

Notification Form for Adverse Events Following Immunization (AEFI)

Patient Information						
Name:			MOH Division:			
Age: <input type="checkbox"/> <input type="checkbox"/> months/years		Sex: Male <input type="checkbox"/> Female <input type="checkbox"/>		Telephone:		
Name & address of the Parent/Guardian:						
Information on the vaccine (primary suspected and other)						
Vaccine (Generic Name)	Vaccine (Trade name)*	Route	Dose (1 st , 2 nd , 3 rd , 4 th)	Batch/Lot Number	Expiry date	VVM Status (I, II, III, IV)
Diluent used: Yes <input type="checkbox"/> No <input type="checkbox"/> If "yes", Diluent batch/lot number						
Expiry date of Diluent						
<i>*Trade name is necessary only in private sector immunization</i>						
Place vaccine administered:					Date:	
Person vaccine administered: Doctor <input type="checkbox"/> PHNS/Nurse <input type="checkbox"/> PHM <input type="checkbox"/> PHI <input type="checkbox"/>					Time: am/pm	
Adverse Events						
Local Adverse Events Requiring investigation	Injection site abscess <input type="checkbox"/> BCG Lymphadenitis <input type="checkbox"/>					
Requiring investigation	Severe local reaction <input type="checkbox"/>					
CNS Adverse Events Requiring Investigation	Vaccine associated paralytic poliomyelitis <input type="checkbox"/> GBS <input type="checkbox"/>					
Requiring Investigation	Encephalopathy <input type="checkbox"/> Encephalitis <input type="checkbox"/> Meningitis <input type="checkbox"/>					
Other Adverse Events Requiring Investigation	Seizures Febrile <input type="checkbox"/> Seizures Afebrile <input type="checkbox"/>					
Other Adverse Events Requiring Investigation	Anaphylaxis <input type="checkbox"/> Persistent screaming <input type="checkbox"/> Osteitis / Osteomyelitis <input type="checkbox"/>					
Other Adverse Events Requiring Investigation	Hypotonic Hyporesponsive Episode <input type="checkbox"/> Toxic Shock Syndrome <input type="checkbox"/>					
Adverse Events Not Requiring Investigation	Allergic reaction <input type="checkbox"/> Arthralgia <input type="checkbox"/>					
Other Adverse Events	High fever (>39°C / 102°F) <input type="checkbox"/> Nodule at the injection site <input type="checkbox"/>					
Other Adverse Events	a)					
Other Adverse Events	b)					
<i>Instruction: Before reporting an AEFI, please refer to the definition for the relevant AEFI given in overleaf and make sure that reporting event agrees with the criteria stipulated in the definition</i>						
Date & Time onset of adverse event:						
Date & Time referring to medical care :						
Medical History/Other	Outcome					
	Hospitalized: Yes No If "Yes": Hospital:					
	BHT: Still in the hospital <input type="checkbox"/> Discharged <input type="checkbox"/>					
	Outcome: Recovered completely <input type="checkbox"/> Partially recovered <input type="checkbox"/> Death <input type="checkbox"/>					
Reporting source						
Date of the notification:		Institution & Designation:			Telephone:	
Name & Signature of the notifying officer/General Practitioner:						

(Medical Officers who attend any patient suffering from Adverse Effects Following Immunization shall notify in this form to the Medical Officer of Health the area of the patients residence)