

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

NATIONAL INCIDENT REPORT FORM (NIRF) NIRF - 01 PERSON

NIMS record Number:

Incident: An event or circumstance which could have, or did lead to unintended and / or unnecessary harm. Please complete this form to the best of your knowledge at the time of reporting the incident.

SECTION A: GENERAL INCIDENT DETAILS	SECTION B: PERSON AFFECTED DETAILS			
Date of incident DDMMYYYY	First name			
Time of incident H H M M Use 24 hour clock	Surname			
Location E.g. Hospital, Health Centre, Residential Centre etc.	Date of birth DDMMYYYY			
Specific Location E.g. Ward, Clients home etc. Offsite?	Female Male			
Description of incident:				
Division (tick one only ✓) Acute Hospital Social Care	Who was involved? (tick one only ✓) Service user – (Resident/Patient/Client) Go to section C			
Health and Wellbeing	Staff member – Go to section D Agency / Panel staff – Go to section D			
Primary Care	Member of public – Proceed to section F			
Mental Health	Volunteer – Go to section D			
Ambulance Service	External Contractor – Go to section E			
National Corporate Services (staff only)	Student – Go to section D			
SECTION C: SERVICE USER DETAILS ONLY	SECTION D: STAFF MEMBER / AGENCY / PANEL STAFF / STUDENT / VOLUNTEER DETAILS ONLY			
Healthcare Record No	Category of person			
Lead Clinician	Employee no.			
This incident involved (tick one only ✓)	Date absence commenced (if known)			
Neonatal Specialties	Date returned to work			
Paediatric Specialties	(if known) Note: For employee incidents reportable to HSA that result in an absence from duty for more than three consecutive days,			
Adolescent Specialties	Work days lost Work days lost Work days lost			
Adult Specialties	the NIMS			
Older Person Specialties	SECTION E: EXTERNAL CONTRACTOR DETAILS ONLY			
Incident Occurred under (Service / Specialty) E.g. Antenatal, Audiology, Radiotherapy, Intellectual Disability, Psychology	Company name Company no.			

SECTION F: WHAT WAS THE OUTCOME AT THE TIME OF THE INCIDENT?				
✓ Outcome			Body Part Affected	
	arly given wrong drug			
occurred	ng drug given but no harm	Category 3		
☐ Injury not requirin	g first aid			
Injury or illness, re	_		E a Aven Sui	no Lynn Other Physiological
	. •	Catagomy 3	E.g. Arm, Spu	ne, Lung, Other Physiological
Injury requiring m		Category 2	\)
	ty / Incapacity (incl. psychosocial)			
Permanent Incapa	city (incl. Psychosocial)	Category 1		
☐ Death				
SECTION G: TYPE O	F INJURY (tick one only ✓)			
SECTION G. TIPE C	☐ Apgar score <5@ 1 min &/or;	HIE Grade 2 - H	ypoxic Ischaemic	Nerve Injury - face
	7@5mins &/or pH ≤ 7.0	Encephalopath		Other unexpected deterioration
	☐ Aspiration	☐ HIE Grade 3 - H	ypoxic Ischaemic \Box	Stillbirth
	Cerebral irritability / neonatal	Encephalopath		Sub-galeal / sub-aponeurotic
Birth Specific Injury	seizure	Hypoglycaemia		haemorrhage
(Baby)	HIE - Hypoxic Ischaemic	☐ Kernicterus		Unknown
	Encephalopathy with Hypoglycaemia	☐ Neonatal death	orachial plexus (incl.	Other
	☐ HIE Grade 1 - Hypoxic Ischaemid		racinai piexus (inci.	
	Encephalopathy	2.25. 3.577		
	□ Death	Perineal tear		Unknown
Birth Specific Injury	Hysterectomy (Perinatal)	Post-Partum Ha		Uterine rupture
(Mother)	☐ Incontinence (faecal)	Rhesus iso-imm	unisation	Other
	☐ Incontinence (urinary)		aecal & urinary)	
Blood Specific Injury	Excessive BleedingFainting	☐ Febrile non-hae reaction	emolytic transfusion \Box	
ыоой эреспіс піјигу	Immunological haemolysis	reaction		Other
	☐ Asbestosis	☐ Hepatitis		Unknown
	Cancer	☐ HIV		Dermatitis
Diagnosed Disease	Acute Radiation Syndrome	Brucellosis		ТВ
Disorder or Cond.	□ Narcolepsy/Cateplexy	Legionnaires		Pleural Plaques
			L	Other
	Clostridium Difficle	Hepatitis		VRE
Diagnosed Infection	☐ COVID-19	☐ MRSA		VRSA
	☐ CPE			Other
	☐ Allergic Reaction (incl. anaphyla		n / Graze / scratch	Malaise / Nausea
	☐ Brain Injury / Concussion	Death		Nerve injury / Loss of Function
	☐ Burn / scald / corrosion	☐ Dental injury &	/or loss	Puncture / bite
General Injuries	☐ Choking / asphyxia	Deterioration		Rash / irritation
	☐ Circulatory / volume depletion	Haemorrhage		Unknown
	☐ Circulatory / volume overload	Blister		Other
	Pain/Discomfort	Tinnitus	П	Othor
Hearing / Sight Injury	Hearing Impairment / lossSight Impairment / loss	☐ Tinnitus ☐ Unknown		Other
NAt-dt	☐ Cancer	☐ Infection		Other
Misdiagnosis	☐ Fracture	Unknown		
	□ Amputation	☐ Fracture		Swelling / Inflammation
	Bruising	Repetitive Strai		Unknown
	Crushing	Slipped / Prola	sed Disc	Whiplash
Musculoskeletal	Dental Fracture / Tooth lossDislocation	□ Sprain / Strain□ Soft tissue injuit	~V	Other
/ Soft Tissue	P. Ulcer Stage 1: Intact skin with	•	•	
	P. Ulcer Stage 2: Part thickness			
	P. Ulcer Stage 3: Full thickness t			
	P. Ulcer Stage 4: Full thickness t	issue loss/necrosis: expo	sed bone/tendon/muscle	
	☐ Additional / Further Surgery	Loss of Wages ,	/ Income /	
Personal Loss	Limb Deformity	Business		Organ Retention
	Defamation of Character	Loss of Consort		- · · · · · · · · · · · · · · · · · · ·
Surgery Specific	Damage to organ / body partDental Damage / Loss	Loss of organ /Nerve injury / L		Unexpected complication / deterioration
Surgery Specific Injury	Foreign body left in situ	Function	U33 UI	Other
yw. y	☐ Unknown	☐ Inadequate ana	iesthesia	
Turning the IF.	☐ Anxiety / Trauma	☐ Stress		Worried Well
Traumatic/Emotional	DISD	Unknown		Other

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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.		
	☐ Birth Specific 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 Process / Treatment / Procedure		
	Procedures	□ Non Invasive		□ Other		
Care	☐ Medication	Route of administration Oral Intravenous Sub Cutaneous Intra Muscular Topical Rectal Inhalation Other / Unknown What medication was involve Medication One				
ical		Medication Two		☐ Wrong Quantity / Duration		
Clinical	☐ Nutrition	☐ 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	 Whole Blood Red Cells Platelet (Apheresis) Platelets (Pooled) Other 	 □ 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 □ Documentation / 	 □ 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 (1 fraction)	□ >1mSv □ <10% □ 10-20% □ >20%		
	☐ Radiotherapy	Records Li Equipment Performing procedure Pregnancy Status Unknown	 ☐ Wrong Drug ☐ Wrong Dose ☐ Wrong Process / Treatment / Intervention ☐ Failure / Malfunction ☐ Inadvertent deterministic effects 			
Bio Hazards	☐ Biological Hazards / Acquired Infections	☐ Bacteria ☐ Fungus / Mould ☐ Prion ☐ Virus ☐ Organism Unknown	Please Specify, if known e.g COVID-19, MRSA, etc	□ 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 □ Physical Assault / Abuse □ Physical Harassment □ Sexual Assault / Abuse □ Sexual Harassment		
	☐ Child Abuse	□ By a Member of the Public□ By a Peer / Student□ By a Prisoner			
	☐ Adult Abuse	□ By a Service User□ 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	 ☐ Human Use / Error ☐ Obstruction / Protruding Object ☐ Physical Training / Sport 		
	☐ Ergonomics (Incl. manual / people handling)	☐ Manual Handling☐ Other☐ Patient Handling☐ Restraint / Intervention	 □ Defective Equipment □ Unsafe / Inappropriate system □ Unknown □ Task 		
	☐ Mechanical Components	 □ Catering equipment □ Door / Gate / Barrier □ Healthcare Equipment □ Lifting Equipment / Accessories □ Office / Business equipment 	 □ Load □ Working Environment □ Individual Capability □ Other 		
	☐ Temperature (Excluding Fire)	☐ Hot ☐ Cold	□ Liquid / Food / Steam □ Equipment / Utensils □ Atmosphere / Environment		
	☐ Fire☐ Vibration☐ Electrical☐ Noise☐ Radiation	□ Please Specify	☐ Defective Equipment ☐ Human Use / Error ☐ Unknown ☐ Unsafe System ☐ Explosion ☐ Exposure ☐ Electrical Wiring / installation		

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SEC	TION H CNTD: WHAT TYPE	OF HAZARD DID THIS INC	CIDENT REL	ATE 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	Le	secticide ead letallic Dust lotor / Gear / Hydraulic Oil atural Gas rganic Dust aint / Paint Product etrol olish adon odenticide oap odium Hydroxide olvents pent / Used Oil Product ulphuric Acid /rong Patient ther	☐ Lack of Supervision ☐ Unknown ☐ Human / User Error ☐ Unsafe System
SEC	CTION I: IMMEDIATE ACTIO	INS TAKEN			
560	MONTH INTERPRETACTION	TAINET			
	CTION K: REPORTED BY: perso wise stated within the organization, this person i		SECTION	N L: WITNESS DETAILS (Name, Contact No. etc.)
Firs	t name				
Suri	name				
Dat	e notified	MYYYY			
Cat	egory of person <u>E.g. Nurse, C</u>	atering Staff, Cleaner			
Loc	al system reference				
no.					
Dat	e D D M	MYYYY			
Cor	tact Details				

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SECTION L: TO BE COMPLETED BY LINE/DEPARTMENT MAN	IAGER			
Has open disclosure happened? * (tick one only ✓)	☐ Yes	□ No		
If No, please specify*:				
CATEGORY 1 INCIDENTS ONLY				
SAO Name:		Date notified	d 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)?	☐ Yes	□ No		
If Yes: Name regulator(s)/agency(ies) reported/notified to:				Date Notified:
1				DDMMYYYY
2				DDMMYYYY
3				DDMMYYYY
Line/Department Manager name:			Title:	
			Date:	DDMMYYYY
SECTION M: TO BE COMPLETED BY QUALITY AND PATIENT	SAFETY (OFFICE		
Is this incident a Serious Reportable Event (SRE)? * (tick one only ✓)	☐ Yes	□ No		
QPS Advisor Name:			Date:	DDMMYYYY

*Mandatory Fields

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