

\*\*\*All PA forms may be found by accessing <https://tnm.providerportal.sxc.com/rxclaim/TNM/PAs.htm>\*\*\*

If the following information is not complete, correct, or legible the PA process can be delayed. **Use one form per member please.**

**Member Information**

 Last Name 

 First Name 

 ID Number 

 Date of Birth  -  - 
**Prescriber Information**

 Last Name 

 First Name 

 NPI# 

 DEA# 

 Phone  -  - 

 Fax  -  - 

### REQUESTED ANTIHYPERKINESIS AGENT

modafinil     Nuvigil     Provigil

Dose \_\_\_\_\_ Directions \_\_\_\_\_ Qty \_\_\_\_\_ Duration of Therapy \_\_\_\_\_

 Request to Backdate PA?  Yes  No

If Yes, Requested PA Start Date \_\_\_\_\_

**Clinical Criteria Documentation**

\*\*\*\*Do not include documentation that is not requested on this form\*\*\*\*

1. What is the diagnosis for this medication?
  - Narcolepsy       Obstructive sleep apnea/hypopnea syndrome
  - ADD/ADHD       Shift work sleep disorder       Other \_\_\_\_\_
2. Has the recipient failed an adequate trial of any other stimulant agent(s)?  Yes (please list)     No
 

Drug 1: \_\_\_\_\_ Strength: \_\_\_\_\_ Quantity: \_\_\_\_\_ Length of trial: \_\_\_\_\_

Reason for discontinuation of the drug: \_\_\_\_\_

Drug 2: \_\_\_\_\_ Strength: \_\_\_\_\_ Quantity: \_\_\_\_\_ Length of trial: \_\_\_\_\_

Reason for discontinuation of the drug: \_\_\_\_\_

Drug 3: \_\_\_\_\_ Strength: \_\_\_\_\_ Quantity: \_\_\_\_\_ Length of trial: \_\_\_\_\_

Reason for discontinuation of the drug: \_\_\_\_\_
3. Has the recipient experienced an adverse event, or been intolerant to, a preferred stimulant?  Yes  No
 

If yes, please list the drug (or drugs) and describe the adverse event or intolerance: \_\_\_\_\_
4. Is the patient currently taking the requested medication?  Yes  No
 

If yes, how long has the recipient been taking the medication? \_\_\_\_\_

How has medication been supplied (other insurance, samples provided, patient discharged from hospital on the medication, etc.)? \_\_\_\_\_
5. If request is for Nuvigil, has the patient tried and failed Provigil?  Yes  No Length of trial: \_\_\_\_\_
 

If no, what is the reason the patient cannot take Provigil? \_\_\_\_\_

Complete this section only if diagnosis is obstructive sleep apnea/hypopnea syndrome.

6. Has the recipient had a sleep study?  Yes  No Date of study: \_\_\_\_\_
7. Does the provider have evidence of documented compliance with a BiPAP or CPAP device?  Yes  No
 

Total length of therapy? \_\_\_\_\_ If no use, why? \_\_\_\_\_

----- continued on next page -----

**TennCare Prior Authorization Form: Provigil®/Nuvigil™**  
**- Page 2 -**

Patient Name: \_\_\_\_\_ DOB \_\_\_\_\_

---

Complete this section only if diagnosis is shift work sleep disorder.

8. Does the patient work a minimum of 6 hours work between the hours of 10 pm and 8 am?  Yes  No

Please note any other information pertinent to this PA request: \_\_\_\_\_  
\_\_\_\_\_  
\_\_\_\_\_

**Prescriber Signature (REQUIRED):** \_\_\_\_\_ **Date:** \_\_\_\_\_  
*(By signature, the physician confirms the above information is accurate and verifiable by patient records.)*

**Fax This Form to: 866-434-5523**  
**Mail requests to: Catamaran PA Department, P.O. Box 3214, Lisle IL 60532-8214**  
**Telephone 866-434-5524**

**Catamaran will provide a response within 24 hours day upon receipt.**

<p>This facsimile transmission contains legally privileged and confidential information intended for the parties identified below. If you have received this transmission in error, please immediately notify us by telephone and return the original message to P.O. Box 3214; Lisle, IL 60532-8214. Distribution, reproduction or any other use of this transmission by any party other than the intended recipient is strictly prohibited.</p>
---