



ADVERSE EVENT FOLLOWING IMMUNISATION REPORTING FORM

MCA-F-305/02

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Patient initials / number: <i>(first, middle, last)</i> Sex: M <input type="checkbox"/> F <input type="checkbox"/> Pregnant <input type="checkbox"/> Lactating <input type="checkbox"/> Date of birth <i>(day/month/year)</i> : __/__/____ OR age group at onset: 0 < 1 year <input type="checkbox"/> 1- 5 years <input type="checkbox"/> > 5years -18 years <input type="checkbox"/> >18 years –60 years <input type="checkbox"/> >60 years <input type="checkbox"/> Date patient notified event to health system <i>(day/month/year)</i> : __/__/____	Reporter's Name: Institution: Designation & Department: Region: Telephone & e-mail: Today's date <i>(day/month/year)</i> : __/__/____
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Health facility (or vaccination centre) name: _____

Vaccine							Diluent		
Name of vaccine (Generic)	*Brand Name and Name of Manufacturer	*Date of vaccination <i>(day/month/year)</i>	*Time of vaccination (24hrs)	Dose (1 st , 2 nd , etc.)	*Batch/ Lot number	Expiry date	*Batch/ Lot number	Expiry date	Time of reconstitution (24hrs)

Adverse event(s): <input type="checkbox"/> Severe local reaction <input type="checkbox"/> >3 days <input type="checkbox"/> beyond nearest joint <input type="checkbox"/> Seizures <input type="checkbox"/> febrile <input type="checkbox"/> afebrile <input type="checkbox"/> Abscess <input type="checkbox"/> Sepsis <input type="checkbox"/> Encephalopathy <input type="checkbox"/> Toxic shock syndrome <input type="checkbox"/> Thrombocytopenia <input type="checkbox"/> Anaphylaxis <input type="checkbox"/> Fever ≥38°C Other (specify) _____ Date <i>(day/month/year)</i> & Time (24 Hr/Min) AEFI started: __/__/____ __/____	Describe AEFI (Signs and symptoms):
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Serious: Yes No If Yes: Death* Life threatening Disability Hospitalisation Congenital anomaly
 Important medical event or Other (Specify) _____
Outcome: Recovering Recovered Recovered with sequelae Not Recovered Unknown
 *Date of death *(day/month/year)*: __/__/____ Autopsy done: Yes No Unknown If yes, provide report

Provide past medical history (including history of similar reaction or other allergies), concomitant medication and dates of administration (exclude those used to treat reaction) other relevant information (e.g. other cases). Use additional sheet if needed:

Date report received by MCA *(day/month/year)*: __/__/____ MCA AEFI number: _____

Comments: _____