

	ADVERSE EVENT RECORDING FORM	UNIQUE MEDICAL INQUIRY / CASE REPORT NUMBER: (to be filled by PV officials)
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<i>Reporter Details</i>			
Name		Phone #	
Address		Fax #	
		Email id	
City		Preferred method to contact	Phone / Fax / Email
State		Consent to contact Reporter	Yes / No
Country		Consent to contact Patient	Yes / No
Qualification		Health Care Professional / Non HCP	

<i>Patient Details</i>			
Patient Initials		Gender (M / F)	
DOB (DD/MMM/YYYY)		Age at the time of event (years)	
Weight (kgs)		Height (cms)	
Ethnicity / Race			



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Suspected Drug(s)

Generic name of the drug

Sr. No.	Suspect Drug Name	Strength	Dosage form	Route	Expiry date	Batch Number	Indication	Start date, Time	Stop date, time	Action Taken*
1										
2										

* Action Taken: 0 - Ongoing; 1 - Dose reduced; 2 - Temporarily stopped; 3 - Drug Withdrawn; 4 - Not Applicable, 5- Unknown

Other Drug Details

Sr. No.	Past / Concomitant Drug	Trade name (Generic name)	Strength	Dosage form	Route	Expiry date	Batch Number	Indication	Start date, Time	Stop date, time
1										
2										
3										

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4										
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Details of Suspected Adverse Drug Reactions

Sr. No.	Adverse Event (Verbatim)	Severity*	Serious (Y/N)	Seriousness Criteria~	Onset Date of Event	Causality¥	Start date, Time	Stop date, time	Outcome±
1									
2									
3									
4									
5									
6									
7									
8									
9									

* 1 - Mild; 2 - Moderate; 3 – Severe

~ 1 - Fatal; 2 - Life-Threatening; 3 - Hospitalization or prolongation of hospitalization; 4 - Persistent or significant disability or Incapacity; 5 - Congenital Anomaly; 6 - Other IME

¥ 1 - Definite; 2 - Probable; 3 - Possible; 4 - Unlikely; 5 – Unclassifiable, 6 - Unascertainable

± 1 - Resolved; 2 - Resolved with sequelae; 3 - Resolving; 4 - Ongoing; 5 – Unknown

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Location where SAE event occurred		
	Other (Please specify)* -	

* 1 - Hospital; 2 - Home; 3 - Nursing home; 4 - Ambulatory Surgical Facility; 5 - Outpatient treatment facility; 6 - Outpatient diagnostic facility; 7 - Other (Please specify)

<i>In case of Hospitalization</i>	
Date of Admission	
Date of Discharge	

For Fatal Outcome			
Date of Death		Time Of Death	
Autopsy Report		Death Certificate	
Cause(s) of Death			



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Laboratory Tests Performed				
Sr. No.	Test Name	Date of Test performed	Result	Reference Range
1				
2				
3				
4				
5				
6				

Medical History / Concurrent Conditions				
Sr. No.	Description	Type*	Start Date	Stop Date
1				
2				
3				
*	1 - Past History (Surgical procedures); 2 - Concurrent Condition			

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Full description of reaction(s) including body site and severity. In addition, description of reported signs and symptoms

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