



## SUSPECTED ADVERSE REACTION REPORTING FORM

MCA-F-305/01

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1 PATIENT'S DETAILS							
1	Patient initials / number <i>(first, middle, last)</i>	Date of Birth <i>(day/month/year)</i>	Age	Sex		Weight (kg)	Reaction start date <i>(day/month/year)</i>
				<b>M</b>	<b>F</b>		
				<input type="checkbox"/>	<input type="checkbox"/>		
2 ADVERSE REACTION							
	Description of reaction <i>(including relevant tests/lab data)</i>					Outcome of reaction <i>(Tick all as appropriate)</i>	
						Patient died	<input type="checkbox"/>
						Life-threatening	<input type="checkbox"/>
		Involvement or prolonged hospitalisation	<input type="checkbox"/>				
		Persistent/ significant disability/incapacity	<input type="checkbox"/>				
		Congenital anomaly/ birth defect	<input type="checkbox"/>				
	Treatment of reaction					Recovered	<input type="checkbox"/>
3 SUSPECTED MEDICINE(S) <i>(including Biologicals, Herbal medicines)</i> <i>(state brand and generic name, batch/lot number, expiry date, name and address of manufacturer and attach product label/ sample if available)</i>							
						Did reaction abate after stopping medicine?	
						Yes <input type="checkbox"/>	No <input type="checkbox"/>
	Indication for use					Did reaction reappear after reintroduction?	
						Yes <input type="checkbox"/>	No <input type="checkbox"/>
	Dosage	Route	Date Started <i>(day/month/year)</i>		Date Stopped <i>(day/month/year)</i>		
4 CONCOMITANT MEDICATION (All medicines taken within the last 3 months including herbal medicines and self-medication, but excluded those used to treat reaction)							
	Brand or Generic Name	Dosage	Route	Date started <i>(day/month/year)</i>		Date stopped <i>(day/month/year)</i>	
4 SOURCE OF REPORT							
	Name and address of reporter					Date of this report <i>(day/month/year)</i>	
	Profession	Tel No/E-mail		Report Type			
				Initial <input type="checkbox"/>	Follow-up <input type="checkbox"/>		