



සෞඛ්‍ය, පෝෂණ සහ දේශීය වෛද්‍ය අමාත්‍යාංශය  
சுகாதார, போசணை மற்றும் சுதேச வைத்திய அமைச்சு  
Ministry of Health, Nutrition & Indigenous Medicine

Strictly Confidential

**Adverse Event / Incident Reporting Form**

Hospital.....

This form is not intended to penalize anyone. Reporting an incident will help us to prevent such events in the future

<b>Part A:</b>	මෙය ඕනෑම සෞඛ්‍ය සේවා වෘත්තීයයෙකුට සම්පූර්ණ කළ හැක; எந்தவொரு சுகாதார பணியாளராலும் நிரப்ப முடியும்; Can be filled by any health care worker.
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Category of person affected: Patient  Staff  Visitor

Name of affected person: .....	Place of occurrence of adverse event: .....
Age: .....	Reporting ward/unit: .....
Gender: Male <input type="checkbox"/> Female <input type="checkbox"/>	BHT No. (If in patient): .....
Address (optional): .....	Date of adverse event/incident: dd/mm/yyyy...
	Time of adverse event/incident: .....

අන්තරායකර/අහිතකර සිදුවීම් (අවස්ථා පැහැදිලි කරන්න) விபரீத விளைவு / சம்பவம் (தயவு செய்து விபரமாக விபரிக்கவும்)  
Adverse event/incident (please describe in detail)

Elaborate (If necessary use a separate paper)

Date and time of occurrence:

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එම අවස්ථාවේ ඒ සම්බන්ධයෙන් ගත් ක්‍රියා මාර්ග; உடனடியாக எடுத்த நடவடிக்கை, **Immediate measures have been taken**

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Directorate of Healthcare Quality & Safety, Ministry of Health, Nutrition & Indigenous Medicine



**Part B: To be filled by Head of the unit (Consultant, MO, Nursing Sister, Chief MLT, Chief Pharmacist, Chief Radiographer, in charges of units etc)**

Describe the risk factors, root causes leading to the Adverse Event / Incident
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**Preventive measures recommended by the unit**

Action recommended	Person responsible for remedial action	Target time period
		dd/mm/yyyy

**Designation of the staff member involved in the adverse event**

Consultant  MO  Nursing sister/NOIC  Nurse  Others:.....

**Outcome of the affected person (Can be more than one. Please tick the relevant outcome/outcomes):**

- a. Completely recovered  Partially disabled  Permanently disabled
- b. Needed hospitalization  Extended hospitalization  Death

**Reference list of adverse events/ Incidents (Please select relevant category/categories)**

Fall/Safety issues	Treatment/diagnosis issues	Drugs/IV/Blood issues	Surgery/Anaesthesia issues	
Fall from bed/ chair/table	Orders/procedures carried out incorrectly	Omitted drug	Wrong patient identifier to call for surgery	
		Wrong drug	Wrong patient/site/site	
		Drug allergy		
Slip & fall	Orders/procedures on wrong patient	Patient/drug/dose/route /time	Left in patient – swab/ instrument	
Found on floor	Order not carried out/ delay	Dispensing error from pharmacy	Discrepancy in swab/ instrument count	
Climbed over bed rail	Plaster allergy/skin tear	Expired drug	Incorrect/no consent	
	Bed sore	Blood transfusion reaction	Diathermy burn	
Injury while transporting	Doctor not notified/ Doctor did not visit	IV site redness/Phlebitis	Equipment not available/ malfunction	
		Wrong diluents		
<b>Laboratory reports</b>	Medication delayed/ notavailable	<b>Miscellaneous issues</b>	Material not available	
Sample lost	Refusal of treatment by patient	Assault to patient/visitor/staff	No PAC	
Label lost	<b>Labour/Delivery issues</b>	Coroners' case	Others (Please specify)	
Wrong sample		Missed medical records		
Reports are not delivered on time		Traumatic birth		Therapeutic procedures
		Forceps injury to infant		Theft
Report lost		Laceration to neonate		Smoking/substance abuse in ward
	Unattended delivery	Wrong drug prescribed, administered		



