



	West Limerick Independent Living CLG Policies				
Title:	INCIDENT MANAGEMENT & REPORTING POLICY				
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Category: Operational Policies
Subject: INCIDENT MANAGEMENT & REPORTING POLICY
Responsible for Review of this Policy: West Limerick Independent Living CLG Board

1. INTRODUCTION

This Incidents Management Policy sets out for the following for Staff and Service Users of West Limerick Independent Living

- how we will enable incidents to be reported
- how these will be investigated
- how we will ensure learning is shared with Staff and Service Users

2. WHAT OUR COMMITMENT MEANS

We are committed to promoting a culture where all incidents and near misses are reported and appropriately investigated. This is achieved by operating an open and just culture which encourages and supports staff and Service Users in reporting incidents so that learning and improvement can take place. The organisation ensures that the different needs in respect of ethnicity, faith, disability, gender age, sexual orientation, and socio-economic group are taken in to account in the reporting and investigation of incidents.

Learning from incidents enables changes to take place in order to:

- improve the safety of staff, Service Users and visitors
- improve the work environment
- improve service users experience

We will ensure that there are appropriate systems in place so that staff and service users are able to report incidents. Go to http://www.limerickcil.com/policies_and_procedures.html for further information.

Supporting the ability to report incidents ensures we:

- can use the information to take appropriate management decisions
- can identify trends in any root causes identified
- can share learning to improve practice within West Limerick Independent Living.

3. SCOPE AND PURPOSE OF THE POLICY

The purpose of this policy is to outline the way in which incidents will be reported. This policy describes procedures which apply to Staff, Service Users, independent contractors and the general public.



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4. WHAT IS AN INCIDENT?

An incident is an event that leads to, or could have caused, loss or harm to an individual or property.

- An incident reported by an individual about their own practice will be known as a reflective incident.
- An incident reported by staff or a service user about another individuals/organisations practice will be known as a notified incident.

5. WHO CAN REPORT INCIDENTS?

Incidents can be reported by West Limerick Independent Living staff and Service Users and the general public.

- about their own practice
- about incidents occurring in our organisation

6. WHEN TO REPORT INCIDENTS

- Incidents should be reported as soon as a concern becomes apparent.
- Staff and Service Users are encouraged to report all incidents in order to gain a true idea of any trends which may be occurring.
- All incidents have to be reported, these could include neglect, self-neglect and any suspicions.
- Incidents which identify concern, allegation, disclosure or suspicion of abuse, MUST also be reported to West Limerick Independent Living Safeguarding Vulnerable Persons. For further information go to http://www.limerickcil.com/policies_and_procedures.html
- Incidents which identify concern, allegation, disclosure or suspicion of abuse involving children MUST also be reported in line with West Limerick Independent Living Child Protection Policy http://www.limerickcil.com/policies_and_procedures.html

Any accident, incident or “near miss” no matter how slight the injury or damage, should be reported to your Service Coordinator before the end of your shift.

- In the event of injury or suspected injury, following a fall or suspected fall go to the Client Falls Management Policy http://www.limerickcil.com/policies_and_procedures.html
- In the event of a service user developing a pressure ulcer or have any risk factors for pressure ulcer development, this should be reported to your Service Coordinator before the end of your shift. Refer to <https://www.hse.ie/eng/about/qavd/incident-management>

Your Service Coordinator is responsible for taking appropriate follow up actions, completing an investigating report and recommending or implementing appropriate corrective actions.

All staff are required to report occurrences that may not have involved injuries or victims but could be potentially dangerous in that respect if repeated. These include but are not limited to;

- Slippery surfaces
- Malfunction of equipment



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- Water or gas leaks
- Inadequate insulation of circuits
- Trailing electrical leads
- Breaking of windows, glass or frames
- Collapses of walls, ceilings etc...

7. MANAGEMENT OF INCIDENTS

The Situation, Background, Assessment, and Recommendation (SBAR) tool has been adopted by West Limerick Independent Living to ensure incidents reported are managed in a consistent and concise manner and that important information is shared clearly, effectively and efficiently.

S = Situation What is going on? A concise statement of the problem or what actually happened?

B = Background What is the background information that is pertinent to the situation?

A = Assessment What did you find? Analysis and considerations of options and risks. What factors contributed to the incident?

R = Recommendation What action/recommendation is needed to correct the problem? What do you want to happen by when? Include any actions taken at the time of the incident.

8. All incidents once logged will be reviewed by the Service Manager

Once investigations have been completed by the Service Manager, all responses will be reviewed by the Manager and Quality and Safety Team and an individual Situation, Background, Assessment, and Recommendation response will be sent to the reporter of the incident as required.

Some no/low harm incidents may not always require an individual Situation, Background, Assessment, and Recommendation response but all responses will be reviewed, themed and shared appropriately with staff and management and using “You said we did” to share any wider learning themes.

9. Reporting of Incidents

Staff will report incidents including all relevant information using the Situation, Background, Assessment, and Recommendation tool.

Incident Management

The Agency Senior Accountable Officer is required to ensure that all incidents relating to service user care and safety; staff safety; accidents, loss or damage to property; incidents involving vehicles are appropriately recorded on the State Claims Agency NIMS system, where the Agency has access, or to the HSE main contact person (Business Managed) named in the Service Arrangement.



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The Agency Senior Accountable Officer should notify the HSE main contact person (Business Managed) who will communicate the information to the relevant personnel within HSE Midwest Community Healthcare, including the Quality, Risk and Patient Safety Business Manager.

A **Serious Incident** is an incident that results in a rating of major or extreme as per the HSE's Risk Impact Table. Serious Reportable Events (SREs) are a defined subset of incidents (i) which are either serious or (ii) that should not occur if the available preventive measures have been effectively implemented by care providers.

HSE Main contact person:	Nuala Kelly
Department/Specific area of responsibility:	Disability Services
Address:	St. Joseph's Hospital, Mulgrave Street, Limerick
Telephone Number:	061-461136
E-mail:	mailto:nuala.kelly1@hse.ie

Appendix:

See Appendix 1. for Incident Management Relevant Policies, Procedures, Protocols, Guidelines (PPPGs)

Appendix 2. A list of Serious Reportable Events is available on the NIMLT page of the Quality Assurance and Verification Division (QAVD) website.

See Appendix 3. Chart of incidents reporting and handling process.

Appendix 4. Use the National Incident Report Form NIRF attached or go to <https://www.hse.ie/eng/about/qavd/incident-management/nirf-01-v10-person.pdf> to download.



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Appendix 1: Incident Management Relevant Policies, Procedures, Protocols, Guidelines (PPPGs)

Key PPPGs:

- Incident Management Framework and guidance (HSE, 2018). <https://www.hse.ie/eng/about/qavd/incident-management/hse-2018-incident-management-framework-guidance-stories.pdf>
- Service User Falls – A practical Guide for Review (HSE, 2018). <https://www.hse.ie/eng/about/qavd/incident-management/service-user-falls-a-practical-guide-for-review.pdf>
- Pressure Ulcers – A practical Guide for Review (HSE, 2018). <https://www.hse.ie/eng/about/qavd/incident-management/pressure-ulcers-a-practical-guide-for-review.pdf>
- Systems Analysis Guidance (HSE (August), 2016). <https://www.hse.ie/eng/about/qavd/incident-management/hse-systems-analysis-investigation-guidelines-part-1-and-part-2.pdf>
- Integrated Risk Management Policy and Guidance (HSE, 2017). <https://www.hse.ie/eng/about/qavd/riskmanagement/risk-management-documentation/hse%20integrated%20risk%20management%20policy%202017.pdf>
 - Part 1: Managing Risk in Everyday Practice (Guidance for Managers) <https://www.hse.ie/eng/about/qavd/riskmanagement/integrated-risk-management-policy-part-1-managing-risk-in-everyday-practice.pdf>
 - Part 2: Risk Assessment and Treatment (Guidance for Managers) <https://www.hse.ie/eng/about/qavd/riskmanagement/risk-management-documentation/hse-integrated-risk-management-policy-part-2-risk-assessment-and-treatment.pdf>
 - Part 3: Managing & Monitoring Risk Registers (Guidance for Managers) <https://www.hse.ie/eng/about/qavd/riskmanagement/integrated-risk-management-policy-part-3-managing-and-monitoring-risk-registers-.pdf>

Risk Management Support Tools: HSE Risk Assessment Tool and Risk Assessment Form.

<https://www.hse.ie/eng/about/who/oqr012-20081210-v4-risk-assessment-tool-and-guidance-incl-guidance-on.pdf>

- Serious Reportable Events (SREs), HSE Implementation Guidance Document (HSE, 2015) <https://www.hse.ie/eng/services/publications/performance-reports/srejan15.pdf>
- Open Disclosure: National Guidelines (HSE, 2013) <https://www.hse.ie/eng/about/who/qid/other-quality-improvement-programmes/opendisclosure/opendiscfiles/opendiscpolicyoct13.pdf>



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Related PPPGs:

- A Board's Role in Improving Quality and Safety (HSE, 2017).
<https://www.hse.ie/eng/about/who/qid/governancequality/board-role-improving-quality-and-safety/a-board-s-role-in-improving-quality-and-safety-guide-final.pdf>
- Quality and Safety Committees Guidance and Sample Terms of Reference (HSE, 2016).
<https://www.hse.ie/eng/about/who/qid/governancequality/boardquality/quality-and-safety-committees-guidance-and-resources-2016.pdf>
- Framework for improving Quality in our Health services: Part 1: Introducing the Framework (HSE, 2016).
<https://www.hse.ie/eng/about/who/qid/framework-for-quality-improvement/framework-for-improving-quality-2016.pdf>
- Guideline on Conducting Look-back Reviews (HSE, 2015). <https://www.hse.ie/eng/about/qavd/incident-management/lookback-review-guideline-final-dec-2015.pdf>
- Policy for Preventing and Managing Critical Incident Stress (HSE, 2012).
<https://www.hse.ie/eng/staff/resources/hrppg/policy-for-preventing-and-managing-critical-incident-stressdecember-2012.pdf>



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APPENDIX 2.

Summary list of Serious Reportable Events 26th January 2015

1.	SURGICAL EVENTS
1A.	Surgery performed on the <i>wrong body part</i> by a healthcare service provider.
1B.	Surgery performed on the <i>wrong patient</i> by a healthcare service provider.
1C.	<i>Wrong surgical procedure</i> performed on patient by a healthcare service provider.
1D.	<i>Unintended retention of a foreign object</i> in an enclosed body cavity in a patient after surgery or other procedure performed by a healthcare service provider.
1E.	<i>Intra-operative or immediately postoperative death</i> in a patient with <i>no known medical problems</i> (ASA Class I) occurring after surgery or other interventional procedure performed by a healthcare service provider.

2.	PRODUCT OR DEVICE EVENTS
2A.	<i>Patient death or serious disability</i> associated with the use of <i>contaminated medications, medical devices, or biologics</i> provided by the healthcare service provider.
2B.	<i>Patient death or serious disability</i> associated with the use or function of a <i>medical device</i> in which the medical device is used or functions other than as intended or anticipated in the care of a patient provided by the healthcare service provider.
2C.	<i>Patient death or serious disability</i> associated with <i>intravascular air embolism</i> that occurs while being cared for by a healthcare service provider but excluding death or serious disability associated with certain neurosurgical procedures or cardiac procedures known to present a high risk of intravascular air embolism.

3.	PATIENT PROTECTION EVENTS
3A.	<i>Child or other dependent person discharged to the wrong person</i> by a healthcare service provider.
3B.	<i>Patient death or serious disability</i> associated with a <i>patient absconding from a healthcare service facility</i> but excluding where a patient advises the healthcare service provider that he or she is leaving against medical advice.
3C.	<i>All sudden unexplained deaths or injuries</i> which result in <i>serious disability of a person who is an inpatient/resident in a mental healthcare facility</i> .

4.	CARE MANAGEMENT EVENTS
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4A.	Patient death or serious disability associated with a <i>medication error</i> by the healthcare service provider but excluding reasonable differences in clinical judgment involving drug selection and/or dose.
4B.	<i>Wrong formulation/route administration of chemotherapy</i> by a healthcare service provider.
4C	<i>Intravenous administration of mis-selected concentrated potassium chloride</i> by a healthcare service provider.
4D	<i>Patient death or serious disability</i> due to the administration of <i>incompatible blood or blood products</i> by a healthcare service provider.
4E	<i>Maternal Death</i> for whom the hospital has accepted medical responsibility, registered with an independent midwife for pregnancy care/ registered with a maternity hospital-during pregnancy or <i>within six weeks of delivery</i> (whether in the hospital or not).
4F(i)	<i>Perinatal death of a neonate</i> occurring in a <i>term infant or an infant weighing more than 2,500g.</i>
4F(ii)	<i>Death or encephalopathy of a normally formed neonate</i> occurring in a <i>term infant or an infant weighing more than 2,500g.</i>
4G	<i>Patient death or serious disability</i> associated with <i>severe hypoglycaemia</i> (excluding neonates), the onset of which occurs while the patient is being cared for in a healthcare service facility.
4H	<i>Death or serious disability (kernicterus)</i> associated with <i>non detection</i> by a healthcare service provider to identify and treat <i>Hyperbilirubinemia in neonates within the first 28 days of life.</i>
4I	<i>Stage 3 or 4 pressure ulcers</i> acquired <i>after admission</i> to a healthcare and social care residential facility.
4J	<i>Patient death or serious disability</i> due to <i>spinal manipulative therapy</i> by a healthcare service provider.
4K	<i>Patient death or serious disability</i> resulting from or associated with the use of <i>restrictive interventions such as physical, mechanical, manual or environmental restraint (e.g. seclusion)</i> to a patient while being cared for in a healthcare service facility.
4L	<i>Diagnostic Error:</i> Death or serious disability associated with a wrong diagnostic result e.g. mislabelled pathology specimen.
4M	The <i>non utilisation of a donor organ</i> deemed suitable for transplantation.
4N	<i>Death of a living organ donor.</i>

5	ENVIRONMENTAL EVENTS
5A	<i>Patient death or serious disability</i> associated with an <i>electric shock</i> while being cared for in a healthcare service facility but excluding events involving planned treatments such as cardioversion.



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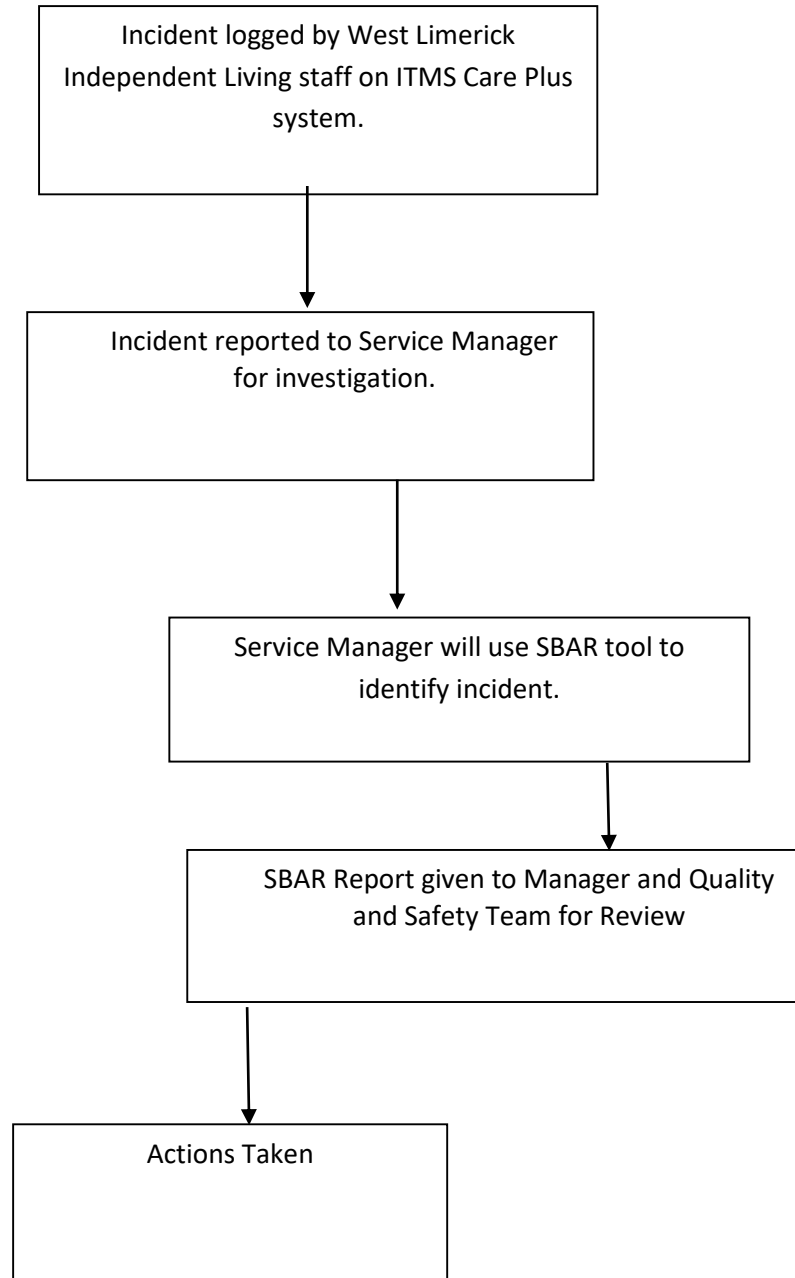
5B	An incident in which a <i>line designated for oxygen or other gas</i> to be delivered to a patient while being cared for by a healthcare service provider <i>contains the wrong gas or is contaminated by toxic substances</i> .
5C	<i>Patient death or serious disability</i> associated with a <i>burn incurred</i> within a healthcare service facility.
5D	Patient death or serious disability associated with a <i>fall</i> – a. while being cared for in a healthcare service facility and/or b. during a clinical intervention from a healthcare professional (includes in the community setting, pre-hospital care and the Ambulance Service).

6	CRIMINAL EVENTS
6A	Any instance of care ordered by or provided by someone <i>impersonating a healthcare professional</i> .
6B	<i>Abduction of a patient</i> of any age while being cared for in a healthcare service facility.
6C	Sexual assault on a patient or other person within or on the grounds of a healthcare service facility.
6D	<i>Death or serious injury/disability</i> of a patient or other person resulting from a <i>physical assault</i> that occurs within or on the grounds of a healthcare service facility.



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APPENDIX. 3.





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APPENDIX 4.

NIMS <small>National Incident Management System</small>	HC NIRF 01 – V10 Date issued: 03/05/2018	NATIONAL INCIDENT REPORT FORM (NIRF) NIRF - 01 PERSON NIMS record Number: <input style="width: 150px;" type="text"/>
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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

Date of incident

Time of incident *Use 24 hour clock*

Location *E.g. Hospital, Health Centre, Residential Centre etc.*

Specific Location *E.g. Ward, Clients home etc.* Offsite?

SECTION B: PERSON AFFECTED DETAILS

First name _____

Surname _____

Date of birth

Female Male

Description of incident:

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

Older Person Specialties

Incident Occurred under (Service / Speciality) *E.g. Antenatal, Audiology, Radiotherapy, Intellectual Disability, Psychology*

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)

Work days lost 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


SECTION E: EXTERNAL CONTRACTOR DETAILS ONLY

Company Name _____

Company no. _____



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SECTION F: WHAT WAS THE OUTCOME AT THE TIME OF THE INCIDENT?		
✓ Outcome	Category	Body Part Affected
<input type="checkbox"/> Near Miss e.g. Nearly given wrong drug	Category 3	 <i>E.g. Arm, Spine, Lung, Other Physiological</i>
<input type="checkbox"/> No Injury e.g. Wrong drug given but no harm occurred		
<input type="checkbox"/> Injury not requiring first aid		
<input type="checkbox"/> Injury or illness, requiring first aid	Category 2	
<input type="checkbox"/> Injury requiring medical treatment		
<input type="checkbox"/> Long-term disability / Incapacity (incl. psychosocial)	Category 1	
<input type="checkbox"/> Permanent Incapacity (incl. Psychosocial)		
<input type="checkbox"/> Death		

SECTION G: TYPE OF INJURY (tick one only ✓)			
Birth Specific Injury (Baby)	<input type="checkbox"/> Apgar score <5@ 1 min &/or; 7@5mins &/or pH ≤ 7.0	<input type="checkbox"/> HIE Grade 2 - Hypoxic Ischaemic Encephalopathy	<input type="checkbox"/> Nerve Injury - face
	<input type="checkbox"/> Aspiration	<input type="checkbox"/> HIE Grade 3 - Hypoxic Ischaemic Encephalopathy	<input type="checkbox"/> Other unexpected deterioration
	<input type="checkbox"/> Cerebral irritability / neonatal seizure	<input type="checkbox"/> Hypoglycaemia - severe	<input type="checkbox"/> Stillbirth
	<input type="checkbox"/> HIE - Hypoxic Ischaemic Encephalopathy with Hypoglycaemia	<input type="checkbox"/> Kernicterus	<input type="checkbox"/> Sub-galeal / sub-aponeurotic haemorrhage
	<input type="checkbox"/> HIE Grade 1 - Hypoxic Ischaemic Encephalopathy	<input type="checkbox"/> Neonatal death	<input type="checkbox"/> Unknown
	<input type="checkbox"/> Nerve Injury - brachial plexus (incl. Erbs Palsy)	<input type="checkbox"/> Other _____	
Birth Specific Injury (Mother)	<input type="checkbox"/> Death	<input type="checkbox"/> Perineal tear	<input type="checkbox"/> Unknown
	<input type="checkbox"/> Hysterectomy (Perinatal)	<input type="checkbox"/> Post-Partum Haemorrhage	<input type="checkbox"/> Uterine rupture
	<input type="checkbox"/> Incontinence (faecal)	<input type="checkbox"/> Rhesus iso-immunisation	<input type="checkbox"/> Other _____
	<input type="checkbox"/> Incontinence (urinary)	<input type="checkbox"/> Incontinence (faecal & urinary)	
Blood Specific Injury	<input type="checkbox"/> Excessive Bleeding	<input type="checkbox"/> Febrile non-haemolytic transfusion reaction	<input type="checkbox"/> Non-immunological haemolysis
	<input type="checkbox"/> Fainting		<input type="checkbox"/> Other _____
	<input type="checkbox"/> Immunological haemolysis		
Diagnosed Disease Disorder or Cond.	<input type="checkbox"/> Asbestosis	<input type="checkbox"/> Hepatitis	<input type="checkbox"/> Unknown
	<input type="checkbox"/> Cancer	<input type="checkbox"/> HIV	<input type="checkbox"/> Dermatitis
	<input type="checkbox"/> Acute Radiation Syndrome	<input type="checkbox"/> Brucellosis	<input type="checkbox"/> TB
	<input type="checkbox"/> Narcolepsy/Cateplexy	<input type="checkbox"/> Legionnaires	<input type="checkbox"/> Pleural Plaques
			<input type="checkbox"/> Other _____
Diagnosed Infection	<input type="checkbox"/> Clostridium Difficile	<input type="checkbox"/> MRSA	<input type="checkbox"/> VRE
	<input type="checkbox"/> ESBL	<input type="checkbox"/> Norovirus	<input type="checkbox"/> VRSA
	<input type="checkbox"/> Hepatitis	<input type="checkbox"/> Unknown	<input type="checkbox"/> Other _____
General Injuries	<input type="checkbox"/> Allergic Reaction (incl. anaphylaxis)	<input type="checkbox"/> Cut / Laceration / Graze / scratch	<input type="checkbox"/> Malaise / Nausea
	<input type="checkbox"/> Brain Injury / Concussion	<input type="checkbox"/> Death	<input type="checkbox"/> Nerve injury / Loss of Function
	<input type="checkbox"/> Burn / scald / corrosion	<input type="checkbox"/> Dental injury &/or loss	<input type="checkbox"/> Puncture / bite
	<input type="checkbox"/> Choking / asphyxia	<input type="checkbox"/> Deterioration	<input type="checkbox"/> Rash / irritation
	<input type="checkbox"/> Circulatory / volume depletion	<input type="checkbox"/> Haemorrhage	<input type="checkbox"/> Unknown
	<input type="checkbox"/> Circulatory / volume overload	<input type="checkbox"/> Blister	<input type="checkbox"/> Other _____
	<input type="checkbox"/> Pain/Discomfort		
Hearing / Sight Injury	<input type="checkbox"/> Hearing Impairment / loss	<input type="checkbox"/> Tinnitus	<input type="checkbox"/> Other _____
	<input type="checkbox"/> Sight Impairment / loss	<input type="checkbox"/> Unknown	
Misdiagnosis	<input type="checkbox"/> Cancer	<input type="checkbox"/> Infection	<input type="checkbox"/> Other _____
	<input type="checkbox"/> Fracture	<input type="checkbox"/> Unknown	
Musculoskeletal / Soft Tissue	<input type="checkbox"/> Amputation	<input type="checkbox"/> Fracture	<input type="checkbox"/> Swelling / Inflammation
	<input type="checkbox"/> Bruising	<input type="checkbox"/> Repetitive Strain Injury (RSI)	<input type="checkbox"/> Unknown
	<input type="checkbox"/> Crushing	<input type="checkbox"/> Slipped / Prolapsed Disc	<input type="checkbox"/> Whiplash
	<input type="checkbox"/> Dental Fracture / Tooth loss	<input type="checkbox"/> Sprain / Strain	<input type="checkbox"/> Other _____
	<input type="checkbox"/> Dislocation	<input type="checkbox"/> Soft tissue injury	
	<input type="checkbox"/> P. Ulcer Stage 1: Intact skin with non-blanchable redness over bony prominence		
	<input type="checkbox"/> P. Ulcer Stage 2: Part thickness dermis loss: blister/open ulcer/no slough		
	<input type="checkbox"/> P. Ulcer Stage 3: Full thickness tissue loss: +/- visible subcutaneous fat		
<input type="checkbox"/> P. Ulcer Stage 4: Full thickness tissue loss/necrosis: exposed bone/tendon/muscle			
Personal Loss	<input type="checkbox"/> Additional / Further Surgery	<input type="checkbox"/> Loss of Wages / Income / Business	<input type="checkbox"/> Unknown
	<input type="checkbox"/> Limb Deformity	<input type="checkbox"/> Loss of Consortium	<input type="checkbox"/> Organ Retention
	<input type="checkbox"/> Defamation of Character	<input type="checkbox"/> Loss of organ / body part	<input type="checkbox"/> Other _____
Surgery Specific Injury	<input type="checkbox"/> Damage to organ / body part	<input type="checkbox"/> Nerve injury / Loss of Function	<input type="checkbox"/> Unexpected complication / deterioration
	<input type="checkbox"/> Dental Damage / Loss	<input type="checkbox"/> Inadequate anaesthesia	<input type="checkbox"/> Other _____
	<input type="checkbox"/> Foreign body left in situ		
	<input type="checkbox"/> Unknown		
Traumatic/Emotional	<input type="checkbox"/> Anxiety / Trauma	<input type="checkbox"/> Stress	<input type="checkbox"/> Worried Well
	<input type="checkbox"/> PTSD	<input type="checkbox"/> Unknown	<input type="checkbox"/> 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.
Clinical Care	<input type="checkbox"/> Birth Specific Procedures	<input type="checkbox"/> Caesarean Section (Elective) <input type="checkbox"/> Caesarean Section (Emergency) <input type="checkbox"/> Instrumental Delivery (Forceps) <input type="checkbox"/> Instrumental Delivery (Vacuum) <input type="checkbox"/> Instrumental Delivery (Multiple Instruments) <input type="checkbox"/> Non Instrumental Delivery	<input type="checkbox"/> Communication / Consent <input type="checkbox"/> Diagnosis / Assessment <input type="checkbox"/> Documentation / Records <input type="checkbox"/> Equipment <input type="checkbox"/> General Care / Management <input type="checkbox"/> Procedure / Treatment / Intervention <input type="checkbox"/> Screening / Prevention <input type="checkbox"/> Specimens / Results <input type="checkbox"/> Tests / Investigations <input type="checkbox"/> Unknown <input type="checkbox"/> Other _____	<input type="checkbox"/> Adverse Effect <input type="checkbox"/> Failure / Malfunction <input type="checkbox"/> Foreign Body left in Situ <input type="checkbox"/> Inappropriate for Task / Wrong device <input type="checkbox"/> Incomplete / Inadequate <input type="checkbox"/> Lack of Availability <input type="checkbox"/> Not performed when indicated / Delay <input type="checkbox"/> Pre Existing Medical Condition <input type="checkbox"/> Shoulder Dystocia <input type="checkbox"/> Unavailable / Mislabeled / Lost <input type="checkbox"/> Wrong Body Part / Site / Side <input type="checkbox"/> Wrong Patient <input type="checkbox"/> Wrong Process / Treatment / Procedure <input type="checkbox"/> Other _____
	<input type="checkbox"/> Clinical Procedures	<input type="checkbox"/> Invasive <input type="checkbox"/> Non Invasive	_____	_____
	<input type="checkbox"/> Medication	<i>Route of administration</i> <input type="checkbox"/> Oral <input type="checkbox"/> Intravenous <input type="checkbox"/> Sub Cutaneous <input type="checkbox"/> Intra Muscular <input type="checkbox"/> Topical <input type="checkbox"/> Rectal <input type="checkbox"/> Inhalation <input type="checkbox"/> Other / Unknown	<input type="checkbox"/> Administration <input type="checkbox"/> Monitoring <input type="checkbox"/> Ordering / Supply / Transport <input type="checkbox"/> Preparation / Dispensing (Pharmacy) <input type="checkbox"/> Prescribing <input type="checkbox"/> Reconciliation <input type="checkbox"/> Storage	<input type="checkbox"/> Adverse Drug Reaction <input type="checkbox"/> Contra-indicated <input type="checkbox"/> Drug Interaction <input type="checkbox"/> Failure / Malfunction of equipment <input type="checkbox"/> Incomplete / Inadequate <input type="checkbox"/> Not performed when indicated / delayed <input type="checkbox"/> Omitted/Delayed Dose <input type="checkbox"/> Wrong Dose / Strength <input type="checkbox"/> Wrong Drug <input type="checkbox"/> Wrong Formulation / Route <input type="checkbox"/> Wrong Frequency <input type="checkbox"/> Wrong Label / Instructions <input type="checkbox"/> Wrong Patient <input type="checkbox"/> Wrong Quantity / Duration
		<i>What medication was involved?</i> Medication One _____ Medication Two _____		
	<input type="checkbox"/> Nutrition	<input type="checkbox"/> Parenteral <input type="checkbox"/> Enteral <input type="checkbox"/> Special Diet <input type="checkbox"/> General Diet <input type="checkbox"/> Other _____	<input type="checkbox"/> Communication / Consent <input type="checkbox"/> Prescribing / Requesting <input type="checkbox"/> Preparation / Dispensing <input type="checkbox"/> Administration <input type="checkbox"/> Storage <input type="checkbox"/> Documentation / Records <input type="checkbox"/> Equipment <input type="checkbox"/> Supply / Ordering / Transport <input type="checkbox"/> Presentation / Packaging <input type="checkbox"/> Transfusing blood <input type="checkbox"/> Other _____	<input type="checkbox"/> Adverse Effect <input type="checkbox"/> Incomplete / Inadequate <input type="checkbox"/> Not performed when indicated / Delay <input type="checkbox"/> Wrong Consistency <input type="checkbox"/> Wrong Diet / Wrong Blood Product <input type="checkbox"/> Wrong Process / Treatment / Procedure <input type="checkbox"/> Wrong Patient <input type="checkbox"/> Lack of Availability <input type="checkbox"/> Wrong dispensing label / instructions <input type="checkbox"/> Inappropriate for task / Wrong device <input type="checkbox"/> Other _____
	<input type="checkbox"/> Blood / Blood Product	<input type="checkbox"/> Whole Blood <input type="checkbox"/> Red Cells <input type="checkbox"/> Platelet (Apheresis) <input type="checkbox"/> Platelets (Pooled) <input type="checkbox"/> Other _____		
	<input type="checkbox"/> Diagnostic Radiology (DR) & Nuclear Medicine (NM)	<input type="checkbox"/> Checking Patient ID procedure <input type="checkbox"/> Clinical Details on Referral <input type="checkbox"/> Communication / Consent <input type="checkbox"/> Documentation / Records <input type="checkbox"/> Equipment <input type="checkbox"/> Performing procedure <input type="checkbox"/> Pregnancy Status <input type="checkbox"/> Unknown	<input type="checkbox"/> Diagnostic Exposure > intended <input type="checkbox"/> X-ray Over Exposure <input type="checkbox"/> Wrong body part / side <input type="checkbox"/> Dose to comforters / carers <input type="checkbox"/> Wrong Patient <input type="checkbox"/> Inadvertent dose to foetus <input type="checkbox"/> Total dose or Volume Variation <input type="checkbox"/> Dose (NM) or Volume Variation (1 fraction)	<input type="checkbox"/> Above Notifiable levels <input type="checkbox"/> Below Notifiable levels <input type="checkbox"/> <1mSv <input type="checkbox"/> >1mSv <input type="checkbox"/> <10% <input type="checkbox"/> 10-20% <input type="checkbox"/> >20%
	<input type="checkbox"/> Radiotherapy		<input type="checkbox"/> Wrong Drug <input type="checkbox"/> Wrong Dose <input type="checkbox"/> Wrong Process / Treatment / Intervention <input type="checkbox"/> Failure / Malfunction <input type="checkbox"/> Inadvertent deterministic effects	
Bio Hazards	<input type="checkbox"/> Biological Hazards / Acquired Infections	<input type="checkbox"/> Bacteria <input type="checkbox"/> Fungus / Mould <input type="checkbox"/> Prion <input type="checkbox"/> Virus <input type="checkbox"/> Organism Unknown		<input type="checkbox"/> Exposure to Bite (Human) <input type="checkbox"/> Exposure to Bite (Insect / Animal) <input type="checkbox"/> Exposure to Bodily Fluids <input type="checkbox"/> Exposure to Ingestion/Food/Water <input type="checkbox"/> Exposure to Needle Stick <input type="checkbox"/> Exposure to Skin Contact <input type="checkbox"/> Inhalation/Airborne <input type="checkbox"/> Equipment, Implements, Facilities, Sharps (Non Needle) <input type="checkbox"/> Unknown <input type="checkbox"/> 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	<input type="checkbox"/> Self-Injurious Behaviour	<input type="checkbox"/> Intentional <input type="checkbox"/> Unintentional	<input type="checkbox"/> Absconsion / Missing <input type="checkbox"/> Attempted Suicide <input type="checkbox"/> Banging Self Against Walls/Furniture/Surfaces <input type="checkbox"/> Hitting Body/Slap/Punch Self incl. Scratching & Picking <input type="checkbox"/> Inappropriate Eating <input type="checkbox"/> Inappropriate Touching <input type="checkbox"/> Self-Harm <input type="checkbox"/> Stripping Clothes in Public Area <input type="checkbox"/> Suicide <input type="checkbox"/> Throwing objects <input type="checkbox"/> Other _____
	<input type="checkbox"/> Violence, Harassment and Aggression	<input type="checkbox"/> By a Family Member / Relative <input type="checkbox"/> By a Member of the Public <input type="checkbox"/> By a Peer / Student <input type="checkbox"/> By a Prisoner <input type="checkbox"/> By a Service User <input type="checkbox"/> By a Staff Member	<input type="checkbox"/> Aggressive towards inanimate object <input type="checkbox"/> Discrimination/Prejudice/Racial <input type="checkbox"/> Intimidation / Threat <input type="checkbox"/> Neglect <input type="checkbox"/> Non-Compliant / Obstructive / Rude <input type="checkbox"/> Physical Assault / Abuse <input type="checkbox"/> Physical Harassment <input type="checkbox"/> Sexual Assault / Abuse <input type="checkbox"/> Sexual Harassment <input type="checkbox"/> Unintentional Aggressive Behaviour <input type="checkbox"/> Bullying <input type="checkbox"/> Verbal Assault / Abuse <input type="checkbox"/> Verbal Harassment <input type="checkbox"/> Other _____
	<input type="checkbox"/> Child Abuse		
	<input type="checkbox"/> Adult Abuse		
Physical Hazards	<input type="checkbox"/> Slip / Trip / Fall	<input type="checkbox"/> From Height <input type="checkbox"/> From Equipment / Furniture <input type="checkbox"/> Same Level / Ground <input type="checkbox"/> On Stairs <input type="checkbox"/> On Steps <input type="checkbox"/> Other _____	<input type="checkbox"/> Unknown <input type="checkbox"/> Pre Existing Medical Condition <input type="checkbox"/> Inadequate supervision gen health / post op <input type="checkbox"/> Obstruction / protruding object <input type="checkbox"/> Surface contaminants <input type="checkbox"/> Rough terrain / irregular surface <input type="checkbox"/> Inappropriate equipment use <input type="checkbox"/> Failure / malfunction of equipment <input type="checkbox"/> Horseplay <input type="checkbox"/> Physical training / sport <input type="checkbox"/> Weather Condition <input type="checkbox"/> Inadequate Lighting / design <input type="checkbox"/> Other _____
	<input type="checkbox"/> Non Mechanical (Incl. Person / Animal)	<input type="checkbox"/> Object / Tools (Non Sharps) <input type="checkbox"/> Sharps (Non Needle) <input type="checkbox"/> Other <input type="checkbox"/> Person	<input type="checkbox"/> Human Use / Error <input type="checkbox"/> Obstruction / Protruding Object <input type="checkbox"/> Physical Training / Sport <input type="checkbox"/> Defective Equipment <input type="checkbox"/> Unsafe / Inappropriate system <input type="checkbox"/> Unknown <input type="checkbox"/> Task <input type="checkbox"/> Load <input type="checkbox"/> Working Environment <input type="checkbox"/> Individual Capability <input type="checkbox"/> Other _____
	<input type="checkbox"/> Ergonomics (Incl. manual / people handling)	<input type="checkbox"/> Manual Handling <input type="checkbox"/> Other <input type="checkbox"/> Patient Handling <input type="checkbox"/> Restraint / Intervention	
	<input type="checkbox"/> Mechanical Components	<input type="checkbox"/> Catering equipment <input type="checkbox"/> Door / Gate / Barrier <input type="checkbox"/> Healthcare Equipment <input type="checkbox"/> Lifting Equipment / Accessories <input type="checkbox"/> Office / Business equipment	
	<input type="checkbox"/> Temperature (Excluding Fire)	<input type="checkbox"/> Hot <input type="checkbox"/> Cold	
	<input type="checkbox"/> Fire <input type="checkbox"/> Vibration <input type="checkbox"/> Electrical <input type="checkbox"/> Noise <input type="checkbox"/> Radiation	<input type="checkbox"/> Please Specify _____	
		<input type="checkbox"/> Liquid / Food / Steam <input type="checkbox"/> Equipment / Utensils <input type="checkbox"/> Atmosphere / Environment <input type="checkbox"/> Defective Equipment <input type="checkbox"/> Human Use / Error <input type="checkbox"/> Unknown <input type="checkbox"/> Unsafe System <input type="checkbox"/> Explosion <input type="checkbox"/> Exposure <input type="checkbox"/> 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	<input type="checkbox"/> Acid / Alkaline	<input type="checkbox"/> Animal Remedy	<input type="checkbox"/> Insecticide	
	<input type="checkbox"/> Agri Chemicals	<input type="checkbox"/> Arsenic	<input type="checkbox"/> Lead	
	<input type="checkbox"/> Gas	<input type="checkbox"/> Asbestos	<input type="checkbox"/> Metallic Dust	
	<input type="checkbox"/> Other Chemical Products	<input type="checkbox"/> Bleach	<input type="checkbox"/> Motor / Gear / Hydraulic Oil	
	<input type="checkbox"/> Particulates	<input type="checkbox"/> Cadmium	<input type="checkbox"/> Natural Gas	
	<input type="checkbox"/> Petroleum / Synthetic Oil Based Products	<input type="checkbox"/> Carbon Dioxide	<input type="checkbox"/> Organic Dust	
	<input type="checkbox"/> Sanitation / Cleaning Chemicals	<input type="checkbox"/> Carbon Monoxide	<input type="checkbox"/> Paint / Paint Product	
	<input type="checkbox"/> Toxic Metals	<input type="checkbox"/> Chemical Fertilizer	<input type="checkbox"/> Petrol	<input type="checkbox"/> Lack of Supervision <input type="checkbox"/> Unknown <input type="checkbox"/> Human / User Error <input type="checkbox"/> Unsafe System
		<input type="checkbox"/> Crystalline Silica	<input type="checkbox"/> Polish	
		<input type="checkbox"/> Detergent	<input type="checkbox"/> Radon	
		<input type="checkbox"/> Diesel / Kerosene	<input type="checkbox"/> Rodenticide	
		<input type="checkbox"/> Disinfectant	<input type="checkbox"/> Soap	
	<input type="checkbox"/> Drain / Oven Cleaner	<input type="checkbox"/> Sodium Hydroxide		
	<input type="checkbox"/> Drugs	<input type="checkbox"/> Solvents		
	<input type="checkbox"/> Fungicide	<input type="checkbox"/> Spent / Used Oil Product		
	<input type="checkbox"/> Glue / Adhesive	<input type="checkbox"/> Sulphuric Acid		
	<input type="checkbox"/> Grease	<input type="checkbox"/> Wrong Patient		
	<input type="checkbox"/> Herbicide	<input type="checkbox"/> Other		
	<input type="checkbox"/> Hydrochloric Acid			

SECTION I: IMMEDIATE ACTIONS TAKEN

SECTION J: REPORTED BY: person who discovers the incident and unless otherwise stated within the organization, this person is responsible for completing the NRP.

First name _____

Surname _____

Date notified

Category of person E.g. Nurse, Catering Staff, Cleaner

Local system reference no. _____

Reporter Signature _____

Date

Contact Details _____

SECTION K: WITNESS DETAILS (Name, Contact No. etc.)



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SECTION L: TO BE COMPLETED BY LINE/DEPARTMENT MANAGER

Has open disclosure happened? (tick one only ✓) Yes No

If No, please specify: _____

CATEGORY 1 INCIDENTS ONLY

SAO Name [Block Capitals]: _____ Date notified to SAO:

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 _____

2 _____

3 _____

Line/Department Manager name [Block Capitals]: _____ Title: _____

Signature of Line/Department Manager: _____ Date:

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 [Block Capitals]: _____

Signature of QPS Advisor: _____ Date: