

HC NIRF 01 – V11 Date issued: 20/03/2020

NATIONAL INCIDENT REPORT FORM (NIRF)

NIRF - 01 PERSON

NIMS record Number:	(

	NINS record number:
dent: An event or circumstance which could have, or did lead to unintended and / or unnecessary SECTION A: GENERAL INCIDENT DETAILS Date of incident Time of incident H H M Use 24 hour clock Location E.g. Hospital, Health Centre, Residential Centre etc. Specific Location E.g. Ward, Clients home etc. Offsite? Description of incident:	harm. Please complete this form to the best of your knowledge at the time of reporting the incident. SECTION B: PERSON AFFECTED DETAILS First name Surname Date of birth Female Male
Division (tick one only ✓) Acute Hospital Social Care Health and Wellbeing Primary Care Mental Health Ambulance Service National Corporate Services (staff only)	Who was involved? (tick one only ✓) Service user – (Resident/Patient/Client) Go to section C Staff member – Go to section D Agency / Panel staff – Go to section D Member of public-Proceed to section F Volunteer – Go to section D External Contractor – Go to section E Student – Go to section D
SECTION C: SERVICE USER DETAILS ONLY Healthcare Record No Lead Clinician This incident involved (tick one only ✓) Neonatal Specialties Paediatric Specialties Adolescent Specialties Adult Specialties	SECTION D: STAFF MEMBER / AGENCY / PANEL STAFF / STUDENT / VOLUNTEER DETAILS ONLY Category of person Employee no. Date absence commenced (if known) Date returned to work (if known) Note: For employee incidents reportable to HSA that result in an absence from duty for more than three consecutive days, excluding the day of the accident, the date absence commenced and the date employee returned to work should be recorded on the NIMS
Older Person Specialties E.g. Antenatal, Audiology, Radiotherapy, Intellectual Disability, Psychology	SECTION E: EXTERNAL CONTRACTOR DETAILS ONLY Company Name Company no.

SECTION F: WHAT WAS THE OUTCOME AT THE TIME OF THE INCIDENT? ✓ Outcome Body Part Affected				
	irly given wrong drug			
	ng drug given but no harm	Category 3		
Injury not requirin	g first aid			
Injury or illness, re	quiring first aid		\)
Injury requiring mo		Category 2		
_	ty / Incapacity (incl. psychosocial)	cutegory 2		
			F - 4 C	the Land Od on District and
Permanent incapa	city (incl. Psychosocial)	Category 1	E.g. Arm, Sp	oine, Lung, Other Physiological
☐ Death				
SECTION G: TYPE C	OF INJURY (tick one only √)			
	☐ Apgar score <5@ 1 min &/or;	☐ HIE Grade 2 - H		Nerve Injury - face
	7@5mins &/or pH ≤ 7.0	Encephalopath HIE Grade 3 - H		Other unexpected deteriorationStillbirth
	☐ Aspiration☐ Cerebral irritability / neonatal	Encephalopath	• •	Sub-galeal / sub-aponeurotic
Birth Specific Injury	seizure	Hypoglycaemia	•	haemorrhage
(Baby)	HIE - Hypoxic Ischaemic	☐ Kernicterus	[Unknown
	Encephalopathy with	Neonatal death	Ĺ	Other
	Hypoglycaemia		rachial plexus (incl.	
	☐ HIE Grade 1 - Hypoxic Ischaemi	c Erbs Palsy)		
	Encephalopathy		d .	
Dinth Cassifis Injum.	☐ Death	Perineal tearPost-Partum Hand		UnknownUterine rupture
Birth Specific Injury (Mother)	☐ Hysterectomy (Perinatal)☐ Incontinence (faecal)	Rhesus iso-imm		Other
(Wother)	☐ Incontinence (urinary)	Incontinence (f		
	☐ Excessive Bleeding		emolytic transfusion	Non-immunological haemolysis
Blood Specific Injury	☐ Fainting	reaction	·	Other
	Immunological haemolysis			
	Asbestosis	Hepatitis	Ĺ	Unknown
Diagnosed Disease	Cancer	☐ HIV	L	Dermatitis
Disorder or Cond.	Acute Radiation SyndromeNarcolepsy/Cateplexy	BrucellosisLegionnaires	L	☐ TB☐ Pleural Plaques
	□ Naicolepsy/Catepiexy	Legioinianes		Uther
	☐ Clostridium Difficle	☐ Hepatitis	-	∨RE
Diagnosed Infection	Covid-19	☐ MRSA		URSA VRE
Diagnosca iniconon	☐ CPE	□ Norovirus	i	Other
	□ ESBL	Unknown		
	Allergic Reaction (incl. anaphyla		n / Graze / scratch	Malaise / Nausea
	☐ Brain Injury / Concussion	Death		Nerve injury / Loss of Function
Canaval Initiation	☐ Burn / scald / corrosion	☐ Dental injury & ☐ Deterioration	or loss	Puncture / bite
General Injuries	☐ Choking / asphyxia☐ Circulatory / volume depletion			Rash / irritation Unknown
	☐ Circulatory / volume depletion	Blister		Other
	☐ Pain/Discomfort			
Hearing / Sight Injury	☐ Hearing Impairment / loss	☐ Tinnitus	Ĺ	Other
ricaring / Signt injury	☐ Sight Impairment / loss	Unknown		
Misdiagnosis	Cancer	☐ Infection ☐ Unknown	[Other
	☐ Fracture			Swalling / Inflammation
	☐ Amputation☐ Bruising	☐ Fracture☐ Repetitive Strain	n Injury (RSI)	Swelling / InflammationUnknown
	☐ Crushing	☐ Slipped / Prola		□ Whiplash
	Dental Fracture / Tooth loss	Sprain / Strain		Other
Musculoskeletal	Dislocation	☐ Soft tissue injur	у	
/ Soft Tissue	P. Ulcer Stage 1: Intact skin with	h non-blanchable rednes	s over bony prominence	
	P. Ulcer Stage 2: Part thickness			
	P. Ulcer Stage 3: Full thickness t			
	P. Ulcer Stage 4: Full thickness t			
Devenuellere	Additional / Further Surgery	Loss of Wages ,		Unknown
Personal Loss	☐ Limb Deformity☐ Defamation of Character	Business Loss of Consort		Organ Retention Other
	☐ Damage to organ / body part	Loss of organ / b		
Surgery Specific	☐ Dental Damage / Loss	☐ Nerve injury / Lo		terioration
Injury	☐ Foreign body left in situ	Function		
	Unknown	Inadequate and	esthesia	Other
Traumatic/Emotional	☐ Anxiety / Trauma	Stress		☐ Worried Well
,	I DISII	LIDINOWN		Lithor

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SEC	SECTION H WHAT TYPE OF HAZARD DID THIS INCIDENT RELATE TO? (Tick one option from Steps 1, 2, 3 & 4)					
	Step 1.	Step 2.	Step 3.	Step 4.		
ē	☐ Birth Specific Procedures ☐ Clinical Procedures	□ Caesarean Section (Elective) □ Caesarean Section (Emergency) □ Instrumental Delivery (Forceps) □ Instrumental Delivery (Vacuum) □ Instrumental Delivery (Multiple Instruments) □ Non Instrumental Delivery	□ Communication / Consent □ Diagnosis / Assessment □ Documentation / Records □ Equipment □ General Care / Management □ Procedure / Treatment / Intervention □ Screening / Prevention □ Specimens / Results □ Tests / Investigations □ Unknown □ Other	□ Adverse Effect □ Failure / Malfunction □ Foreign Body left in Situ □ Inappropriate for Task / Wrong device □ Incomplete / Inadequate □ Lack of Availability □ Not performed when indicated / Delay □ Pre Existing Medical Condition □ Shoulder Dystocia □ Unavailable / Mislabelled / Lost □ Wrong Body Part / Site / Side □ Wrong Patient □ Wrong Process / Treatment / Procedure □ Other		
	☐ Medication	Route of administration Oral Intravenous Sub Cutaneous Intra Muscular Topical Rectal Inhalation Other / Unknown What medication was involve		□ Adverse Drug Reaction □ Contra-indicated □ Drug Interaction □ Failure / Malfunction of equipment □ Incomplete / Inadequate □ Not preformed when indicated / delayed □ Omitted/Delayed Dose □ Wrong Dose / Strength □ Wrong Drug □ Wrong Formulation / Route □ Wrong Frequency		
Clinical Care		Medication One		☐ Wrong Label / Instructions☐ Wrong Patient☐ Wrong Quantity / Duration		
Clini	□ Nutrition	Medication Two □ Parenteral □ Enteral □ Special Diet □ General Diet □ Other	 ☐ Communication / Consent ☐ Prescribing / Requesting ☐ Preparation / Dispensing ☐ Administration ☐ Storage 	 □ Adverse Effect □ Incomplete / Inadequate □ Not performed when indicated / Delay □ Wrong Consistency □ Wrong Diet / Wrong Blood Product 		
	□ Blood / Blood Product		 □ Documentation / Records □ Equipment □ Supply / Ordering / Transport □ Presentation / Packaging □ Transfusing blood □ Other 	 □ Wrong Process / Treatment / Procedure □ Wrong Patient □ Lack of Availability □ Wrong dispensing label / instructions □ Inappropriate for task / Wrong device □ Other 		
	☐ Diagnostic Radiology (DR) & Nuclear Medicine (NM)	 □ Checking Patient ID procedure □ Clinical Details on Referral □ Communication / Consent 	□ Diagnostic Exposure > intended □ X-ray Over Exposure □ Wrong body part / side □ Dose to comforters / carers □ Wrong Patient □ Inadvertent dose to foetus □ Total dose or Volume Variation □ Dose (NM) or Volume Variation			
	☐ Radiotherapy	 □ Documentation / Records □ Equipment □ Performing procedure □ Pregnancy Status □ Unknown 	(1 fraction) Wrong Drug Wrong Dose Wrong Process / Treatment / Intervention Failure / Malfunction Inadvertent deterministic effects	>20%		
Bio Hazards	☐ Biological Hazards / Acquired Infections	☐ Bacteria☐ Fungus / Mould☐ Prion☐ Virus☐ Organism Unknown	☐ Please specify, if known: ————————————————————————————————————	□ Exposure to Bite (Human) □ Exposure to Bite (Insect / Animal) □ Exposure to Bodily Fluids □ Exposure to Ingestion/Food/Water □ Exposure to Needle Stick □ Exposure to Skin Contact □ Inhalation/Airborne □ Equipment, Implements, Facilities, □ Sharps (Non Needle) □ Unknown □ Other		

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SECTION H CNTD: WHAT TYPE OF HAZARD DID THIS INCIDENT RELATE TO? (Tick one option from Steps 1, 2 & 3)				
	Step 1.	Step 2.	Step 3.	
Behavioural Hazards	└ Self-Injurious Behaviour	☐ Intentional ☐ Unintentional		
	☐ Violence, Harassment and Aggression	☐ By a Family Member / Relative	 □ Aggressive towards inanimate object □ Discrimination/Prejudice/Racial □ Intimidation / Threat □ Neglect □ Non-Compliant / Obstructive / Rude 	
	☐ Child Abuse	By a Peer / Student By a Prisoner By a Service User	 ☐ Non-Compliant / Obstructive / Rude ☐ Physical Assault / Abuse ☐ Physical Harassment ☐ Sexual Assault / Abuse ☐ Sexual Harassment 	
	□ Adult Abuse	□ By a Staff Member	 Unintentional Aggressive Behaviour Bullying Verbal Assault / Abuse Verbal Harassment Other 	
Physical Hazards	□ Slip / Trip / Fall	☐ From Height ☐ From Equipment / Furniture ☐ Same Level / Ground ☐ On Stairs ☐ On Steps ☐ Other	 Unknown Pre Existing Medical Condition Inadequate supervision gen health / post op Obstruction / protruding object Surface contaminants Rough terrain / irregular surface Inappropriate equipment use Failure / malfunction of equipment Horseplay Physical training / sport Weather Condition Inadequate Lighting / design Other 	
	☐ Non Mechanical (Incl. Person / Animal)	 □ Object / Tools (Non Sharps) □ Sharps (Non Needle) □ Other □ Person □ Manual Handling 	 ☐ Human Use / Error ☐ Obstruction / Protruding Object ☐ Physical Training / Sport ☐ Defective Equipment 	
	Ergonomics(Incl. manual / people handling)	☐ Other ☐ Patient Handling ☐ Restraint / Intervention	 ☐ Unsafe / Inappropriate system ☐ Unknown ☐ Task 	
	☐ Mechanical Components	 □ Catering equipment □ Door / Gate / Barrier □ Healthcare Equipment □ Lifting Equipment / Accessories □ Office / Business equipment 		
	☐ Temperature (Excluding Fire)	☐ Hot ☐ Cold	□ Liquid / Food / Steam □ Equipment / Utensils □ Atmosphere / Environment	
	☐ Fire☐ VibrationElectrical☐ Noise☐ Radiation	☐ Please Specify ————————————————————————————————————	 □ Defective Equipment □ Human Use / Error □ Unknown □ Unsafe System □ Explosion □ Exposure □ Electrical Wiring / installation 	

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SECTION H CNTD: WHAT TYPE OF HAZARD DID THIS INCIDENT RELATE TO? (Tick one option from Steps 1, 2, & 3)				
	Step 1.		Step 2.	Step 3.
Chemical Hazards	 □ Acid / Alkaline □ Agri Chemicals □ Gas □ Other Chemical Products □ Particulates □ Petroleum / Synthetic Oil Based Products □ Sanitation / Cleaning Chemicals □ Toxic Metals 	Animal Remedy Arsenic Asbestos Bleach Cadmium Carbon Dioxide Carbon Monoxide Chemical Fertilizer Crystalline Silica Detergent Diesel / Kerosene Disinfectant Drain / Oven Cleaner Drugs Fungicide Glue / Adhesive Grease Herbicide Hydrochloric Acid	Insecticide Lead Metallic Dust Motor / Gear / Hydraulic Oil Natural Gas Organic Dust Paint / Paint Product Petrol Polish Radon Rodenticide Soap Sodium Hydroxide Solvents Spent / Used Oil Product Sulphuric Acid Wrong Patient Other	☐ Lack of Supervision☐ Unknown☐ Human / User Error☐ Unsafe System
SEC	TION I: IMMEDIATE ACTIO	NS TAKEN		
-				
otherv	TION J: REPORTED BY: person is name		SECTION K: WITNESS DETAILS	(Name, Contact No. etc.)
Surn	ame			
Date	notified DDM	MYYYY		
	gory of person $E.g. Nurse, C$ I system	Catering Staff, Cleaner		
refe	rence no.			
Date	orter Signature	MVVVV		
	tact Details			

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SECTION L: TO BE COMPLETED BY LINE/DEPARTMENT MANAGER				
Has open disclosure happened? (tick one only ✓)	□ No			
If No, please specify:				
CATEGORY 1 INCIDENTS ONLY				
SAO Name [Block Capitals]:	Date notified to SAO:	DDMMYYYY		
SAO Email and Contact Details:				
Is there a requirement to report this incident to any external regulators/agencies/insurers (other than the State Claims Agency)?				
If Yes: Name regulator(s)/agency(ies) reported/notified to:		Date Notified:		
1		DDMMYYYY		
2		DDMMYYYY		
3		DDMMYYYY		
Line/Department Manager name [Block Capitals]:	Title:			
Signature of Line/Department Manager:	Date:	DDMMYYYY		
SECTION M: TO BE COMPLETED BY QUALITY AND PATIENT SAFETY C	OFFICE			
Is this incident a Serious Reportable Event (SRE)? (tick one only ✓)	□ No			
QPS Advisor Name [Block Capitals]:				
Signature of QPS Advisor:	Date:	DDMMYYYY		

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