

SUSPECTED ADVERSE DRUG REACTION REPORTING FORM

For VOLUNTARY reporting of Adverse Drug Reactions by healthcare professionals

						AMC/ NCC Use only				
						AMC Report No.				
						Worldwide Unique No.				
A Patient Information						12. Relevant tests / laboratory data with dates				
1. Patient Initials _____		2. Age at time of Event or date of birth _____		3. Sex <input type="checkbox"/> M <input type="checkbox"/> F _____						
				4. Weight _____ Kgs						
B Suspected Adverse Reaction						13 Other relevant history including pre-existing medical conditions (e.g. allergies, race, pregnancy, smoking, alcohol use, hepatic/ renal dysfunction etc)				
5. Date of reaction stated (dd/mm/yyyy)										
6. Date of recovery (dd/mm/yyyy)										
7. Describe reaction or problem _____										
						14 Outcomes <input type="checkbox"/> Fatal <input type="checkbox"/> Recovering <input type="checkbox"/> Unknown <input type="checkbox"/> Continuing <input type="checkbox"/> Recovered <input type="checkbox"/> Other (specify)_____				
C. Suspected medication(s)										
S.No	8. Name (brand and /or generic name)	Manufacturer (if known)	Batch No./ Lot No. (if known)	Exp. Date (if known)	Dose used	Route used	Frequency	Therapy dates (if known give duration)		Reason for use of prescribed for
								Date started	Date stopped	
i										
ii										
iii										
iv										
Sl.No As per C	9. Reaction abated after drug stopped or dose reduced					10. Reaction reappeared after reintroduction				
	Yes	No	Unknown	NA	Reduced dose	Yes	No	Unknown	NA	If reintroduced dose
i										
ii										
iii										
iv										
11 Concomitant medical product including self medication and herbal remedies with therapy dates (exclude those used to treat reaction)						D. Reporter (see confidentiality section in first page)				
						15. Name and Professional Address : _____				
						Pin code _____ E-mail _____				
						Tel. No. (with STD code) _____				
						Occupation _____ Signature _____				
16. Causality Assessment					17. Date of this report (dd/mm/yyyy)					
18. Professional identification number _____										
19. Approving authority _____										