to a client on school premises are only permitted when delivered before or after school hours. The only PT, OT, and ST services that can be delivered during school hours are therapy services provided by school districts as School Health and Related Services (SHARS).

Clients who are eligible for PT, OT, and ST through the public school system (SHARS), may only receive additional therapy through Medicaid if medical necessity criteria is met as outlined in this handbook.

Refer to: Subsection 2.8, “Early Childhood Intervention (ECI) Services” in the Children’s Services Handbook (Vol. 2, Provider Handbooks) for the specific guidelines for therapy services that are provided through Early Childhood Intervention (ECI).

Section 3, “School Health and Related Services (SHARS)” in the Children’s Services Handbook (Vol. 2, Provider Handbooks) for information about therapy services provided through SHARS.

Therapy services must be performed by one of the following: a licensed physical therapist, licensed occupational therapist, licensed speech-language pathologist, a physician within their scope of practice, or one of the following under the supervision of a licensed therapist of the specific discipline:

- Licensed therapy assistant
- Licensed speech-language pathology intern (Clinical Fellow)

Note: An advanced practice registered nurse (APRN) or a physician assistant (PA) may sign all documentation related to the provision of therapy services on behalf of the client’s physician when the physician delegates this authority to the APRN or PA.

PT, OT, and ST services are provided in one of the following places of service by setting and provider:

<table>
<thead>
<tr>
<th>Place of Service</th>
<th>Provider Types</th>
</tr>
</thead>
<tbody>
<tr>
<td>Office</td>
<td>Physician, physical therapy group, independently enrolled therapist, podiatrist, SHARS and ECI</td>
</tr>
<tr>
<td>Home</td>
<td>Home health agency, independently enrolled therapist, physical therapy group, ECI, SHARS</td>
</tr>
<tr>
<td>Outpatient</td>
<td>Physician: podiatrist, outpatient hospital/clinic, outpatient rehabilitation center (includes comprehensive outpatient rehabilitation facility [CORF]/outpatient rehabilitation facility [ORF]) PPECC: home health agency, independently enrolled therapist, physical therapy group, ECI</td>
</tr>
<tr>
<td>Other</td>
<td>ECI, SHARS, independently enrolled therapist, home health agency, and physical therapy group</td>
</tr>
</tbody>
</table>

In determining whether a service requires the skill of a licensed physical and occupational therapist or speech language pathologist, consideration must be given to the inherent complexity of the service, the condition of the client, the accepted standards of medical and therapy practice guidelines, with consideration of the following:

- If the service could be performed by the average nonmedical person, the absence of a competent person (such as a family member or medical assistant) to perform it does not cause it to be a skilled therapy service.
- If the nature of a service is such that it can be safely and effectively performed by the average nonmedical person without direct supervision of a licensed therapist, the services cannot be regarded as skilled therapy.

Refer to: Subsection 2.1, “CCP Overview” in the Children’s Services Handbook (Vol. 2, Provider Handbooks) for additional information about CCP.
5.1.1 Acute PT, OT, and ST Services

Acute PT, OT, and ST services are benefits of Texas Medicaid for the medically necessary short term treatment of an acute medical condition or an acute exacerbation of a chronic medical condition.

Treatments are expected to significantly improve, restore or develop physical functions diminished or lost as a result of a recent trauma, illness, injury, disease, surgery, or change in medical condition, in a reasonable and generally predictable period of time (60 days), based on the prescribing provider’s and therapist’s assessment of the client’s restorative potential.

Note: Recent is defined as occurring within the past 90 days of the prescribing provider’s evaluation of condition.

Treatments are directed towards restoration of or compensation for lost function.

Services do not duplicate those provided concurrently by any other therapy.

Services must meet acceptable standards of medical practice and be specific and effective treatment for the client’s condition.

Services are provided within the provider’s scope of practice, as defined by state law.

Acute is defined as an illness or trauma with a rapid onset and short duration.

A medical condition is considered chronic when 120 days have passed from the start of therapy or the condition is no longer expected to resolve or may be slowly progressive over an indefinite period of time.

With documentation of medical need physical, occupational, and speech therapy may continue for a maximum of 120 days for an acute medical condition or an acute exacerbation of a chronic medical condition.

Once the client’s condition is no longer considered acute, continued therapy for a chronic condition will only be considered for clients who are 20 years of age or younger.

5.1.2 Chronic Services

Chronic physical, occupational, and speech therapy services are benefits of Texas Medicaid for the medically necessary treatment of chronic medical conditions and developmental delay when a medical need is established for the developmental delay as indicated in this handbook. All eligible clients who are birth through 20 years of age may continue to receive all medically necessary therapy services, with documentation proving medical necessity.

The goals of the services provided are directed at maintaining, improving, adapting, or restoring functions which have been lost or impaired due to a recent illness, injury, loss of body part, congenital abnormality, degenerative disease, or developmental delay.

Services do not duplicate those provided concurrently by any other therapy.

Services must meet acceptable standards of medical practice and be specific and effective treatment for the client’s condition.

Services are provided within the provider’s scope of practice, as defined by state law.

Treatment for chronic medical conditions and developmental delay will only be considered for clients who are birth through 20 years of age.

5.2 Authorization Requirements for PT, OT, and ST Services

Prior authorization requests may be submitted to the TMHP Prior Authorization Department via mail, fax, or the electronic portal. Prescribing or ordering providers, dispensing providers, clients’ responsible adults, and clients may sign prior authorization forms and supporting documentation using electronic or wet signatures.
Refer to: Subsection 5.5.1.2, “Document Requirements and Retention” in “Section 5: Fee-for-Service Prior Authorizations” (Vol. 1, General Information) for additional information about electronic signatures.

Providers must list all relevant procedure codes on the Texas Medicaid Physical, Occupational, or Speech Therapy (PT, OT, ST) Prior Authorization Form when requesting prior authorization for therapy services.

Therapy services performed in the acute care inpatient setting do not require prior authorization.

Coverage periods do not coincide necessarily with calendar weeks or months, but cover a number of services to be scheduled between a start and end date that is assigned during the prior authorization period.

5.2.1 Initial Evaluation and Considerations for Prior Authorization for Treatment

Initial evaluations do not require prior authorization (procedure codes 92521, 92522, 92523, 92524, 92610, 97161, 97162, 97163, 97165, 97166, and 97167); however, documentation kept in the client’s record must include a signed and dated prescribing provider’s order for the evaluation, support a medical need for the therapy evaluation, and be available when requested.

A therapy evaluation is considered current when it is performed within 60 days before the prior authorization request is received.

To complete the prior authorization process by paper, the provider must complete and submit the prior authorization requirements documentation through fax or mail, and must maintain a copy of the prior authorization request and all submitted documentation in the client’s medical record at the therapy provider’s place of business.

To complete the prior authorization process electronically, the provider must complete and submit the prior authorization requirements documentation through any approved electronic method, and must maintain a copy of the prior authorization request and all submitted documentation in the client’s medical record at the therapy provider’s place of business.

To avoid unnecessary denials, the prescribing provider must provide correct and complete information, including documentation of medical necessity for the service(s) requested. The prescribing provider must maintain documentation of medical necessity in the client’s medical record. The requesting therapy provider may be asked for additional information to clarify or complete a request.

Therapy services, regardless of place or provider, occurring after the initial evaluation, require prior authorization. PT, OT, or ST services may be prior authorized to be provided in the following locations: home of the client, home of the caregiver or guardian, client’s daycare facility or the client’s school.

5.2.1.1 Initial Evaluation for Acute and Chronic Therapy Services

For acute therapy services, i.e. acute services billed with an AT modifier, prior authorization requests may not exceed a 60 day period per each request. After two 60 day authorized periods, any continued requests for therapy services must be considered under the chronic sections of this handbook.

For chronic therapy services, prior authorization may be granted for up to 180 days with documentation of medical necessity and additional prior authorizations.

Initial prior authorization (PA) requests must be received no later than five business days from the date therapy treatments are initiated. Requests received after the five-business-day period will be denied for dates of service that occurred before the date that the PA request was received.
All of the following documentation is required when submitting an initial request for therapy services initiated after the completion of the evaluation for acute or chronic services:

- A completed Texas Medicaid Physical, Occupational, or Speech Therapy (PT, OT, ST) Prior Authorization Form signed and dated by both the therapist and by the prescribing provider is required. When the request form is unsigned by the prescribing provider, it must be accompanied by a signed and dated written order or prescription or a documented verbal order delineating the prescribed therapy services.

- The prescribing provider must certify that the Texas Health Steps (THSteps) checkup is current or that a developmental screening has been performed within the last 60 days. Signature of prescribing provider on PA form will attest that this service has been provided. If prescribing provider provides verbal order or written order separate from PA form, staff member who conveys the verbal or written order must communicate that prescribing provider attests that THSteps checkup is current or that a developmental screening has been performed within the last 60 days.

- For acute services: Documentation from the prescribing provider that a visit for the acute or acute exacerbation of the medical condition requiring therapy has occurred within the last 90 days.

- Evaluation and Treatment Plan or Plan of Care (POC) with all of the following required elements:
  - Client’s medical history and background
  - All medical diagnoses related to the client’s condition
  - Date of onset of the client’s condition requiring therapy or exacerbation date as applicable
  - Date of evaluation
  - Time in and time out
  - Baseline objective measurements based on standardized testing performed or other standard assessment tools

Refer to: Subsection 5.3, “Developmental Delay Criteria” in this handbook for information about chronic services.

- Safety risks
- Client-specific, measurable short and long-term functional goals within the length of time the service is requested
- Interpretation of the results of the evaluation, including recommendations for therapy amount, frequency per week and duration of services
- Therapy treatment plan/POC to include specific modalities and treatments planned
- Documentation of client’s primary language
- Documentation of client’s age and date of birth
- Adaptive equipment or assistive devices, as applicable
- Prognosis for improvement
- Requested dates of service for planned treatments after the completion of the evaluation
- Responsible adult’s expected involvement in client’s treatment
- History of prior therapy and referrals as applicable
- Signature and date of treating therapist
5.2.2 Additional Evaluation and Documentation Requirements for Speech Therapy

Additional evaluation and documentation requirements for speech therapy include one or more of the following:

- Language evaluations—Oral-peripheral speech mechanism examination and formal or informal assessment of hearing, articulation, voice and fluency skills;
- Speech production (voice)—Formal screening of language skills, and formal or informal assessment of hearing, voice and fluency skills;
- Speech production (fluency and articulation)—Formal screening of language skills, formal or informal assessment of hearing, voice and fluency skills;
- Oral Motor/Swallowing/Feeding—In addition to formal screening of language skills, formal or informal assessment of hearing, voice, and fluency skills, if swallowing problems and/or signs of aspiration are noted, then a statement indicating that a referral has been made to the client’s prescribing provider to consider a video fluoroscopic swallow study must be included.

5.2.2.1 Bilingual Testing Requirements

Bilingual and multilingual speakers are frequently misclassified as developmentally delayed. Equivalent proficiency in both languages should not be expected.

Criterion-referenced assessment tools can be used to identify and evaluate a client’s strengths and weaknesses, as opposed to norm-referenced testing, which assesses an individual relative to a group.

When possible, use culturally and linguistically adapted test equivalents in both languages to compare potential deficits and include in the documentation. The therapist will show the highest score of the two languages to determine whether the child qualifies and which language will be used for the child’s therapy. Testing for all subsequent re-evaluations should only be conducted in the language used in therapy.

5.2.3 Written and Verbal Orders

For new authorizations and recertifications of therapies, if the submitted request form is not signed and dated by the prescribing provider, the request must be accompanied by a verbal or written order.

The request form or written or verbal order must be signed and dated within the 60-day period before the initiation of services. A prescribing physician’s order to evaluate and treat is acceptable for the evaluation or re-evaluation, but is not acceptable for the therapy treatment. The therapy treatment order must contain the prescribing provider’s ordered frequency, duration, and affirmation that the client’s THSteps checkup is current or that a developmental screening has been performed within the last 60 days.

The documentation for a verbal order must meet the following criteria:

- It must be signed and dated by the licensed professional who by state and federal law may take a verbal order.
- It must have the name and credentials of the licensed professional who took the order and who is responsible for furnishing or supervising the ordered services.
- The verbal order must include the date on which the verbal order was taken.
- The verbal order must include the services, frequency, and duration that was prescribed by the ordering provider.
5.2.4 Requests for Recertification—Acute Therapy Services

A recertification for prior authorization of acute therapy services may be considered up to a maximum of 60 day increments, when services continue to meet authorization criteria. Re-evaluation codes (procedure codes 97164, 97168, and S9152) require authorization for acute therapy services and must be submitted with the recertification request. Therapy for clients who are birth through 20 years of age who do not meet the acute therapy services criteria may be considered for chronic therapy services.

Recertification for an acute or acute exacerbation of medical conditions includes a progress summary and a Texas Medicaid Physical, Occupational or Speech Therapy (PT, OT, ST) Prior Authorization Form.

A complete recertification request must be received no earlier than 30 days before the current authorization period expires. Requests for recertification services received after the current authorization expires will be denied for dates of service that occurred before the date the submitted request was received.

Prior authorization for recertification requests may be considered for increments up to 60 days for each therapy service request, with documentation supporting the medical necessity including all of the following:

- Texas Medicaid Physical, Occupational or Speech Therapy (PT, OT, ST) Prior Authorization Form or electronic equivalent signed and dated by the therapist and signed and dated by the prescribing provider. When the request form is unsigned by the prescribing provider, it must be accompanied by a written order or prescription or a verbal order for the prescribed therapy services.

- A progress summary (see progress summary documentation requirements), and

- A revised treatment plan or plan of care for the recertification dates of service requested, including all of the following:
  - Date therapy services started
  - Changes in the treatment plan, the rationale and the requested change in frequency of visits for changing the plan
  - Documentation of reasons continued therapy services are medically needed
  - Documentation of client’s participation in treatment, as well as client or responsible adult’s participation or adherence with a home treatment program
  - New treatment plan or plan of care for the recertification dates of service requested
  - Updated or new functional and measurable short and long-term treatment goals with new time frames, as applicable
  - Adaptive equipment or assistive devices, as applicable
  - Prognosis with clearly established discharge criteria
  - Documentation of consults with other professionals and services or referrals made and coordination of service when applicable (e.g., for school aged clients, documentation of the coordination of care and referrals made for school therapies).
  - The updated treatment plan or POC must be signed and dated by the therapist responsible for the therapy services.

A progress summary, which may be contained in the last treatment note, must be included with the recertification request and contains all of the following:

- Date therapy started
- Date the summary completed
• Time period (dates of service) covered by the summary
• Client’s medical and treatment diagnoses
• A summary of client’s response to therapy and current treatment plan, to include:
  • Documentation of any issues limiting the client’s progress
  • Documentation of objective measures of functional progress related to each treatment goal established on the initial evaluation
  • An assessment of the client’s therapy prognosis and overall functional progress
  • Documentation of client’s participation in treatment as well as client or responsible adult’s participation or adherence with a home treatment program
  • Updated or new functional and measurable short and long-term treatment goals with time frames, as applicable
  • Documentation of client’s continued need for therapy
  • Clearly established discharge criteria
  • Documentation of consults with other professionals and services or coordination of service when applicable.
• The progress summary must be signed and dated by the therapist responsible for the therapy services.

5.2.5 Requests for Recertification - Chronic Therapy Services

5.2.5.1 Re-evaluation (every 180 days)
A re-evaluation is a comprehensive evaluation and must take place every 180 days and contains all the elements of an initial evaluation, including affirmation that the client’s THSteps checkup is current or that a developmental screening was performed by the prescribing provider within the last 60 days. It may be used to make a determination whether or not skilled therapy is medically necessary, or when determining the effectiveness of the current plan, or when the current plan requires significant modification and revision of the interventions and goals due to changes in the client’s medical status or lack of progress with the current treatment. A re-evaluation requires authorization and must be submitted with the recertification request (procedure codes 97164, 97168, and S9152).

Routine reassessments that occur during each treatment session or visit or for a progress report required for an extension of services or discharge summary are not considered a comprehensive re-evaluation.

Tests used must be norm-referenced, standardized, and specific to the therapy provided.

Refer to: Subsection 5.3, “Developmental Delay Criteria” in this handbook for information about documentation about developmental delay criteria.

A recertification request may be considered when services will be medically needed after the previously approved authorization period ends.

A complete request must be received no earlier than 30 days before the current authorization period expires.

Requests for recertification services received after the current authorization expires will be denied for dates of service that occurred before the date the request is received.

A re-evaluation may occur as early as 60 days prior to the end of the current authorization period.

A therapy re-evaluation is considered current when it is performed within 60 days before the current authorization period expires.

The re-evaluation must occur within 30 days of the signed and dated order from the referring provider.
Prior authorization for recertification requests may be considered for increments up to 180 days for each request with documentation supporting the medical necessity including all of the following:

- Texas Medicaid Physical, Occupational, or Speech Therapy (PT, OT, ST) Prior Authorization Form or electronic equivalent signed and dated by the therapist and by the prescribing provider. When the request form is unsigned by the prescribing provider, it must be accompanied by a written order or prescription or a verbal order for the prescribed therapy services.

- A re-evaluation must include a revised treatment plan or plan of care including all of the following:
  - Documentation that the THSteps checkup is current or that a developmental screening was performed by the prescribing provider within the last 60 days
  
  Note: Additional documentation is not necessary if the prescribing provider signs the Texas Medicaid Physical, Occupational, or Speech Therapy (PT, OT, ST) Prior Authorization Form.
  
  - Date therapy services started
  - Changes in the treatment plan, the rationale, and the requested change in frequency of visits
  - Documentation of reasons continued therapy services are medically needed
  - Documentation of developmental delay

Refer to: Subsection 5.3, “Developmental Delay Criteria” in this handbook for information about documentation about developmental delay criteria.

- Documentation of client’s participation in treatment, as well as client or responsible adult’s participation or adherence with a home treatment program

- New treatment plan or POC for the recertification dates of service requested

- Updated or new functional and measurable short and long-term treatment goals with new time frames, as applicable. Previous authorization period’s goals and progress must be included.

- Prognosis with clearly established discharge criteria. The discharge plan must reflect realistic expectations from the episode of therapy.

- Documentation of consults with other professionals and services or referrals made and coordination of service when applicable (e.g., for school aged clients, documentation of the coordination of care and referrals made for school therapies)

- The updated treatment plan or POC must be signed and dated by the therapist responsible for the therapy services.

5.2.6 Requests for Revisions to Existing Prior Authorizations or Recertification for Acute and Chronic Therapy Services

A revision to an existing authorization/recertification must be documented in the client’s record when significant changes occur in the frequency or treatment plan. When frequency is increased, or services requiring separate authorization are added, a request for revision must be submitted for prior authorization.

Requests for revisions must be received no later than five business days from the date the revised therapy treatments are initiated. Requests for revisions received after the five business day period will be denied for dates of service that occurred before the date the request was received.

A prior authorization request for revisions to services may be considered up to the end of the current approved prior authorization.
Requests for revision must be submitted with the following documentation:

- Texas Medicaid Physical, Occupational, or Speech Therapy (PT, OT, ST) Prior Authorization Form, including the date the revision was initiated, signed and dated by the therapist and signed and dated by the prescribing provider. When the request form is not signed and dated by the prescribing provider, it must be accompanied by a written order or prescription or a verbal order for the prescribed services.
- Progress summary for acute services indicating the medical rationale for the change requested, and
- Updated treatment plan or POC addressing all the elements of the previous plan and addressing all revisions to the services planned, including updated or new functional and measurable short and long-term treatment goals with new time frames, as applicable. Previous authorization period’s goals and progress must be included.
- The updated treatment plan or POC must be signed and dated by the therapist responsible for the therapy services.

5.2.7 Change of Therapy Provider

If a provider or client discontinues therapy during an existing prior authorized period and the client requests services through a new provider outside the current group or agency, they must start a new request for authorization and submit all documentation required for an initial evaluation, and also the following:

- A change-of-therapy provider letter, signed by the client or responsible adult
- The letter must document the date that the client ended therapy (effective date of change) with the previous provider, or last date of service
- The name of the new provider and previous provider

When a provider or client discontinues therapy during an existing prior authorization period and the client requests services through a new provider located within the same enrolled group of providers or within a group of independently enrolled providers collaboratively working together, the new provider can use the same evaluation and plan of care.

The authorization period will not change when the provider changes.

5.2.8 Treatment Note

The following documentation must be kept on file by the treating provider and be available when requested:

- Client’s name
- Date of service
- Time in and out of each therapy session
- Objectives addressed (should coincide with plan of care) and progress noted, if applicable
- A description of specific therapy services provided and the activities rendered during each therapy session, along with a form of measurement.
- Assessments of client’s progress or lack of progress
- Treatment notes must be legible
- Therapist must sign each date of entry with full signature and credentials

All documentation for evaluations, re-evaluations, progress summaries, treatment notes, and discharge summaries must show client’s name, date of service, time in and time out of each therapy session.
5.3 Developmental Delay Criteria

To establish a developmental delay, all of the following criteria must be met:

- Tests used must be norm-referenced, standardized, and specific to the therapy provided.
- Retesting with norm-referenced standardized test tools for re-evaluations must occur every 180 days. Tests must be age appropriate for the child being tested and providers must use the same testing instrument as used in the initial evaluation. If reuse of the initial testing instrument is not appropriate, i.e. due to change in client status or restricted age range of the testing tool, provider should explain the reason for the change.
- Eligibility for therapy will be based upon a score that falls 1.5 standard deviations (SD) or more below the mean in at least one subtest area of composite score on a norm-referenced, standardized test. Raw scores must be reported along with score reflecting SD from mean.
- When the client’s test score is less than 1.5 SD below the mean, a criterion-referenced test along with informed evidenced-based clinical opinion must be included to support the medical necessity of services and will be sent to physician review to determine medical necessity.
- If a child cannot complete norm-referenced standardized assessments, then a functional description of the child’s abilities and deficits must be included. Measurable functional short and long term goals will be considered along with test results. Documentation of the reason a standardized test could not be used must be included in the evaluation.

Specific developmental delay criteria requirements for speech diagnoses are as follows:

- Language—at least one norm-referenced, standardized test with good reliability and validity, a score that falls 1.5 SD or more below the mean, and clinical documentation of an informal assessment that supports the delay
- Articulation—at least one norm-referenced, standardized test with good reliability and validity, a score that falls 1.5 SD or more below the mean, and clinical documentation of an informal assessment that supports the delay
- Apraxia—at least one norm-referenced, standardized test with good reliability and validity, a score that falls 1.5 SD or more below the mean, and clinical documentation of an informal assessment that supports the delay
- Fluency—at least one norm-referenced, standardized test with good reliability, a score that falls 1.5 SD or more below the mean, and clinical documentation of an informal assessment that supports the delay
- Voice—a medical evaluation is required for eligibility and based on medical referral
- Oral Motor/Swallowing/Feeding—an in-depth, functional profile of oral motor structures and function

If the client’s test score is less than 1.5 SD below the mean, additional documentation supporting the client’s medical need for therapy will be considered and the request will be sent to physician review to determine medical necessity.

Additional speech therapy visits or sessions may be considered for moderate speech language, articulation, voice and dysphagia developmental delays when documentation submitted supports medical necessity as delineated in the frequency criteria in this handbook.

5.4 Age Adjustment for Children Born Prematurely

Age is adjusted for children born before 37 weeks gestation and is based on a 40-week term. The developmental age must be measured against the adjusted age rather than chronological age until the child is 24 months old. The age adjustment cannot exceed 16 weeks.
5.5 **PT, OT, and ST Procedure Codes**

PT, OT, and ST treatment procedure codes are either time-based and billable in units or untimed and billable per daily encounter.

5.5.1 **Timed PT and OT Treatment Procedure Codes**

All time-based PT and OT treatment procedure codes are cumulatively limited to one hour per date of service per discipline (4 units).

The following time-based PT and OT treatment procedure codes must be billed in 15 minute increments and are limited to a combined total of 2 units (thirty minutes) per date of service per discipline:

<table>
<thead>
<tr>
<th>Procedure Codes</th>
</tr>
</thead>
<tbody>
<tr>
<td>97034</td>
</tr>
<tr>
<td>97035</td>
</tr>
</tbody>
</table>

The following time-based PT and OT treatment procedure code must be billed in 15 minute increments, is limited to a combined total of 3 units (45 minutes) per date of service per discipline, and is not payable in the home or other setting:

<table>
<thead>
<tr>
<th>Procedure Codes</th>
</tr>
</thead>
<tbody>
<tr>
<td>97036</td>
</tr>
</tbody>
</table>

The following time-based PT and OT treatment procedure codes must be billed in 15 minute increments and are limited to a combined total of 4 units (one hour) per date of service per discipline:

<table>
<thead>
<tr>
<th>Procedure Codes</th>
</tr>
</thead>
<tbody>
<tr>
<td>97032 97033 97110 97112 97113 97116 97124 97140 97530 97535</td>
</tr>
<tr>
<td>97537 97542 97750 97760 97761 97763</td>
</tr>
</tbody>
</table>

*Note: Procedure code 97113 is not payable to home health agencies.*

5.5.2 **Untimed PT and OT Treatment Procedure Codes**

The following supervised modality PT and OT treatment procedure codes are limited to once per date of service per procedure code and must be delivered on the same date of service as one or more time-based PT and OT procedure codes and are subject to CMS NCCI relationships:

<table>
<thead>
<tr>
<th>Procedure Codes</th>
</tr>
</thead>
<tbody>
<tr>
<td>97012 97014 97016 97018 97022 97024 97026 97028</td>
</tr>
</tbody>
</table>

The following PT and OT group therapy treatment procedure code is limited to once per date of service:

<table>
<thead>
<tr>
<th>Procedure Codes</th>
</tr>
</thead>
<tbody>
<tr>
<td>97150</td>
</tr>
</tbody>
</table>

Separate prior authorization is required for medically necessary therapeutic procedures not addressed by procedure codes outlined in this handbook.

The following procedure code requires supporting documentation indicating why an unlisted procedure code is required and is limited to once per date of service:

<table>
<thead>
<tr>
<th>Procedure Codes</th>
</tr>
</thead>
<tbody>
<tr>
<td>97799</td>
</tr>
</tbody>
</table>
If the therapy treatment services that are billed exceed one hour (four units per day), the claim will be denied, and it may be appealed. On appeal, the provider must meet the following conditions:

- The appeal must document the prior authorization period week or month for the date of service appealed.
- The appeal must include an attestation that the provider has billed all therapy services for the week or month in question.

For clients who are 20 years of age and younger, when physical or occupational group therapy is administered, providers can bill procedure code 97150 for each member of the group.

A client may receive therapy in more than one discipline (physical, occupational, or speech) in more than one setting (outpatient, office or home setting) in one day.

### 5.5.3 ST Treatment Procedure Codes

Individual speech treatment is limited to one encounter per date of service per provider. Only one of the following individual speech treatment procedure codes will be reimbursed per date of service:

<table>
<thead>
<tr>
<th>Procedure Codes</th>
</tr>
</thead>
<tbody>
<tr>
<td>92507</td>
</tr>
</tbody>
</table>

An encounter for speech therapy individual treatment is defined as face-to-face time with the patient and/or caregiver for a length of time compliant with nationally recognized professional speech-language pathology standards for a typical session.

The following group speech therapy procedure code is limited to once per date of service:

<table>
<thead>
<tr>
<th>Procedure Codes</th>
</tr>
</thead>
<tbody>
<tr>
<td>92508</td>
</tr>
</tbody>
</table>

### 5.5.4 PT, OT, and ST Evaluation and Re-evaluation Codes

Evaluation and re-evaluation procedure codes in the following table are untimed:

<table>
<thead>
<tr>
<th>Procedure Codes</th>
</tr>
</thead>
<tbody>
<tr>
<td>92521</td>
</tr>
<tr>
<td>97166</td>
</tr>
</tbody>
</table>

### 5.5.5 PT, OT, and ST Reimbursement Guidelines

If a therapy evaluation or re-evaluation procedure code and like therapy procedure code are billed for the same date of service by any provider, the like therapy evaluation or re-evaluation will be denied.

An evaluation or re-evaluation performed on the same day as therapy from a different therapy type must be performed at distinctly separate times to be considered for reimbursement.

Physical therapy provided in the nursing home setting is limited to the nursing facility because it must be made available to nursing home residents on an “as needed” basis and must be provided directly by the staff of the facility or furnished by the facility through arrangements with outside qualified resources. Nursing home facilities should refrain from admitting clients who need goal directed therapy if the facility is unable to provide these services.

Procedure codes for PT, OT, and ST evaluations are payable once every three years to the same rendering provider.

For acute services, PT, OT, and ST re-evaluations may be reimbursed once every 60 days to any provider when a recertification of services is planned.
For chronic services, PT, OT, and ST re-evaluations are reimbursed once every 180 days to any provider when a recertification of services is planned.

Additional PT, OT, or ST evaluations or re-evaluations exceeding the limits outlined in this handbook may be considered for with documentation of one of the following:

- A significant change in the client’s medical condition as documented in the plan of care or treatment plan
- A change of provider has occurred and a change of provider letter is submitted with the appeal.
- The re-evaluation is required for recertification of an existing authorization.

Therapy services may be billed when rendered in a PPECC even if the provider would typically be restricted to a home setting. Home health providers rendering therapy services in a PPECC must include the PPECC’s NPI number on their institutional claim form, in addition to their own NPI. Therapy providers who bill using a professional claim form must include the PPECC name and NPI number on the claim and indicate outpatient hospital as the place of service. The therapy provider and PPECC must have a written agreement for each client related to the provision of therapy services provided at the PPECC. The written agreement must address responsibilities of both parties, and how the parties will coordinate related to the client’s plan of care. The written agreement must be maintained in the client’s medical record.

A modifier must be used to indicate when treatment services have been rendered by a licensed therapist/physician or a therapy assistant under supervision of a licensed therapist.

The following modifiers are not required for evaluation or re-evaluation codes because those services may not be rendered by therapy assistants.

<table>
<thead>
<tr>
<th>Modifier</th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td>UB</td>
<td>Services delivered by a licensed therapy assistant under supervision of a licensed therapist</td>
</tr>
<tr>
<td>U5</td>
<td>Services delivered by a licensed therapist or physician</td>
</tr>
</tbody>
</table>

### 5.5.6 Therapy Co-Treatment

Claims for co-treatment services must be submitted with modifier U3:

<table>
<thead>
<tr>
<th>Modifier</th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td>U3</td>
<td>Therapy Co-Treatment Modifier</td>
</tr>
</tbody>
</table>

### 6 Adult Services

#### 6.1 Services, Benefits, and Limitations

Unless otherwise specified, “days” refers to calendar days.

Physical therapy (PT), occupational therapy (OT), and speech therapy (ST) services are benefits of Texas Medicaid for the medically necessary short term treatment of an acute medical condition or an acute exacerbation of a chronic medical condition for clients who are 21 years of age and older.

Treatments are expected to significantly improve, restore, or develop physical functions diminished or lost as a result of a recent trauma, illness, injury, disease, surgery, or change in medical condition, in a reasonable and generally predictable period of time (60 days), based on the prescribing provider’s and therapist’s assessment of the client’s restorative potential.

**Note:** Recent is defined as occurring within the past 90 days of the prescribing provider's evaluation of condition.
Treatments are directed towards restoration of or compensation for lost function.

Services do not duplicate those provided concurrently by any other therapy.

Services must meet acceptable standards of medical practice and be specific and effective treatment for the client's condition.

Services are provided within the provider's scope of practice, as defined by state law.

Acute is defined as an illness or trauma with a rapid onset and short duration.

Adult therapy services are limited to a maximum of 120 days per identified acute medical condition or acute exacerbation of a chronic medical condition requiring therapy or whenever the maximum benefit from therapy has been achieved, whichever comes first.

A medical condition is considered chronic when 120 days have passed from the start of therapy, or the condition is no longer expected to resolve or may be slowly progressive over an indefinite period of time.

Physical and occupational therapy services for acute conditions are benefits of Texas Medicaid for adult clients in the office, outpatient, and home settings.

Speech therapy services for acute conditions are benefits of Texas Medicaid for adult clients in the office and outpatient setting only.

Therapy services must be performed by one of the following:

- Licensed physical therapist
- Licensed occupational therapist
- Licensed speech-language pathologist
- Physician within their scope of practice

Therapy services may also be performed by one of the following under the supervision of a licensed therapist of the specific discipline:

- Licensed therapy assistant
- Licensed speech-language pathology intern (Clinical Fellow)

### 6.1.1 Adult Acute Therapy Place of Service

PT and OT providers render acute therapy services for adult clients who are 21 years of age and older in one of the following places of service:

<table>
<thead>
<tr>
<th>Place of Service</th>
<th>Provider types</th>
</tr>
</thead>
<tbody>
<tr>
<td>Office</td>
<td>Physician; podiatrist; independently enrolled therapist; physical therapy group</td>
</tr>
<tr>
<td>Home</td>
<td>Home health agency; independently enrolled physical therapist; and physical therapy group</td>
</tr>
<tr>
<td>Outpatient</td>
<td>Physician; podiatrist; outpatient hospital/clinic</td>
</tr>
<tr>
<td>Other</td>
<td>Independently enrolled therapist; home health agency; physical therapy group</td>
</tr>
</tbody>
</table>

ST providers render acute therapy services for adult clients who are 21 years of age and older in one of the following places of service:

<table>
<thead>
<tr>
<th>Place of Service</th>
<th>Provider types</th>
</tr>
</thead>
<tbody>
<tr>
<td>Office</td>
<td>Physician; independently enrolled speech-language pathologist</td>
</tr>
<tr>
<td>Outpatient</td>
<td>Physician; outpatient hospital/clinic</td>
</tr>
</tbody>
</table>
In determining whether a service requires the skill of a licensed therapist, consideration must be given to the inherent complexity of the service, the condition of the client, and the accepted standards of medical and therapy practice guidelines.

If the service could be performed by the average nonmedical person, the absence of a competent person to perform it does not cause it to be a skilled therapy service.

If the nature of a service is such that it can safely and effectively be performed by the average nonmedical person without direct supervision of a licensed therapist, the services cannot be regarded as skilled therapy.

6.2 Authorization Requirements for Outpatient and Home Health—PT, OT, and ST Services

Prior authorization requests may be submitted to the TMHP Prior Authorization Department via mail, fax, or the electronic portal. Prescribing or ordering providers, dispensing providers, clients’ responsible adults, and clients may sign prior authorization forms and supporting documentation using electronic or wet signatures.

Refer to: Subsection 5.5.1.2, “Document Requirements and Retention” in “Section 5: Fee-for-Service Prior Authorizations” (Vol. 1, General Information) for additional information about electronic signatures.

Coverage periods do not coincide necessarily with calendar weeks or months, but cover a number of services to be scheduled between a start and end date that is assigned during the prior authorization period.

6.2.1 Initial Evaluation and Considerations for Prior Authorization for Treatment

Initial evaluations do not require prior authorization (procedure codes 92521, 92522, 92523, 92524, 92610, 97161, 97162, 97163, 97165, 97166, and 97167); however, documentation kept in the client’s record must include a signed and dated prescribing provider’s order for the evaluation, support a medical need for the therapy evaluation and be available when requested.

To complete the prior authorization process by paper, the provider must complete and submit the prior authorization requirements documentation through fax or mail, and must maintain a copy of the prior authorization request and all submitted documentation in the client’s medical record at the therapy provider’s place of business.

To complete the prior authorization process electronically, the provider must complete and submit the prior authorization requirements documentation through any approved electronic method, and must maintain a copy of the prior authorization request and all submitted documentation in the client’s medical record at the therapy provider’s place of business.

To avoid unnecessary denials, the physician must provide correct and complete information, including documentation of medical necessity for the service(s) requested. The physician must maintain documentation of medical necessity in the client’s medical record. The requesting therapy provider may be asked for additional information to clarify or complete a request.

Therapy services, regardless of place or provider, occurring after the initial evaluation, require prior authorization. Prior authorization requests may not exceed a 60 day period.

Prior authorization (PA) requests must be received no later than five business days from the date therapy treatments following the evaluation are initiated. Requests received after the five-business-day period will be denied for dates of service that occurred before the date that the PA request is received.
6.2.1.1 Documentation

All of the following documentation is required when submitting an initial request for therapy services initiated after the completion of the evaluation:

- A completed Texas Medicaid Physical, Occupational, or Speech Therapy (PT, OT, ST) Prior Authorization Form signed and dated by the therapist and signed and dated by the prescribing provider is required. When the request form is unsigned by the prescribing provider, it must be accompanied by a signed and dated written order, or prescription, or a documented verbal order delineating the prescribed therapy services.

- Documentation of the acute or acute exacerbation of the medical condition requiring therapy. Evaluation and Treatment Plan or Plan of Care (POC) with all of the following required elements:
  - Client’s medical history and background
  - All medical diagnoses related to the client’s condition
  - Date of onset of the client’s condition requiring therapy, or exacerbation date as applicable
  - Date of evaluation
  - Baseline objective measurements documented based on any testing performed
  - Explanation of how identified limitations impair the overall function of the client
  - Safety risks
  - Client-specific, measurable short and long-term functional goals within the length of service time requested
  - Interpretation of the results of the evaluation, including recommendations for therapy amount, frequency per week and duration of services
  - When a client also receives PPECC services, indicate if therapy services will be delivered in a PPECC setting, the amount, frequency, and duration of the therapy services to be delivered in a PPECC setting in contrast with other locations (e.g., the home)
  - Therapy treatment plan/POC to include specific modalities and treatments planned
  - Documentation of client’s primary language
  - Documentation of client’s age and date of birth
  - Adaptive equipment or assistive devices, as applicable
  - Prognosis for improvement
  - Time in and time out on evaluation
  - Requested dates of service for planned treatments after the completion of the evaluation
  - Responsible adult’s expected involvement in client’s treatment
  - History of prior therapy and referrals as applicable
  - Signature and date of treating therapist

Additional requirements for speech therapy include one or more of the following:

- Language evaluations—oral-peripheral speech mechanism examination and formal or informal assessment of hearing, articulation, voice and fluency skills.

- Speech production (voice)—formal screening of language skills, and formal or informal assessment of hearing, voice and fluency skills.
• Speech production (fluency)—formal screening of language skills, formal or informal assessment of hearing, voice and fluency skills.

• Oral Motor/Swallowing/Feeding—If swallowing problems and/or signs of aspiration are noted, then include a statement indicating that a referral for a video fluoroscopic swallow study has been made; formal screening of language skills, formal or informal assessment of hearing, voice and fluency skills.

6.2.2 Written and Verbal Orders

For new authorizations and recertifications of therapies, if the submitted request form is not signed and dated by the physician, the request must be accompanied by a verbal or written order.

The request form or written or verbal order must be signed and dated within the 60-day period before the initiation of services. A prescribing physician’s order to evaluate and treat is acceptable for the evaluation or re-evaluation, but is not acceptable for the therapy treatment. The therapy treatment order must contain the prescribing provider’s ordered frequency and duration.

The documentation for a verbal order must meet the following criteria:

• It must be signed and dated by the licensed professional who by state and federal law may take a verbal order.

• It must have the name and credentials of the licensed professional who took the order and who is responsible for furnishing or supervising the ordered services.

• The verbal order must include the date on which the verbal order was taken.

• The verbal order must include the services and the frequency and duration that was prescribed by the ordering physician.

6.2.3 Requests for Recertification -Up to an Additional 60 days for Acute Services

A recertification request may be considered when services will be medically needed after the previously approved authorization period ends.

Re-evaluation codes (procedure codes 97164, 97168, and S9152) require prior authorization and must be submitted with the recertification request. Required documentation for recertifications includes a progress summary and Texas Medicaid Physical, Occupational, or Speech Therapy (PT, OT, ST) Prior Authorization Form.

A complete recertification request must be received no earlier than 30 days before the current authorization period expires. Requests for recertification services received after the current authorization expires will be denied for dates of service that occurred before the date the submitted request was received.

One recertification request may be considered for an additional 60 days for each therapy service request with documentation supporting the medical necessity including all of the following:

• Texas Medicaid Physical, Occupational, or Speech Therapy (PT, OT, ST) Prior Authorization Form or electronic equivalent signed and dated by the therapist and signed and dated by the ordering physician. When the request form is unsigned by the physician, it must be accompanied by a written order or prescription or a verbal order for the prescribed therapy services.

• A progress summary (see progress summary documentation requirements), and

• An updated treatment plan or POC for the recertification dates of service requested, including all of the following:
  • Date therapy services started
• Changes in the treatment plan, the rationale and the requested change in frequency of visits for changing the plan
• Documentation of reasons continued therapy services are medically needed
• Documentation of client’s participation in treatment, as well as client and responsible adult’s participation or adherence with a home treatment program
• Updated or new functional and measurable short and long-term treatment goals with new time frames, as applicable
• Adaptive equipment or assistive devices, as applicable
• Prognosis with clearly established discharge criteria
• Documentation of consults with other professionals and services or referrals made and coordination of service when applicable
• The updated treatment plan or plan of care must be signed and dated by the therapist responsible for the therapy services.

A progress summary which may be contained in the last treatment note, must be included with the recertification request and contains all of the following:
• Date therapy started
• Date the summary completed
• Time period (dates of service) covered by the summary
• Client’s medical and treatment diagnoses
• A summary of client’s response to therapy and current treatment plan, to include:
  • Documentation of any issues limiting the client’s progress
  • Documentation of objective measures of functional progress related to each treatment goal established on the initial evaluation
  • An assessment of the client’s therapy prognosis and overall functional progress
  • Documentation of client’s participation in treatment as well as client and responsible adult’s participation or adherence with a home treatment program
  • Updated or new functional and measurable short and long-term treatment goals with time frames, as applicable
  • Documentation of client’s continued need for therapy
  • Clearly established discharge criteria
  • Documentation of consults with other professionals and services or coordination of service when applicable.
• The progress summary must be signed and dated by the therapist responsible for the therapy services.

6.2.4 Requests for Revisions to Existing Prior Authorization or Recertification

A revision to an existing authorization/recertification must be documented in the client’s record when significant changes occur in the frequency or treatment plan. When the frequency is increased or services requiring separate authorization are added, a request for revision must be submitted for prior authorization.

Requests for revisions must be received no later than five business days from the date the revised therapy treatments are initiated.
Requests for revisions received after the five business day period will be denied for dates of service that occurred before the date the request was received.

- A prior authorization request for revisions to services may be considered up to the end of the current approved prior authorization.
- Requests for revision must be submitted with the following documentation:
  - Texas Medicaid Physical, Occupational, or Speech Therapy (PT, OT, ST) Prior Authorization Form, including the date the revision was initiated, signed and dated by the therapist and signed and dated by the physician. When the request form is not signed and dated by the physician, it must be accompanied by a written order or prescription or a verbal order for the prescribed services.
  - Progress summary including the medical rationale for the change requested, and
  - Updated treatment plan or POC addressing all the elements of the previous plan and addressing all revisions to the services planned, including updated or new functional and measurable short and long-term treatment goals with new time frames, as applicable. Previous authorization period’s goals and progress must be included.
  - The updated treatment plan or POC must be signed and dated by the therapist responsible for the therapy services.

### 6.2.5 Change of Therapy Provider

If a provider or client discontinues therapy during an existing prior authorized period and the client requests services through a new provider outside the current group or agency, they must start a new request for authorization and submit all documentation required for an initial evaluation, and also all of the following:

- A change-of-therapy provider letter signed by the client or responsible adult,
- The letter must document the date that the client ended therapy (effective date of change) with the previous provider, or last date of service,
- The name of the new provider and previous provider

When a provider or client discontinues therapy during an existing prior authorization period and the client requests services through a new provider located within the same enrolled group of providers or within a group of independently enrolled providers collaboratively working together, the new provider can use the same evaluation and plan of care.

The authorization period will not change when the provider changes.

### 6.2.6 Treatment Note

The following documentation must be kept on file by the treating provider and available when requested:

- Client’s name
- Date of service
- Time in and out of each therapy session
- Objectives addressed (should coincide with plan of care) and progress noted, if applicable
- A description of specific therapy services provided and the activities rendered during each therapy session, along with a form of measurement.
- Assessments of client’s progress or lack of progress
- Treatment notes must be legible
• Therapist must sign each date of entry with full signature and credentials

All documentation for evaluations, re-evaluations, progress summaries, treatment notes, and discharge summaries must show client’s name, date of service, time in and time out for each therapy session.

6.3  PT, OT, and ST Procedure Codes

PT, OT, and ST treatment procedure codes are either time-based and billable in units or untimed and billable per daily encounter.

6.3.1  PT and OT Treatment Procedure Codes

All time-based PT and OT treatment procedure codes are cumulatively limited to one hour per date of service per discipline (4 units).

The following time-based PT and OT treatment procedure codes must be billed in 15 minute increments and are limited to a combined total of 2 units (thirty minutes) per date of service per discipline:

<table>
<thead>
<tr>
<th>Procedure Codes</th>
</tr>
</thead>
<tbody>
<tr>
<td>97034</td>
</tr>
<tr>
<td>97035</td>
</tr>
</tbody>
</table>

The following time-based PT and OT treatment procedure code must be billed in 15 minute increments, is limited to a combined total of 3 units (45 minutes) per date of service per discipline, and is not payable in the home or other setting:

<table>
<thead>
<tr>
<th>Procedure Codes</th>
</tr>
</thead>
<tbody>
<tr>
<td>97036</td>
</tr>
</tbody>
</table>

The following time-based PT and OT treatment procedure codes must be billed in 15 minute increments and are limited to a combined total of 4 units (one hour) per date of service per discipline:

<table>
<thead>
<tr>
<th>Procedure Codes</th>
</tr>
</thead>
<tbody>
<tr>
<td>97032 97033 97110 97112 97113 97116 97124 97140 97530 97535</td>
</tr>
<tr>
<td>97537 97542 97750</td>
</tr>
</tbody>
</table>

Note: Procedure code 97113 is not payable to home health agencies.

6.3.2  Untimed PT and OT Treatment Procedure Codes

The following supervised modality PT and OT treatment procedure codes are limited to once per date of service per procedure code and must be delivered on the same date of service as one or more time-based PT and OT procedure code(s) and are subject to CMS NCCI relationships:

<table>
<thead>
<tr>
<th>Procedure Codes</th>
</tr>
</thead>
<tbody>
<tr>
<td>97012 97014 97016 97018 97022 97024 97026 97028</td>
</tr>
</tbody>
</table>

The following PT and OT group therapy treatment procedure code is limited to once per date of service:

<table>
<thead>
<tr>
<th>Procedure Codes</th>
</tr>
</thead>
<tbody>
<tr>
<td>97150</td>
</tr>
</tbody>
</table>

Separate prior authorization is required for medically necessary therapeutic procedures not addressed by procedure codes outlined in this handbook.
The following procedure code requires supporting documentation indicating why an unlisted procedure code is required and is limited to once per date of service:

<table>
<thead>
<tr>
<th>Procedure Codes</th>
<th>97799</th>
</tr>
</thead>
</table>

If the therapy treatment services that are billed exceed one hour (four units per day), the claim will be denied, and it may be appealed. On appeal, the provider must meet the following conditions:

- The appeal must document the prior authorization period week or month for the date of service appealed.
- The appeal must include an attestation that the provider has billed all therapy services for the week or month in question.

For clients who are 21 years of age and older, when physical or occupational group therapy is administered, providers should bill procedure code 97150 for each member of the group.

A client may receive therapy in more than one discipline (physical, occupational, or speech) in the outpatient, office or home setting in one day.

### 6.3.3 ST Treatment Procedure Codes

Individual speech treatment is limited to one encounter per date of service per provider. Only one of the following individual speech treatment procedure codes will be reimbursed per date of service:

<table>
<thead>
<tr>
<th>Procedure Codes</th>
<th>92507 92526</th>
</tr>
</thead>
</table>

An encounter for speech therapy individual treatment is defined as face-to-face time with the patient and/or caregiver for a length of time compliant with nationally recognized professional speech-language pathology standards for a typical session.

The following group speech therapy procedure code is limited to once per date of service:

<table>
<thead>
<tr>
<th>Procedure Codes</th>
<th>92508</th>
</tr>
</thead>
</table>

### 6.3.4 PT, OT, and ST Evaluation and Re-evaluation Procedure Codes

Evaluation and re-evaluation procedure codes in the following table are untimed:

<table>
<thead>
<tr>
<th>Procedure Codes</th>
<th>92521 92522 92523 92524 92610 97161 97162 97163 97164 97165 97166 97167 97168 S9152</th>
</tr>
</thead>
</table>

### 6.3.5 PT, OT, and ST Reimbursement Guidelines

If a therapy evaluation or re-evaluation procedure code and like therapy procedure codes are billed for the same date of service by any provider, the like therapy evaluation or re-evaluation will be denied.

An evaluation or re-evaluation performed on the same day as therapy from a different therapy type must be performed at distinctly separate times to be considered for reimbursement.
Physical therapy provided in the nursing home setting is limited to the nursing facility because it must be made available to nursing home residents on an “as needed” basis and must be provided directly by the staff of the facility or furnished by the facility through arrangements with outside qualified resources. Nursing home facilities should refrain from admitting clients who need goal directed therapy if the facility is unable to provide these services.

Procedure codes for PT, OT, ST evaluations are payable once every three years to the same rendering provider.

For acute services, PT, OT, ST re-evaluations may be reimbursed once every 60 days to any provider when a recertification of services is planned.

Additional PT, OT, or ST evaluations or re-evaluations exceeding the limits outlined in this handbook may be considered for reimbursement on appeal with documentation of one of the following:

- A significant change in the client’s medical condition as documented in the plan of care or treatment plan,
- A change of provider has occurred and a change of provider letter is submitted with the appeal.

The re-evaluation is required for recertification of an existing authorization.

A modifier must be used to indicate when treatment services have been rendered by a licensed therapist/physician or a therapy assistant under supervision of a licensed therapist.

The following modifiers are not required for evaluation or re-evaluation codes because those services may not be rendered by therapy assistants:

<table>
<thead>
<tr>
<th>Modifier</th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td>UB</td>
<td>Services delivered by a licensed therapy assistant under supervision of a licensed therapist</td>
</tr>
<tr>
<td>U5</td>
<td>Services delivered by a licensed therapist or physician</td>
</tr>
</tbody>
</table>

This modifier is to be utilized as indicated with all physical, occupational, and speech therapy treatment procedure codes.

Claims for co-treatment services must be submitted with modifier U3:

<table>
<thead>
<tr>
<th>Modifier</th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td>U3</td>
<td>Therapy Co-Treatment Modifier</td>
</tr>
</tbody>
</table>

## 7 Claims Filing and Reimbursement

### 7.1 Claims Information

Therapy services must be submitted to TMHP in an approved electronic format or on a CMS-1500 paper claim form. Providers may purchase CMS-1500 paper claim forms from the vendor of their choice. TMHP does not supply the forms. Claims may be filed electronically in a CMS-1500 format as long as the nine-digit prior authorization number is reflected in the equivalent electronic field.

CORF and ORF providers must submit services in an approved electronic claims format or on the UB-04 CMS-1450 paper claim form from the vendor of their choice. TMHP does not supply the forms. Revenue and Current Procedural Terminology (CPT) procedure codes are used when submitting claims for CORF and ORF services. The only POS is outpatient facility (POS 5).

When completing a CMS-1500 or a UB-04 CMS-1450 paper claim form, all required information must be included on the claim, as TMHP does not key information from attachments.
Refer to: “Section 3: TMHP Electronic Data Interchange (EDI)” (Vol. 1, General Information) for information on electronic claims submissions.

“Section 6: Claims Filing” (Vol. 1, General Information) for general information about claims filing.

Subsection 6.5, “CMS-1500 Paper Claim Filing Instructions” in “Section 6: Claims Filing” (Vol. 1, General Information). Blocks that are not referenced are not required for processing by TMHP and may be left blank.


7.2 **Reimbursement/Billing Guidelines**

Physical, occupational, and speech therapy services are reimbursed in accordance with 1 TAC § 355. See the OFL or the applicable fee schedule on the TMHP website at [www.tmhp.com](http://www.tmhp.com) for reimbursement rates.

Therapy providers are reimbursed in accordance with 1 TAC §355.8085.

Refer to: “Section 3: TMHP Electronic Data Interchange (EDI)” (Vol. 1, General Information) for information on electronic claims submissions.

Subsection 6.4, “Claims Filing Instructions” in “Section 6: Claims Filing” (Vol. 1, General Information) for general information about claims filing.

Subsection 6.5, “CMS-1500 Paper Claim Filing Instructions” in “Section 6: Claims Filing” (Vol. 1, General Information) in “Section 6: Claims Filing” (Vol. 1, General Information). Blocks that are not referenced are not required for processing by TMHP and may be left blank.

Texas Medicaid implemented mandated rate reductions for certain services. The OFL and static fee percentage reductions applied. Additional information about rate changes is available on the TMHP website at [www.tmhp.com/resources/rate-and-code-updates/rate-changes](http://www.tmhp.com/resources/rate-and-code-updates/rate-changes).

When there is a change of provider, or a change in the client’s medical condition requiring therapy, a denied claim for a therapy (PT, OT, or ST) evaluation, re-evaluation or swallowing function evaluation that exceeded the limits outlined in this handbook may be considered on appeal for reimbursement with documentation of one of the following:

- A change in the client’s medical condition or new therapy related diagnosis with date of onset documented in the plan of care or treatment plan
- A change of provider letter signed and dated by the client or responsible adult documenting all of the following:
  - The date the client ended therapy (effective date of change) with the previous provider
  - The name of the new provider and previous provider

7.2.1 **Method for Counting Minutes for Timed Procedure Codes in 15-Minute Units**

Modifiers GP, GO, and GN are required on all claims except when billing evaluation and re-evaluation procedure codes. The AT modifier must be included on claims for acute therapy services.

All claims for reimbursement of procedure codes paid in 15 minute increments are based on the actual amount of billable time associated with the service. For those services for which the unit of service is 15 minutes (1 unit = 15 minutes), partial units should be rounded up or down to the nearest quarter hour.
To calculate billing units, count the total number of billable minutes for the calendar day for the client, and divide by 15 to convert to billable units of service. If the total billable minutes are not divisible by 15, the minutes are converted to one unit of service if they are greater than seven and converted to zero units of service if they are seven or fewer minutes.

For example, 68 total billable minutes/15 = 4 units + 8 minutes. Since the 8 minutes are more than 7 minutes, those 8 minutes are converted to one unit. Therefore, 68 total billable minutes = 5 units of service.

Time intervals for 1 through 8 units are as follows:

<table>
<thead>
<tr>
<th>Units</th>
<th>Number of Minutes</th>
</tr>
</thead>
<tbody>
<tr>
<td>0 units</td>
<td>0 minutes through 7 minutes</td>
</tr>
<tr>
<td>1 unit</td>
<td>8 minutes through 22 minutes</td>
</tr>
<tr>
<td>2 units</td>
<td>23 minutes through 37 minutes</td>
</tr>
<tr>
<td>3 units</td>
<td>38 minutes through 52 minutes</td>
</tr>
<tr>
<td>4 units</td>
<td>53 minutes through 67 minutes</td>
</tr>
<tr>
<td>5 units</td>
<td>68 minutes through 82 minutes</td>
</tr>
<tr>
<td>6 units</td>
<td>83 minutes through 97 minutes</td>
</tr>
<tr>
<td>7 units</td>
<td>98 minutes through 112 minutes</td>
</tr>
<tr>
<td>8 units</td>
<td>113 minutes through 127 minutes</td>
</tr>
</tbody>
</table>

### 7.3 Claims Resources

Providers may refer to the following sections or forms when filing claims:

<table>
<thead>
<tr>
<th>Resource</th>
<th>Location</th>
</tr>
</thead>
<tbody>
<tr>
<td>Appendix A: State, Federal, and TMHP Contact Information</td>
<td>Appendix A (Vol. 1, General Information)</td>
</tr>
<tr>
<td>Appendix D: Acronym Dictionary</td>
<td>Appendix D (Vol. 1, General Information)</td>
</tr>
<tr>
<td>Automated Inquiry System (AIS)</td>
<td>Appendix A: State, Federal, and TMHP Contact Information (Vol. 1, General Information)</td>
</tr>
<tr>
<td>CMS-1500 Paper Claim Filing Instructions</td>
<td>Subsection 6.5 (Vol. 1, General Information)</td>
</tr>
<tr>
<td>Section 3: TMHP Electronic Data Interchange (EDI)</td>
<td>Section 3 (Vol. 1, General Information)</td>
</tr>
<tr>
<td>TMHP Electronic Claims Submission</td>
<td>Subsection 6.2 (Vol. 1, General Information)</td>
</tr>
<tr>
<td>UB-04 CMS-1450 Paper Claim Filing Instructions</td>
<td>Subsection 6.6 (Vol. 1, General Information)</td>
</tr>
</tbody>
</table>

### 7.4 Contact TMHP

The TMHP Contact Center at 1-800-925-9126 is available Monday-Friday from 7 a.m. to 7 p.m., Central Time.

### 8 Forms

The following linked forms can also be found on the Forms page of the Provider section of the TMHP website at [www.tmhp.com](http://www.tmhp.com):

<table>
<thead>
<tr>
<th>Forms</th>
</tr>
</thead>
<tbody>
<tr>
<td>Texas Medicaid Physical, Occupational, or Speech Therapy (PT, OT, ST) Prior Authorization Form</td>
</tr>
</tbody>
</table>
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   2.3 Documentation Requirements ....................................................... 23
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1 General Information

This information is intended for Texas Medicaid independent (freestanding) laboratories, radiological laboratories, and physiological laboratories. The handbook provides information about Texas Medicaid’s benefits, policies, and procedures applicable to these providers.

Important: All providers are required to read and comply with “Section 1: Provider Enrollment and Responsibilities” (Vol. 1, General Information). In addition to required compliance with all requirements specific to Texas Medicaid, it is a violation of Texas Medicaid rules when a provider fails to provide health-care services or items to Medicaid clients in accordance with accepted medical community standards and standards that govern occupations, as explained in Title 1 Texas Administrative Code (TAC) §371.1659. Accordingly, in addition to being subject to sanctions for failure to comply with the requirements that are specific to Texas Medicaid, providers can also be subject to Texas Medicaid sanctions for failure, at all times, to deliver health-care items and services to Medicaid clients in full accordance with all applicable licensure and certification requirements including, without limitation, those related to documentation and record maintenance.

Refer to: “Section 1: Provider Enrollment and Responsibilities” (Vol. 1, General Information).

1.1 Payment Window Reimbursement Guidelines for Services Preceding an Inpatient Admission

According to three-day and one-day payment window reimbursement guidelines, most professional and outpatient diagnostic and nondiagnostic services that are rendered within the designated timeframe of an inpatient hospital stay and are related to the inpatient hospital admission will not be reimbursed separately from the inpatient hospital stay if the services are rendered by the hospital or an entity that is wholly owned or operated by the hospital.

These reimbursement guidelines do not apply in the following circumstances:

- The professional services are rendered in the inpatient hospital setting.
- The hospital and the physician office or other entity are both owned by a third party, such as a health system.
- The hospital is not the sole or 100-percent owner of the entity.

Refer to: Subsection 3.7.4.14, “Payment Window Reimbursement Guidelines” in the Inpatient and Outpatient Hospital Services Handbook (Vol. 2, Provider Handbooks) for additional information about the payment window reimbursement guidelines.

2 Independent Laboratory

The requirements in this section apply to all providers who bill laboratory services.

2.1 Enrollment

Providers must meet the following requirements and submit a complete application in order to enroll as independent (freestanding) laboratory providers:

- The provider must be actively enrolled in Medicare as an independent laboratory.
- The independent laboratory must be independent from a physician’s office or hospital.
- The independent laboratory must meet staff, equipment, and testing capability standards for certification by the Health and Human Services Commission (HHSC).
2.1.1 Clinical Laboratory Improvement Amendments (CLIA)

CLIA regulations set standards that are designed to improve quality in all laboratory testing and include specifications for quality control (QC), quality assurance (QA), patient test management, personnel, and proficiency testing.

The regulations concern all laboratory testing that is used for the assessment of human health or the diagnosis, prevention, or treatment of disease. Under CLIA 88, all clinical laboratory providers (including those located in physicians’ offices), regardless of location, size, or type of laboratory, must meet certain standards based on the complexity of the tests they perform.

Providers must hold the appropriate CLIA certificates to perform certain tests as indicated in this handbook. Providers that are certified only to perform waived tests must use modifier QW as indicated on the CMS website.

Refer to: The CLIA Regulations and Federal Register Documents page of the Centers for Medicare & Medicaid Services (CMS) website at www.cms.gov for the CLIA rules and regulations. The regulations are found at Title 42 Code of Federal Regulations, Part 493.

2.1.2 CLIA Requirements

To be eligible for reimbursement by Medicare and Medicaid, all providers that perform laboratory tests must do the following:

• Pay the applicable fee to CMS.

• Contact HHSC at 1-512-834-6650 to receive a CLIA registration or certification number. Submit CLIA applications to the following address:

  Health Facility Licensing and Certification Division  
  HHSC  
  1100 West 49th Street  
  Austin, TX 78756

• Notify the Texas Medicaid & Healthcare Partnership (TMHP) of the assigned CLIA number at the following address:

  Texas Medicaid & Healthcare Partnership  
  Provider Enrollment  
  PO Box 200795  
  Austin, TX 78720-0795

TMHP monitors claims that are submitted by clinical laboratory providers to verify that the clinical laboratory has a CLIA number on file. If the provider does not have a CLIA number on file with TMHP, the laboratory services claims may be denied.

2.2 Services, Benefits, Limitations, and Prior Authorization

Texas Medicaid only covers professional and technical services that an independent laboratory is certified by CLIA to perform.

Provider documentation must be maintained in the client’s medical record and must delineate the medical need for administering the laboratory test.

The physician is responsible for providing to the performing laboratory the clinical diagnosis code that is associated with the individual test so that the performing laboratory may bill Texas Medicaid directly for the analysis of the specimen.

Refer to: The Current Procedural Terminology (CPT) manual for information regarding examples of laboratory codes. Correct use of CPT coding requires using the most specific laboratory code that matches the services provided, based on the code’s description.
2.2.1 CLIA Certificates
Texas Medicaid follows the Medicare categorization of tests for CLIA certificate holders.

Refer to: The Categorization of Tests page of the CMS website at www.cms.gov for additional information.

Providers that are certified only to perform waived tests must use modifier QW as indicated on the CMS website.

2.2.2 Laboratory Handling Fees and Reference Laboratories

2.2.2.1 Independent Laboratory Providers
An independent laboratory provider may be reimbursed for tests performed in the laboratory and for laboratory handling fees for tests that are forwarded to another laboratory (i.e., reference laboratory).

An independent laboratory that forwards a specimen to another laboratory without performing any tests on that specimen may not bill for any laboratory tests.

An independent laboratory may only bill Texas Medicaid for tests referred to another independent or hospital laboratory if it performs at least one test that it is certified by CLIA to perform, and forwards a portion of the same specimen to the other laboratory to have one or more tests performed. The referring laboratory may then bill for tests it has performed on the specimen. When billing, the following information must be on the claim:

- Block 20: “Yes” box must be checked.
- Block 32: The name, address, and ZIP Code of the reference laboratory (i.e., the laboratory to which the specimen was referred).
- Block 24-J: The provider number of the reference laboratory must be included next to each procedure to be performed by the reference laboratory.

An independent laboratory that forwards a specimen to another laboratory (independent or hospital) may bill a handling fee (procedure code 99001) for collecting and forwarding the specimen to the other laboratory if the specimen is collected by routine venipuncture or catheterization.

2.2.2.2 Physician Providers
A physician may bill only one laboratory handling charge (procedure code 99000) per client visit when the specimen is collected by drawing a blood sample through venipuncture or collecting a urine specimen by catheterization, unless the specimen is divided and sent to different laboratories or there are different specimens collected and sent to different laboratories.

The claim must indicate the name and address of each laboratory where a specimen is sent for more than one laboratory handling charge to be paid.

Refer to: Subsection 9.2.39, “Laboratory Services” in the Medical and Nursing Specialists, Physicians, and Physician Assistants Handbook (Vol. 2, Provider Handbooks) for more information about laboratory services reimbursed to physician providers.

2.2.2.3 Outpatient Hospital Providers
An outpatient hospital may be reimbursed for a laboratory handling charge (procedure code 99001) for each independent laboratory to where it sends specimens when the laboratory handling charge is not being billed through other methods.
2.2.2.4 **Family Planning Laboratory Tests**
Family planning agencies must use procedure code 99000 with a family planning diagnosis code to bill their laboratory handling charges for laboratory specimens sent out; modifier FP must be omitted. Providers may refer to the appropriate section in the provider manual for instructions for billing family planning services. As with procedure code 99000, only one handling fee may be charged for each laboratory to where the agency sends specimens, regardless of the number of specimens taken.

When family planning test specimens, such as Pap smears, are collected, providers must direct the laboratory to indicate that the claim for the test is to be billed as a family planning service.


2.2.3 **Nonclinical Laboratory Procedures**
The reimbursement for nonclinical laboratory procedure codes can be found on the appropriate Texas Medicaid fee schedules on the TMHP website at [www.tmhp.com](http://www.tmhp.com).

2.2.4 **Clinical Laboratory Procedures**
The reimbursement for clinical laboratory procedures can be found on the appropriate Texas Medicaid fee schedule. Fee schedules are available on the TMHP website at [www.tmhp.com](http://www.tmhp.com).

2.2.4.1 **Repeat Procedures**
Modifier 91 should be used for repeat clinical diagnostic tests as follows:

- Modifier 91 must not be used when billing the initial procedure. It must be used to indicate the repeated procedure.
- If more than two services are billed on the same day by the same provider, regardless of the use of modifier 91, the claim or detail is denied.
- If a repeated procedure performed by the same provider on the same day is billed without modifier 91, it is denied as a duplicate procedure.
- If a claim is denied for a quantity more than two or as a duplicate procedure, the times of these procedures and services must be documented on appeal.

Providers may appeal claims that have been denied for documentation of time. Most procedure codes that initially required modifier 91 will continue to be audited for modifier 91.

When appealing claims with modifier 91 for repeat procedures, providers must separate the details. One detail should be appealed without the modifier and one detail with the modifier, including documentation of times for each repeated procedure.

2.2.5 **Automated Laboratory Tests and Laboratory Paneling**
The reimbursement for the complete panel procedure code represents the total payment for all automated laboratory tests that are covered under that panel combined with any other automated tests that are billed for the client for the same date of service. The Texas Medicaid allowable fee for the individual components of the complete laboratory panel will not exceed the automated test panel (ATP) fee for the total number of automated tests that are billed for the client for the same date of service.

When all of the components of the panel are performed, the complete panel procedure code must be billed. When only two or more components of the panel are performed, the individual procedure codes for each laboratory test performed may be billed.
2.2.5.1 Fee Calculations for Automated Tests and Laboratory Panels

Automated test and laboratory panel procedure codes may be reimbursed according to the appropriate ATP level payment based on the total number of automated tests that are performed on the same day for the same client.

Refer to: The “Clinical Laboratory, Automated Test Panels - Insert” Texas Medicaid fee schedule on the TMHP website at www.tmhp.com for the ATP level payment for automated test and lab panel procedure codes.

ATP Level Pricing

The amount that is allowed for each automated test and lab panel procedure code that is billed with the same date of service for the same client will be a percentage of the total ATP level payment. To calculate the automated test pricing, the following information is necessary:

- The number of automated tests that are billed for the client for the same date of service (including individual automated tests and all automated tests that are represented by the laboratory panels.) Procedure codes that are duplicated between panels are not counted more than once.
- The ATP pricing fee that corresponds to the number of automated tests that are billed for the client for the same date of service.
- The total billed amount for all automated test and laboratory panel procedure codes that are billed for the client for the same date of service.

The automated test pricing may be calculated as follows:

Step 1

A percentage for each automated test or lab panel detail is derived from dividing the billed amount (B/A) for each procedure by the total billed amount (TB/A) for all automated test and laboratory panel procedure codes with the same date of service for the same client.

Example:

<table>
<thead>
<tr>
<th>Automated Test</th>
<th>B/A</th>
<th>Percentage</th>
</tr>
</thead>
<tbody>
<tr>
<td>Detail 1 Automated Test</td>
<td>$50.00</td>
<td>29%</td>
</tr>
<tr>
<td>Detail 2 Automated Test</td>
<td>$25.00</td>
<td>14%</td>
</tr>
<tr>
<td>Detail 3 Lab Panel</td>
<td>$100.00</td>
<td>57%</td>
</tr>
<tr>
<td>Detail 4 Clinical Lab Test</td>
<td>$20.00</td>
<td>0%</td>
</tr>
<tr>
<td>TB/A = $175.00</td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

Note: The TB/A is for automated test and laboratory panel procedure codes (details 1, 2, and 3 only). Detail 4 is not included in the calculations for the automated tests because it is a clinical lab procedure code and may be reimbursed as indicated on the fee schedule.

Calculations:

<p>| | | | |</p>
<table>
<thead>
<tr>
<th></th>
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</tr>
</thead>
<tbody>
<tr>
<td>Detail 1</td>
<td>50.00 / 175.00</td>
<td>= .285714285714…</td>
<td>= 29%</td>
</tr>
<tr>
<td>Detail 2</td>
<td>25.00 / 175.00</td>
<td>= .142857142857…</td>
<td>= 14%</td>
</tr>
<tr>
<td>Detail 3</td>
<td>100.00 / 175.00</td>
<td>= .571428571428…</td>
<td>= 57%</td>
</tr>
</tbody>
</table>

Step 2

The detail allowed amount for each automated test (AT) procedure code will be the calculated percentage of the ATP level payment.
Example:

<table>
<thead>
<tr>
<th>Automated Test</th>
<th>Number of Automated Tests</th>
<th>Allowed Amount</th>
</tr>
</thead>
<tbody>
<tr>
<td>Detail 1 Automated Test</td>
<td>= 1</td>
<td>= $3.10</td>
</tr>
<tr>
<td>Detail 2 Automated Test</td>
<td>= 1</td>
<td>= $1.55</td>
</tr>
<tr>
<td>Detail 3 Lab Panel</td>
<td>= 4</td>
<td>= $6.19</td>
</tr>
<tr>
<td>Detail 4 Clinical Lab Test</td>
<td>= 0</td>
<td>= Fee Schedule</td>
</tr>
<tr>
<td>ATP = 6 = $10.84</td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

Note: The total number of automated tests includes the individual automated test procedure codes and the number of automated tests that are represented by each panel procedure code that is billed. Automated tests that are duplicated between panels are not counted more than once.

Calculations:

<p>| | | | |</p>
<table>
<thead>
<tr>
<th></th>
<th></th>
<th></th>
<th></th>
</tr>
</thead>
<tbody>
<tr>
<td>Detail 1</td>
<td>= 29% of 10.84</td>
<td>= (.285714285714…)X(10.84)</td>
<td>= $3.10</td>
</tr>
<tr>
<td>Detail 2</td>
<td>= 14% of 10.84</td>
<td>= (.142857142857…)X(10.84)</td>
<td>= $1.55</td>
</tr>
<tr>
<td>Detail 3</td>
<td>= 57% of 10.84</td>
<td>= (.571428571428…)X(10.84)</td>
<td>= $6.19</td>
</tr>
</tbody>
</table>

$10.84

Note: If a clinical laboratory procedure code is included in a panel, the fee schedule rate for the clinical laboratory procedure is added to the ATP rate, and the resulting sum is divided among the automated test and laboratory panel procedure codes that are billed for the date of service.

The total allowed amount for all laboratory services that are billed for the client for the same date of service will represent the ATP level pricing combined with any clinical laboratory test fee schedule pricing.

2.2.6 Breast Cancer Gene 1 and 2 (BRCA) Testing

Breast Cancer Gene 1 and 2 (BRCA) testing services are benefits of Texas Medicaid when billed with the following procedure codes:

| Procedure Codes |
|-----------------|-----------------|-----------------|-----------------|-----------------|-----------------|-----------------|-----------------|-----------------|-----------------|
| 81162 | 81163 | 81164 | 81165 | 81166 | 81167 | 81212 | 81215 | 81216 | 81217 |

Breast cancer gene 1, early onset (BRCA1) and breast cancer gene 2, susceptibility protein (BRCA2) are tumor repressor genes responsible for keeping breast cells from growing too rapidly or in an uncontrolled way. Mutations within the gene interrupt this regulatory function and increase the risk of breast and ovarian cancer, have been linked to other types of cancer such as pancreatic and prostate, and can be inherited from a person’s mother or father.

Note: Coverage of BRCA mutation analysis testing, including large rearrangement gene mutation analysis testing, are based on the National Comprehensive Cancer Network (NCCN) guidelines.

Interpretation of gene mutation analysis results is not separately reimbursable. Interpretation is part of the physician E/M service.

BRCA gene mutation analysis testing must be ordered based on familial medical history and the availability of previous familial gene mutation analysis testing results and only if the test results will affect treatment decisions or provide prognostic information.
Unaffected male or females with a family history of breast cancer, including diagnosis of ductal carcinoma in situ (DCIS), should only be considered for testing when the appropriate affected family member is unavailable. Clinical judgment should be used to determine if the client has a reasonable likelihood of a mutation, considering the client’s current age and the age of the unaffected female relatives who link the client with the affected relatives.

It is mandatory that a client who is at risk for BRCA1, BRCA2, or BRCA large rearrangement genetic mutation receive genetic counseling before and after BRCA gene mutation testing.

BRCA gene mutation analysis testing (procedure codes 81162, 81163, 81164, 81165, 81166, 81167, 81212, 81215, 81216, 81217) is limited to once per lifetime.

The ordering provider is responsible for ordering the appropriate BRCA test (e.g., BRCA1 versus BRCA2) based on medical necessity of the testing criteria and genetic counseling results. If a client’s BRCA test has a positive result, any further BRCA testing services requested with documentation of medical necessity will be considered on a case-by-case basis by Medical Director review for prior authorization.

Procedure codes 81162 and 81163 are for comprehensive BRCA gene mutation analysis testing. Procedure 81162 may only be used for clients who meet the criteria for both BRCA comprehensive and BRCA large rearrangement gene mutation analysis testing.

2.2.6.1 Genetic Counseling
Genetic counseling must be provided by a trained genetic counselor, nurse specialist in genetics, or other medical provider possessing expertise in genetic counseling that is not affiliated with the genetic testing laboratory for clients before and after BRCA gene mutation testing.

Both pre- and post-test counseling must provide the depth of content and time for the client to make an informed testing decision. The genetic counseling must be nondirective, and information about the purpose and nature of the tests must be provided to the client.

Pre-test genetic counseling must include:
- The risks and benefits of the specific genetic testing.
- The limitations of the specific genetic testing to be performed and the limitations of interpreting test results for an unaffected individual.

2.2.6.2 Documentation
Providers must maintain the following gene mutation analysis testing documentation in the client’s medical record:
- The appropriateness of the genetic testing
- The client’s specific high-risk criteria
- The benefit of the specific genetic testing to be performed
- Pre-testing genetic counseling, including all of the following:
  - Pre-test counseling date with the name and qualifications of the counseling professional
  - The risks, benefits, and limitations discussed with the client
  - The client’s ability to understand the risks, benefits, and limitations and the client’s informed choice to proceed with the specific gene mutation analysis testing as evidenced by the client’s signature on a consent form specific to the genetic mutation testing to be performed.
  - The client’s previous BRCA comprehensive gene mutation analysis testing results to support medical necessity for ordering BRCA gene mutation analysis testing
- Post-testing genetic counseling, including all the following:
• Post-test counseling date with the name and qualifications of the counseling professional
• The client’s ability to understand the results of the gene mutation analysis testing and the appropriate medical treatment resulting from the test results.
• The client’s treatment plan and any treatment plan changes based on interpretation of the test results.

The medical record is subject to retrospective review.

2.2.6.3 Prior Authorization for BRCA Testing

Prior authorization is required for initial BRCA testing (procedure codes 81162, 81163, 81164, 81165, 81166, 81167, 81212, 81215, 81216, 81217).

For clients with a known familial BRCA variant, targeted testing for BRCA1 (procedure code 81215) and BRCA2 (procedure 81217) for the specific variant must be performed before utilizing more comprehensive tests.

If the client is of Ashkenazi Jewish, Icelandic, Swedish, or Hungarian descent, testing for the three known founder variants (procedure code 81212) should be performed first.

The prior authorization request must include documentation that indicates that the client meets one or more of the criteria below:

• An individual (male or female) from a family with a known deleterious BRCA1/BRCA2 mutation
• A female with a personal history of breast cancer, including invasive or ductal carcinoma in situ (DCIS), diagnosed at age 45 years or younger
• A female with a personal history of breast cancer, including DCIS, diagnosed at any age and of an ethnicity associated with higher mutation frequency, such as: Ashkenazi Jewish, Icelandic, Swedish, or Hungarian descent
• A female with a personal history of epithelial ovarian cancer, including fallopian tube and primary peritoneal cancers, diagnosed at any age
• A male with a personal history of breast cancer, including DCIS, diagnosed at any age
• A female with a personal history of breast cancer, including DCIS, diagnosed at age 50 years or younger, and has one of the following:
  • An additional primary (2 primary sites, including bilateral or clearly separate ipsilateral) tumors occurring either synchronously or asynchronously
  • At least one close blood relative with breast cancer at any age
  • An unknown or limited family history
• A female with a personal history of breast cancer, including DCIS, diagnosed at age 60 years or younger with triple negative breast cancer
• A female with a personal history of breast cancer, including DCIS, diagnosed at any age, and has one of the following:
  • At least one close blood relative with breast cancer diagnosed at age 50 years or younger
  • At least two close blood relatives with breast cancer at any age
  • At least one close blood relative with epithelial ovarian cancer, including fallopian tube and primary peritoneal cancers
  • At least two close blood relatives with pancreatic cancer or prostate cancer (Gleason score 7 or greater) at any age
• A close male blood relative with breast cancer at any age

• A male or female with a personal history of pancreatic cancer or prostate cancer (Gleason score 7 or greater) at any age regardless of ancestry with at least two close blood relatives with one of the following:
  • Breast cancer
  • Ovarian cancer, including fallopian tube and primary peritoneal cancers
  • Prostate cancer (Gleason score 7 or greater)
  • Pancreatic cancer

• A male or female with a personal history of pancreatic cancer at any age and of an ethnicity associated with higher mutation frequency, such as Ashkenazi Jewish, Icelandic, Swedish, or Hungarian descent and one or more close blood relatives with pancreatic cancer

• A male or female with a family history of breast or ovarian cancer, including DCIS with one of the following:
  • At least one first or second degree blood relative meeting any of the criteria above; or
  • At least one third degree blood relative who has breast cancer, including DCIS, or ovarian cancer, including fallopian tube and primary peritoneal cancers, and at least two close blood relatives with one of the following:
    • Breast cancer, including DCIS, of which at least one with breast cancer was diagnosed at 50 years of age or younger
    • Ovarian cancer, including fallopian tube and primary peritoneal cancers

Note: The term “close blood relative” includes first-degree male or female relatives (e.g., parents, siblings), second-degree relatives (e.g., aunts, uncles, grandparents), and third-degree relatives (e.g., first cousins, great grandparents), from the same side of the family as the client.

Prior authorization for additional BRCA testing may be considered on a case-by-case basis by Medical Director review when testing criteria for these studies are met for clients who:

• Have previously been tested for BRCA sequencing gene mutation analysis testing and received negative results. Documentation of negative results for all previous BRCA1 sequencing gene mutation analysis testing is required.

• Every reasonable effort and documentation of the specific efforts made to obtain the previous BRCA sequencing gene mutation analysis test results from the client’s genetic testing physician or the testing laboratory.

A completed Hereditary Breast and Ovarian Cancer (HBOC) Genetic Testing Prior Authorization Request Form that has been signed and dated by the ordering provider must be submitted.

A provider’s signature on a submitted document indicates that the provider certifies, to the best of the provider’s knowledge, the information in the document is true, accurate, and complete.

For comprehensive sequencing (procedure codes 81162 or 81163), the ordering physician must indicate one of the following on the prior authorization request form:

• The client’s familial genetic history that supports medical necessity for the requested BRCA1 and BRCA2 comprehensive sequencing gene mutation analysis testing.

• Every reasonable effort was made to obtain the client’s familial genetic history and have been unable to obtain BRCA1 and BRCA2 comprehensive sequencing gene mutation analysis testing results for the affected family member(s). Documentation of the specific efforts made to obtain the client’s familial genetic history must be submitted with the request.
To facilitate a determination of medical necessity and avoid unnecessary denials, the ordering physician must provide correct and complete information, including accurate medical necessity of the services requested. Medical documentation that is submitted by the ordering physician must verify the client’s diagnosis or family history. Requisition forms from the laboratory are not sufficient for the establishment of a client’s personal and family history.

To complete the prior authorization process, the ordering provider must mail or fax the request to the TMHP Special Medical Prior Authorization Unit and include documentation of medical necessity.

A request for retroactive authorization must be submitted no later than seven calendar days beginning the day after the lab draw is performed.

### 2.2.7 Complete Blood Count (CBC)

A CBC and its components may be reimbursed by Texas Medicaid without prior authorization. The medical necessity for all laboratory services must be documented in the client’s medical record, and the services must be referenced to an appropriate diagnosis code.

Texas Medicaid considers a baseline CBC appropriate for the evaluation and management of existing and suspected disease processes. CBC tests should be individualized and based on client history, clinical indications, or proposed therapy, and will not be reimbursed for screening purposes.

When related CBC procedure codes are billed for the same date of service by the same provider, the first procedure code will be reimbursed and all other procedure codes will be denied.

Reticulocyte procedure codes may be reimbursed in addition to the CBC, hemogram, differential analysis, and platelet procedure codes indicated above.

**Referto:** The appropriate Texas Medicaid fee schedule on the TMHP website at [www.tmhp.com](http://www.tmhp.com) for CBC procedure codes that may be reimbursed.

### 2.2.8 Drug Testing and Therapeutic Drug Assays

The following procedure codes for drug testing and therapeutic drug assays may be reimbursed by Texas Medicaid:

<table>
<thead>
<tr>
<th>Procedure Codes</th>
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</thead>
<tbody>
<tr>
<td>80143 80150 80151 80155 80156 80157 80158 80159 80161 80162</td>
</tr>
<tr>
<td>80163 80164 80165 80167 80168 80169 80170 80171 80173 80175</td>
</tr>
<tr>
<td>80176 80177 80178 80179 80180 80181 80183 80184 80185 80186</td>
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<td>80360 80361 80362 80363 80364 80365 80366 80367 80368 80369</td>
</tr>
<tr>
<td>80370 80371 80372 80373 80374 80375 80376 80377 G0480 G0481</td>
</tr>
<tr>
<td>G0482 G0483 G0659</td>
</tr>
</tbody>
</table>

**Note:** The procedure codes above do not require prior authorization.

Providers are encouraged to reference the American Board of Internal Medicine (ABIM) Foundation’s “Choosing Wisely” lists to determine appropriateness of laboratory tests.
Procedure codes G0480, G0481, G0482, G0483, and G0659 are limited to once per day by any provider.

Chemistry procedure codes used for specimen validity testing (procedure codes 82540, 82550, 82552, 82553, 82554, 83986, and 84311) will be denied when submitted on the same date of service by the same provider, as procedure code G0480, G0481, G0482, G0483, or G0659.

The following CPT Drug Assay procedure codes will be denied when billed on the same date of service, by the same provider with the corresponding HCPCS Drug Assay procedure codes identified with an “X”:

<table>
<thead>
<tr>
<th>CPT Drug Assay Procedure Codes</th>
<th>G0480</th>
<th>G0481</th>
<th>G0482</th>
<th>G0483</th>
<th>G0659</th>
</tr>
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<tr>
<td>80305^</td>
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(X) The “80000” CPT procedure code will be denied if billed with the HCPCS “G” procedure code indicated with an “X.”

(Blank) There is no relationship between the “80000” CPT procedure code and the HCPCS “G” procedure code. Both procedure codes may be reimbursed if billed with the same date of service.

^QW Modifier
2.2.8.1 Documentation Requirements

All services outlined in this section are subject to retrospective review. Documentation in the client’s medical record must be maintained by the physician and support the medical necessity for the services provided.

2.2.9 Evocative and Suppression Testing

The following procedure codes for evocative and suppression testing may be reimbursed by Texas Medicaid:

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Note: The procedure codes above do not require prior authorization.

Providers are encouraged to reference the American Board of Internal Medicine (ABIM) Foundation’s “Choosing Wisely” lists to determine appropriateness of laboratory tests.

2.2.10 Genetic Testing for Colorectal Cancer

Genetic testing is provided to clients who have a first- or second-degree relative who has or has had colorectal cancer in order to determine if the client may have increased risk for developing colorectal cancer.

Note: A first-degree relative is defined as: sibling, parent, or offspring. A second-degree relative is defined as: uncle, aunt, grandparent, nephew, niece, or half-sibling.

Interpretation of gene mutation analysis results are part of the evaluation and management service and will not be reimbursed separately.

Genetic test results, when informative, may influence clinical management decisions. The documentation that is maintained in the client’s medical record must reflect that the client or family member has been given information on the nature, inheritance, and implications of genetic disorders to help them make informed medical and personal decisions prior to the genetic testing. The testing must be medically necessary and supported by documentation with a clear rationale for testing, which must be retained in the client’s medical record and made available upon request.

2.2.10.1 Documentation Requirements

Providers must maintain the following documentation in the client’s medical record for genetic testing for colorectal cancer:

- Documentation of formal pre-test counseling, including assessment of the client’s ability to understand the risks and limitations of the test.
- The client’s informed choice to proceed with the genetic testing for colorectal cancer.

The ordering provider must select the test based on the familial medical history and the availability of previous family testing results. The medical record is subject to retrospective review.
Refer to: The appropriate Texas Medicaid fee schedule on the TMHP website at www.tmhp.com for genetic testing procedure codes that may be reimbursed.

2.2.10.2 Authorization Requirements

Prior authorization is required for gene mutation analysis. A completed Special Medical Authorization (SMPA) Request Form, signed and dated by the provider rendering direct care, must be submitted to the TMHP Prior Authorization Department via mail, fax, or the electronic portal. Prior authorized services may be reimbursed once per lifetime when billed by any provider. Additional services will not be prior authorized.

Prior authorization requests may be considered for Familial Adenomatous Polyposis (FAP) testing for clients of any age with well defined hereditary cancer syndromes and for which either a positive or negative result will change medical care. The client for whom the request is made must have more than 20 polyps or a first-degree relative with FAP and a documented mutation.

Note: Clients who are seven years of age or younger must have clear rationale for testing and documentation of medical necessity from the client’s medical record must be submitted with the prior authorization request.

Prior authorization requests may be considered for Hereditary Nonpolyposis Colorectal Cancer (HNPCC) testing for clients of any age. Testing for HNPCC is used to determine whether an individual has an increased risk for colorectal cancer or other HNPCC-associated cancers. Results of the test may influence clinical management decisions. The request must include one or more of the following criteria for testing:

- The client has three or more family members (at least one must be a first-degree relative) who have colorectal cancer, and FAP has been ruled out. Two successive generations were affected, and one or more of the relatives was diagnosed with colorectal cancer at 50 years of age or younger.
- The client has had two HNPCC cancers.
- The client has colorectal cancer and a first-degree relative who also has colorectal cancer or HNPCC extracolonic cancer at 50 years of age or younger or colorectal adenoma at 40 years of age or younger.
- The client has had colorectal cancer or endometrial cancer at 50 years of age or younger.
- The client has had right-sided colorectal cancer with an undifferentiated pattern on histology at 50 years of age or younger.
- The client has had signet-cell type colorectal cancer at 50 years of age or younger.
- The client has had colorectal adenoma at 40 years of age or younger.
- The client is an asymptomatic individual with a first- or second-degree relative with a documented HNPCC mutation.

Note: Clients who are 20 years of age or younger must have clear rationale for testing and documentation of medical necessity from the client’s medical record must be submitted with the prior authorization request.

The ordering provider’s signature on a submitted document indicates that the provider certifies, to the best of the provider’s knowledge, the information in the document is true, accurate, and complete.

Requisition forms from the laboratory are not sufficient for verification of the personal and family history.

To facilitate a determination of medical necessity and avoid unnecessary denials, the physician must provide correct and complete information, including accurate medical necessity of the services requested. Medical documentation that is submitted by the physician must verify the client’s diagnosis or family history.
Guidelines for MLH1 and MLH2 mutation testing are based on guidelines established by the American College of Medical Genetics and the American Gastroenterological Association.

A request for retroactive authorization must be submitted no later than seven calendar days beginning the day after the lab draw is performed.

### 2.2.11 Hematology and Coagulation

The following hematology and coagulation procedure codes may be reimbursed by Texas Medicaid:

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**Note:** The procedure codes above do not require prior authorization.

The following procedure codes are limited to one per day by the same provider:

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^ QW Modifier

Procedure codes 85460 and 85461 may be reimbursed for female clients who are 10 through 55 years of age.

Procedure code 85004 will deny if billed on the same day by the same provider as procedure code 85007.

Procedure code 85660 is limited to once per lifetime, any provider. An additional test may be considered on appeal with documentation indicating the provider was unaware the client was tested previously or was unable to obtain client’s medical records.

Providers are encouraged to reference the American Board of Internal Medicine (ABIM) Foundation’s “Choosing Wisely” lists to determine appropriateness of laboratory tests.

### 2.2.12 Human Immunodeficiency Virus (HIV) Drug Resistance Testing

Standard treatment regimens for HIV therapy require a combination of three or more drugs. Standard therapy continues if a reduction in viral load is achieved. Incomplete virus suppression favors the development of a drug resistance and jeopardizes the success of future therapy. Testing for drug resistance as a prerequisite to further therapy is indicated under such circumstances.

To ensure accurate testing results, the client must be on appropriate antiretroviral therapy at the time of testing or have discontinued the drug regimen within the past four weeks.
Testing for antiretroviral drug resistance is indicated in certain clinical situations. These indications include any of the following:

- Individuals who have an initial (new onset) acute HIV infection, to determine if a drug-resistant viral strain was transmitted and to plan a drug regimen accordingly.
- Individuals who have virological failure during antiretroviral therapy, laboratory results showing HIV RNA levels greater than 500 and less than 1,000 copies/ml.
- HIV-infected pregnant women before initiation of therapy.
- HIV-infected pregnant women entering pregnancy with HIV RNA levels at or below 400 copies/ml while the women are on therapy.

Documentation must be maintained in the client’s medical record to support medical necessity for the HIV drug-resistance testing. Specific documentation requirements are dependent on the rationale for the testing. Documentation must include, but is not limited to, the date the drug regimen was initiated, the dosage and frequency of the prescribed medication, and laboratory tests that support all of the following:

- Acute HIV infection, with identification of the specific viral strain
- Virological failure during antiretroviral therapy with HIV RNA levels greater than 500 and less than 1,000 copies/ml
- Positive pregnancy results in an HIV positive female client
- HIV RNA levels of 400 copies/ml or less during pregnancy

HIV drug-resistance testing is not recommended when one of the following criteria is met:

- The drug regimen has been discontinued for more than four weeks
- The viral load is less than 500 copies/ml


### 2.2.13 Microbiology

The following procedure codes may be reimbursed by Texas Medicaid:

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### Note: The procedure codes above do not require prior authorization.

The following procedure codes are limited to one per day by the same provider:

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Procedure code G0433 will be denied if billed on the same day by the same provider as procedure code 86703.

Providers are encouraged to reference the American Board of Internal Medicine (ABIM) Foundation’s “Choosing Wisely” lists to determine appropriateness of laboratory tests.

#### 2.2.13.1 Zika Virus Testing

Procedure codes 86794 and 87662 may be used to bill for Zika virus testing.

Procedure code 87662 may be reimbursed up to two times on the same day by the same provider.

#### 2.2.14 Organ or Disease-Oriented Panels

The following organ or disease-oriented panel procedure codes may be reimbursed by Texas Medicaid:

<table>
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</tr>
</thead>
<tbody>
<tr>
<td>80047^</td>
<td>80048^</td>
<td>80050</td>
<td>80051^</td>
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<td>80081</td>
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</tbody>
</table>
Note: The procedure codes above do not require prior authorization.

Procedure codes 80055 and 80081 are limited to female clients who are 10 through 55 years of age. Only one service for procedure code 80055 or one service for procedure code 80081 will be reimbursed per pregnancy to the same provider.

Procedure code 80061 is limited to once per rolling year, by any provider, when performed as part of a preventative care medical checkup.

The reimbursement for the complete panel procedure code represents the total payment for all automated laboratory tests that are covered under that panel combined; including any other automated tests billed for the client for the same date of service (DOS). The Texas Medicaid allowable fee for the individual components of the complete laboratory panel will not exceed the automated test panel (ATP) fee for the total number of automated tests that are billed for the client for the same DOS.

When all of the components of the panel are performed, the complete panel procedure code must be billed. When only two or more components of the panel are performed, the individual procedure codes for each laboratory test may be billed.

Providers are encouraged to reference the American Board of Internal Medicine (ABIM) Foundation’s “Choosing Wisely” lists to determine appropriateness of laboratory tests.

### 2.2.15 Pharmacogenetics


### 2.2.16 Urinalysis and Chemistry

The following urinalysis and chemistry procedure codes may be reimbursed by Texas Medicaid for once per day:

<table>
<thead>
<tr>
<th>Procedure Codes</th>
<th>Urinalysis</th>
<th>Chemistry</th>
</tr>
</thead>
<tbody>
<tr>
<td>81000</td>
<td>81001</td>
<td>82009</td>
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<td>81020</td>
<td>82016</td>
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<td>82024</td>
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<td>81099</td>
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<td>82040^</td>
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<td>82658</td>
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</table>

*CLIA Waived test
+Add-on code
^QW Modifier
Texas Medicaid follows the Medicare categorization of tests for CLIA certificate holders.


For waived tests, providers must use modifier QW as indicated on the CMS website.

<table>
<thead>
<tr>
<th>Procedure Codes</th>
<th>83006</th>
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</thead>
<tbody>
<tr>
<td>Molecular Testing</td>
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<tr>
<td>Ophthalmology and Optometry</td>
<td></td>
</tr>
</tbody>
</table>

 Texas Medicaid follows the Medicare categorization of tests for CLIA certificate holders.


For waived tests, providers must use modifier QW as indicated on the CMS website.
Texas Medicaid limits reimbursement for the procedure codes listed in the table above to one per day without a modifier and one per day with a modifier when billed by the same provider.

Procedure code 84583 will be denied if it is billed on the same day by the same provider as procedure code 81000, 81001, 81002, 81003, 81005, or 81020.

Procedure codes 82013, 82105, 82106, 82677, 83080, 84163, and 84704 are limited to one per 210 days when billed by any provider.

Procedure code 83698 is limited to the following diagnosis codes:

<table>
<thead>
<tr>
<th>Procedure</th>
<th>Diagnosis Codes</th>
</tr>
</thead>
<tbody>
<tr>
<td>81025</td>
<td>E1165 E119 E139 E7800 E7801 E781 E782 E783</td>
</tr>
<tr>
<td>82016</td>
<td>E7841 E7849 E785 I2584</td>
</tr>
</tbody>
</table>

Procedure code 83698 is limited to two per rolling year when billed by any provider. Claims submitted for procedure code 83698 that are in excess of two per year may be considered on appeal with documentation of any of the following:

- Medical necessity for the additional test.
- The provider was unable to obtain the previous records from a different provider.
- The provider was new to treating the client and was not aware the client had received the test.

Procedure codes 82757, 84066, 84152, and 84154 are limited to male clients.

Procedure codes 82120, 84135, and 84138 are limited to female clients.

The following procedure codes are restricted to females who are 10 through 55 years of age:

<table>
<thead>
<tr>
<th>Procedure Codes</th>
</tr>
</thead>
<tbody>
<tr>
<td>81025*</td>
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<td>83664</td>
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<tr>
<td>84112</td>
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<tr>
<td>84163</td>
</tr>
</tbody>
</table>

*CLIA waived tests


2.2.17 Additional Laboratory Services

2.2.17.1 Colorectal Cancer Screening

Referto: Subsection 4.2.8, “Colorectal Cancer Screening” in the Inpatient and Outpatient Hospital Services Handbook (Vol. 2, Provider Handbooks).


2.2.17.2 Cytopathology Studies


2.2.17.3 Helicobacter pylori Testing

2.2.17.4 Laboratory Services for Clients on Dialysis
Refer to: Subsection 6.2.9, “Laboratory and Radiology Services” in the Clinics and Other Outpatient Facility Services Handbook (Vol. 2, Provider Handbooks).

2.2.17.5 Prognostic Breast and Gynecological Cancer Studies
Refer to: Subsection 6.2.9, “Laboratory and Radiology Services” in the Clinics and Other Outpatient Facility Services Handbook (Vol. 2, Provider Handbooks).

2.2.17.6 THSteps Outpatient Laboratory Services
Refer to: Subsection 5.3.11.6, “Laboratory Test” in the Children’s Services Handbook (Vol. 2, Provider Handbooks).

2.2.17.7 Authorization Requirements
Prior authorization is not required for most laboratory services. Providers may refer to the specific sections for those services that require authorizations.

2.3 Documentation Requirements
All services require documentation to support the medical necessity of the service rendered, including independent laboratory services. Independent laboratory services are subject to retrospective review and recoupment if documentation does not support the service billed.

Independent laboratory documentation must include the physician’s signed and dated order for the laboratory tests. The specific tests ordered by the physician must be listed on the order. The test results must also be included in the documentation.

2.4 Claims Filing and Reimbursement

2.4.1 Claims Information
When family planning test specimens, such as Pap smears, are collected, providers must direct the laboratory to indicate that the claim for the test is to be billed as a family planning service using a family planning diagnosis code.


A National Provider Identifier (NPI) is required for all claims. In addition, for paper claims, the Texas Provider Identifier (TPI) is required for the billing and performing provider only. NPI-only is required for all other fields.

Providers must submit independent laboratory services to TMHP in an approved electronic format or on a CMS-1500 paper claim form. Providers must purchase CMS-1500 claim forms from the vendor of their choice. TMHP does not supply the forms.

When completing a CMS-1500 claim form, all required information must be included on the claim, as information is not keyed from attachments. Superbills, or itemized statements, are not accepted as claim supplements.
Referto: “Section 3: TMHP Electronic Data Interchange (EDI)” (Vol. 1, General Information) for information on electronic claims submissions.

“Section 6: Claims Filing” (Vol. 1, General Information) for general information about claims filing.

Subsection 6.5, “CMS-1500 Paper Claim Filing Instructions” in “Section 6: Claims Filing” (Vol. 1, General Information). Blocks that are not referenced are not required for processing by TMHP and may be left blank.

2.4.1.1 Electronic Filing for Laboratory Providers

Referring provider information is always required on laboratory claims. Failure to submit this data will result in a claim rejection on the TMHP Electronic Data Interchange (EDI).

When the place of service is 6, and the billing provider identifier belongs to a laboratory, there is no need to submit the same provider identifier in the facility ID field. This notation causes the claim to suspend processing unnecessarily, and may cause a delay in the disposition of the claim. For questions about the electronic fields, contact the commercial software vendor or the TMHP EDI Help Desk at 1-888-863-3638.

2.4.2 Reimbursement

The Medicaid rates for independent laboratories are calculated in accordance with 1 TAC §355.8085 and §355.8610, and the Deficit Reduction Act (DEFRA) of 1984. By federal law, Medicaid payments for clinical laboratory services cannot exceed the Medicare payment for that service.

As the result of the Tax Equity and Fiscal Responsibility Act (TEFRA) of 1982, independent laboratories are not directly reimbursed by Texas Medicaid when providing tests to clients who are registered as hospital inpatients. Hospital reimbursements (i.e., inpatient DRG reimbursement) include payment for all pathology and laboratory services, including those sent to referral laboratories. Hospital-based and referral laboratory providers must obtain reimbursement for the technical portion from the hospital. The technical portion includes the handling of specimens and the automated or technician-generated reading and reporting of results. These services are not billable to Medicaid-covered clients.

Refereto: Subsection 2.2, “Fee-for-Service Reimbursement Methodology” in “Section 2: Texas Medicaid Fee-for-Service Reimbursement” (Vol. 1, General Information).

Texas Medicaid pays up to the amount allowed for the total component for the same procedure, same client, same date of service, any provider.

- Providers who perform the technical service and interpretation must bill for the total component.
- Providers who perform only the technical service must bill for the technical component.
- Providers who perform only the interpretation must bill for the interpretation component.

Claims filed in excess of the amount allowed for the total component for the same procedure, same dates of service, same client, any provider, are denied. Claims are paid based on the order in which they are received.

For example, if a claim is received for the total component and TMHP has already made payment for the technical or interpretation component for the same procedure, same dates of service, same client, any provider, the claim for the total component will be denied as previously paid to another provider. The same is true if a total component has already been paid and claims are received for the individual components. Texas Medicaid implemented mandated rate reductions for certain services. The Online Fee Lookup (OFL) and static fee schedules include a column titled “Adjusted Fee” to display the individual fees with all mandated percentage reductions applied. Additional information about rate changes is available on the TMHP website at www.tmhp.com/resources/rate-and-code-updates/rate-changes.
2.4.2.1 National Correct Coding Initiative (NCCI) and Medically Unlikely Edit (MUE) Guidelines

The Healthcare Common Procedure Coding System (HCPCS) and Current Procedural Terminology (CPT) codes included in the Texas Medicaid Provider Procedures Manual and the Texas Medicaid Bulletin are subject to NCCI relationships, which supersede any exceptions to NCCI code relationships that may be noted in the manuals and bulletins. Providers should refer to the CMS NCCI web page for correct coding guidelines and specific applicable code combinations.

3 Radiological and physiological laboratory services

3.1 Enrollment

To enroll in Texas Medicaid, physiological laboratory, portable X-ray supplier, independent diagnostic testing facility (IDTF), and radiological laboratory providers must be actively enrolled in Medicare.

3.1.1 Enrollment Criteria for Mammography Providers

All mammography providers, including those providing stereotactic biopsies, must be certified by the Bureau of Radiation Control (BRC).

Additionally, the Department of State Health Services (DSHS) issues mammography certification to providers who render mammography services. Providers can submit this certification to the TMHP Provider Enrollment Department in lieu of certification issued by the Food and Drug Administration (FDA), because a mammography certification issued by DSHS is recognized by the FDA. TMHP will also accept mammography certification issued by the FDA. The certificate will contain the BRC certification number, dates of issue and expiration, type of service, and Texas Medicaid and Children with Special Health Care Needs (CSHCN) Services Program provider identifiers.

Providers must check the expiration date of their mammography certification and submit an updated mammography certification prior to the expiration date. The certifications can be uploaded to PIMS, faxed to 1-512-514-4214, or mailed to:

Texas Medicaid & Healthcare Partnership
Provider Enrollment
PO Box 200795
Austin, TX 78720-0795

3.2 Services, Benefits, Limitations, and Prior Authorization

The following high-technology radiology services may be reimbursed by Texas Medicaid with prior authorization:

- Cardiac nuclear imaging
- Computed tomography (CT)
- Computed tomography angiography (CTA)
- Functional MRI (fMRI)
- Magnetic resonance imaging (MRI)
- Magnetic resonance angiography (MRA)
- Positron emission tomography (PET) scan imaging

  **Note:** Providers and facilities are required to use the lowest possible radiation dose that is consistent with acceptable image quality for cardiac nuclear imaging, CT, and PET examinations of all clients. It is recommended that providers and facilities use national standards such as those established by the American College of Radiology in their ACR Practice Guidelines and Technical Standards manual.

Radiology interpretations in any place of service will be denied if they are billed by the attending physician. Services that are billed by the attending physician are included in the facility fee and are not reimbursed separately.

  **Note:** The 3-dimensional (3-D) obstetric ultrasound is not a benefit of Texas Medicaid.

  **Referto:** PA section for exceptions to prior authorization.

### 3.2.1 Cardiac Nuclear Imaging

Cardiac nuclear imaging is a benefit of Texas Medicaid and may be reimbursed using the following procedure codes:

<table>
<thead>
<tr>
<th>Procedure Codes</th>
</tr>
</thead>
<tbody>
<tr>
<td>78414 78428 78451 78452 78453 78454 78466 78468 78472 78473</td>
</tr>
<tr>
<td>78481 78483 78494 78496</td>
</tr>
</tbody>
</table>

The cardiac nuclear imaging study may be reimbursed separately from the diagnostic radiopharmaceutical.

  **Referto:** The [online fee lookup (OFL)](www.tmhp.com) or the applicable fee schedules on the TMHP website at [www.tmhp.com](http://www.tmhp.com) to review the diagnostic radiopharmaceuticals that are reimbursed by Texas Medicaid.

#### 3.2.1.1 Authorization Requirements

Authorization is required for cardiac nuclear imaging.

  **Referto:** Subsection 3.2.6, “Authorization Requirements for CT, CTA, MRI, fMRI, MRA, PET, and Cardiac Nuclear Imaging Services” in this handbook.

### 3.2.2 Computed Tomography and Magnetic Resonance Imaging

CT, CTA, MRI, fMRI, and MRA services are benefits of Texas Medicaid.

The following procedure codes may be reimbursed with prior authorization for CT, CTA, MRI, fMRI, and MRA radiology services:

<table>
<thead>
<tr>
<th>Procedure Codes</th>
</tr>
</thead>
<tbody>
<tr>
<td>70336 70450 70460 70470 70480 70481 70482 70486 70487 70488</td>
</tr>
<tr>
<td>70490 70491 70492 70496 70498 70540 70542 70543 70544 70545</td>
</tr>
<tr>
<td>70546 70547 70548 70549 70551 70552 70553 70554 70555 71250</td>
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<td>71260 71270 71271 71275 71550 71551 71552 71555 72125 72126</td>
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<tr>
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<tr>
<td>74174 74175 74176 74177 74178 74181 74182 74183 74185 75557</td>
</tr>
</tbody>
</table>
Texas Medicaid may reimburse the total component for procedure codes 76497 and 76498 when the service is rendered in the office and outpatient hospital setting by radiation treatment center providers.

The professional component may be reimbursed when the service is rendered in the office, inpatient hospital, or outpatient hospital setting by physician providers.

The technical component will be a benefit when rendered in the office setting by physician, radiation treatment center, portable X-ray supplier, radiological laboratory, and physiological laboratory providers.

Procedure codes 76497 and 76498 will be a benefit when rendered in the outpatient hospital setting by radiation treatment center providers.

The following revenue codes must be billed with the most appropriate corresponding procedure code for CT, CTA, MRI, fMRI, and MRA radiology services rendered by outpatient hospital providers:

<table>
<thead>
<tr>
<th>Revenue Codes</th>
</tr>
</thead>
<tbody>
<tr>
<td>350</td>
</tr>
</tbody>
</table>

The addition of post 3-D reconstruction (procedure codes 76376 and 76377) CT, CTA, MRI, and MRA studies must be prior authorized. No additional payment will be made in absence of prior authorization.

### Functional MRI (fMRI)

Texas Medicaid considers fMRI medically necessary when it is being used as part of a preoperative evaluation for a planned craniotomy and is required for localization of eloquent areas of the brain, such as those responsible for speech, language, motor function, and senses, which might potentially be put at risk during the proposed surgery.

Neurofunctional testing procedure code 96020 must be reported in conjunction with brain fMRI procedure code 70555. Procedure code 96020 is informational and will not be reimbursed separately.

### Intraoperative MRI (iMRI)

Indications for intracranial neurosurgical procedures using intraoperative MRI (iMRI) include, but are not limited to, the following:

- Oncologic neurosurgical procedures
- Epilepsy
- Chiari surgery
- Deep brain stimulators

Only one iMRI procedure code may be billed per operative session. Procedure codes 70557, 70558, and 70559 must not be billed in conjunction with procedure code 61751, 77021, or 77022.

Intraoperative MRI procedure codes 70557, 70558, and 70559 that are billed with modifier 26 may be reimbursed to physician providers for interpretation.

Procedure codes 75559 and 75563 must be billed in conjunction with stress testing procedure codes 93015, 93016, 93017, and 93018.
3.2.2.3 Authorization Requirements and Flexibility

Authorization is required for CT, CTA, MRI, fMRI, and MRA procedures.

**Note:** Intraoperative MRI (iMRI) does not require prior authorization.

**Refer to:** Subsection 3.2.6, “Authorization Requirements for CT, CTA, MRI, fMRI, MRA, PET, and Cardiac Nuclear Imaging Services” in this handbook.

If the ordering physician or radiologist determines that a CT, CTA, MRI, fMRI, or MRA procedure that is different from the authorized procedure is required or that additional procedures are required, the following will apply:

- The procedure performed is less complex than the procedure authorized but of the same modality (e.g., an MRI with contrast is prior authorized and the actual procedure performed is an MRI without contrast). Full reimbursement is allowed for the billed procedure.

- The authorized procedure is performed and an additional higher-level procedure of the same modality is deemed medically necessary within the same authorization period. A separate authorization is required. The additional procedure must be prior authorized separately and submitted on a separate claim.

- The procedure billed is more complex than the procedure authorized but of the same modality. No authorization update will result in reimbursement according to the rate of the lesser authorized code. For full reimbursement of the more complex procedure, the authorization requires an update.

The following table includes the recognized relationships for authorization flexibility:

<table>
<thead>
<tr>
<th>Level 1 (High)</th>
<th>Level 2 (Moderate)</th>
<th>Level 3 (Low)</th>
</tr>
</thead>
<tbody>
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<td>72127</td>
<td>72126</td>
<td>71550</td>
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<tr>
<td>72130</td>
<td>72129</td>
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<td>72133</td>
<td>72132</td>
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<td>72156</td>
<td>72142</td>
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<td>72157</td>
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<td>72158</td>
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<td>72194</td>
<td>72193</td>
<td>72148</td>
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<td>72197</td>
<td>72196</td>
<td>72192</td>
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<tr>
<td>73202</td>
<td>73201</td>
<td>72195</td>
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<td>73220</td>
<td>73219</td>
<td>73200</td>
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<tr>
<td>73223</td>
<td>73222</td>
<td>73218</td>
</tr>
<tr>
<td>73702</td>
<td>73701</td>
<td>73221</td>
</tr>
</tbody>
</table>
3.2.3 Positron Emission Tomography (PET) Scan Imaging

PET scan imaging services are benefits of Texas Medicaid and may be reimbursed using the following procedure codes:

<table>
<thead>
<tr>
<th>Procedure Codes</th>
</tr>
</thead>
<tbody>
<tr>
<td>78608</td>
</tr>
<tr>
<td>78811</td>
</tr>
<tr>
<td>78812</td>
</tr>
<tr>
<td>78813</td>
</tr>
<tr>
<td>78814</td>
</tr>
<tr>
<td>78815</td>
</tr>
<tr>
<td>78816</td>
</tr>
</tbody>
</table>

Procedure codes 78459, 78491, and 78492 are not reimbursed by Texas Medicaid.

The PET scan procedure may be reimbursed separately from the diagnostic radiopharmaceutical.

Refer to: The online fee lookup (OFL) or the applicable fee schedules on the TMHP website at www.tmhp.com to review the diagnostic radiopharmaceuticals that are reimbursed by Texas Medicaid.

3.2.3.1 Authorization Requirements

Prior authorization is required for PET imaging services.

Refer to: Subsection 3.2.6, “Authorization Requirements for CT, CTA, MRI, fMRI, MRA, PET, and Cardiac Nuclear Imaging Services” in the Radiology and Laboratory Services Handbook (Vol. 2, Provider Handbooks).

3.2.4 Radiology/Diagnostic Imaging Policy

Radiography and fluoroscopy radiology/diagnostic imaging may be reimbursed by Texas Medicaid using the following procedure codes:

<table>
<thead>
<tr>
<th>Procedure Codes</th>
</tr>
</thead>
<tbody>
<tr>
<td>74775</td>
</tr>
<tr>
<td>75956</td>
</tr>
<tr>
<td>75957</td>
</tr>
<tr>
<td>75958</td>
</tr>
<tr>
<td>75959</td>
</tr>
</tbody>
</table>

Procedure code 74775 may be reimbursed for services rendered to clients who are 20 years of age and younger.

The procedure code in Column A must be billed with the procedure codes in Column B by the same provider with the same date of service to be reimbursed:

<table>
<thead>
<tr>
<th>Column A</th>
<th>Column B</th>
</tr>
</thead>
<tbody>
<tr>
<td>75956</td>
<td>33880</td>
</tr>
<tr>
<td>75957</td>
<td>33881</td>
</tr>
<tr>
<td>75958</td>
<td>33883 or 33884</td>
</tr>
<tr>
<td>33884</td>
<td>33883</td>
</tr>
<tr>
<td>75959</td>
<td>33886</td>
</tr>
</tbody>
</table>
Texas Medicaid may reimburse the professional interpretation component only when the physician bills procedure codes 75956, 75957, 75958, or 75959. The professional and technical service may be reimbursed to the inpatient hospital through the DRG reimbursement.

### 3.2.4.1 Authorization Requirements

Prior authorization is not required for the radiology/diagnostic imaging procedure codes in this section.

### 3.2.5 Physician-Performed Radiology Services


### 3.2.6 Authorization Requirements for CT, CTA, MRI, fMRI, MRA, PET, and Cardiac Nuclear Imaging Services

Prior authorization is not required for emergency department services, outpatient observation services, or inpatient hospital radiology services.

Prior authorization is required for outpatient nonemergent services (i.e., those that are planned or scheduled). Prior authorization must be obtained before the service is rendered.

The following table summarizes the authorization requirements for CT, CTA, MRI, fMRI, MRA, PET, and cardiac nuclear imaging services:

<table>
<thead>
<tr>
<th>Condition</th>
<th>Authorization Requirements</th>
</tr>
</thead>
<tbody>
<tr>
<td>Emergency department visit</td>
<td>Authorization is not required for emergency department radiology services that are rendered during an emergency department visit.</td>
</tr>
<tr>
<td></td>
<td>• For professional claims, the appropriate radiology procedure code must be billed with modifier U6.</td>
</tr>
<tr>
<td></td>
<td>• The facility may be reimbursed using the appropriate, corresponding emergency services revenue code.</td>
</tr>
<tr>
<td>Outpatient observation</td>
<td>Authorization is not required for radiology services rendered during outpatient observation.</td>
</tr>
<tr>
<td></td>
<td>• For professional claims, the appropriate radiology procedure code must be billed with modifier U6.</td>
</tr>
<tr>
<td></td>
<td>• The facility may be reimbursed using the appropriate, corresponding outpatient observation revenue code.</td>
</tr>
<tr>
<td>Nonemergent condition: planned or scheduled</td>
<td>Texas Medicaid defines a nonemergent condition as a symptom or condition that is neither acute nor severe and can be diagnosed and treated immediately, or that allows adequate time to schedule an office visit for a history, physical, or diagnostic studies prior to diagnosis and treatment.</td>
</tr>
<tr>
<td>radiology service</td>
<td>Prior authorization is required for outpatient nonemergent (i.e., those studies that are planned or scheduled) CT, CTA, MRI, fMRI, MRA, PET scan, and cardiac nuclear imaging services.</td>
</tr>
</tbody>
</table>
### Condition | Authorization Requirements
---|---
#### Outpatient urgent condition

*Important:* The authorization number must be on the claim when it is submitted to TMHP for reimbursement. Only one authorization is allowed per claim. For the most accurate and efficient claims processing, TMHP recommends that the procedure code that is submitted on the claim match the procedure code that is authorized. Providers are encouraged to contact TMHP and update the prior authorization if the ordering physician or radiologist changes the actual procedure that is performed. Providers have 14 calendar days after the day on which the study was completed to update the prior authorization.

Additional or alternate studies identified and ordered by the radiologist at the time of a prior-authorized study meet the definition of urgent condition and require retroactive authorization.

*Refer to:* Subsection 3.2.6.1, “Retroactive Authorization” in this handbook.

*Note:* Additional or alternate studies identified and ordered by the radiologist at the time of a prior-authorized study meet the definition of urgent condition and require retroactive authorization.

#### Outpatient emergent condition

Retroactive authorization is required for unplanned radiology procedures performed during other planned or scheduled outpatient visits or procedures.

Texas Medicaid defines an emergent condition as a medical condition (including emergency labor and delivery) manifesting itself by acute symptoms of sufficient severity (including severe pain, psychiatric disturbances, or symptoms of substance use) such that a prudent layperson with an average knowledge of health and medicine could reasonably expect the absence of immediate medical attention to result in at least one of the following:

- Placing the recipient’s health (or, with respect to a pregnant woman, the health of the woman or her unborn child) in serious jeopardy
- Serious impairment to bodily functions
- Serious dysfunction to any bodily organ or part

The physician must determine that a medical emergency, which imminently threatens life or limb exists and that the medical emergency requires advanced diagnostic imaging.

*Refer to:* Subsection 3.2.6.1, “Retroactive Authorization” in this handbook.

#### Inpatient hospital

Authorization is not required for inpatient hospital radiology services.
Prior authorization of nonemergent services is considered on an individual basis, adhering to standard clinical evidence-based guidelines. Documentation must support medical necessity for the service and must be maintained in the client’s medical record, both by the ordering physician (i.e., the physician who orders the study) and the performing facility.

Nationally-accepted guidelines and radiology protocols based on medical literature are used in the authorization processes for urgent, emergent, and nonemergent services. These include, but are not limited to:

- American College of Radiology (specifically, their Appropriateness Criteria)
- American Academy of Neurology
- American Academy of Orthopedic Surgeons
- American College of Cardiology
- American Heart Association
- National Comprehensive Cancer Care Network

Refer to: Subsection 3.2.2.3, “Authorization Requirements and Flexibility” in this handbook for information about authorization flexibility for CT, CTA, MRI, fMRI, and MRA procedures.

### 3.2.6.1 Retroactive Authorization

A request for retroactive authorization for an emergent or urgent CT, MR, PET, or cardiac nuclear imaging service must be submitted no later than 14 calendar days after the day on which the study was completed.

Retroactive authorization of urgent or emergent services is considered on an individual basis, adhering to standard clinical evidence-based guidelines. Documentation must support medical necessity for the study and must be maintained in the client’s medical record, both by the ordering physician (i.e., the physician ordering the study) and the performing facility.

Retroactive authorization for outpatient urgent services is considered when all of the following criteria are met:

- The physician who renders the imaging service determines, during the provision of prior-authorized services, that additional or alternate procedures are medically indicated.
- The urgent condition requires additional or alternate advanced diagnostic imaging.

Retroactive authorization for outpatient emergent services is considered when all of the following criteria are met:

- The physician determines that a medical emergency that imminently threatens life or limb exists.
- The medical emergency requires advanced diagnostic imaging.

Retroactive authorization is not required when a prior-authorized CT or MR procedure is changed by the ordering physician or radiologist to a lesser procedure of the same modality (e.g., MRI with contrast is authorized and the actual procedure performed is MRI without contrast).

### 3.2.6.2 Request Form and Documentation

Regardless of method of submission, the ordering physician must complete and retain the Radiology Contractor Prior Authorization Request Form with an original signature in the client’s medical record.

Providers must submit the form with the request information related to the medical necessity for the service, including all of the following:

- Diagnosis
• Treatment history
• Treatment plan
• Medications that the client is currently taking
• Previous imaging results

Providers may also be asked to provide additional documentation as necessary during the authorization process.

Section 1 of the Radiology Contractor Prior Authorization Request Form must be completed, signed, and dated by the ordering physician before requesting prior authorization, regardless of the method of request for authorization.

Section 2 of the Radiology Contractor Prior Authorization Request Form must be completed, signed, and dated by the physician who performs the service prior to requesting retroactive authorization for urgent or emergent studies.

Residents, physician assistants (PAs), and nurse practitioners (NPs) may order radiological procedures; however, the ordering or referring clinician must sign the authorization form and provide the group or supervising provider’s provider identifier.

The Radiology Decision Support Tool is provided by eviCore as a resource for providers. eviCore uses the evidence-based guidelines to authorize advanced imaging services for TMHP, and these guidelines help providers determine the most appropriate treatment option for the client related to advanced imaging services. The documents include the recognized clinical guidelines for CT, CTA, MR, MRA, PET, and cardiac nuclear imaging studies.

Refer to:
Section 3, “Inpatient Hospital (Medical/Surgical Acute Care Inpatient Facility)” in the Inpatient and Outpatient Hospital Services Handbook (Vol. 2, Provider Handbooks).
Section 9, “Physician” in the Medical and Nursing Specialists, Physicians, and Physician Assistants Handbook (Vol. 2, Provider Handbooks) for more information on MRI and contrast material.

3.2.6.3 Methods of Submission

Authorization requests for CT, CTA, MRI, fMRI, MRA, PET, and cardiac nuclear imaging studies for Texas Medicaid clients must be submitted to eviCore. eviCore is the TMHP subcontractor for high-tech radiology services. Providers can submit authorizations to eviCore:

• On the eviCore website at www.evicore.com/provider
• By telephone to 1-800-572-2116
• By fax to 1-800-572-2119
• By mail to:

Texas Medicaid & Healthcare Partnership
400 Buckwalter Place Blvd
Bluffton, SC 29910

All prior authorization requests for outpatient urgent or emergent radiology services should be made by telephone in order to ensure a timely response. Requests for retroactive authorization may be submitted online using the eviCore prior authorization portal, or by telephone, fax, or mail.
3.2.7 Additional Radiology and Physiological Laboratory Services

3.2.7.1 Ambulatory Electroencephalogram


3.2.7.2 Brachytherapy


3.2.7.3 Diagnostic Doppler Sonography


3.2.7.4 Electrocardiograms


3.2.7.5 Electromyography (EMG) and Nerve Conduction Studies (NCS)


3.2.7.6 Esophageal pH Probe Monitoring


3.2.7.7 Mammography Services

The following procedure codes will be denied if the provider does not have a BRC mammography certification on file:

<table>
<thead>
<tr>
<th>Procedure Codes</th>
</tr>
</thead>
<tbody>
<tr>
<td>77032</td>
</tr>
<tr>
<td>77053</td>
</tr>
<tr>
<td>77054</td>
</tr>
</tbody>
</table>

Refer to: Subsection 9.2.15.5, “Prognostic Breast and Gynecological Cancer Studies” in the Medical and Nursing Specialists, Physicians, and Physician Assistants Handbook (Vol. 2, Provider Handbooks) for more information about mammography services.

3.2.7.8 Nonsurgical Vision Services


3.2.7.9 Obstetric Services


3.2.7.10 Radiation Therapy Services

Refer to: Subsection 9.2.60, “Radiation Therapy” in the Medical and Nursing Specialists, Physicians, and Physician Assistants Handbook (Vol. 2, Provider Handbooks) for information about radiation therapy, including brachytherapy and stereotactic radiosurgery.
3.2.7.11 Screening and Diagnostic Studies of the Breast

Refer to: Subsection 9.2.15.4, “Mammography (Screening and Diagnostic Studies of the Breast)” in the Medical and Nursing Specialists, Physicians, and Physician Assistants Handbook (Vol. 2, Provider Handbooks).

3.2.7.12 Sleep Studies


3.3 Documentation Requirements

All services require documentation to support the medical necessity of the service rendered, including radiological and physiological laboratory services. Radiological and physiological laboratory services are subject to retrospective review and recoupment if documentation does not support the service billed.

3.4 Claims Filing and Reimbursement

3.4.1 Claims Information

Claims for radiological and physiological laboratory services and portable X-ray supplier services must include the referring or ordering provider. Radiological and physiological laboratory services and portable X-ray supplier services must be submitted to TMHP in an approved electronic format or on a CMS-1500 paper claim form. Providers may purchase CMS-1500 claim forms from the vendor of their choice. TMHP does not supply the forms.

When completing a CMS-1500 claim form, all required information must be included on the claim, as information is not keyed from attachments. Superbills, or itemized statements, are not accepted as claim supplements.

Refer to: “Section 3: TMHP Electronic Data Interchange (EDI)” (Vol. 1, General Information) for information on electronic claims submissions.

“Section 6: Claims Filing” (Vol. 1, General Information) for general information about claims filing.

Subsection 6.5, “CMS-1500 Paper Claim Filing Instructions” in “Section 6: Claims Filing” (Vol. 1, General Information). Blocks that are not referenced are not required for processing by TMHP and may be left blank.

3.4.1.1 Diagnosis Requirements

A diagnosis is not required with a provider’s request for payment except when providing the following services:

- Ambulatory Electroencephalogram (A/EEG)
- Arteriogram
- Cardiac nuclear imaging
- Chest X-ray
- Computed tomography imaging (CT)
- Echography
- Electrocardiogram (ECG)
- Functional MRI (fMRI)
- Magnetic resonance angiography (MRA)
• Magnetic resonance imaging (MRI)
• Mammographies, noninvasive diagnostic studies
• Positron emission tomography (PET) scan
• Polysomnographies
• Venographies

Claims for all services provided to clients who are eligible for “Emergency Care Only” must have a diagnosis to be considered for reimbursement. As with all procedures billed to Texas Medicaid, most baseline screening or comparison studies are not a benefit.

Refer to: Section 9, “Physician” in the Medical and Nursing Specialists, Physicians, and Physician Assistants Handbook (Vol. 2, Provider Handbooks) for more information on these services.

3.4.1.2 Modifier Requirements for Type of Service Assignment

For the radiology, physiological lab, and X-ray procedures in this chapter, providers must bill modifier 26 for the interpretation component or modifier TC for the technical component. No modifier is necessary for the total component.

Refer to: Subsection 6.2.5, “Modifier Requirements for TOS Assignment” in “Section 6: Claims Filing” (Vol. 1, General Information).

Subsection 6.3.2, “Type of Service (TOS)” in “Section 6: Claims Filing” (Vol. 1, General Information).

3.4.2 Reimbursement

Radiological and physiological laboratory and portable X-ray supplier providers are reimbursed in accordance with 1TAC §355.8085. Providers can refer to the Online Fee Lookup (OFL) or the applicable fee schedule on the TMHP website at www.tmhp.com.

Refer to: Subsection 2.2, “Fee-for-Service Reimbursement Methodology” in “Section 2: Texas Medicaid Fee-for-Service Reimbursement” (Vol. 1, General Information) for more information about reimbursement.

Texas Medicaid pays only up to the amount allowed for the total component for the same procedure submitted for reimbursement by the same provider for the same client with the same date of service. Providers who perform the technical service and the interpretation must bill the total component. Providers who perform only the technical service must bill the technical component, and those who perform only the interpretation must bill the interpretation component. The total component and the technical or interpretation component for the same procedure are not reimbursed separately when billed by any provider with the same date of service; the first claim may be reimbursed and the additional claim(s) will be denied. Claims are considered for reimbursement based on the order in which they are received.

For example, if a claim is received for the total component and TMHP has already made payment for the technical or interpretation component for the same procedure with the same date of service for the same client, regardless of provider, the claim for the total component is denied. The same is true if a total component has already been paid and claims are received for the individual components.

Radiology and physiological laboratory and portable X-ray services are not payable when the client is in an inpatient setting. The reimbursement for these services are included in the diagnosis-related group (DRG) payment.

Imaging services submitted by outpatient hospital providers may be reimbursed a flat fee.

Imaging services procedure codes can be found on the TMHP fee schedule website titled, “Hospital Outpatient Imaging Services.”
Texas Medicaid implemented mandated rate reductions for certain services. The Online Fee Lookup (OFL) and static fee schedules include a column titled “Adjusted Fee” to display the individual fees with all mandated percentage reductions applied. Additional information about rate changes is available on the TMHP website at www.tmhp.com/resources/rate-and-code-updates/rate-changes.

### 3.4.2.1 NCCI and MUE Guidelines

The HCPCS and CPT codes included in the *Texas Medicaid Provider Procedures Manual* and the *Texas Medicaid Bulletin* are subject to NCCI relationships, which supersede any exceptions to NCCI code relationships that may be noted in the manuals and bulletins. Providers should refer to the CMS NCCI web page for correct coding guidelines and specific applicable code combinations.

In instances when Texas Medicaid limitations are more restrictive than NCCI MUE guidance, Texas Medicaid limitations prevail.

## 4 Claims Resources

<table>
<thead>
<tr>
<th>Resource</th>
<th>Location</th>
</tr>
</thead>
<tbody>
<tr>
<td>Acronym Dictionary</td>
<td>&quot;Appendix C: Acronym Dictionary&quot; <em>(Vol. 1, General Information)</em></td>
</tr>
<tr>
<td>State, federal, and TMHP contact information</td>
<td>“Appendix A: State, Federal, and TMHP Contact Information” <em>(Vol. 1, General Information)</em></td>
</tr>
<tr>
<td>TMHP Electronic Data Interchange (EDI) information</td>
<td>“Section 3: TMHP Electronic Data Interchange (EDI)” <em>(Vol. 1, General Information)</em></td>
</tr>
</tbody>
</table>

## 5 Contact TMHP

The TMHP Contact Center at 1-800-925-9126 is available Monday through Friday from 7 a.m. to 7 p.m., Central Time.

## 6 Forms

The following linked forms can also be found on the Forms page of the Provider section of the TMHP website at www.tmhp.com:

<table>
<thead>
<tr>
<th>Forms</th>
</tr>
</thead>
<tbody>
<tr>
<td>Hereditary Breast and Ovarian Cancer (HBOC) Genetic Testing</td>
</tr>
</tbody>
</table>
7  **Claim Form Examples**

The following linked claim form examples can also be found on the [Claim Form Examples](https://www.tmhp.com) page of the Provider section of the TMHP website at [www.tmhp.com](http://www.tmhp.com):

<table>
<thead>
<tr>
<th>Claim Form Examples</th>
</tr>
</thead>
<tbody>
<tr>
<td>Independent Laboratory</td>
</tr>
<tr>
<td>Office Visit with Lab and Radiology</td>
</tr>
<tr>
<td>Radiological/Physiological Laboratory and Portable X-Ray Supplier</td>
</tr>
</tbody>
</table>
# TELECOMMUNICATION SERVICES HANDBOOK

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1 General Information

This handbook contains information about Texas Medicaid fee-for-service benefits. For information about managed care benefits, refer to the Medicaid Managed Care Handbook (Vol. 2, Provider Handbooks).

Managed care carve-out services are administered as fee-for-service benefits. A list of all carve-out services is available in section 8, “Carve-Out Services” in the Medicaid Managed Care Handbook (Vol. 2, Provider Handbooks).

The information in this handbook is intended for home health agencies, hospitals, nurse practitioners (NP), clinical nurse specialists (CNS), certified nurse midwives (CNM), licensed professional counselors (LPC), licensed marriage and family therapists (LMFT), licensed clinical social workers (LCSW), physicians, physician assistants (PA), psychologists, licensed psychological associates, provisionally licensed psychologists, and licensed dieticians.

Important: All providers are required to read and comply with “Section 1: Provider Enrollment and Responsibilities” (Vol. 1, General Information). In addition to required compliance with all requirements specific to Texas Medicaid, it is a violation of Texas Medicaid rules when a provider fails to provide health-care services or items to Medicaid clients in accordance with accepted medical community standards and standards that govern occupations, as explained in Title 1 Texas Administrative Code (TAC) §371.1659. Accordingly, in addition to being subject to sanctions for failure to comply with the requirements that are specific to Texas Medicaid, providers can also be subject to Texas Medicaid sanctions for failure, at all times, to deliver health-care items and services to Medicaid clients in full accordance with all applicable licensure and certification requirements including, without limitation, those related to documentation and record maintenance.

Refer to: “Section 1: Provider Enrollment and Responsibilities” (Vol. 1, General Information).

2 Enrollment

Providers may provide telecommunication services for Texas Medicaid clients under the provider’s Texas Medicaid provider identifier. No additional enrollment is required to provide telemedicine medical service or telehealth services.

Home health agency and hospital providers who wish to provide telemonitoring services must notify the Texas Medicaid & Healthcare Partnership (TMHP) as follows:

• Current providers must use the Provider Information Management System (PIMS) to indicate that they provide telemonitoring services.
• Newly enrolling or re-enrolling home health agency or outpatient hospital providers must indicate whether they provide telemonitoring services during the enrollment process.
Refer to:


Subsection 2.10.1, “Enrollment” in the Children’s Services Handbook (Vol. 2, Provider Handbooks) for information about licensed dietitian enrollment.

3 Services, Benefits, Limitations, and Prior Authorization

Telemedicine medical service and telehealth services must be provided in compliance with standards established by the respective licensing or certifying board of the professional providing the services.

The use of telemedicine medical services within intermediate care facilities for individuals with intellectual disabilities (ICD-IID) and State Supported Living Centers is subject to the policies established by the Health and Human Services Commission (HHSC).

More than one medically necessary telemedicine medical service or telehealth service may be reimbursed for the same date and same place of service if the services are billed by providers of different specialties.

All confidentiality and Health Insurance Portability and Accountability Act (HIPAA) standards apply to telemedicine medical service and telehealth transmissions.

Refer to: Subsection 1.7.6, “Release of Confidential Information” in “Section 1: Provider Enrollment and Responsibilities” (Vol. 1, General Information) for more information about confidentiality standards.

3.1 Patient Health Information Security

The software system used by the distant site provider must allow secure authentication of the distant site provider and the client.

The physical environments of the client and the distant site provider must ensure that the client’s protected health information remains confidential. A parent or legal guardian may be physically located in the patient site or distant site environment during a telehealth or telemedicine medical service with a child.
A parent or legal guardian must provide written or verbal consent to the distant site provider to allow any other individual, other than the health professional as required by Texas Government Code §531.0217(c-4)(4) for school-based telemedicine medical services, to be physically present in the distant or patient site environment during a telehealth or telemedicine medical service with a child.

An adult client must also provide written or verbal consent to the distant site provider to allow any other individual to be physically present in the distant or patient site environment during a telehealth or telemedicine medical service.

Providers of telehealth or telemedicine medical services must maintain the confidentiality of protected health information (PHI) as required by Federal Register 42, Code of Federal Regulations (CFR) Part 2, 45 CFR Parts 160 and 164, Chapters 111 and 159 of the Texas Occupations Code, and other applicable federal and state law.

Providers of telehealth or telemedicine medical services must also comply with the requirements for authorized disclosure of PHI relating to clients in state mental health facilities and residents in state supported living centers, which are included in, but not limited to, 42 CFR Part 2, 45 CFR Parts 160 and 164, Texas Health and Safety Code §611.004, and other applicable federal and state law.

All client health information generated or utilized during a telehealth or telemedicine medical service must be stored by the distant site provider in a client health record. If the distant site provider stores the patient health information in an electronic health record, the provider should use software that complies with Health Insurance Portability and Accountability Act (HIPAA) confidentiality and data encryption requirements, as well as with the United States Department of Health and Human Services (HHS) rules implementing HIPAA.

3.2 General Guidelines for Texas Medicaid MCOs

Texas Medicaid managed care organizations (MCOs) are prohibited from denying reimbursement for covered services solely because they are delivered remotely. MCOs must consider reimbursement for all medically necessary Medicaid-covered services that are provided using telemedicine or telehealth and must consider clinical effectiveness and cost-effectiveness to determine whether a telemedicine or telehealth visit is appropriate.

Texas Medicaid MCOs must determine whether to reimburse for a telemedicine or telehealth service based on the following considerations:

- Is it medically necessary?
- Is it clinically effective?
- Is the telemedicine or telehealth service cost-effective?
- Is the telemedicine or telehealth service provided in accordance with the law and in accordance with contract requirements that are applicable to the provision of the same health-care service provided in person?
- Does the use of telemedicine or telehealth promote and support patient-centered medical homes?

Texas Medicaid MCOs must consider reimbursement for all services that are currently a Medicaid benefit when they are provided using telemedicine or telehealth, including the procedure codes that are identified in the tables of subsection 3.3.4, “Telemedicine Benefits for FQHCs” and subsection 3.4.8, “Distant-Site Telehealth Benefits for FQHCs” in this handbook.

All other medically necessary Medicaid-covered services that are provided using telemedicine or telehealth must also be considered for reimbursement.

Texas Medicaid MCOs cannot deny, limit, or reduce reimbursement for a covered health-care service or procedure based on the provider’s choice of telecommunications platform to provide the service or procedure using telemedicine or telehealth.
Providers should refer to individual MCO policies for additional coverage information.

### 3.3 Telemedicine Services

Telemedicine medical services are defined as health-care services delivered by a physician licensed in Texas or a health professional who acts under the delegation and supervision of a health professional licensed in Texas and within the scope of the health professional’s license to a patient at a different physical location using telecommunications or information technology.

#### 3.3.1 Distant Site

A distant site is the location of the provider rendering the service. Distant-site telemedicine benefits include services that are performed by the following providers, who must be enrolled as a Texas Medicaid provider:

- Physician
- CNS
- NP
- PA
- CNM
- Federally Qualified Health Center (FQHC)

An out-of-state physician who is a distant site provider may provide episodic telemedicine medical services without a Texas medical license as outlined in Texas Occupations Code §151.056 and Title 22 Texas Administrative Code (TAC) §172.2(g)(4) and 172.12(f).

Distant site providers that provide mental health services must be appropriately licensed or certified in Texas, or be a qualified mental health professional-community services (QMHP-CS), as defined in 26 TAC §301.303(48).

A valid practitioner-patient relationship must exist between the distant site provider and the patient receiving telemedicine services. A valid practitioner-patient relationship exists between the distant site provider and the patient if:

- The distant site provider meets the same standard of care required for an in-person service.
- The relationship can be established through:
  - A prior in-person service.
  - A prior telemedicine medical service that meets the delivery modality requirements specified in Texas Occupations Code §111.005(a)(3).
  - The current telemedicine medical service.

The relationship can be established through a call coverage agreement established in accordance with Texas Medical Board (TMB) administrative rules in 22 TAC §177.20.

The distant site provider must obtain informed consent to treatment from the patient, patient’s parent, or the patient’s guardian prior to rendering a telemedicine medical service.
Distant site providers that communicate with clients using electronic communication methods other than phone or facsimile must provide clients with written notification of the physician’s privacy practices prior to evaluation and treatment. A good faith effort must be made to obtain the client’s written acknowledgment of the notice, including by email response.

A distant site provider should provide patients who receive a telemedicine medical service with guidance on the appropriate follow-up care.

Procedure codes that indicate remote (telemedicine medical services) delivery in the description do not need to be billed with the 95 modifier.

The following procedure codes, when billed with the 95 modifier, are a benefit for distant-site telemedicine providers:

<table>
<thead>
<tr>
<th>Procedure Codes</th>
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</thead>
<tbody>
<tr>
<td>90791</td>
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<td>90792</td>
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<td>90832</td>
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<td>90833</td>
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<td>90836</td>
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<td>G0407*</td>
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<td>G0408*</td>
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<td>G0425</td>
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<td>G0426</td>
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<tr>
<td>G0427</td>
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<tr>
<td>G0459</td>
</tr>
</tbody>
</table>

Note: Procedure codes for behavioral health services are subject to the benefits and limitations outlined in Subsection 4.2, “Services, Benefits, Limitations” in the Behavioral Health and Case Management Services Handbook (Vol. 2, Provider Handbooks). Procedure codes 90833, 90836, and 90838 are add-on codes and must be billed with a primary E/M procedure code in order to be reimbursed.

Preventive health visits under Texas Health Steps (THSteps) are not benefits if performed using telemedicine medical services. Health care or treatment using telemedicine medical services after a THSteps preventive health visit for conditions identified during a THSteps preventive health visit is a benefit. Medical services provided through telemedicine for abnormalities identified during these preventive health visits may be reimbursed separately to the distant site provider if an acute care evaluation and management procedure code is billed.

Refer to: Subsection 5.3.6, “THSteps Medical Checkups” in the Children’s Services Handbook (Vol. 2, Provider Handbooks) for information about THSteps preventive health visits.

### 3.3.2 Telemedicine Medical Services Delivery Modalities

The following modalities may be used to deliver telemedicine medical services within fee-for-service (FFS) Medicaid:

- Synchronous audiovisual interaction between the distant site provider and the client in another location
- Asynchronous store and forward technology, including asynchronous store and forward technology in conjunction with synchronous audio interaction between the distant site provider and the client in another location. The distant site provider would need to use one of the following:
  - Clinically relevant photographic or video images, including diagnostic images
  - The client’s relevant medical records, such as medical history, laboratory and pathology results, and prescriptive histories
- Other forms of audiovisual telecommunication technologies that allow the distant site provider to meet the in-person visit standard of care
A health benefit plan, including a Texas Medicaid managed care organization (MCO), is not required to provide reimbursement for telemedicine medical services that are provided through only synchronous or asynchronous audio interactions including:

- An audio-only telephone consultation
- A text-only email message
- A facsimile transmission

Texas Medicaid MCOs may optionally provide reimbursement for telemedicine medical services that are provided through only synchronous or asynchronous audio interactions. Distant site providers should contact each MCO to determine whether an MCO provides reimbursement for a specified modality.

### 3.3.2.1 Prescriptions Generated from a Telemedicine Medical Service

A distant site provider may issue a valid prescription as part of a telemedicine medical service. An electronic prescription (e-script) may be used as permitted by applicable federal and state statues and rules.

The same standards that apply for the issuance of a prescription during an in-person setting apply to prescriptions issued by a distant site provider.

The prescription must be issued for a legitimate medical purpose by the distant site provider as part of a valid practitioner-patient relationship.

The prescribing physician must be licensed in Texas. If the prescription is for a controlled substance, the prescribing physician must have a current valid U.S. Drug Enforcement Administration (DEA) registration number.

A licensed health professional acting under the delegation and supervision of a physician licensed in Texas may also issue a valid prescription. Prescribing must be in accordance with the required prescriptive authority agreement or other forms of delegation.

If the prescription is for a controlled substance, the health professional must have a current valid DEA registration number. If the prescription is for a schedule II controlled substance, the health professional must comply with DEA regulations regarding the use of electronic prescriptions. The health professional may also use the official prescription forms issued with their name, address, phone number, DEA registration number, delegating physician’s name, and delegating physician’s DEA registration number.

As applicable, all drug prescriptions must meet the requirements of the Texas Controlled Substance Act (Texas Health and Safety Code §481), the Texas Dangerous Drug Act (Texas Health and Safety Code §483), and any other federal or state statutes or rules.

Treatment of a client for chronic pain with scheduled drugs using telemedicine medical services is prohibited, as provided by 22 TAC §174.5(e). Chronic pain is defined in 22 TAC §170.2(2).

Treatment of a client for acute pain with scheduled drugs using telemedicine medical services is permitted, as provided by 22 TAC §174.5(e). Acute pain is defined by 22 TAC §170.2(2).

### 3.3.3 Patient Site

A patient site is the place where the client is physically located. A client’s home may be the patient site for telemedicine medical services.

Patient-site providers that are enrolled in Texas Medicaid may only be reimbursed for the facility fee using procedure code Q3014. Procedure code Q3014 is payable to NP, CNS, PA, physicians, and outpatient hospital providers. Charges for other services that are performed at the patient site may be submitted separately. Procedure code Q3014 is not a benefit if the patient site is the client’s home.
3.3.3.1 School-Based Setting

Telemedicine medical services provided in a school-based setting by a physician, even if the physician is not the client’s primary care physician or provider, are benefits if all of the following criteria are met:

- The physician is an authorized health-care provider enrolled in Texas Medicaid.
- The client is a child who is receiving the service in a primary or secondary school-based setting.
- The parent or legal guardian of the client provides consent before the service is provided.

Telemedicine medical services provided in a school-based setting are also a benefit if the physician delegates provision of services to a nurse practitioner, clinical nurse specialist, or physician assistant, as long as the nurse practitioner, clinical nurse specialist, or physician assistant is working within the scope of their professional license and within the scope of their delegation agreement with the physician.

3.3.4 Telem medicine Benefits for FQHCs

3.3.4.1 Distant Site

FQHCs that provide distant-site telemedicine services may be reimbursed for the following procedure codes:

<table>
<thead>
<tr>
<th>Procedure Codes</th>
</tr>
</thead>
<tbody>
<tr>
<td>G0466</td>
</tr>
</tbody>
</table>

*Note: Telemedicine services should be billed using modifier 95.*

FQHCs may be reimbursed the distant-site provider fee for telemedicine services at the Prospective Payment System (PPS) rate or Alternative Prospective Payment System (APPS) rate.

3.3.4.2 Patient Site

FQHCs may be reimbursed the facility fee (procedure code Q3014) as an add-on procedure code that should not be included in any cost reporting that is used to calculate a PPS or APPS per visit encounter rate.

To receive reimbursement for more than one facility fee for the same client on the same date of service, an FQHC must submit documentation of medical necessity that indicates that the client needed multiple distant-site provider consultations. An FQHC can use a signed letter from the client’s treating health-care provider at the FQHC to document the client’s medical need for receiving multiple distant-site provider consultations on the same date of service. The letter must state that the client suffered an illness or injury that required additional diagnosis or treatment by a distant-site provider.

If an FQHC is eligible for payment of both an encounter fee and a facility fee for the same client on the same date of service, the FQHC must submit a claim for the facility fee separate from the claim that was submitted for the encounter.

3.3.5 Documentation Requirements for Telemedicine Medical Services

Medical records must be maintained for all telemedicine medical services.

Documentation for a service provided via telemedicine must be the same as for a comparable in-person service.

If a patient has a primary care provider who is not the distant site provider and the patient or their parent or legal guardian provides consent to a release of information, a distant site provider must provide the patient’s primary care provider with the following information:

- A medical record or report with an explanation of the treatment provided by the distant site provider
- The distant site provider’s evaluation, analysis, or diagnosis of the patient
Unless the telemedicine medical services are rendered to a child in a school-based setting, distant site providers of mental health services are not required to provide the patient’s primary care provider with a treatment summary.

For telemedicine medical services provided to a child in a school-based setting, a notification provided by the telemedicine medical services physician to the child’s primary care provider must include a summary of the service, exam findings, prescribed or administered medications, and patient instructions.

If the child does not have a primary care provider, the notification must be provided to the child’s parent or legal guardian. In addition to providing treatment information, the notification must include a list of primary care providers from which the child’s parent or legal guardian may select a primary care provider.

### 3.4 Telehealth Services

Telehealth services are a benefit of Texas Medicaid. Telehealth services are defined as health-care services, other than telemedicine medical services, delivered by a health professional licensed, certified or otherwise entitled to practice in Texas and acting within the scope of the health professional’s license, certification or entitlement to a patient at a different physical location other than the health professional using telecommunications or information technology.

Telehealth services are reimbursed in accordance with 1 TAC §355.

#### 3.4.1 Distant Site

A distant site is the location of the provider rendering the service. Distant-site telehealth benefits include services that are performed by the following providers, who must be enrolled as a Texas Medicaid provider:

- Early Childhood Intervention (ECI)
- Licensed professional counselor
- LMFT
- LCSW
- Psychologist
- Licensed psychological associate
- Provisionally licensed psychologist
- Licensed dietitian
- CCP providers (occupational therapist, speech-language pathologist)
- Home health agency
- School Health and Related Services (SHARS)
- FQHC

Post-doctoral psychology fellows and pre-doctoral psychology interns under a psychologist supervision may deliver telehealth services. A distant site provider is the health professional that is licensed, certified, or otherwise entitled to practice in Texas who uses telehealth services to provide health-care services to a patient in Texas.

Distant site providers who provide mental health services must be appropriately licensed or certified in Texas or be a QMHP-CS as defined in 26 Texas Administrative Code §301.303(48).

The distant site provider must obtain informed consent to treatment from the patient, patient’s parent or the patient’s legal guardian prior to rendering a telehealth service.
Distant site providers should meet all other telehealth service requirements specified in Texas Occupations Code §111.

Procedure codes that indicate remote (telehealth service) delivery in the description do not need to be billed with the 95 modifier.

The following procedure codes, when billed with the 95 modifier, are a benefit for distant-site telehealth providers:

<table>
<thead>
<tr>
<th>Procedure Codes</th>
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<tbody>
<tr>
<td>90791 90792 90832* 90833* 90834* 90836* 90837* 90838* 90951 90952</td>
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<tr>
<td>90954 90955 90957 90958 90960 90961 97802 97803 97804 99202</td>
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<td>99203 99204 99205 99211 99212 99213 99214 99215 99241 99242</td>
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<td>99243 99244 99245 99251 99252 99253 99254 99255 S9470</td>
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</tbody>
</table>

*Services may be performed for or include family members.

**Note:** Procedure codes for behavioral health services are subject to the benefits and limitations outlined in subsection 4.2, “Services, Benefits, Limitations” in the Behavioral Health and Case Management Services Handbook (Vol. 2, Provider Handbooks).

**Note:** Procedure codes 90833, 90836, and 90838 are add-on codes and must be billed with a primary E/M procedure code in order to be reimbursed.

Preventive health visits under Texas Health Steps (THSteps) are not benefits if performed using telehealth medical services. Health care or treatment using telehealth medical services after a THSteps preventive health visit for conditions identified during a THSteps preventive health visit is a benefit. Telehealth services for abnormalities identified during these preventive health visits may be reimbursed separately to the distant site provider if an acute care evaluation and management procedure code is billed.

**Refer to:** Subsection 5.3.6, “THSteps Medical Checkups” in the Children’s Services Handbook (Vol. 2, Provider Handbooks) for information about THSteps preventive health visits.

### 3.4.2 School-Based Telehealth Services

Occupational therapist and speech-language pathologist providers may be reimbursed for telehealth services delivered to children in school-based settings with the following criteria:

- Reimbursement for occupational therapist and speech-language pathologist providers is only available when the patient site is a school-based setting.
- Children receiving telehealth services rendered by occupational therapist and speech-language pathologist providers must be eligible for these services through Texas Health Steps-Comprehensive Care Program (CCP).
- All medical necessity criteria and prior authorization requirements for in-person OT and ST services apply when services are delivered to children in school-based settings.
- Services provided to a client on public school or open-enrollment charter school premises are only permitted when delivered before or after school hours.

All other prior authorization, reimbursement, and billing guidelines that are applicable to in-person services will also apply when OT and ST services are delivered as telehealth services.

Licensed clinical social workers (LCSW), licensed professional counselors (LPC), licensed marriage and family therapists (LMFT), and psychologist providers may be reimbursed for telehealth services in school-based settings.
Children receiving telehealth services rendered by LCSW, LPC, LMFT, and psychologist providers must be eligible for these services through Texas Health Steps-CCP or through SHARS.

### 3.4.3 SHARS Telehealth Services

Schools that participate in the SHARS program may be reimbursed for telehealth OT and ST services delivered to children in school-based settings with the following criteria:

- Children who are eligible for OT and ST services through SHARS may receive additional therapy through Texas Health Steps-CCP if medical necessity criteria is met.
- OT and ST services provided by school districts through SHARS can be delivered during school hours.

### 3.4.4 Prior Authorization Requirements

The following initial evaluation and re-evaluation procedure codes do not require prior authorization for telehealth services:

<table>
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<tr>
<th>Procedure Codes</th>
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<tbody>
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<td>97167</td>
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</table>

The following evaluation and re-evaluation procedure codes require prior authorization for telehealth services:

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<th>Procedure Codes</th>
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<td>92507</td>
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<td>97168</td>
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<tr>
<td>97530</td>
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<tr>
<td>S9152</td>
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</tbody>
</table>

Prior authorization is not required for procedure codes 92507, 92508, or 97150 when services are provided through SHARS.

### 3.4.5 Early Childhood Intervention Telehealth Services

Telehealth services delivered to children who are eligible for the Early Childhood Intervention (ECI) Program are a benefit of Texas Medicaid.

The following procedure codes may be reimbursed for services rendered as telehealth services through the ECI program:

<table>
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<th>Procedure Codes</th>
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<tbody>
<tr>
<td>92507</td>
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<td>97530</td>
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<td>97535</td>
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<tr>
<td>S9152</td>
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<tr>
<td>T1027</td>
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</tbody>
</table>

Prior authorization is not required for the following procedure codes when services are provided through the ECI program:

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<tr>
<th>Procedure Codes</th>
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<td>97535</td>
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<tr>
<td>S9152</td>
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<tr>
<td>T1027</td>
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</tbody>
</table>
3.4.6 Distant Site Billing Requirements

Telehealth services should be billed using modifier 95. The following OT and ST procedure codes may be reimbursed when rendered as telehealth services to children eligible through Texas Health Steps-CCP:

<table>
<thead>
<tr>
<th>Procedure Codes</th>
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</thead>
<tbody>
<tr>
<td>92507</td>
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<tr>
<td>97168</td>
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</tbody>
</table>

The following OT and ST procedure codes may be reimbursed when rendered as telehealth services to children eligible for services through SHARS:

<table>
<thead>
<tr>
<th>Procedure Codes</th>
</tr>
</thead>
<tbody>
<tr>
<td>92507</td>
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<tr>
<td>97530</td>
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</tbody>
</table>

The following procedure codes should be billed with the modifiers listed in the table below:

<table>
<thead>
<tr>
<th>Procedure Codes</th>
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</thead>
<tbody>
<tr>
<td>92507</td>
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<tr>
<td>97150</td>
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</table>

Procedure code T1027 should be billed with the U1 modifier.

<table>
<thead>
<tr>
<th>Modifier</th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td>GO</td>
<td>OT services</td>
</tr>
<tr>
<td>GN</td>
<td>ST services</td>
</tr>
<tr>
<td>AT</td>
<td>Acute OT or ST services</td>
</tr>
</tbody>
</table>

Modifiers GO and GN are required on all claims except when billing evaluation and re-evaluation procedure codes. The AT modifier must be included on claims for acute therapy services.

3.4.7 Patient Site

A patient site is where the client is physically located while the service is rendered. A client’s home may be the patient site for telehealth services.

The facility fee (procedure code Q3014) is not a benefit for telehealth services.

3.4.8 Distant-Site Telehealth Benefits for FQHCs

FQHCs providing distant-site telehealth services may be reimbursed for the following procedure codes:

<table>
<thead>
<tr>
<th>Procedure Codes</th>
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</thead>
<tbody>
<tr>
<td>G0466</td>
</tr>
</tbody>
</table>

Note: Telehealth services should be billed using modifier 95.

FQHCs may be reimbursed the distant-site provider fee for telehealth services at the Prospective Payment System (PPS) rate or Alternative Prospective Payment System (APPS) rate.

FQHC practitioners may be employees of the FQHC or contracted with the FQHC.
3.4.9 Telehealth Service Delivery Modalities

The following modalities may be used to deliver telehealth services within FFS Medicaid:

- Synchronous audiovisual interaction between the distant site provider and the client in another location
- Asynchronous store and forward technology, including asynchronous store and forward technology in conjunction with synchronous audio interaction between the distant site provider and the client in another location. The distant site provider would need to use one of the following:
  - Clinically relevant photographic or video images, including diagnostic images
  - The client’s relevant medical records, such as medical history, laboratory and pathology results, and prescriptive histories
  - Other forms of audiovisual telecommunication technologies that allow the distant site provider to meet the in-person visit standard of care

A health benefit plan, including a Texas Medicaid MCO, is not required to provide reimbursement for telehealth services that are provided through only synchronous or asynchronous audio interactions including:

- An audio-only telephone consultation
- A text-only email message
- A facsimile transmission

Texas Medicaid MCOs may optionally provide reimbursement for telehealth services that are provided through only synchronous or asynchronous audio interactions. Distant site providers should contact each MCO to determine whether an MCO provides reimbursement for a specified modality.

3.4.10 Telehealth Service Documentation Requirements

Medical records must be maintained for all telehealth services.

Documentation for a telehealth service must be the same as a comparable in-person service.

If a client has a primary care provider, and the client or their parent or legal guardian provides consent to release information, a distant site provider must provide the client’s primary care provider with the following information:

- A medical record or report with an explanation of the treatment provided by the distant site provider
- The distant site provider’s evaluation, analysis, or diagnosis of the client

Providers of mental health services are not required to provide a client’s primary care provider with a treatment summary.

ECI providers are not required to provide the client’s primary care physician with a treatment summary.

3.5 Telemonitoring Services

Home telemonitoring is a health service that requires scheduled remote monitoring of data related to a client’s health, and transmission of the data from the client’s home to a licensed home health agency or a hospital. The data transmission must comply with standards set by HIPAA.

Data parameters are established as ordered by a physician’s plan of care.

Data must be reviewed by a registered nurse (RN), NP, CNS, or PA, who is responsible for reporting data to the prescribing physician in the event of a measurement outside the established parameters.
Scheduled periodic reporting of the client data to the physician is required at least once every 30 days, even when there have been no readings outside the parameters established in the physician’s orders. The RN, NP, CNS, or PA in a licensed home health agency or a hospital is responsible for reporting data to the prescribing physician. Telemonitoring providers must be available 24 hours a day, 7 days a week. Although transmissions are generally at scheduled times, they can occur any time of the day or any day of the week, according to the client’s plan of care.

Collection and interpretation of a client’s data for home telemonitoring services (procedure code 99091) is a benefit in the office or outpatient hospital setting when services are provided by a physician or other qualified health care professional. Procedure code 99091 is limited to once in a 30-day period.

Home telemonitoring is a benefit for clients who have been diagnosed with either diabetes or hypertension or both.

Home telemonitoring services is also a benefit for clients who are 20 years of age and younger, with one or more of the following conditions:

- End-stage solid organ disease
- Organ transplant recipient
- Requiring mechanical ventilation

The physician who orders home telemonitoring services has a responsibility to ensure the following:

- The client has a choice of home telemonitoring providers.
- The client has the right to discontinue home telemonitoring services at any time.

Although Texas Medicaid supports the use of home telemonitoring, clients are not required to use this service.

### 3.5.1 Facility Services

The provision and maintenance of home telemonitoring equipment is the responsibility of the home health agency or the hospital. The one-time initial setup and installation (procedure code S9110 with modifier U1) of the equipment in the client’s home is a benefit when services are provided by a home health agency or an outpatient hospital. Monthly home monitoring services (procedure code S9110 with the appropriate modifier) are a benefit when services are provided by a home health agency or an outpatient hospital. Hospital providers must submit revenue code 780 with procedure code S9110 and one of the appropriate modifiers listed in the table within this section.

Use one of the following modifiers with monthly home monitoring services procedure code S9110 to indicate the number of transmission days per month:

<table>
<thead>
<tr>
<th>Modifier</th>
<th>Number of Days Per Month</th>
</tr>
</thead>
<tbody>
<tr>
<td>U2</td>
<td>1 through 5 days per month</td>
</tr>
<tr>
<td>U3</td>
<td>6 through 10 days per month</td>
</tr>
<tr>
<td>U4</td>
<td>11 through 15 days per month</td>
</tr>
<tr>
<td>U7</td>
<td>16 through 20 days per month</td>
</tr>
<tr>
<td>U8</td>
<td>21 through 25 days per month</td>
</tr>
<tr>
<td>U9</td>
<td>26 through 30 days per month</td>
</tr>
</tbody>
</table>

The unit of reimbursement for procedure code S9110 and the appropriate modifier is a rolling month. Providers must bill the appropriate modifier to indicate the number of days that transmissions of data were received and reviewed for the client within a rolling month.
Providers are not required to submit modifiers U2, U3, U4, U7, U8, or U9 for telemonitoring on the prior authorization request, but are required to submit the appropriate modifier on the claim for reimbursement based on the number of days as outlined in the table.

Documentation supporting medical necessity for telemonitoring services must be maintained in the client’s medical record by the entity providing the service (home health agency or hospital) and is subject to retrospective review. All paid telemonitoring services not supported by documentation of medical necessity are subject to recoupment.

3.6 Prior Authorization

Prior authorization is not required for telemedicine or telehealth services; however, it may be required for the individual procedure codes billed.

3.6.1 Prior Authorization of Telemonitoring Services

Procedure code S9110 with or without modifier U1 requires prior authorization. Home telemonitoring services may be approved for up to 180 days per prior authorization request.

Procedure code S9110 with modifier U1 can only be prior authorized once per episode of care even if monitoring parameters are added after initial setup and installation, unless the provider submits documentation that extenuating circumstances require another installation of telemonitoring equipment.

Procedure code S9110 for the transmission of client data will be prior authorized no more than once per month for the duration of the prior authorization period.

Prior authorization requests may be submitted to the TMHP Prior Authorization Department by mail, fax, or the electronic portal. Prescribing or ordering providers, dispensing providers, clients’ responsible adults, and clients may sign prior authorization forms and supporting documentation using electronic or wet signatures.

The initial request for prior authorization must be received no more than three business days from the date the home telemonitoring services are initiated. Requests received after the three-business day period will be denied for dates of service that occurred before the date the request was received.

The request must include the physician-ordered frequency of the clinical data transmission and the client’s diagnoses and risk factors that qualify the client for home telemonitoring services.

Requests for additional home telemonitoring services received after the current prior authorization period ends will be denied for dates of service provided before the date the request was received.

A completed Home Telemonitoring Services Prior Authorization Request form must be submitted to request home telemonitoring services.

The Home Telemonitoring Services Prior Authorization Request form must be signed and dated within 30 days before the start of care. An RN, NP, CNS, or PA may sign the prior authorization request form on behalf of the client’s physician when the physician delegates this authority to the RN, NP, CNS, or PA. The RN, NP, CNS, or PA name, TPI or NPI, signature, and date must appear on the form.

If the prior authorization form is not signed and dated by the physician or an authorized delegate, the prior authorization request must be accompanied by a written order or prescription that is signed and dated by the physician, or a complete verbal order from the physician. A complete written order or verbal order must include all of the following:

- Physician ordered home telemonitoring transmission frequency.
- The client’s qualifying condition(s) and risk factors for home telemonitoring services.
- The ordered services including applicable procedure codes or descriptions.
- Dates of service matching or greater than those on the prior authorization request form.
• If prior authorization is being requested for the initial setup and installation, orders for initial

**Note:** A verbal order is considered current when the date received is on, or no more than, 30 days before the start of home telemonitoring services for the requested authorization period. A written order or prescription is considered current when it is signed and dated on, or no more than, 30 days before the start of home telemonitoring services.

A request received without a physician’s or an authorized delegate’s signature, documented verbal order, or written prescription will not be approved and may be denied.

Telemonitoring services will not be approved for clients of any age who have diabetes or hypertension unless they have two or more of the following risk factors:

• Two or more hospitalizations in the previous 12-month period
• Frequent or recurrent emergency department visits
• A documented history of poor adherence to medication regimens
• Documented history of falls in the previous 6-month period
• Limited or absent informal support systems
• Living alone or being home alone for extended periods of time
• A documented history of care access challenges

Prior authorization will be considered for clients who are 20 years of age and younger and have at least one of the following conditions:

• End-stage solid organ disease
• Organ transplant recipient
• Mechanical ventilation

**Refer to:** Subsection 5.5.1.2, “Document Requirements and Retention” in “Section 5: Fee-for-Service Prior Authorizations” (Vol. 1, General Information) for additional information about electronic signatures.

### 3.7 Documentation Requirements

Documentation for a telecommunication service must be the same as for a comparable in-person service.

#### 3.7.1 Documentation Requirements for Telemonitoring Providers

The home health agency or hospital must maintain documentation in the client’s medical record that shows:

• The telemonitoring equipment is:
  • Capable of monitoring any data parameters included in the plan of care.
  • A Food and Drug Administration (FDA) Class II hospital-grade medical device.
  • Capable of measuring and transmitting the client’s weight, oxygen levels in blood, glucose levels in blood, or blood pressure data.

• The monitoring equipment is being used, which must be demonstrated with data transmission information such as the:
  • Date of transmission.
  • Frequency of transmission.
  • Clinical data that was provided to the client’s primary care physician or the physician’s designee.
• The provider’s staff is qualified to install the telemonitoring equipment and to monitor the client’s data, which will be transmitted according to the client’s care plan.
• No other provider is also monitoring the client’s clinical data.
• Whether the client is able to operate the equipment or has a willing and able person to assist in completing electronic transmission of data. (Not required if the equipment does not require active participation from the client.)
• There are written protocols, policies, and procedures about the provision of the home telemonitoring services that are available to HHSC or its designee upon request and that they address the:
  • Authentication and authorization of users.
  • Authentication of the origin of client data transmitted.
  • Prevention of unauthorized access to the system or information.
  • System security, including the integrity of information that is collected, program integrity, and system integrity.
  • Maintenance of documentation about system and information usage.
  • Information storage, maintenance, and transmission.
  • Synchronization and verification of patient profile data.

4 Claims Filing and Reimbursement

4.1 Claims Information
Claims for telecommunication services must be submitted to TMHP in an approved electronic format or on the CMS-1500 paper claim form or the UB-04 CMS-1450 paper claim form. Providers may purchase CMS-1500 paper claim forms or UB-04 CMS-1450 paper claim forms from the vendor of their choice. TMHP does not supply the forms.

When completing a CMS-1500 paper claim form or a UB-04 CMS-1450 paper claim form, all required information must be included on the claim, as information is not keyed from attachments. Superbills and itemized statements are not accepted as claim supplements.

Refer to: “Section 3: TMHP Electronic Data Interchange (EDI)” (Vol. 1, General Information) for information on electronic claims submissions.

“Section 6: Claims Filing” (Vol. 1, General Information) for general information about claims filing.

Subsection 6.5, “CMS-1500 Paper Claim Filing Instructions” in “Section 6: Claims Filing” (Vol. 1, General Information) for instructions on completing paper claims. Blocks that are not referenced are not required for processing by TMHP and may be left blank.

Subsection 6.6, “UB-04 CMS-1450 Paper Claim Filing Instructions” in “Section 6: Claims Filing” (Vol. 1, General Information) for instructions on completing paper claims. Blocks that are not referenced are not required for processing by TMHP and may be left blank.

4.1.1 Telemonitoring Services
Providers submitting claims for clients who have dual eligibility for Medicaid and Medicare must first submit their claims to Medicare for procedure code 99091. Claims for procedure code S9110 with any modifier should not be submitted to Medicare. Procedure code S9110 is not payable by Medicare.
4.2 Reimbursement

For fee information, providers can refer to the Online Fee Lookup (OFL) on the TMHP website at www.tmhp.com.

Texas Medicaid implemented mandated rate reductions for certain services. The OFL and static fee schedules include a column titled “Adjusted Fee” to display the individual fees with all mandated percentage reductions applied. Additional information about rate changes is available on the TMHP website at www.tmhp.com/resources/rate-and-code-updates/rate-changes.

Referto: Subsection 2.2, “Fee-for-Service Reimbursement Methodology” in “Section 2: Texas Medicaid Fee-for-Service Reimbursement” (Vol. 1, General Information) for more information about reimbursement.

5 Claims Resources

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</tr>
</thead>
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</tr>
<tr>
<td>Automated Inquiry System (AIS)</td>
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</tr>
<tr>
<td>CMS-1500 Paper Claim Filing Instructions</td>
<td>Subsection 6.5, “CMS-1500 Paper Claim Filing Instructions” in “Section 6: Claims Filing” (Vol. 1, General Information)</td>
</tr>
<tr>
<td>State, federal, and TMHP contact information</td>
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</tr>
<tr>
<td>TMHP electronic claims submission information</td>
<td>Subsection 6.2, “TMHP Electronic Claims Submission” in “Section 6: Claims Filing” (Vol. 1, General Information)</td>
</tr>
<tr>
<td>TMHP Electronic Data Interchange (EDI)</td>
<td>“Section 3: TMHP Electronic Data Interchange (EDI)” (Vol. 1, General Information)</td>
</tr>
</tbody>
</table>

6 Contact TMHP

The TMHP Contact Center at 1-800-925-9126 is available Monday–Friday from 7 a.m. to 7 p.m., Central Time.

7 Forms

The following linked form can also be found on the Forms page of the Provider section of the TMHP website at www.tmhp.com:

Form

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1 General Information

The information in this handbook is intended for optometrists (doctors of optometry), ophthalmologists, and opticians who render services related to the eye and vision and for hearing aid professionals (fitters and dispensers, physicians, and audiologists) who provide hearing evaluations or fitting and dispensing services. The handbook provides information about Texas Medicaid’s benefits, policies, and procedures applicable to these providers.

**Important:** All providers are required to read and comply with subsection 4.1, “Enrollment” in this handbook. In addition to required compliance with all requirements specific to Texas Medicaid, it is a violation of Texas Medicaid rules when a provider fails to provide healthcare services or items to Medicaid clients in accordance with accepted medical community standards and standards that govern occupations, as explained in Title 1 Texas Administrative Code (TAC) §371.1659. Accordingly, in addition to being subject to sanctions for failure to comply with the requirements that are specific to Texas Medicaid, providers can also be subject to Texas Medicaid sanctions for failure, at all times, to deliver healthcare items and services to Medicaid clients in full accordance with all applicable licensure and certification requirements including, without limitation, those related to documentation and record maintenance.

**Refer to:** “Section 1: Provider Enrollment and Responsibilities” (Vol. 1, General Information).

This handbook contains information about Texas Medicaid fee-for-service benefits. For information about managed care benefits, refer to the Medicaid Managed Care Handbook (Vol. 2, Provider Handbooks).

Managed care carve-out services are administered as fee-for-service benefits. A list of all carve-out services is available in section 8, “Carve-Out Services” in the Medicaid Managed Care Handbook (Vol. 2, Provider Handbooks).

1.1 Payment Window Reimbursement Guidelines for Services Preceding an Inpatient Admission

According to the three-day and one-day payment window reimbursement guidelines, most professional and outpatient diagnostic and nondiagnostic services that are rendered within the designated timeframe of an inpatient hospital stay and are related to the inpatient hospital admission will not be reimbursed separately from the inpatient hospital stay if the services are rendered by the hospital or an entity that is wholly owned or operated by the hospital.

These reimbursement guidelines do not apply in the following circumstances:

- The professional services are rendered in the inpatient hospital setting.
- The hospital and the physician office or other entity are both owned by a third party, such as a health system.
- The hospital is not the sole or 100-percent owner of the entity.

**Refer to:** Subsection 3.7.4.14, “Payment Window Reimbursement Guidelines” in the Inpatient and Outpatient Hospital Services Handbook (Vol. 2, Provider Handbooks) for additional information about the payment window reimbursement guidelines.
2 Nonimplantable Hearing Aid Devices and Related Services

2.1 Enrollment
To enroll in Texas Medicaid, hearing aid professionals (physicians, audiologists, and hearing aid fitters and dispensers) who provide hearing evaluations or fitting and dispensing services must be licensed by the licensing board of their profession to practice in the state where the service is performed. Hearing aid providers are eligible to enroll as individuals and facilities.

Providers cannot be enrolled if their license is due to expire within 30 days. A current license must be submitted.

2.1.1 School Districts, State Agencies, and Inpatient Facilities
To be reimbursed for audiology and audiometry evaluation and diagnostic services for suspected and confirmed hearing loss (other than audiology evaluation and therapy services reimbursed to School Health and Related Services [SHARS] providers), audiologists employed by or contracted with school districts, state agencies, and inpatient hospitals must enroll as individual practitioners or group practitioners by choosing “Audiologist” on the enrollment application.

To be reimbursed for hearing aid devices and accessories, and fitting and dispensing visits and revisits, audiologists and hearing aid fitters and dispensers employed by or contracted with school districts, state agencies, and inpatient hospitals must enroll as individual practitioners or facilities by choosing “Hearing Aid” on the enrollment application.

Note: A SHARS Texas Provider Identifier (TPI) cannot be used to bill for these services.

2.2 Services, Benefits, Limitations, and Prior Authorization
The Texas Medicaid hearing services benefit includes those services that are medically necessary for clients of any age who have suspected or identified hearing loss that can be improved or ameliorated using a hearing aid device. Such services may be reimbursed to audiologists or hearing aid fitters and dispensers.

Note: Hearing-related services that are medically necessary because of a medical condition that cannot be improved or ameliorated using a nonimplantable hearing aid device are not considered part of the Texas Medicaid hearing services benefit. Providers may refer to the other Texas Medicaid Provider Procedures Manual Handbooks for benefit and limitation information about other hearing-related services.

Texas Medicaid clients of any age are eligible to receive medically necessary hearing aid devices and services through the hearing services benefit outlined in the following sections. The Texas Medicaid hearing services benefit includes a broad range of hearing services for clients of all ages and reimburses providers who are appropriately enrolled with Texas Medicaid in accordance with their licensure and scope of practice. Prior authorization is not necessary for benefits within program limitations unless specifically addressed in the sections below.

The following hearing services are benefits of Texas Medicaid to appropriately-enrolled audiologists, hearing aid fitters and dispensers, and physicians according to their licensure, scope of practice, and enrollment as indicated:

- Audiologists and physicians may be reimbursed for audiology and audiometry evaluation and diagnostic services for suspected and confirmed hearing loss.
- Hearing aid fitters and dispensers may be reimbursed for hearing aid devices and accessories and fitting and dispensing visits and revisits.
- Physicians may be reimbursed for physician otology and otorhinolaryngology (ENT) services.
Texas Medicaid clients whose jobs are contingent on their possessing a hearing aid or who appear to have vocational potential and who need a hearing aid may be referred to the Texas Workforce Commission (TWC) for hearing aids.

### 2.2.1 Limitations and Required Forms

All services provided to Texas Medicaid clients must be medically necessary. Unless otherwise specified, services may be reimbursed without prior authorization within the set limitations. In addition to services that always require prior authorization, providers may request prior authorization for medically necessary services that exceed benefit limitations.

Required forms, which are indicated in the specific sections below, are not required to be submitted with the claim, but the forms must be completed and maintained in the client’s medical record and made available upon request by the Texas Health and Human Services Commission (HHSC) or the Texas Medicaid & Healthcare Partnership (TMHP) for retrospective review.

### 2.2.2 Hearing Screenings

Hearing screening provided due to client concern, or at the provider’s discretion, is a benefit for clients of any age when the client is referred by a Medicaid-enrolled physician, and the screening is provided by a Medicaid-enrolled provider licensed to perform these services.

**Note:** A nurse practitioner, clinical nurse specialist, or a physician assistant under physician supervision and delegation may also refer the client for hearing screening.

Routine newborn hearing screenings and Texas Health Steps (THSteps) medical checkup hearing screenings are benefits for Texas Medicaid clients, and are included in the reimbursement for the routine service or visit.

#### 2.2.2.1 Routine Hearing Screenings

Routine hearing screenings that are required as part of the newborn hospital stay and as part of a THSteps medical checkup are included in the Texas Medicaid hearing services benefit. These routine screenings are not reimbursed to audiologists, hearing aid fitters and dispensers, or physicians.

**Newborn Hearing Screen**

The newborn hearing screening is included in the reimbursement to the hospital for the newborn hospital stay and is not reimbursed separately. A newborn hearing screening must be offered to each newborn by the facility where the birth occurs, through a program mandated by the Texas State Legislature and certified by the Texas Department of State Health Services (DSHS). The screening is covered as part of the newborn delivery. An infant born outside a birthing facility and not admitted to a birthing facility shall be referred to a facility that provides newborn hearing screening. If a facility is not required by legislative mandate to perform newborn hearing screening, a referral must be made to a facility that offers the screening.

**Refer to:** Subsection 5.3.9, “Newborn Examination” in the *Children’s Services Handbook (Vol. 2, Provider Handbooks)* for more information about the newborn hearing screening.

**THSteps Medical Checkup Hearing Screen**

Hearing screening is a required component of the THSteps medical checkup, and a standardized audiometric hearing screening is required at specific ages according to the periodicity schedule.

**Refer to:** The THSteps Medical Checkups Periodicity Schedule including the footnotes, which is available on the DSHS website at [www.dshs.state.tx.us/thsteps/providers.shtm](http://www.dshs.state.tx.us/thsteps/providers.shtm), for coverage criteria when performed as part of a THSteps medical checkup.

Subsection 5.3.11.2.3, “Hearing Screening” in the *Children’s Services Handbook (Vol. 2, Provider Handbooks)* for more information on THSteps checkup hearing screening.
2.2.2.2 Additional Hearing Screenings

A hearing screening requested outside of a routine newborn or THSteps medical checkup may be reimbursed as medically necessary without prior authorization using procedure code 92551.

Further diagnostic testing may also be reimbursed using the appropriate procedure code as indicated in subsection 2.2.3, “Audiology and Audiometry Evaluation and Diagnostic Services” in this handbook.

2.2.2.3 Abnormal Hearing Screening Results

If the screening returns abnormal results, the client must be referred to a Texas Medicaid-enrolled provider who is a licensed audiologist or physician who provides audiology services. Clients who are 20 years of age or younger and have abnormal screening results must be referred to a Texas Medicaid-enrolled provider who is an audiologist or physician who is experienced with the pediatric population and who offers auditory services.

The referring physician who performs the screening must complete the Physician’s Examination Report, which is maintained in the client’s medical record. A new Physician’s Examination Report must be completed whenever there is a change in the client’s hearing or a new hearing aid is needed. Retrospective review may be performed to ensure documentation supports the medical necessity of the service.

In addition to being referred to an appropriate provider for further testing, clients who are 35 months of age and younger and have suspected hearing loss must be referred to Early Childhood Intervention (ECI) as soon as possible but no longer than 7 days after identification, even if the client was referred to an appropriate provider for further testing.

The client’s responsible adult may refuse to permit the referral or decline ECI services at any time. The provider must document the client’s responsible adult’s decision in the client’s medical record.

Refer to: Subsection 2.8, “Early Childhood Intervention (ECI) Services” in the Children’s Services Handbook (Vol. 2, Provider Handbooks) for more information about ECI.

2.2.3 Audiology and Audiometry Evaluation and Diagnostic Services

Audiometry is a benefit of Texas Medicaid for clients of any age. Physicians must recommend hearing evaluations based on examination of the client. Only physicians or licensed audiologists will be reimbursed for hearing evaluations. Hearing aid fitters and dispensers are not reimbursed for hearing evaluations.

Important: The date of service for audiology and audiometry evaluations and diagnostic services is the date the service is rendered to the client. The date of service that is billed on the claim must match the date of service that is documented in the client’s medical record.

The following audiometry procedure codes are benefits of Texas Medicaid for a basic comprehensive audiometry survey:

<table>
<thead>
<tr>
<th>Procedure Codes</th>
</tr>
</thead>
<tbody>
<tr>
<td>92550</td>
</tr>
</tbody>
</table>

The following additional procedure codes may be benefits for audiometric testing:

<table>
<thead>
<tr>
<th>Procedure Codes</th>
</tr>
</thead>
<tbody>
<tr>
<td>92558</td>
</tr>
<tr>
<td>92576</td>
</tr>
<tr>
<td>92652</td>
</tr>
</tbody>
</table>
Automated auditory brainstem response (AABR) and otoacoustic emissions (OAE) are benefits for clients of any ages when performed to identify and diagnose hearing loss and for newborns when performed for the purpose of a newborn hearing screening.

**Note:** AABR and OAE tests performed as part of the newborn hearing screen are reimbursed as part of the hospital visit and are not reimbursed separately.

### 2.2.3.1 Otological Examinations

Otological examinations are a benefit when medically necessary and provided by a Medicaid-enrolled physician licensed to perform this service.

Procedure codes 92504 and 92505 are benefits for otological examinations.

An otological examination may also include physician evaluation and management (E/M) services provided to diagnose or treat medical conditions.

**Refer to:** Subsection 9.2.56.2, “Group Clinical Visits” in the *Medical and Nursing Specialists, Physicians, and Physician Assistants Handbook* (Vol. 2, Provider Handbooks) for information about medically necessary physician E/M services.

### 2.2.3.2 Vestibular Evaluations

Vestibular evaluations are a benefit when medically necessary and provided by a Medicaid-enrolled physician or nonphysician provider licensed to perform this service.

The following procedure codes for vestibular evaluations are benefits:

<table>
<thead>
<tr>
<th>Procedure Codes</th>
</tr>
</thead>
<tbody>
<tr>
<td>92531</td>
</tr>
<tr>
<td>92545</td>
</tr>
</tbody>
</table>

### 2.2.3.3 Evaluative and Therapeutic Services

The following procedure codes may be reimbursed for evaluative and therapeutic services:

<table>
<thead>
<tr>
<th>Procedure Codes</th>
</tr>
</thead>
<tbody>
<tr>
<td>92620</td>
</tr>
</tbody>
</table>

Audiology providers may be reimbursed for services rendered in the office setting for procedure code 92621.

### 2.2.3.4 Intraoperative Neurophysiology Monitoring (IONM)

Audiology providers may be reimbursed for performing intraoperative neurophysiology monitoring (IONM) with the following procedure codes:

<table>
<thead>
<tr>
<th>Procedure Codes</th>
</tr>
</thead>
<tbody>
<tr>
<td>95940</td>
</tr>
</tbody>
</table>

**Refer to:** Section 9.2.27.3, “Evoked Potential Testing” in the *Medical and Nursing Specialists, Physicians, and Physician Assistants Handbook* (Vol. 2, Provider Handbooks) for more information about intraoperative neurophysiology monitoring.
2.2.3.5 Forms and Documentation

Providers of hearing evaluations must have a report in the client’s record. Providers must include in the report hearing evaluation test data. The Hearing Evaluation, Fitting, and Dispensing Report (Form 3503) must be completed by the physician or audiologist who conducts the diagnostic testing. The provider who signs the report must maintain it in the client’s file. The report includes audiometric assessment results of the hearing evaluation and must provide objective documentation that amplification improves communication ability. Retrospective review may be performed to ensure documentation supports the medical necessity of the service.

For physician diagnostic hearing services (procedure codes 92502, 92504, 92540, 95940, and 95941), providers must maintain documentation of medical necessity in the client’s medical record. Retrospective review may be performed to ensure that the documentation supports medical necessity for the service.

2.2.3.6 Prior Authorization

Hearing screening and testing services do not require prior authorization. Documentation of medical necessity must be maintained by the provider in the client’s medical record. Retrospective review may be performed to ensure that the documentation supports medical necessity for the service.

Prior authorization requests may be submitted to the TMHP Prior Authorization Department via mail, fax, or the electronic portal. Prescribing or ordering providers, dispensing providers, clients’ responsible adults, and clients may sign prior authorization forms and supporting documentation using electronic or wet signatures.

Providers should use the Special Medical Prior Authorization (SMPA) Request Form for all prior authorization requests.

2.2.3.7 Limitations

Newborn hearing screenings provided during the birth admission are considered part of the newborn delivery payment to the facility and are not reimbursed as separate procedures.

An otological examination is a benefit of Texas Medicaid when medically necessary and provided by a Medicaid-enrolled physician licensed to perform this service.

An otological examination may also include physician E/M services provided to diagnose or treat medical conditions.

Refer to: Subsection 9.2.56.2, “Group Clinical Visits” in the Medical and Nursing Specialists, Physicians, and Physician Assistants Handbook (Vol. 2, Provider Handbooks) for information about medically necessary physician E/M services.

Audiometry survey procedure codes and evoked potential and otoacoustic emissions screening procedure codes may be reimbursed once per day.

Procedure codes 92551, 92552, and 92553 for pure tone audiometry are limited to one of any of these procedure codes per day, same provider, same client.

Procedure codes 92553 and 92556 are not reimbursed on the same day by the same provider for the same client. If both procedure codes are billed for the same date of service, same provider, and same client, they are denied with instructions to bill with the more appropriate, comprehensive audiometry procedure code 92557.

Procedure codes 92558, 92587, 92588, 92650, 92651, 92652, and 92653 may be reimbursed once per day, same provider, same client, assuming testing is done in both ears and must be billed with modifier 52 if the testing is only performed on one ear.
Procedure codes 92620, 92621, and 92625 may be reimbursed to the same provider four times each per rolling year. Providers must submit a prior authorization request for additional reimbursement of either procedure code.

2.2.3.7.1 Tympanometry
Tympanometry (procedure code 92567) must be limited to selected individual cases where its use demonstrably adds to the provider’s ability to establish a diagnosis and provide appropriate treatment. Tympanometry is limited to three services per rolling year when billed by any provider and is based on medical necessity, which must be documented in the client’s medical record.

2.2.3.7.2 Electrical Testing
Electrical testing may be reimbursed for services rendered to clients of any age.

Electrical testing (procedure code 92547) must be billed with the same date of service by the same provider as procedure codes 92540, 92541, 92542, 92544, 92545, or 92546.

2.2.3.7.3 Vestibular Evaluation
Vestibular evaluation is a benefit of Texas Medicaid when medically necessary and provided by a provider who is licensed to provide this service.

Hearing pathway tests such as audiometry, AABR, and electrocochleography (ECoG) can also be used for the same purpose and are frequently combined with vestibular tests.

2.2.3.7.4 AABR and OAE Hearing Screening Services
An electroencephalogram (EEG) may be reimbursed for the same date of service as evoked response testing by any provider.

Procedure code 92591 may be reimbursed as often as is medically necessary.

Texas Medicaid may reimburse physicians for ear and throat examination procedure codes 92502, 92504, and 92540. Audiologists will not be reimbursed for these services.

Refer to: Subsection 9.2.56, “Physician Evaluation and Management (E/M) Services” in the Medical and Nursing Specialists, Physicians, and Physician Assistants Handbook (Vol. 2, Provider Handbooks) for more information about these services.

Procedure codes 95940 and 95941 may be reimbursed in addition to each evoked potential test. Procedure codes 95940 and 95941 are limited to a maximum of 2 hours per day, per client, any provider, without documentation of medical necessity. Procedure codes 95940 and 95941 cannot be reported by the surgeon or anesthesiologist.

2.2.3.8 SHARS Audiology Services
Audiology evaluation and therapy services procedure codes 92507, 92508, 92521, 92522, 92523, 92524, and 92620 may be reimbursed to school districts and state agencies that are enrolled with Texas Medicaid as SHARS providers.

Refer to: Section 3, “School Health and Related Services (SHARS)” in the Children’s Services Handbook (Vol. 2, Provider Handbooks) for more information about SHARS services.

Other hearing evaluation, diagnostic, and hearing aid services may be reimbursed to appropriately-enrolled audiologists, hearing aid fitters and dispensers, and physicians as outlined in this section.

2.2.3.9 Noncovered Services
Texas Medicaid does not reimburse for a hearing screening completed in a day care, Head Start location, or a school, unless it is performed in a school-based health clinic as follow-up to an acute care medical visit. Separate procedure codes must not be billed for these services.
### 2.2.4 Hearing Aid Devices and Accessories (Nonimplantable)

Texas Medicaid may reimburse hearing aid fitters and dispensers for the following devices and accessories:

<table>
<thead>
<tr>
<th>Service</th>
<th>Limitation</th>
</tr>
</thead>
</table>
| Hearing aid devices   | **Limitation:**  
  - For clients who are 20 years of age and younger, 1 hearing aid device per ear may be reimbursed every 5 rolling years from the month it is dispensed.  
  - For clients who are 21 years of age and older, if the client has at least a 35 dB hearing loss in both ears, 1 hearing aid device may be reimbursed every 5 years from the month it is dispensed. Either the left or the right may be reimbursed, but not both in the same 5 year period.  
  
  Refer to: Subsection 2.2.4.1, “Forms and Documentation” in this handbook for additional medical necessity criteria.  

Replacement hearing aid devices that are required within the same 5-year period must be prior authorized.  
Repairs or modifications may be reimbursed without prior authorization once per rolling year after the 1-year warranty period has lapsed if the requested repair or modification is a better alternative than a new purchase.  
Procedure codes: See below for monaural and binaural procedure codes.  
Procedure code V5014 may be reimbursed for repairs and modifications.  
**Date of service:** The date of service for the initial hearing aid device is the date the client successfully completes the 30-day trial period and accepts the hearing aid device.  

**Note:** During the warranty period, Texas Medicaid may reimburse providers for a replacement hearing aid and replacement hearing aid batteries. Texas Medicaid will not reimburse hearing aid repairs or modifications that are rendered during the 12-month manufacturer’s warranty period. Providers must follow the manufacturer’s repair process as outlined in their warranty contract.

<table>
<thead>
<tr>
<th>Service</th>
<th>Limitation</th>
</tr>
</thead>
</table>
| Hearing aid accessories | **Limitation:** As often as is medically necessary for clients who are 20 years of age and younger with prior authorization.  
  
  **Note:** Hearing aid accessories include, but are not limited to, chin straps, clips, boots, and headbands.  
  
  **Procedure code:** V5267  

**Date of service:** The date of service is the date the client successfully completes the 30-day trial period and accepts the hearing aid device or the date the client receives the replacement accessory item.

<table>
<thead>
<tr>
<th>Service</th>
<th>Limitation</th>
</tr>
</thead>
</table>
| Ear impression        | **Limitation:** 1 each per hearing aid device as follows:  
  - For one impression, bill a quantity of 1.  
  - For two impressions, bill a quantity of 2.  
  
  **Procedure codes:** V5275  

**Date of service:** The date of service for the ear impression is the date the ear impression is taken.
The following monaural procedure codes may be reimbursed for medically necessary hearing aid devices and replacements that are rendered to clients of any age when they are billed with the appropriate modifier LT or RT to indicate for which ear the hearing aid device was purchased and fitted:

<table>
<thead>
<tr>
<th>Service</th>
<th>Limitation</th>
</tr>
</thead>
<tbody>
<tr>
<td>Ear mold</td>
<td><strong>Limitation:</strong> As medically necessary for clients who are 20 years of age and younger.</td>
</tr>
<tr>
<td></td>
<td>For clients who are 21 years of age and older:</td>
</tr>
<tr>
<td></td>
<td>• 3 ear molds per rolling year for custom ear molds</td>
</tr>
<tr>
<td></td>
<td>• 4 ear molds per 30 days for disposable ear molds</td>
</tr>
<tr>
<td></td>
<td>Ear molds must be billed using the appropriate LT or RT modifier.</td>
</tr>
<tr>
<td></td>
<td>Replacement ear molds may be reimbursed as often as is medically necessary without prior authorization.</td>
</tr>
<tr>
<td></td>
<td>Documentation of medical necessity must be maintained in the client’s medical record.</td>
</tr>
<tr>
<td></td>
<td><strong>Procedure codes:</strong> V5264 and V5265</td>
</tr>
<tr>
<td></td>
<td><strong>Date of service:</strong> The date of service for the ear mold is the date the ear mold is dispensed to the client.</td>
</tr>
</tbody>
</table>

Batteries (Replacement only)

<table>
<thead>
<tr>
<th>Limitation</th>
<th>Replacement batteries may be reimbursed as often as is medically necessary when a hearing aid device has been previously reimbursed by Texas Medicaid.</th>
</tr>
</thead>
<tbody>
<tr>
<td>Note</td>
<td><em>If a hearing aid has not been reimbursed by Texas Medicaid in the last 5 years, the replacement batteries may be reimbursed on appeal with a statement that documents medical necessity.</em></td>
</tr>
<tr>
<td>Procedure code</td>
<td>V5266</td>
</tr>
<tr>
<td>Date of service</td>
<td>The date of service is the date the client receives the replacement batteries.</td>
</tr>
</tbody>
</table>

The following binaural procedure codes may be reimbursed for medically necessary hearing aid devices and replacements that are rendered to clients who are 20 years of age and younger only.

<table>
<thead>
<tr>
<th>Procedure Codes</th>
</tr>
</thead>
<tbody>
<tr>
<td>V5030 V5040 V5244 V5245 V5246 V5247 V5254 V5255 V5256 V5257 V5298</td>
</tr>
</tbody>
</table>

Procedure codes V5171, V5172, and V5181 may be reimbursed for monaural hearing aids that are rendered to clients who are 20 years of age and younger only.

The following binaural procedure codes may be reimbursed for medically necessary hearing aid devices and replacements that are rendered to clients who are 20 years of age and younger:

<table>
<thead>
<tr>
<th>Procedure Codes</th>
</tr>
</thead>
<tbody>
<tr>
<td>V5100 V5211 V5212 V5213 V5214 V5215 V5221 V5249 V5250 V5251 V5252 V5253 V5258 V5259 V5260 V5261</td>
</tr>
</tbody>
</table>

Binaural hearing aid procedure codes must be submitted with a quantity of 1 per procedure code. Providers can refer to the Online Fee Lookup (OFL) or the applicable fee schedule on the TMHP website at www.tmhp.com for reimbursement rates.

**Refer to:** Subsection 2.4.2, “Reimbursement” in this handbook for more information about manual pricing.
2.2.4.1 Forms and Documentation

Monaural hearing aids may be reimbursed for clients who have no medical contraindication for using a hearing aid and who have documentation of medical necessity. The following documentation of medical necessity must be maintained in the client’s medical record:

- Hearing loss in the better ear of 35 dB or greater for the pure tone average of 500, 1000, 1500, and 2000 Hz, or a spondee threshold in the better ear of 35 dB or greater when pure tone thresholds cannot be established
- Documentation of communication need and a statement that the patient is alert and oriented and able to use the device appropriately by themselves or with assistance

Clients who are 21 years of age and older must meet the medical necessity criteria outlined above and have at least a 35 dB hearing loss in both ears to qualify for the purchase of a monaural hearing aid device.

Clients who are 20 years of age and younger must meet the medical necessity criteria outlined above and have at least a 35 dB hearing loss in both ears to qualify for the purchase of binaural hearing aid devices.

Claims for non-implantable hearing aid devices must be submitted with a manufacturer invoice showing the net acquisition cost of the non-implantable hearing aid device.

An invoice printed from an email or the Internet will not be accepted and should not be submitted with the claim as documentation to show the net acquisition cost of the hearing aid device unless the invoice reflects the actual price the provider paid for the hearing aid device.

Note: The requirement to submit the net acquisition cost of the hearing aid device applies only to non-implantable monaural and binaural hearing aid devices including, but not limited to, procedure code V5298.

Refer to: Subsection 6.3.1.1, “Place of Service (POS) Coding” in “Section 6: Claims Filing” (Vol. 1, General Information) for more information about coding place of service for other locations.

2.2.4.2 Prior Authorization

Prior authorization is not required for medically necessary hearing aid devices and supplies that are provided within the limitations outlined in the table above.

Prior authorization is required for the following:

- Replacement hearing aid devices that are required within the same 5-year period. A replacement hearing aid device may be considered for prior authorization when loss or irreparable damage has occurred. A copy of the police or fire report, when appropriate, and measures to be taken to prevent reoccurrence must be submitted with the prior authorization request. Replacements will not be authorized when the equipment has been abused or neglected by the client, the client's family, or the caregiver.

- Hearing aid repair in excess of one per rolling year. The prior authorization request must include documentation supporting the need for the requested repair.

- Hearing aid accessories for clients who are birth through 20 years of age. Requests for prior authorization for children’s hearing aid accessories including, but not limited to, chin straps, clips, boots, and headbands will be considered when the requests are submitted with documentation that shows that the client is birth through 20 years of age and that the requested supply is medically necessary for the proper use or functioning of the hearing aid device.

- Hearing aid devices that are not currently a benefit of Texas Medicaid but that are medically necessary for clients who are birth through 20 years of age (using procedure code V5298).

The prior authorization request must include:

- The medical necessity for the requested hearing aid device.
• The name of the manufacturer.
• The model number, serial number, and the dates that the warranty is in effect for the requested hearing aid.
• Additional medically necessary repairs or modifications beyond 1 per year. For additional repairs or modifications, requests for prior authorization must include documentation that supports the need for the requested repair.

For services that require prior authorization, prior authorization must be obtained before the services are rendered. The prior authorization number must be included on the claim form when the claim is submitted to TMHP.

Prior authorization requests must be submitted to the TMHP Special Medical Prior Authorization (SMPA) Department with documentation that supports medical necessity for the requested device, service, or supply. Authorization may be submitted on the TMHP website at www.tmhp.com or by fax to 1-512-514-4213.

**Important:** For clients who are birth through 20 years of age, if the authorization request is denied because it does not meet benefit criteria, the TMHP SMPA Department will refer the request to the TMHP Comprehensive Care Program (CCP) Department for consideration under CCP. The provider is not required to complete additional forms or request referral to the TMHP CCP Department.

Providers may use the form of their choice to submit the required information to the TMHP SMPA Department. No specific request form is required.

**Refer to:** “Section 6: Claims Filing” (Vol. 1, General Information) for more information about the authorizations and claims filing processes.

### 2.2.4.3 Limitations

The following services and supplies must be provided to Texas Medicaid clients if a nonimplantable hearing aid device is medically necessary:

• An individual client assessment to identify the appropriate type of device
• The fitting/implantation of the device
• The re-assessment to determine whether the device allows for adequate hearing
• Expendable supplies that are necessary to keep the device functioning properly, such as batteries and accessories

Hearing devices are a benefit for clients of any age. Some types of hearing devices are age restricted.

A hearing aid dispensed through Texas Medicaid must meet the following criteria:

• Be a new and current model
• Meet the performance specifications indicated by the manufacturer
• Include, at minimum, a standard 12-month warranty that begins on the dispensing date of the hearing aid.

Providers must dispense each hearing aid reimbursed through Texas Medicaid with all necessary hearing aid accessories and supplies, including a 1-month supply of batteries. The reimbursement for monaural and binaural procedure codes includes the required hearing aid package as follows, and no separate reimbursement will be made for these items:

• Acquisition cost of the hearing aid (the actual cost or net cost of the hearing aid after any discounts or rebates have been deducted)
• Manufacturer’s postage and handling charges, including shipping insurance
• All necessary hearing aid accessories or supplies
• Instructions for care and use
• A 1-month supply of batteries

Note: TMHP does not supply the hearing aid devices, supplies, and accessories. Providers must purchase equipment directly from manufacturers and submit claims to TMHP for reimbursement using the appropriate procedure codes.

Procedure code V5298 may be reimbursed with prior authorization for hearing aid devices that are not currently a benefit of Texas Medicaid but that are medically necessary for clients who are birth through 20 years of age.

Services for residents in a skilled nursing facility (SNF), intermediate care facility (ICF), or extended care facility (ECF) must be ordered by the attending physician. The order must be on the client’s chart, must state the condition that necessitates the hearing aid services, and must be signed by the attending physician.

2.2.5 Hearing Aid Services

The following additional hearing aid related procedures are benefits for services that are rendered to clients of any age:

<table>
<thead>
<tr>
<th>Procedure Codes</th>
</tr>
</thead>
<tbody>
<tr>
<td>92590 92592 92594 V5010 V5011 V5264 V5265 V5266 V5275</td>
</tr>
</tbody>
</table>

The following additional hearing aid related procedures are benefits for services that are rendered to clients who are 20 years of age and younger only:

<table>
<thead>
<tr>
<th>Procedure Codes</th>
</tr>
</thead>
<tbody>
<tr>
<td>92591 92593 92595</td>
</tr>
</tbody>
</table>

Texas Medicaid may reimburse hearing aid fitters and dispensers for the following services:

<table>
<thead>
<tr>
<th>Service</th>
<th>Limitation</th>
</tr>
</thead>
<tbody>
<tr>
<td>Hearing test for sensitivity</td>
<td><strong>Limitation:</strong> As often as is medically necessary</td>
</tr>
<tr>
<td></td>
<td><strong>Procedure code:</strong> 92564 (SISI hearing test)</td>
</tr>
<tr>
<td>Fitting and dispensing visits</td>
<td><strong>Limitation:</strong> 1 fitting per hearing aid procedure code per 5 rolling year period, regardless of the number of times a device is returned as unacceptable during a 30-day trial period</td>
</tr>
<tr>
<td></td>
<td><strong>Procedure code:</strong> V5011</td>
</tr>
<tr>
<td></td>
<td><strong>Limitation:</strong> 1 dispensing fee each time a hearing aid is dispensed and a new 30-day trial period begins</td>
</tr>
<tr>
<td></td>
<td><strong>Procedure codes:</strong> V5090 and V5241 (for clients of any age) and V5110, V5160, V5200, and V5240 (for clients who are 20 years of age and younger)</td>
</tr>
<tr>
<td></td>
<td>The dispensing fee may be reimbursed separately from the fitting of the hearing aid.</td>
</tr>
<tr>
<td></td>
<td>The post-fitting check is included in the reimbursement for the dispensing procedure and is not reimbursed separately.</td>
</tr>
</tbody>
</table>
2.2.5.1 Forms and Documentation

The forms and documentation required for the fitting and dispensing visits are as follows:

- Physician’s Examination Report—The referring physician who performs the screening must complete the Physician’s Examination Report, which is maintained in the client’s medical record.

  Note: An advanced practice registered nurse (APRN) or a physician assistant (PA) under physician supervision and delegation may also complete the Physician’s Examination Report.

- Hearing Evaluation, Fitting, and Dispensing Report (Form 3503)—The Hearing Evaluation, Fitting, and Dispensing Report (Form 3503) must be completed by the fitter/dispenser that conducts the fitting and dispensing visit. The provider who signs the report must maintain it in the client’s file. The report includes audiometric assessment results of the hearing evaluation and must provide objective documentation to support improved communication ability with amplification. Retrospective review may be performed to ensure documentation supports the medical necessity of the device, service, or supply.

- Client Acknowledgement Statement (created by the provider)—At the time the hearing aid device and supplies are dispensed, the client must sign a client acknowledgement statement to verify the client was evaluated and offered an appropriate hearing aid that meets the client’s hearing need. The acknowledgement statement must include language that indicates the client is responsible for paying any hearing aid rental fees if charged. The provider must obtain the signed acknowledgment statement before dispensing the hearing aid device and supplies and must keep the signed acknowledgment statement in the client’s file. Retrospective review may be performed to ensure documentation supports the medical necessity of the device, service, or supply.

- Prior to dispensing a hearing instrument, a provider must enter into a written contract with the client that meets the Texas Department of Licensing and Regulation (TDLR) rule requirements in 16 TAC §112.140. The signed contract should verify that:
  - The client has a 30-day trial period for the hearing aid.
  - If the client is not satisfied with the purchased hearing aid, the client may return it to the provider within the 30-day trial period. If the device is returned within the 30-day trial period, the provider may charge the client a rental fee not to exceed $2.00 per day. This fee is not a benefit of Texas Medicaid. The 30-day trial period and any charged rental fee must meet TDLR rule requirements in 16 TAC §112.140.
  - The contract must be executed prior to dispensing and must be maintained in the client’s file.
  - The client must receive a copy of the executed contract.
  - All charges and fees associated with the trial period must be stated in the contract, which shall also include the name, address, and telephone number of TDLR.

After at least 30 days and the successful completion of the trial period, the provider must update the statement to indicate that the trial was successful and the client accepted the dispensed hearing aid device. The updated statement must be maintained in the client’s file. Retrospective review may be performed to ensure documentation supports the medical necessity of the device, service, or supply.
For hearing aids that are dispensed in a provider’s office, if a client fails to return by the end date of the trial period, the provider must contact the client. After 3 attempts have been made, if the client does not return to the provider’s office, the provider must document all attempts to contact the client and must maintain this documentation in the client’s file. Retrospective review may be performed to ensure documentation supports the contact attempts and the client’s failure to return to the provider’s office. This requirement does not apply for services that are rendered to clients who receive hearing aids in other places of service (i.e., nursing homes).

2.2.5.2 Prior Authorization

Prior authorization is not required for fitting and dispensing visits and revisits.

2.2.5.3 Limitations

The following hearing aid visits may be reimbursed by Texas Medicaid:

- The fitting and dispensing visits that encompass a 30-day trial period and include a post-fitting check 5 weeks after the trial period has been successfully completed
- A first revisit as needed after the post-fitting check
- A second revisit as needed after the first revisit

The fitting visit includes the fitting, dispensing, and post-fitting check of the hearing aid. A trial period of up to 30 days is authorized by Texas Occupations Code §402.401. The 30-day trial period, and any charged rental fee, must meet the Texas Department of Licensing and Regulation (TDLR) rule requirements in 16 TAC §112.140.

Providers must allow each Texas Medicaid client a 30-consecutive-day trial period that begins with the dispensing date. This trial period gives the client time to determine whether the hearing aid device meets the client’s needs. If the client is not satisfied with the purchased hearing aid, the client may return it to the provider, who must accept it. If the device is returned within 30 days of the date it was dispensed, the provider may charge the client a rental fee not to exceed $2 per day. This fee is not a benefit of Texas Medicaid and will not be reimbursed. The client is responsible for paying the hearing aid rental fees if the provider chooses to charge a fee for the rental of returned hearing aid devices.

During the trial period, providers may dispense additional hearing aids as medically necessary until either the client is satisfied with the results of the hearing aid or the provider determines that the client cannot benefit from the dispensing of another hearing aid. The dispensing date of each additional hearing aid starts a new trial period.

The licensed audiologist or fitter/dispenser must perform a post-fitting check of the hearing aid within 5 weeks of the initial fitting.

The first and second revisits are available if additional visits are required after the post-fitting check.

- **First revisit.** The first revisit must include a hearing aid check.
- **Second revisit.** The second revisit is available as needed after the post-fitting check and first revisit. The second revisit must include either a real ear measurement or aided sound field testing according to the guidelines specified for the hearing evaluation. If the aided sound field test scores suggest a decrease in hearing acuity, the provider must include puretone and speech audiometry readings from the first evaluation.

Services for residents in an SNF, ICF, or ECF must be ordered by the attending physician. The order must be on the client’s chart, must state the condition that necessitates the hearing aid services, and must be signed by the attending physician.

Home visit hearing evaluations and fittings are permitted only with documentation of client need in the physician’s, advanced practice registered nurse (APRN) or a physician assistant (PA) written order.

**Note:** APRN or a PA under physician supervision and delegation may also perform the evaluation.
2.3 Documentation Requirements

All services, including hearing services, require documentation to support the medical necessity of the service rendered. Hearing services are subject to retrospective review and recoupment if documentation does not support the service billed.

Required forms for nonimplantable hearing devices and services, which are indicated in the specific sections above, are not submitted with the claim to TMHP, but the forms must be completed and maintained in the client’s medical record and made available upon request by HHSC or TMHP for retrospective review.

2.4 Claims Filing and Reimbursement

2.4.1 Claims Filing

Hearing services must be submitted to TMHP in an approved electronic format or on the CMS-1500 paper claim form. Providers may purchase CMS-1500 paper claim forms from the vendor of their choice. TMHP does not supply the forms.

When completing a CMS-1500 paper claim form, all required information must be included on the claim, as information is not keyed from attachments. Superbills, or itemized statements, are not accepted as claim supplements.

Refer to: Subsection 2.2, “Fee-for-Service Reimbursement Methodology” in “Section 2: Texas Medicaid Fee-for-Service Reimbursement” (Vol. 1, General Information) for more information about reimbursement.

Subsection 1.7.11, “Billing Clients” in “Section 1: Provider Enrollment and Responsibilities” (Vol. 1, General Information).

“Section 3: TMHP Electronic Data Interchange (EDI)” (Vol. 1, General Information) for information on electronic claims submissions.

Subsection 6.1, “Claims Information” in “Section 6: Claims Filing” (Vol. 1, General Information) for general information about claims filing.

Subsection 6.5, “CMS-1500 Paper Claim Filing Instructions” in “Section 6: Claims Filing” (Vol. 1, General Information). Blocks that are not referenced are not required for processing by TMHP and may be left blank.

Providers must file all claims electronically or on the appropriate Centers for Medicare & Medicaid Services (CMS) paper claim form after providing the services.

Exception: Claims for non-implantable hearing aid devices must be submitted on the CMS-1500 paper claim form because electronic claim submissions do not allow for the submission of attachments.

Claims must include the following information:

- The most appropriate International Classification of Diseases, Tenth Revision, Clinical Modification (ICD-10-CM) diagnosis code that represents the purpose for the service.
- The most appropriate Current Procedural Terminology (CPT) or Healthcare Common Procedure Coding System (HCPCS) procedure code(s) that represent the service(s) provided.
- The appropriate information as indicated on the provider enrollment letter (Electronic claims must also include the most appropriate attested taxonomy code.)

Note: For Texas Medicaid managed care clients, all hearing aid benefits and otology, and audiology services are administered by the client’s Medicaid managed care organization (MCO).
2.4.1.1 Non-implantable Hearing Aid Devices

To be reimbursed for a non-implantable hearing aid device, providers must submit documentation with the paper claim showing their cost for the hearing aid device. The Texas Health and Human Services Commission (HHSC) requires providers to submit non-implantable hearing aid claims using the CMS-1500 paper claim form because electronic claim submissions do not allow for the submission of attachments.

Providers must use the net acquisition cost as the amount billed on the claim. The net acquisition cost is the actual price the provider paid for the device, including the wholesale cost plus sales tax, shipping and handling, and any reductions resulting from discounts or rebates. Providers must not use usual and customary fees as the amount billed.

The documentation submitted with the claim must be a manufacturer invoice showing the net acquisition cost of the non-implantable hearing aid device. An original invoice that includes the manufacturer name, model number, serial number, and warranty expiration date must be submitted. Invoice templates will not be accepted.

An invoice printed from an email or the Internet will not be accepted and should not be submitted with the claim as documentation to show the net acquisition cost of the hearing aid device unless the invoice reflects the actual price the provider paid for the hearing aid device.

2.4.1.2 Third Party Liability

Standard third party liability (TPL) rules apply to all hearing services claims.

Refer to: “Section 8: Third Party Liability (TPL)” (Vol. 1, General Information).

2.4.2 Reimbursement

Hearing aid devices and all hearing and audiological services are reimbursed in accordance with 1 TAC §355.8141. To be reimbursed for both audiology services and hearing aid fitting and dispensing services, audiologists must enroll with Texas Medicaid as audiologists and also as hearing aid fitters and dispensers. Audiology services must be billed using the audiologist provider number and benefit code (for electronic claims only) as indicated on the provider enrollment letter that indicates “Audiologist,” and hearing aid fitting and dispensing services must be billed with the hearing aid provider number and benefit code (for electronic claims only) as indicated on the provider enrollment letter that indicates “Hearing Aid.”

Requested items that are not represented by a specific procedure code must be prior authorized and are priced manually during the authorization process. Manually priced items for clients who are birth through 20 years of age require prior authorization that must be obtained through the TMHP SMPA Department. The reimbursement will be determined based on either the MSRP less 18 percent or based on the provider’s documented invoice cost if there is no MSRP available.

Manually priced items are indicated with “Note Code 5” in the Texas Medicaid fee schedule.

Texas Medicaid implemented mandated rate reductions for certain services. The OFL and static fee schedules include a column titled “Adjusted Fee” to display the individual fees with all mandated percentage reductions applied. Additional information about rate changes is available on the TMHP website at www.tmhp.com/resources/rate-and-code-updates/rate-changes.

 Providers may refer to the OFL or the applicable fee schedule on the TMHP website at www.tmhp.com.

2.4.2.1 National Correct Coding Initiative (NCCI) and Medically Unlikely Edit (MUE) Guidelines

The HCPCS and CPT codes included in the Texas Medicaid Provider Procedures Manual are subject to NCCI relationships, which supersede any exceptions to NCCI code relationships that may be noted in the manuals. Providers should refer to the CMS NCCI web page for correct coding guidelines and specific applicable code combinations.
In instances when Texas Medicaid limitations are more restrictive than NCCI MUE guidance, Texas Medicaid limitations prevail.

3  Implantable Hearing Devices and Related Services

3.1  Enrollment
To enroll in Texas Medicaid, hearing services professionals who provide implantable hearing devices and services must be appropriately enrolled according to their licensure and scope of practice.

Providers cannot be enrolled if their license is due to expire within 30 days. A current license must be submitted.

3.2  Services, Benefits, Limitations and Prior Authorization
Implantable hearing devices, including the cochlear implant device, the auditory brainstem implant (ABI), and the bone anchored hearing device (BAHD), are benefits of Texas Medicaid for clients of all ages.

The following services and supplies must be provided to Texas Medicaid clients if an implantable hearing aid device is medically necessary:

• An individual client assessment to identify the appropriate type of device
• The fitting of the device
• The reassessment to determine whether the device allows for adequate hearing
• Expendable supplies that are necessary to keep the device functioning properly, such as batteries and accessories

Hearing devices are a benefit for clients of any age. Some types of hearing devices are age restricted.

3.2.1  Cochlear Implants
The following procedure codes may be reimbursed for the cochlear implant device, separate components, and services:

<table>
<thead>
<tr>
<th>Procedure Codes</th>
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</thead>
<tbody>
<tr>
<td>69930</td>
</tr>
<tr>
<td>L7368</td>
</tr>
<tr>
<td>L8499</td>
</tr>
<tr>
<td>L8614</td>
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<tr>
<td>L8615</td>
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<tr>
<td>L8616</td>
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<td>L8617</td>
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<td>L8618</td>
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<td>L8619</td>
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<td>L8621</td>
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<td>L8622</td>
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<td>L8623</td>
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<td>L8624</td>
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<tr>
<td>L8627</td>
</tr>
<tr>
<td>L8628</td>
</tr>
<tr>
<td>L8629</td>
</tr>
</tbody>
</table>

The following procedure codes may be reimbursed for diagnostic analysis of the cochlear implant:

<table>
<thead>
<tr>
<th>Procedure Codes</th>
</tr>
</thead>
<tbody>
<tr>
<td>92601</td>
</tr>
<tr>
<td>92602</td>
</tr>
<tr>
<td>92603</td>
</tr>
<tr>
<td>92604</td>
</tr>
</tbody>
</table>

3.2.1.1  Prior Authorization
Prior authorization is required for the following:

• Cochlear implant surgery, device, and replacement parts
• Sound processor repair or replacement
• Battery recharger unit
• Replacement batteries beyond the limitations outlined in the sections below
Requests for prior authorization must be submitted by the provider to the SMPA Department with documentation supporting the medical necessity for the requested device, service, or supply.

**Note:** Requests for clients who are 20 years of age or younger who do not meet the medical necessity criteria may be considered through Comprehensive Care Program (CCP).

Documentation submitted for review must indicate who will be providing the cochlear implant device (i.e., the facility or the Durable Medical Equipment (DME) or medical supplier). The supplier’s provider number must be included on the prior authorization request.

Prior authorization for a unilateral or bilateral cochlear implant may be granted for clients who are 12 months of age and older with documentation of all of the following criteria:

- Cognitive ability to use auditory cues and written documentation of agreement by the client or the client’s parent or guardian that the client will participate in a program of post-implantation auditory rehabilitation. This documentation must be maintained in the client’s medical record.
- Postlingual deafness or prelingual deafness.
- Freedom from middle-ear infection, an accessible cochlear lumen that is structurally suited to implantation, and freedom from lesions in the auditory nerve and acoustic areas of the central nervous system.
- No contraindications to surgery.
- Inability to derive benefit from appropriately fitted hearing aid devices.
- Documentation of poor speech discrimination and a recommendation for cochlear implant candidacy and the most appropriate ICD-10-CM diagnoses for severe-to-profound bilateral sensorineural hearing loss.

The initial lithium ion battery recharger unit, additional medically necessary units, and additional replacement batteries beyond the limitations indicated in the following sections may be reimbursed with prior authorization. Documentation must be submitted with the prior authorization request to support medical necessity for the request.

**Refer to:** Subsection 3.2.4, “Sound Processor Replacement and Repair” in this handbook for more information about sound processor repair or replacement.

Prior authorization requests may be submitted to the TMHP Prior Authorization Department by mail, fax, or the electronic portal. Prescribing or ordering providers, dispensing providers, clients’ responsible adults, and clients may sign prior authorization forms and supporting documentation using electronic or wet signatures.

### 3.2.1.2 Limitations

#### Surgery

Procedure code 69930 with the appropriate modifier LT or RT may be reimbursed for unilateral cochlear implantation. Procedure code 69930 with modifier 50 may be reimbursed for bilateral cochlear implantation performed simultaneously.

#### Device and Components

Procedure codes L8627, L8628, and L8629 for the cochlear implant device and components may be reimbursed for clients who are 12 months of age and older as follows:

- The device must be approved by the Food and Drug Administration (FDA) and be age-appropriate for the client.
- One per day may be reimbursed with prior authorization.

The cochlear implant device and the surgery to implant the device may be reimbursed separately.
Replacement Batteries and Related Items

Replacement batteries and related items for the cochlear implant device include non-rechargeable batteries, rechargeable batteries, and recharger units as follows:

<table>
<thead>
<tr>
<th>Procedure Code</th>
<th>Prior Authorization</th>
<th>Limitation</th>
</tr>
</thead>
<tbody>
<tr>
<td>L8621 (Zink air non-rechargeable)</td>
<td>Not required</td>
<td>Maximum of 50 per month</td>
</tr>
<tr>
<td>L8622 (Alkaline non-rechargeable)</td>
<td>Not required</td>
<td>Maximum of 31 per month</td>
</tr>
<tr>
<td>L8623 (Lithium ion rechargeable)</td>
<td>Not required</td>
<td>2 batteries per calendar year</td>
</tr>
<tr>
<td>L8624 (Lithium ion rechargeable)</td>
<td>Not required</td>
<td>2 batteries per calendar year</td>
</tr>
<tr>
<td>L7368 (Battery recharger unit for lithium ion rechargeable batteries)</td>
<td>Required</td>
<td>1 replacement unit every 5 rolling years</td>
</tr>
</tbody>
</table>

Replacement batteries for clients with bilateral cochlear implants and two sound processors may be reimbursed when billed with the applicable battery procedure code and the appropriate LT or RT modifier.

Replacement batteries for the cochlear device are limited to clients with a previously paid cochlear implant procedure, device, or supply. Replacement batteries for clients who did not receive the cochlear implant through Texas Medicaid will be considered for reimbursement on appeal with a physician’s statement documenting medical necessity.

Additional batteries and lithium ion battery recharger units beyond these limitations may be reimbursed with prior authorization.

3.2.1.3 Auditory Rehabilitation

Auditory rehabilitation is a benefit of Texas Medicaid when it is medically necessary for clients who have received a surgically implanted hearing device, or who have prelingual or postlingual hearing loss when the treating physician has determined that auditory rehabilitation would be beneficial.

The following procedure codes may be reimbursed for auditory rehabilitation:

<table>
<thead>
<tr>
<th>Procedure Codes</th>
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<tr>
<td>92626</td>
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</table>

One auditory rehabilitation evaluation and 12 visits per 180 day period may be reimbursed without prior authorization. Additional visits during a six rolling month period for clients who are 12 months of age through 20 years of age require prior authorization.

Procedure code 92627 is an add-on procedure, and must be billed with the primary procedure code 92626 to be considered for reimbursement.

Procedure code 92627 may be reimbursed up to four times per day to the same provider.

**Note:** Additional therapy services may be a benefit through the Texas Medicaid speech therapy benefit.

**Refer to:** The Physical Therapy, Occupational Therapy, and Speech Therapy Services Handbook (Vol. 2, Provider Handbooks) for information about speech therapy.

Personal frequency modulated (FM) systems are not benefits of Texas Medicaid.
3.2.2 Auditory Brainstem Implant (ABI)

The following procedure codes may be reimbursed for the ABI, related components, and services:

<table>
<thead>
<tr>
<th>Procedure Codes</th>
</tr>
</thead>
<tbody>
<tr>
<td>92640</td>
</tr>
</tbody>
</table>

3.2.2.1 Prior Authorization

The following implantable hearing devices and services require prior authorization:

- ABI surgery, device, and replacement parts
- Sound processor repair or replacement
- Replacement batteries beyond the limitations outlined in the sections below

Requests for prior authorization must be submitted to the SMPA Department with documentation supporting the medical necessity for the requested device, service, or supply.

Prior authorization requests and claims for ABI is limited to clients with a condition of neurofibromatosis, type II or schwannomatosis.

Refer to:

Subsection 2.2.1, “Limitations and Required Forms” in this handbook for additional information about replacement batteries.

Subsection 3.2.4, “Sound Processor Replacement and Repair” in this handbook for more information about sound processor repair or replacement.

3.2.2.2 Limitations

ABI is a benefit for clients who are 12 years of age and older.

Diagnostic analysis of the ABI (procedure code 92640) is limited to 2 hours per day when billed by any provider.

3.2.3 Bone-Anchored Hearing Device (BAHD)

The following procedure codes must be submitted for the BAHD and related components:

<table>
<thead>
<tr>
<th>Procedure Codes</th>
</tr>
</thead>
<tbody>
<tr>
<td>69714</td>
</tr>
</tbody>
</table>

3.2.3.1 Prior Authorization

The following implantable hearing devices and services require prior authorization:

- BAHD implant surgery, device, and replacement parts
- Sound processor repair or replacement

Requests for prior authorization must be submitted to the SMPA Department with documentation supporting the medical necessity for the requested device, service, or supply.

Providers should use the Special Medical Prior Authorization (SMPA) Request Form for all prior authorization requests.

Prior authorization requests may be granted for clients who are 5 years of age and older with all of the following:

- Documentation of previous attempts at hearing aid devices and why these devices are inadequate or have failed
- Documentation of scores on hearing tests for bone conduction thresholds and on maximum speech discrimination
• Documentation of audiological testing showing good inner ear function
• Documentation of a multidisciplinary assessment including physical, cognitive, communicative, and behavioral limitations describing the client’s auditory disability and expected benefit with use of the BAHD implant
• Documentation of an appropriate diagnosis.

Benefit-eligible conditions may include, but are not limited to the following:
• Conductive hearing loss
• Sensorineural hearing loss
• Other anomalies of external ear with impairment of hearing
• Anomalies of skull and face bones

Refer to: Subsection 3.2.4, “Sound Processor Replacement and Repair” in this handbook for more information about sound processor repair or replacement.

3.2.3.2 Limitations
Implanted BAHDs are a benefit for clients who are 5 years of age and older.

A BAHD sound processor that is specifically worn on a soft headband is a benefit for clients five years old or younger. The BAHD sound processor is not implanted. A BAHD sound process worn on a soft headband may be reimbursed using procedure code L8692.

Replacement batteries for the BAHD (procedure code V5266) do not require prior authorization. The replacement batteries are limited to clients with a previously paid hearing device. Replacement batteries for clients who did not receive the hearing device through Texas Medicaid will be considered for reimbursement on appeal with a physician’s statement documenting the medical necessity.

Procedure codes L8691, L8692, L8693, and L8694 will be denied as part of another service when billed by any provider with the same date of service as procedure code L8690.

Procedure code L8692 for the BAHD device and components may be reimbursed once per day with prior authorization.

Bilateral BAHD procedures are not benefits of Texas Medicaid.

3.2.4 Sound Processor Replacement and Repair

3.2.4.1 Prior Authorization
Replacement and repair of a sound processor require prior authorization.

Documentation by the provider must explain the need for the replacement of the sound processor. The processor must be used for a minimum of 12 months before replacement of the unit will be considered.

The prior authorization request must include evidence of the purchase, such as the manufacturer’s warranty.

Repair of a sound processor will be considered for prior authorization with documentation of medical necessity for the requested repair. Repair of a sound processor will be manually priced at the time the prior authorization is reviewed and granted. If the actual cost of the repair differs from the prior authorized fee, the provider must contact the SMPA Department to update the authorization before filing a claim for the repair services.

3.2.4.2 Limitations
Procedure code L8499 with modifier RB may be reimbursed for sound processor repair.

Repair or replacement of a sound processor is not a benefit during the manufacturer’s warranty period.
3.2.5 Electromagnetic Bone Conduction Hearing Device - Removal Only
The removal of the electromagnetic bone conduction hearing aid may be reimbursed by Texas Medicaid using procedure code 69711.

The removal or repair of an electromagnetic bone conduction hearing device is limited to two procedures per lifetime when billed by any provider.

The implantation of the device is not a benefit of Texas Medicaid.

3.3 Documentation Requirements
All implantable hearing aid services require documentation to support the medical necessity of the service rendered. Hearing services are subject to retrospective review and recoupment if documentation does not support the service billed.

3.4 Claims Filing and Reimbursement

3.4.1 Claims Filing
Hearing services must be submitted to TMHP in an approved electronic format or on the CMS-1500 paper claim form. Providers may purchase CMS-1500 paper claim forms from the vendor of their choice. TMHP does not supply the forms.

When completing a CMS-1500 paper claim form, all required information must be included on the claim, as information is not keyed from attachments. Superbills, or itemized statements, are not accepted as claim supplements.

Refer to:
Subsection 2.2, “Fee-for-Service Reimbursement Methodology” in “Section 2: Texas Medicaid Fee-for-Service Reimbursement” (Vol. 1, General Information) for more information about reimbursement.

Subsection 1.7.11, “Billing Clients” in “Section 1: Provider Enrollment and Responsibilities” (Vol. 1, General Information).

“Section 3: TMHP Electronic Data Interchange (EDI)” (Vol. 1, General Information) for information on electronic claims submissions.

Subsection 6.1, “Claims Information” in “Section 6: Claims Filing” (Vol. 1, General Information) for general information about claims filing.

Subsection 6.5, “CMS-1500 Paper Claim Filing Instructions” in “Section 6: Claims Filing” (Vol. 1, General Information). Blocks that are not referenced are not required for processing by TMHP and may be left blank.

Note: For Texas Medicaid managed care clients, all implantable hearing devices and services are administered by the client’s Medicaid MCO.

3.4.1.1 Third Party Liability
Standard TPL rules apply to all hearing services claims.

Refer to: “Section 8: Third Party Liability (TPL)” (Vol. 1, General Information).

3.4.2 Reimbursement
Implantable hearing aids and related services are reimbursed in accordance with 1 TAC §355.8141.

Implantable hearing aids and related services are reimbursed at the lesser of the billed charges or the published Texas Medicaid fee. Unless otherwise indicated, providers may not make additional charges to the client for covered services; such charges constitute a breach of the Texas Medicaid contract.
Requested items that are not represented by a specific procedure code must be prior authorized and are priced manually during the authorization process. Manually priced items for clients who are birth through 20 years of age require prior authorization that must be obtained through the TMHP SMPA Department. The reimbursement will be determined based on either the MSRP less 18 percent or based on the provider’s documented invoice cost. Manually priced items are indicated with “MP” in the reimbursement rate table at the end of this article.

Texas Medicaid implemented mandated rate reductions for certain services. The OFL and static fee schedules include a column titled “Adjusted Fee” to display the individual fees with all mandated percentage reductions applied. Additional information about rate changes is available on the TMHP website at www.tmhp.com/resources/rate-and-code-updates/rate-changes.

Providers may refer to the OFL or the applicable fee schedule on the TMHP website at www.tmhp.com.

### 3.4.2.1 NCCI and MUE Guidelines

The HCPCS and CPT codes included in the Texas Medicaid Provider Procedures Manual are subject to NCCI relationships, which supersede any exceptions to NCCI code relationships that may be noted in the manuals. Providers should refer to the CMS NCCI web page for correct coding guidelines and specific applicable code combinations.

In instances when Texas Medicaid limitations are more restrictive than NCCI MUE guidance, Texas Medicaid limitations prevail.

## 4 Vision Care Professionals

### 4.1 Enrollment

To enroll in Texas Medicaid, optometrists (doctors of optometry [ODs]) and ophthalmologists must be licensed by the licensing board of their profession to practice in the state where the service is performed, at the time the service is performed, and be enrolled as Medicare providers.

An optometrist or ophthalmologist cannot be enrolled if their license is due to expire within 30 days; a current license must be submitted.

### 4.2 Provider Responsibilities

Suppliers of eyewear must comply with all Medicaid provider responsibilities and adhere to the following guidelines:

- Do not delay the ordering of eyewear or the dispensing of eyeglasses to the client while payment is pending from TMHP.
- Deliver the eyewear in a reasonable amount of time (usually two or three weeks from the date the order is placed by the client).
- Check the client’s eligibility
- Visit TexMedConnect or access the Medicaid Client Portal for Providers to determine whether eyeglasses have been reimbursed by Texas Medicaid within the last 24 months. Providers should ask clients if they recently received vision care services that may not be documented in My Account yet.
- Submit claims for eyewear services as soon as possible so the client’s record indicates that eyewear or eyeglasses have been dispensed.
- Have the client, parent, or guardian sign and date the Vision Care Eyeglass Patient (Medicaid Client) Certification Form and retain it in their records. When a client chooses an eyeglass or contact lens option beyond the program limitations, or if nonprosthetic eyeglasses or contact lenses
are replaced because of loss or destruction, the client must acknowledge their choice and his/her liability for the cost difference by signing the Vision Care Eyeglass Patient (Medicaid Client) Certification Form. The form must remain in the provider’s records.

- Do not charge a Medicaid client more than a patient not enrolled in Texas Medicaid for noncovered services (e.g., tints, oversized lenses, or frames).
- Keep invoices on file for a minimum of five years.
- Submit claims using the date eyeglasses were ordered as the date of service (DOS) (the start of the 95-day filing period), not the date the eyewear was dispensed.

4.3 Services, Benefits, Limitations, and Prior Authorization

Examination and treatment of eye conditions, including prescribing and dispensing of medically necessary eyeglasses or contact lenses, are benefits of Texas Medicaid and may be reimbursed to optometrist, ophthalmologist, and optician providers as is within the scope of practice for each.

The following services are included in other services and will not be considered for separate reimbursement:

- Vision screening conducted to meet State screening requirements, such as the DSHS School Vision and Hearing Screening Program.
- Expenses for medical supplies, equipment, and other items that are not specifically made-to-order for the client are considered to have been incurred on the date the item is delivered.

**Ophthalmologist and Optometrist**

Examination and treatment services rendered by an ophthalmologist or optometrist are not limited to the procedure codes included in this handbook.

**Refer to:** The Texas Medicaid fee schedules on the TMHP web site at www.tmhp.com for a complete list of procedure codes that may be reimbursed by Texas Medicaid.

**Optician**

Services rendered by an optician are limited to fitting and dispensing of medically necessary eyeglasses and contact lenses.

**Note:** In accordance with the Omnibus Reconciliation Act of 1986, Section 9336, a Doctor of Optometry is considered a physician, with respect to the provision of any item or service the optometrist is authorized to perform by state law or regulation.

4.3.1 Services Performed in Long-Term Care Facilities

Ophthalmological, optometric, and eyeglass or contact lens services provided in a skilled or intermediate care facility may be reimbursed when the client’s attending physician has ordered the service and the signed order is included in the client’s medical record at the nursing facility.

The ordering physician’s name and provider identifier must be documented on the claim when ophthalmological, optometric, or eyeglasses or contact lenses services are performed in a skilled or intermediate care facility.

4.3.2 Services Performed in Federally Qualified Healthcare Centers (FQHC)

Vision services rendered by FQHC providers may be reimbursed based on an all-inclusive rate per visit.

**Refer to:** Subsection 2.2, “Services, Benefits, Limitations, and Prior Authorization” in the Clinics and Other Outpatient Facility Services Handbook (Vol. 2, Provider Handbooks) for information about vision services that may be reimbursed to FQHC providers.
4.3.3 THSteps Medical Checkup Vision Screening

A vision screening must be completed during each THSteps medical checkup with standardized screenings performed at specific ages, as listed in the THSteps Periodicity Schedule. Providers may perform a vision screening during an acute care visit with the appropriate screening tools or refer at-risk infants and children to an optometrist or ophthalmologist who is experienced with the pediatric population and who can perform further testing, diagnosis, and treatment.

Refer to: Subsection 5.3.11.2.4, “Vision Screening” in the Children’s Services Handbook (Vol. 2, Provider Handbooks) for information about THSteps medical checkup vision screenings.

4.3.3.1 Vision Screening Outside of a THSteps Preventive Care Medical Checkup

Vision screening for clients who are birth through 20 years of age may be completed at any primary care provider’s office visit upon the following:

- Request from a parent
- Referral from a school vision screening program
- Referral from a school nurse

Clients who are birth through 20 years of age must be screened for eye abnormalities by history, observation, and physical exam. Clients who are identified as high risk must be referred to an appropriate Medicaid-enrolled optometrist or ophthalmologist.

4.3.4 Noncovered Services

The following services and supplies are not a benefit of Texas Medicaid:

- Artificial eyes for clients who are 21 years of age and older.
  
  Note: Artificial eyes for clients who are birth through 20 years may be considered under Texas Health Steps-Comprehensive Care Program (THSteps-CCP).

- Eyeglasses for residents of institutions where the reimbursement formula and vendor reimbursement include this service.

- Eyeglasses or contact lenses prescribed or dispensed to clients at a hospital or nursing facility without documented orders of the attending physician in the client’s medical records.

- Low vision aids.
  
  Note: Clients may be referred to TWC for low vision aids.

- Optional eyeglass features that are requested by the client but that do not increase visual acuity (e.g., lens tint, industrial hardening, and decorative accessories or lettering).

- Plano sunglasses.

- Extended color vision examination (procedure code 92283), dark adaptation examination (procedure code 92284), and vision screening (procedure codes 99172 or 99173).

- Spectacle (eyeglass) fitting services when billed separately.

Clients may be billed for noncovered frames and other items beyond Medicaid benefits. Providers must have the client sign and date the Vision Care Eyeglass Patient (Medicaid Client) Certification Form and retain it in the provider’s records. The client payment amount is not considered other insurance and must not be entered as a credit amount in the electronic field.

Example: Texas Medicaid may reimburse providers a total of $32.55 for eyeglass frames that are within the provider’s selection for Medicaid reimbursement plus the allowed cost per lens. If the client chooses a pair of frames (such as $200 frames) that are outside of the provider’s selections for Medicaid reimbursement and if the client chooses other items or services that are not a benefit of Texas Medicaid (such as tinted lenses for an extra $10
The provider may withhold the noncovered eyewear, contacts, or eyeglasses until the client pays for those items. If the client fails to pay for the noncovered items or has not returned for finished eyewear within a reasonable length of time (two to three months), the provider may return any reusable items to stock. Any payment made by TMHP for frames or lenses must be refunded to Texas Medicaid. If a client requests eyewear that is beyond program benefits (for example, scratch-resistant coating), Medicaid allows reimbursement up to the maximum fee. The provider may charge the client the difference between the Medicaid payment and the customary charge for the eyewear requested, when the client has been shown the complete selection of Medicaid-covered eyewear and when the following conditions are met:

- The client rejects the Medicaid-covered eyewear and wants eyewear that complies with Texas Medicaid specifications, but is not included in the selection of Medicaid-covered eyewear.
- The client indicates a willingness to pay the difference between the Medicaid payment and the actual charge. The provider must have the client sign the Vision Care Eyeglass Patient (Medicaid Client) Certification Form and retain it in the provider’s records.

**Note:** Children and young adults between the ages of birth and 22 years who have been determined by their provider to have a vision impairment that may require intensive or comprehensive vision impaired related services should be referred to the HHSC Blind Children’s Vocational Discovery and Development Program. Providers should also refer adults, youths, and students that would like to prepare for employment or post-secondary opportunities to TWC.

### 4.3.5 Vision Testing

Vision testing and examination and treatment of eye conditions are benefits of Texas Medicaid and may be reimbursed to ophthalmologist or optometrist providers.

Eye examinations with refraction testing may be reimbursed using the following procedure codes:

<table>
<thead>
<tr>
<th>Procedure Codes</th>
</tr>
</thead>
<tbody>
<tr>
<td>S0620</td>
</tr>
<tr>
<td>S0621</td>
</tr>
</tbody>
</table>

Medical evaluation and examination may be reimbursed using the following procedure codes:

<table>
<thead>
<tr>
<th>Procedure Codes</th>
</tr>
</thead>
<tbody>
<tr>
<td>92002</td>
</tr>
<tr>
<td>92004</td>
</tr>
<tr>
<td>92012</td>
</tr>
<tr>
<td>92014</td>
</tr>
<tr>
<td>92015</td>
</tr>
</tbody>
</table>

Procedure codes 92002, 92004, 92012, and 92014 are limited to one service per day by any provider.

**Refer to:** Subsection 9.2.56.1.1, “New and Established Patient Services” in the Medical and Nursing Specialists, Physicians, and Physician Assistants Handbook (Vol. 2, Provider Handbooks) for information about new patient and established patient E/M services.

Vision testing procedure codes are subject to the CMS NCCI relationships. Claims that are submitted by physicians with the same specialty who are in the same group practice are processed as if they were the same provider. Providers should refer to the Current Procedural Terminology (CPT) Manual for additional information about intermediate and comprehensive ophthalmological services.

### 4.3.5.1 Routine Vision Testing

Procedure codes S0620 and S0621 may be reimbursed for routine vision testing with refraction when they are billed with diagnosis code Z0100 or Z0101.
Procedure codes S0620 and S0621 will be denied if billed with the same date of service as procedure codes 92020, 92273, and 92274.

Clients who are birth through 20 years of age are eligible for a routine eye examination with refraction testing for the purpose of obtaining eyeglasses or contact lenses once every 12 months. The limitation for refraction testing can be exceeded for clients who are birth through 20 years of age only when:

- The parent, teacher, or school nurse requests the refraction testing and it is medically necessary.
- There is a significant change in vision, and documentation supports a diopter (d) change of 0.5d or greater in the sphere, cylinder, prism measurements, or axis changes.

Clients who are 21 years of age and older are eligible for a routine eye examination with refraction testing for the purpose of obtaining eyeglasses or contact lenses once every 24 months. The limitation for refraction testing can be exceeded for clients who are 21 years of age and older only when there is a significant change in vision, and documentation supports a diopter change of 0.5d or more in the sphere, cylinder, prism measurements, or axis changes.

### 4.3.5.2 Medically Necessary Eye Examinations

An eye examination with or without refraction (procedure code 92002, 92004, 92012, 92014, or 92015) may be reimbursed for medical evaluations and examinations of the eye. Procedure codes 92002, 92004, 92012, 92014, and 92015 will not be reimbursed for routine exams.

Documentation in the client’s medical record must support the medical necessity of the service performed.

Procedure codes 92002, 92004, 92012, and 92014 may be reimbursed when it is medically necessary to ophthalmologist or optometrist providers for medically necessary eye examinations without refraction.

Procedure code 92015 may be reimbursed to ophthalmologist or optometrist providers for refraction in addition to the eye examination procedure code 92002, 92004, 92012, or 92014. A refractive state (procedure code 92015) will be denied as part of another service if it is billed with the same date of service by the same provider as procedure code S0620 or S0621.

### 4.3.5.3 Ophthalmological Examination and Evaluation with General Anesthesia

An ophthalmological examination and evaluation under general anesthesia (procedure codes 92018 and 92019) performed by an ophthalmologist may be medically necessary when a client has significant injury or cannot otherwise tolerate the procedure while conscious.

Procedure codes 92018 and 92019 may be reimbursed once per service, per day, when billed by any provider.

### 4.3.5.4 Ophthalmic Ultrasound

Ophthalmic ultrasound is an ultrasonic diagnostic test that uses high frequency sound waves that are used to provide additional information about the interior of the eye and surrounding areas. The following procedure codes may be reimbursed for ophthalmic ultrasound services:

<table>
<thead>
<tr>
<th>Procedure Codes</th>
</tr>
</thead>
<tbody>
<tr>
<td>76510</td>
</tr>
</tbody>
</table>

Procedure codes 76510, 76511, 76512, 76513, 76516, and 76519 are limited to two services per calendar year by any provider.

Procedure code 76999 requires prior authorization.
Procedure code 76514 may be reimbursed once per lifetime, when billed by any provider, for the following diagnosis codes:

<table>
<thead>
<tr>
<th>Diagnosis Codes</th>
</tr>
</thead>
<tbody>
<tr>
<td>H40001</td>
</tr>
<tr>
<td>H40023</td>
</tr>
<tr>
<td>H40052</td>
</tr>
<tr>
<td>H4010X3</td>
</tr>
<tr>
<td>H401121</td>
</tr>
<tr>
<td>H401134</td>
</tr>
<tr>
<td>H401212</td>
</tr>
<tr>
<td>H401230</td>
</tr>
<tr>
<td>H401313</td>
</tr>
<tr>
<td>H401331</td>
</tr>
<tr>
<td>H401511</td>
</tr>
<tr>
<td>H401524</td>
</tr>
<tr>
<td>H4020X2</td>
</tr>
<tr>
<td>H402212</td>
</tr>
<tr>
<td>H402230</td>
</tr>
<tr>
<td>H40233</td>
</tr>
<tr>
<td>H4031X4</td>
</tr>
<tr>
<td>H4033X2</td>
</tr>
<tr>
<td>H4042X0</td>
</tr>
<tr>
<td>H4043X3</td>
</tr>
<tr>
<td>H4052X1</td>
</tr>
<tr>
<td>H4053X4</td>
</tr>
<tr>
<td>H4062X2</td>
</tr>
<tr>
<td>H40811</td>
</tr>
<tr>
<td>H40833</td>
</tr>
</tbody>
</table>

Procedure code 76999 may be reimbursed with prior authorization.

Ophthalmic ultrasounds may be reimbursed when they are billed with the same date of service by the same provider as an eye examination visit or consultation.

Ophthalmic ultrasounds (procedure codes 76514 and 76516) are limited to one service, per day, by any provider. Procedure codes 92002, 92004, 92012, 92014, and 92015 will not be reimbursed for routine exams.

Procedure code 76519 may be reimbursed as follows:
- The professional interpretation component may be reimbursed when procedure code 76519 is billed with modifier LT or RT to identify the eye on which the service was performed.
- The technical component may be reimbursed once when the service is performed on one or both eyes on the same date of service by any provider.
• The total component may be reimbursed along with an additional professional service when the service is performed on both eyes on the same date of service by the any provider. The claim for the additional interpretation component must include modifier LT or RT.

Ophthalmic ultrasound procedure codes are subject to CMS NCCI relationships, except for procedure code 76511, which will be denied when it is billed with the same date of service by the same provider as procedure code 76506.

Refer to: The CMS NCCI web page for the published correct coding guidelines and specific applicable code combinations.

Prior Authorization Requirements
Procedure code 76999 requires prior authorization. The provider must submit the following documentation with the request:

• A clear, concise description of the ophthalmic ultrasound being performed.
• A procedure code that is comparable to the ophthalmic ultrasound being requested or the provider’s intended fee for performing the ophthalmic ultrasound.

Note: Services and procedures that are investigational or experimental are not a benefit of Texas Medicaid.

4.3.5.5 Corneal Topography
Procedure code 92025 may be reimbursed for corneal topography. Procedure code 92025 is considered medically necessary to diagnose, monitor, and treat various visual conditions such as, but not limited to the following:

• Corneal abrasion
• Corneal irregularities
• Corneal disease
• Corneal injury
• Keratoconus

Corneal topography may be reimbursed when it is billed with the same date of service by the same provider as an eye examination visit or consultation.

Corneal topography (procedure code 92025) is limited to one service, per day, by any provider and two services per calendar year.

4.3.5.6 Gonioscopy
A gonioscopy consists of an eye examination to look at the front part of the eye (anterior chamber) between the cornea and the iris. A microscope (slit lamp) is used to look inside the eye.

Procedure code 92020 may be reimbursed for gonioscopy when billed with one of the diagnosis codes in the following table:

<table>
<thead>
<tr>
<th>Diagnosis Codes</th>
</tr>
</thead>
<tbody>
<tr>
<td>H40001</td>
</tr>
<tr>
<td>H40023</td>
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<tr>
<td>H40052</td>
</tr>
<tr>
<td>H4010X3</td>
</tr>
<tr>
<td>H401121</td>
</tr>
<tr>
<td>H401134</td>
</tr>
</tbody>
</table>
4.3.5.7 Sensorimotor Examination

A sensorimotor examination is an evaluation of the function of the ocular neuro-muscular system. This exam includes interpretation and reporting of multiple ocular deviation measurements and includes, but is not limited to, visual motor integration, reversal frequency (letters and numbers), motor speed and precision, visual memory, and visualization to test eye movement and control, focusing ability, eye teaming ability, depth perception, and visual perception skills.

Sensorimotor examination (procedure code 92060) may be reimbursed once per day and twice per calendar year by any provider. Procedure code 92060 may be reimbursed in addition as an eye examination visit.

4.3.5.8 Orthoptic or Pleoptic Training

Orthoptics, a component of vision training or vision therapy, are exercises designed to improve the function of the eye muscles with an emphasis on binocular vision and eye movements. Pleoptics are exercises designed to improve impaired vision when there is no evidence of organic eye diseases.

Procedure code 92065 may be reimbursed for orthoptic or pleoptic training when it is billed with one of the following diagnosis codes:

<table>
<thead>
<tr>
<th>Diagnosis Codes</th>
</tr>
</thead>
<tbody>
<tr>
<td>H401212</td>
</tr>
<tr>
<td>H401230</td>
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<tr>
<td>H401313</td>
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<tr>
<td>H401331</td>
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<tr>
<td>H401511</td>
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<tr>
<td>H401524</td>
</tr>
<tr>
<td>H4020X2</td>
</tr>
<tr>
<td>H402212</td>
</tr>
<tr>
<td>H402230</td>
</tr>
<tr>
<td>H40233</td>
</tr>
<tr>
<td>H4031X4</td>
</tr>
<tr>
<td>H4033X2</td>
</tr>
<tr>
<td>H4042X0</td>
</tr>
<tr>
<td>H4043X3</td>
</tr>
<tr>
<td>H4052X2</td>
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<tr>
<td>H4053X4</td>
</tr>
<tr>
<td>H4062X2</td>
</tr>
<tr>
<td>H40811</td>
</tr>
<tr>
<td>H42</td>
</tr>
</tbody>
</table>
Orthoptic or pleoptic training may be reimbursed one service per day for up to 6 services when it is billed with one of the diagnosis codes in the above diagnosis table. Up to an additional 6 services may be reimbursed with prior authorization for a total of 12 services per lifetime.

The provider must attest that current therapy has resulted in an improvement with presenting symptomatology over the course of treatment, including, but not limited to:

- Blurred vision
- Double vision
- Amblyopia
- Accommodation or near point of convergence measurements

**Note:** Orthoptic or pleoptic training services over the 12 per lifetime limit may be considered with prior authorization through CCP for clients who are birth through 20 years of age. Documentation for medical necessity must be submitted with the prior authorization request.

Procedure code 92065 may be reimbursed in addition to an eye examination visit.

### 4.3.5.9 Special Ophthalmological Services

The following procedure codes may be reimbursed for special ophthalmological services:

<table>
<thead>
<tr>
<th>Procedure Codes</th>
</tr>
</thead>
<tbody>
<tr>
<td>92081</td>
</tr>
</tbody>
</table>

*Total component only

Procedure codes 92081, 92082, and 92083 may be reimbursed once per day and twice per calendar year by any provider of any combination.

Procedure codes 92100, 92132, 92133, and 92134 may be reimbursed once per day and twice per calendar year by any provider.

Procedure code 92136 may be reimbursed twice per calendar year by any provider as follows:

- The professional interpretation component may be reimbursed when billed with modifier LT or RT to identify the eye on which the service was performed.
- The technical component may be reimbursed once when the service is performed on one or both eyes on the same date of service by any provider.
- The total component may be reimbursed along with an additional professional service when the service is performed on both eyes on the same date of service by the any provider.
4.3.5.10 Ophthalmoscopy and Extended Ophthalmoscopy

Ophthalmoscopy and extended ophthalmoscopy may be reimbursed using the following procedure codes:

<table>
<thead>
<tr>
<th>Procedure Codes</th>
</tr>
</thead>
<tbody>
<tr>
<td>92201</td>
</tr>
<tr>
<td>* Total component only</td>
</tr>
</tbody>
</table>

Ophthalmoscopy, fluorescein angiography, indocyanine-green angiography and fluorescein angiography (procedure codes 92230, 92235, and 92240) may be reimbursed for a quantity of two if both the left and right eyes are evaluated. If two services are billed for the same date of service, one may be reimbursed at the full rate, and the other may be reimbursed at half rate.

Procedure codes 92230, 92235, and 92240 are limited to one service per eye per day and two services per eye per calendar year when billed by any provider.

Procedure codes 92230, 92235, and 92240 must be billed with modifier LT or RT to identify the eye on which the service was performed.

Procedure codes 92201 and 92202 are limited to one service per day and two services per calendar year by any provider.

Procedure codes 92242, 92250 and 92260 are limited to one service per day and two services per calendar year by any provider.

Ophthalmoscopy, angiography, and angiography procedure codes are subject to CMS NCCI relationships.

Fundus photography (procedure code 92250) are considered medically necessary when a clinical condition exists that is subject to change in extent, appearance or size and where such change would directly affect the management of client care. These conditions include, but are not limited to the following:

- Macular degeneration
- Glaucoma
- Hypertension
- Neoplasms of the retina
- Choroid (benign or malignant)
- Retinal hemorrhages
- Ischemia
- Retinal detachment
- Choroid disturbances
- Diabetic retinopathy
- Assessment of recently performed retinal laser surgery

Note: Fundus photography performed for a routine screen of a normal eye, in the absence of a clinical condition, that is subject to change in extent, appearance or size is not a benefit of Texas Medicaid.
### 4.3.5.11 Other Specialized Vision Services

The following procedure codes may be reimbursed by Texas Medicaid when the services are medically necessary:

<table>
<thead>
<tr>
<th>Procedure Codes</th>
</tr>
</thead>
<tbody>
<tr>
<td>92227*</td>
</tr>
<tr>
<td>92228</td>
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<tr>
<td>92229</td>
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<tr>
<td>92265</td>
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<tr>
<td>92270</td>
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<tr>
<td>92273</td>
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<tr>
<td>92274</td>
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<tr>
<td>92285</td>
</tr>
<tr>
<td>92286</td>
</tr>
<tr>
<td>92287</td>
</tr>
</tbody>
</table>

*Total component only

Procedure codes 92265, 92270, 92273, 92274, 92285, 92286, and 92287 may each be reimbursed once per day and twice per calendar year by any provider.

External ocular photography (procedure code 92285) may be reimbursed once per day, when it is billed by any provider.

For other professional services, fitting services are included in the reimbursement for prosthetic eyeglasses or contact lenses.

Microfluidic analysis is a CLIA-waived lab test and may be performed by optometrists in the office setting.

**Refer to:** Subsection 2.2.16, “Urinalysis and Chemistry” in the Radiology and Laboratory Services Handbook (Vol. 2, Provider Handbooks) for information about microfluidic analysis (procedure code 83861).

### 4.3.6 Nonprosthetic Eyeglasses or Contact Lenses

Nonprosthetic eyeglasses or contact lenses are lenses that are medically necessary to correct defects in vision when the eye’s organic lens is present. Providers may refer to TAC §354.1015 for more information.

Nonprosthetic eyeglasses or contact lenses may be reimbursed for clients of any age when there is no other option available to correct or ameliorate a visual defect. Contact lenses require prior authorization with documentation of medical necessity. Prescribing and dispensing medically necessary eyeglasses or contact lenses are benefits of Texas Medicaid as follows:

- Nonprosthetic eyeglasses or contact lenses may be reimbursed once every 24 months. Additional services within the 24-month period may be considered when documentation in the client’s medical record supports medical necessity that includes a diopter change of 0.5d or more in the sphere, cylinder, prism measurements, or axis changes. A new 24 month benefit period for eyewear begins with the placement of the new nonprosthetic eyewear.

- Replacement of nonprosthetic eyeglasses or contact lenses because of loss or destruction is a benefit of Texas Medicaid for clients who are birth through 20 years of age. If the eyeglasses or contact lenses are lost or destroyed, the provider must have the client sign the Vision Care Eyeglass Patient (Medicaid Client) Certification Form and the signed form must be maintained in the client’s medical record.

- For clients who have had insertion of an intraocular lens (IOL), one pair of eyeglasses or contact lenses may be reimbursed. Additional eyeglasses or contact lenses may be considered when documentation in the client’s medical record supports medical necessity that includes a diopter change of 0.5d or more in the sphere, cylinder, prism measurements, or axis changes.

  **Note:** Because the IOL is considered the prosthetic device, the eyeglasses or contact lenses, and any replacements, are considered nonprosthetic.

**Refer to:** Subsection 4.3.5.1, “Routine Vision Testing” in this handbook for information about vision testing for the purposes of prescribing eyewear.
The prescription for eyeglasses must be given to the client upon request. A provider may not withhold a prescription for eyeglasses from a client even if Medicaid reimbursement for the eye examination has not been received.

To be considered by Texas Medicaid, the eyeglasses or contact lenses must be:

- Medically necessary.
- Prescribed by a doctor of medicine, optometry, or osteopathy.
- Prescribed to significantly improve vision or correct a medical condition.
- In compliance with eyeglass program specifications for frames and lenses as stated in TAC Rule 354.1017, Specifications for Eyewear and Rule 363.503, Specifications for Eyewear.

Note: Contact lenses require prior authorization unless provided in a medical emergency.

### 4.3.6.1 Eyeglass Lenses and Frames

The following eyeglass lens procedure codes may be billed with frame procedure codes V2020 and V2025 for reimbursement of a pair of eyeglasses:

<table>
<thead>
<tr>
<th>Procedure Codes</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>Single Vision Lenses</strong></td>
</tr>
<tr>
<td>V2100 V2101 V2102 V2103 V2104 V2105 V2106 V2107 V2108 V2109</td>
</tr>
<tr>
<td>V2110 V2111 V2112 V2113 V2114 V2115 V2118 V2121</td>
</tr>
<tr>
<td><strong>Bifocal Lenses</strong></td>
</tr>
<tr>
<td>V2200 V2201 V2202 V2203 V2204 V2205 V2206 V2207 V2208 V2209</td>
</tr>
<tr>
<td>V2210 V2211 V2212 V2213 V2214 V2215 V2218 V2219 V2220 V2221</td>
</tr>
<tr>
<td><strong>Trifocal Lenses</strong></td>
</tr>
<tr>
<td>V2300 V2301 V2302 V2303 V2304 V2305 V2306 V2307 V2308 V2309</td>
</tr>
<tr>
<td>V2310 V2311 V2312 V2313 V2314 V2315 V2318 V2319 V2320 V2321</td>
</tr>
</tbody>
</table>

For the purpose of Texas Medicaid, high-powered lenses are lenses with a sphere greater than 7.00d or a cylinder greater than 4.00d.

Providers must bill a quantity of two when billing for bilateral lenses with the same prescription.

The following procedure codes may be reimbursed for add-on services:

<table>
<thead>
<tr>
<th>Add-On Procedure Codes</th>
</tr>
</thead>
<tbody>
<tr>
<td>V2410 V2430 V2700 V2710* V2715* V2718* V2730 V2755 V2770 V2780</td>
</tr>
</tbody>
</table>

* Procedure codes for prism correction: prism, slab off, ground in, Fresnell, press-on

Add-on procedure codes will not be reimbursed unless they are billed with the appropriate lens procedure code by the same provider for the same date of service.

The fitting of eyeglasses (procedure codes 92340, 92341, 92342, and 92370) is considered part of the dispensing procedure and is not separately reimbursed.

**Polycarbonate Lens**

Polycarbonate lenses (procedure code V2784) may be reimbursed for clients with a medical or physical condition such as, but not limited to the following:

- Cerebral palsy
- Multiple sclerosis
• Muscular dystrophy
• Epilepsy
• Autism
• Down’s syndrome
• Brain trauma
• Balance disorders
• Parkinson’s disease
• Seizure disorder
• Motor ataxia
• Marfan’s syndrome
• Ocular prostheses
• Amblyopia

In addition to the medical or physical conditions identified above, polycarbonate lenses also may be reimbursed when the client meets the following criteria:

• Lens power in at least one meridian of -5.25/+4.00 diopters or more and the eyeglasses are not functional in regular standard glass or plastic lens materials due to weight, thickness or aberration
• Monocular vision with functional vision in one eye
• Retinal detachment or risk for retinal detachment (e.g., lattice degeneration, history of retinal detachment in the family, posterior vitreous detachment)

Procedure code V2784 may be reimbursed when it is billed with one of the following diagnosis codes:

<table>
<thead>
<tr>
<th>Diagnosis Codes</th>
</tr>
</thead>
<tbody>
<tr>
<td>F840</td>
</tr>
<tr>
<td>G2119</td>
</tr>
<tr>
<td>G40009</td>
</tr>
<tr>
<td>G40211</td>
</tr>
<tr>
<td>G40A19</td>
</tr>
<tr>
<td>G40419</td>
</tr>
<tr>
<td>G4089</td>
</tr>
<tr>
<td>G7109</td>
</tr>
<tr>
<td>G71220</td>
</tr>
<tr>
<td>G7241</td>
</tr>
<tr>
<td>G802</td>
</tr>
<tr>
<td>H33012</td>
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<td>H33041</td>
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<td>H33193</td>
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<td>H33312</td>
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<tr>
<td>H3341</td>
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<tr>
<td>H53012</td>
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<tr>
<td>H53041</td>
</tr>
</tbody>
</table>
Polycarbonate lens claims must include a lens procedure code with lens power in at least one meridian of -5.25/+4.00 diopters or more, and the eyeglasses are not functional in regular standard glass or plastic lens material due to weight, thickness or aberration.

For diagnoses not listed in the above table or for lens power other than those listed in this section, providers must submit documentation of medical necessity. If documentation is not submitted with the claim, the polycarbonate lenses will be denied.

**Undeliverable Eyeglasses**

The provider may be reimbursed for the lenses based on the services furnished and the materials used up to the time the provider learned that the eyeglasses were undeliverable due to any of the following:

- The client cancels an order for eyeglasses prior to their completion and delivery.
- The prescription changes prior to completion and delivery of the eyeglasses.
- The client dies prior to completion and delivery of the eyeglasses.

Reimbursement will not be made for the frames.

**4.3.6.2 Contact Lenses**

Prior authorization is required for all contact lenses. The following procedure codes may be reimbursed for prosthetic and nonprosthetic contact lenses:
The following procedure codes may be reimbursed for the fitting or modification of a contact lens:

<table>
<thead>
<tr>
<th>Procedure Codes</th>
</tr>
</thead>
<tbody>
<tr>
<td>V2500 V2501 V2502 V2510 V2511 V2512 V2513 V2520 V2521 V2522</td>
</tr>
<tr>
<td>V2523 V2530 V2531 V2599</td>
</tr>
</tbody>
</table>

**Contact Fitting for Corneal Bandage**

A contact lens fitting for the placement of a corneal bandage lens may be medically necessary for eye protection and pain control due to a disease process or an injury. Procedure codes 92071 and 92072 may be reimbursed for the fitting of the corneal bandage for treatment and management.

Procedure code 92071 is limited to once per eye by any provider. Modifier LT or RT must be included on the claim to identify the eye on which the service was performed. When procedure code 92071 is performed on both eyes on the same date of service, one procedure may be reimbursed at the full rate and the second procedure may be reimbursed at half rate.

Procedure code 92072 is limited to one service per lifetime by the same provider.

Procedure code 92072 will be denied if billed on the same date of service by the same provider as procedure code 92071.

**Prior Authorization Requirements**

Nonprosthetic contact lenses and corneal plano bandage lenses must be prior authorized. The following documentation must be submitted with a request for nonprosthetic contact lenses and must be signed and dated by the prescribing physician or optometrist:

- Diagnosis causing the refractive error (such as keratoconus)
- Include the current and new prescriptions supporting a change of 0.5d or more in the sphere, cylinder, or prism measurements
- Indicate which eyes to be treated
- Specify the procedure codes requested
- Include a brief statement addressing the medical necessity for vision correction by contact lens(es) and specify why eyeglasses are inappropriate or contraindicated for this client

For the contact fitting of the corneal bandage lens (procedure code 92071 or 92072), nonprosthetic contact lenses for nonemergency placement require prior authorization that must be obtained before the lenses are dispensed. Documentation submitted with the request must include the information listed above.

Nonprosthetic contact lenses for emergency placement do not require prior authorization. The emergency condition necessitating a corneal bandage must be documented on the claim.

Additional nonprosthetic contact lenses may be considered more frequently than the limitations outlined in this handbook when documentation in the client’s medical record supports medical necessity for a diopter change of 0.5d or more in the sphere, cylinder, prism measurements, or axis changes.
4.3.6.3 Dispensing Requirements

Providers must be able to dispense standard size frames at no cost to the eligible client. The following criteria must be met for the dispensed frames:

- Providers must offer each client who is 20 years of age or younger a choice of six styles in three colors for each type of frame: metal, zylonite, or combination of metal and zylonite.
- Providers must offer each client who is 21 years of age or older a choice of three styles in three colors for each type of frame: metal, zylonite, or combination of metal and zylonite.

When a client chooses eyeglass or contact lens options that are beyond program limitations, the client must acknowledge their choice and his or her liability for the cost difference by signing the Vision Care Eyeglass Patient (Medicaid Client) Certification Form.

Dispensing of contact lenses include the fabrication, ordering, adjustment, dispensing, sale, and delivery to the client of the contact lenses prescribed by and dispensed in accordance with a prescription from a licensed physician or optometrist.

Dispensing of eyeglasses includes the design, verification, fitting, adjustment, sale, and delivery to the client of (1) fabricated and finished spectacle lenses, (2) frames, or (3) other ophthalmic devices, prescribed by and dispensed in accordance with a prescription from a licensed physician or optometrist.

4.3.6.4 Repair

The eyeglass supplier is required to perform minor repairs on request (without charge) on eyeglasses that they have dispensed regardless of the client’s age. Minor repairs are those that cost $2 or less. The minor repairs are included in the reimbursement for the eyeglasses and are not reimbursed separately.

For clients who are birth through 20 years of age, repairs that cost $2 or more may be reimbursed using procedure code V2799. The following criteria apply:

- The cost of repair supplies cannot exceed the cost of replacement eyeglasses.
- All repair supplies must be new and at least equivalent to the original item.
- The provider must maintain in the client’s medical record an itemized list of repairs and the replacement cost to determine whether criteria are met for repair.

For clients who are 21 years of age and older, repair of nonprosthetic eyeglasses or contact lenses is not a benefit when the actual cost of materials exceeds $2.

The provider must make the client’s medical record available for review upon request.

4.3.6.5 Nonprosthetic Replacement

Clients who are birth through 20 years of age may obtain replacement nonprosthetic eyeglasses if the first pair is lost or destroyed. There are no limitations on the number of replacements a client who is birth through 20 years of age may receive. If the eyewear is lost or destroyed, the provider must have the client sign the Vision Care Eyeglass Patient (Medicaid Client) Certification Form. Claims for replacement lenses must be submitted with the RB modifier to ensure accurate processing. Prior authorization is not required for the replacement of nonprosthetic eyeglasses.

Replacement of eyeglasses or contact lenses is also allowed with a change in axis. A new prescription must have at least one of the following changes:

- A change of 0.50 diopters or more in any corresponding meridian.
- A cylinder axis change of at least 20 degrees for a cylinder power of 0.50-0.62 diopters.
- A cylinder axis change of at least 15 degrees for a cylinder power of 0.75-0.87 diopters.
- A cylinder axis change of at least 10 degrees for a cylinder power of 1.00-1.87 diopters.
• A cylinder axis change of at least 5 degrees for a cylinder power of 2.00 diopters or greater.

   **Note:** Replacement glasses will not be reimbursed for a cylinder power of 0.12-0.37 diopters with a change in axis.

Prior authorization is required for replacement of non-prosthetic contact lenses.

If the client is diagnosed with aphakia, procedure code 92326 may be reimbursed for the replacement of a contact lens.

### 4.3.6.6 Medicare Coverage for Nonprosthetic Eyewear

Eye examinations for the purpose of prescribing, fitting, or changing eyeglasses or contact lenses because of refractive errors are not a benefit of Medicare. These services must be filed directly to Texas Medicaid when performed for a Medicaid Qualified Medicare Beneficiary (MQMB) client. Medicare coverage is limited to eye examinations for treatment of eye disease or injury and for a diagnosis of aphakia. When performing an eye examination with refraction for an MQMB client diagnosed with aphakia or disease or injury to the eye, the following procedures must be followed:

• Procedure code 92015 must be used to bill Texas Medicaid for the refractive portion of the examination and is payable with a diagnosis of aphakia or ocular disease only.

• The medical portion of the eye examination (procedure code 92002, 92004, 92012, or 92014) is covered by Medicare and must be billed to Medicare first. Medicare forwards this portion of the examination automatically to TMHP for deductible and coinsurance payment consideration according to current guidelines.

**Refer to:** Subsection 2.7, “Medicare Crossover Claim Reimbursement” in “Section 2: Texas Medicaid Fee-for-Service Reimbursement” (Vol. 1, General Information) for more information about current coinsurance and deductible payment guidelines.

**Important:** Providers performing eye exams for refractive errors on Medicaid Qualified Medicare Beneficiary (MQMB) clients must bill TMHP. Do not send the refraction (procedure code 92015) to Medicare first. Texas Medicaid will not waive the 95-day filing deadline if the claim is billed to Medicare in error, nor will Medicare transfer the refraction to Texas Medicaid for payment.

Medicare allows payment of one pair of conventional eyewear (contact lens or glasses) for clients who have had cataract surgery with insertion of an IOL. Medicare considers the IOL the prosthetic device. Texas Medicaid providers must bill Medicare for the conventional (nonprosthetic) eyewear provided following an IOL insertion and bill Texas Medicaid for any replacements of the conventional (nonprosthetic) eyewear using the procedure codes in subsection 4.3.6, “Nonprosthetic Eyeglasses or Contact Lenses” in this handbook.

### 4.3.7 Vision Services for Prosthetic Eyewear

Prosthetic eyeglasses or contact lenses are lenses that replace the eye’s organic lens when it is absent due to congenital or acquired aphakia. Aphakia may be the result of a congenital abnormality or defect or an acquired condition as a result of trauma or cataract removal without intraocular lens (IOL) insertion.

Prosthetic eyeglasses or contact lenses may be provided based on medical necessity. Eye examinations and prosthetic eyewear may be reimbursed as follows:

• Eye examinations for aphakia (including congenital aphakia) and disease or injury to the eye may be reimbursed as often as medically necessary.

• One pair of permanent prosthetic eyeglasses or contact lenses is a benefit during a client’s lifetime.
• Replacement of prosthetic eyeglasses or contact lenses may be reimbursed for clients of any age due to loss or destruction of the eyewear or due to a significant change in visual acuity with a diopter change of 0.5d or more in the sphere, cylinder, prism measurements, or axis changes. The provider must maintain in the client’s medical record documentation that supports the medical necessity for the replacement eyeglasses or contact lenses.

Prosthetic contact lenses may be provided, with prior authorization, for clients of any age with congenital or acquired aphakia.

Note: Fitting services are included in the reimbursement for prosthetic eyeglasses or contact lenses.

Providers must use modifier VP with a diagnosis code of aphakia when billing for prosthetic eyeglasses or contact lenses.

Refer to: Subsection 4.3.6, “Nonprosthetic Eyeglasses or Contact Lenses” in this handbook for the eyeglass lens, frame, and contact lens procedure codes and dispensing requirements that apply to prosthetic and nonprosthetic eyewear.

The date of cataract surgery is not required on the claim for conventional eyeglasses or contact lenses after surgery, or prosthetic eyeglasses or contact lenses after cataract surgery without IOL insertion.

4.3.7.1 Eyeglasses or Contact Lenses Following Cataract Surgery
Temporary eyeglasses or contact lenses after cataract surgery may be reimbursed when they are billed with the appropriate lens and frame procedure codes and diagnosis code Z961.

Temporary eyeglasses or contact lenses may be reimbursed for up to 4 months after surgery until the client is ready for conventional eyeglasses or contact lenses, when it is medically necessary. The date of surgery is used to determine the convalescence period for temporary eyeglasses or contact lenses. Temporary eyeglasses or contact lenses will be denied if they are dispensed more than 4 months after the date of surgery.

Temporary eyeglasses or contact lenses may be reimbursed as often as is medically necessary during the postsurgical convalescence period.

4.3.7.2 Repair
The eyeglass supplier is required to perform minor repairs on request (without charge) on eyeglasses that they have dispensed regardless of the client’s age. Minor repairs are those that cost $2 or less. The minor repairs are included in the reimbursement for the eyeglasses and are not reimbursed separately.

Repairs that cost $2 or more may be reimbursed using procedure code V2799. The following criteria apply:

• The cost of repair supplies cannot exceed the cost of replacement eyeglasses.
• All repair supplies must be new and at least equivalent to the original item.
• The provider must maintain in the client’s medical record an itemized list of repairs and the replacement cost to determine whether criteria are met for repair.

The provider must make the client’s medical record available for review upon request.

4.3.7.3 Prosthetic Replacement
Replacement prosthetic eyeglasses or contact lenses may be reimbursed as often as is medically necessary if the replacement is due to loss, destruction, or a significant change in visual acuity.

Replacement of eyeglasses or contact lenses is also allowed with a change in axis. A new prescription must have at least one of the following changes:

• A change of 0.50 diopters or more in any corresponding meridian.
• A cylinder axis change of at least 20 degrees for a cylinder power of 0.50-0.62 diopters.
• A cylinder axis change of at least 15 degrees for a cylinder power of 0.75-0.87 diopters.
• A cylinder axis change of at least 10 degrees for a cylinder power of 1.00-1.87 diopters.
• A cylinder axis change of at least 5 degrees for a cylinder power of 2.00 diopters or greater.

**Note:** Replacement glasses will not be reimbursed for a cylinder power of 0.12-0.37 diopters with a change in axis.

The appropriate eyeglass and frame or contact lens procedure codes must be billed with modifier RB to indicate replacement.

**Refer to:** Subsection 4.3.6, “Nonprosthetic Eyeglasses or Contact Lenses” in this handbook for the eyeglass lens, frame, and contact lens procedure codes and dispensing requirements that apply to nonprosthetic eyewear.

### 4.3.7.4 Intraocular Lens (IOL) and Additional Eyewear

Intraocular lenses are benefits of Texas Medicaid. If conventional eyewear is medically necessary in addition to the IOL, the IOL is considered the prosthetic device, and the eyewear and any replacements are considered nonprosthetic.

**Refer to:** Subsection 9.2.46.4, “Intraocular Lens (IOL)” in the Medical and Nursing Specialists, Physicians, and Physician Assistants Handbook (Vol. 2, Provider Handbooks) for more information about IOL benefits.

Subsection 4.3.6, “Nonprosthetic Eyeglasses or Contact Lenses” in this handbook for more information about nonprosthetic eyewear.

### 4.3.7.5 Artificial Eyes

For clients who are birth through 20 years of age, artificial eyes may be considered under CCP.

### 4.3.7.6 Ultraviolet (U-V) Protection

Procedure code V2755 may be reimbursed for U-V protection when billed with one of the following diagnosis codes:

<table>
<thead>
<tr>
<th>Diagnosis Codes</th>
</tr>
</thead>
<tbody>
<tr>
<td>H2701</td>
</tr>
<tr>
<td>H27123</td>
</tr>
</tbody>
</table>

UV lens procedure code V2755 will be denied when billed with the same date of service by the same provider as polycarbonate lens procedure code V2784.

UV and polycarbonate lens procedure codes are subject to CMS NCCI relationships.

**Refer to:** The CMS NCCI web page for the published correct coding guidelines and specific applicable code combinations.

### 4.3.8 Surgical Vision Services

**Refer to:** Subsection 7.29, “Fluocinolone Acetonide (Retisert)” in the Outpatient Drug Services Handbook (Vol. 2, Provider Handbooks) for more information about fluocinolone acetonide benefits.

4.3.9 **Prior Authorization**

Prior authorization is required for the following:

- Orthoptic or pleoptic training beyond the maximum limits outlined in subsection 4.3.5.8, “Orthoptic or Pleoptic Training” in this handbook
- The unlisted ultrasound procedure
- All contact lenses, except corneal bandage lens(es) for emergency placement

A completed Special Medical Prior Authorization (SMPA) Request Form must be submitted by fax or mail to the Special Medical Prior Authorization department. The form must be signed and dated by a physician familiar with the client before requesting prior authorization. The completed Special Medical Prior Authorization (SMPA) Request Form must include the procedure codes and numerical quantities for services requested. The completed, signed, and dated Special Medical Prior Authorization (SMPA) Request Form must be maintained by the provider and the prescribing physician in the client’s medical record.

Prior authorization requests may be submitted to the TMHP Prior Authorization Department by mail, fax, or the electronic portal. Prescribing or ordering providers, dispensing providers, clients’ responsible adults, and clients may sign prior authorization forms and supporting documentation using electronic or wet signatures.

To avoid unnecessary denials, the physician must provide correct and complete information, including documentation for medical necessity of the vision service requested. The physician must maintain documentation of medical necessity in the client’s medical record. The requesting provider may be asked for additional information to clarify or complete a request for the vision service.

4.4 **Documentation Requirements**

All services require documentation to support the medical necessity of the service rendered, including vision services. Vision services are subject to retrospective review and recoupment if documentation does not support the service billed.

The client must sign and date the Vision Care Eyeglass Patient (Medicaid Client) Certification Form, and the provider must retain it in the provider’s records.

When a client chooses an eyeglasses or contact lens option beyond the program limitations, or nonprosthetic eyeglasses or contact lenses are replaced because of loss or destruction the client must acknowledge their choice and liability for the cost difference by signing the Vision Care Eyeglass Patient (Medicaid Client) Certification Form and retain it in the provider’s records.

The current and previous prescriptions must be documented in the client’s medical record.

The provider must make the client’s medical record available for review upon request by the following:

- HHSC
- Office of the Attorney General
- TMHP

4.5 **Claims Filing and Reimbursement**

4.5.1 **Claims Filing**

Vision care service claims must be submitted to TMHP in an approved electronic format or on a CMS-1500 paper claim form. Providers may purchase CMS-1500 paper claim forms from the vendor of their choice. TMHP does not supply the forms. When completing a CMS-1500 paper claim form, all required information must be included on the claim, as information is not keyed from attachments. Superbills, or itemized statements, are not accepted as claim supplements.
When submitting the client’s old and new prescriptions to show an axis change or a diopter change of 0.5 or more, enter the new prescription in Block 24D, line 5, and the old prescription in Block 24D, line 6 of the CMS-1500 paper claim form.

Claims for eye examination services require a diagnosis. Claims for eye examinations that lack a diagnosis are listed as an incomplete claim on the Remittance and Status (R&S) report and must be resubmitted for payment consideration. Electronic claims that lack a diagnosis will be rejected. A letter with the reason for rejection and instructions for resubmission will be mailed the following business day.

When the eye exam limitation is exceeded for clients who are 20 years of age and younger, identify one of the following situations in Block 19 of the CMS-1500 paper claim form:

- A school nurse, teacher, or parent requests the eye examination.
- The eye examination is medically necessary.

### 4.5.2 Reimbursement

Providers must reflect the highest level of specificity for vision related diagnoses on claims or other documentation. Professional services by an optometrist for contact lenses and prosthetic eyewear are reimbursed in accordance with 1 TAC §355.8001 and §355.8085.

FQHCs are paid an all-inclusive rate per visit for payable services in accordance with 1 TAC, §355.8261.

Suppliers of nonprosthetic lenses and frames are reimbursed the lesser of their billed amount or of the established maximum allowable fee in accordance with 1 TAC, §355.8001. See the OFL or the applicable fee schedule on the TMHP website at www.tmhp.com.

**Refer to:** Subsection 2.2, “Fee-for-Service Reimbursement Methodology” in “Section 2: Texas Medicaid Fee-for-Service Reimbursement” (Vol. 1, General Information) for more information about reimbursement.

**Vision Services** on the TMHP website at www.tmhp.com for a claim form example.

The nonsurgical vision procedure codes included in this handbook may be subject to the CMS NCCI relationships.

**Refer to:** The CMS website at www.cms.gov for more information about NCCI relationships.

Texas Medicaid implemented mandated rate reductions for certain services. The OFL and static fee schedules include a column titled “Adjusted Fee” to display the individual fees with all mandated percentage reductions applied. Additional information about rate changes is available on the TMHP website at www.tmhp.com/resources/rate-and-code-updates/rate-changes.

### 4.5.2.1 NCCI and MUE Guidelines

The HCPCS and CPT codes included in the Texas Medicaid Provider Procedures Manual are subject to NCCI relationships, which supersede any exceptions to NCCI code relationships that may be noted in the manuals. Providers should refer to the CMS NCCI web page for correct coding guidelines and specific applicable code combinations.

In instances when Texas Medicaid limitations are more restrictive than NCCI MUE guidance, Texas Medicaid limitations prevail.

If applicable and consistent with CMS billing guidelines, procedure codes must be billed with modifier LT (left side) or RT (right side) to identify the eye on which the service was performed.
5 Claims Resources

Refer to the following sections and forms when filing claims:

<table>
<thead>
<tr>
<th>Resource</th>
<th>Location</th>
</tr>
</thead>
<tbody>
<tr>
<td>Acronym Dictionary</td>
<td>“Appendix C: Acronym Dictionary” (Vol. 1, General Information)</td>
</tr>
<tr>
<td>Automated Inquiry System (AIS)</td>
<td>Subsection A.10, “TMHP Telephone and Fax Communication” in “Appendix A: State, Federal, and TMHP Contact Information” (Vol. 1, General Information)</td>
</tr>
<tr>
<td>CMS-1500 Paper Claim Filing Instructions</td>
<td>Subsection 6.5, “CMS-1500 Paper Claim Filing Instructions” in “Section 6: Claims Filing” (Vol. 1, General Information)</td>
</tr>
<tr>
<td>State, federal, and TMHP contact information</td>
<td>“Appendix A: State, Federal, and TMHP Contact Information” (Vol. 1, General Information)</td>
</tr>
<tr>
<td>TMHP electronic claims submission information</td>
<td>Subsection 6.2, “TMHP Electronic Claims Submission” in “Section 6: Claims Filing” (Vol. 1, General Information)</td>
</tr>
<tr>
<td>TMHP Electronic Data Interchange (EDI)</td>
<td>“Section 3: TMHP Electronic Data Interchange (EDI)” (Vol. 1, General Information)</td>
</tr>
</tbody>
</table>

6 Contact TMHP

The TMHP Contact Center at 1-800-925-9126 is available Monday through Friday from 7 a.m. to 7 p.m., Central Time.

7 Forms

The following linked forms can also be found on the Forms page of the Provider section of the TMHP website at www.tmhp.com:

<table>
<thead>
<tr>
<th>Forms</th>
</tr>
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<tbody>
<tr>
<td>Hearing Evaluation, Fitting, and Dispensing Report (Form 3503)</td>
</tr>
<tr>
<td>Physician’s Examination Report</td>
</tr>
<tr>
<td>Vision Care Eyeglass Patient (Medicaid Client) Certification Form</td>
</tr>
<tr>
<td>Vision Care Eyeglass Patient (Medicaid Client) Certification Form (Spanish)</td>
</tr>
</tbody>
</table>

8 Claim Form Examples

The following linked claim form examples can also be found on the Claim Form Examples page of the Provider section of the TMHP website at www.tmhp.com:

<table>
<thead>
<tr>
<th>Claim Form Examples</th>
</tr>
</thead>
<tbody>
<tr>
<td>Hearing Aid Assessments</td>
</tr>
<tr>
<td>Vision Services</td>
</tr>
</tbody>
</table>