

## STANDARD REPORTING FORM FOR ADVERSE EVENTS FOLLOWING IMMUNIZATION (AEFI)

<p><b>*Patient name or initials:</b></p> <p><b>*Patient's full Address:</b></p> <p>Telephone:</p> <p>Sex: <input type="checkbox"/> M <input type="checkbox"/> F (Pregnant - Trimester I <input type="checkbox"/> II <input type="checkbox"/> III <input type="checkbox"/> /Lactating <input type="checkbox"/>)</p> <p><b>*Date of birth (DD/MM/YYYY):</b> __/__/____</p> <p>OR Age at onset: <input type="checkbox"/><input type="checkbox"/> Years <input type="checkbox"/><input type="checkbox"/> Months <input type="checkbox"/><input type="checkbox"/><input type="checkbox"/> Days</p> <p>OR Age Group: <input type="checkbox"/> 0 &lt; 1 year <input type="checkbox"/> 1-5 years <input type="checkbox"/> &gt; 5 years - 18 years</p> <p><input type="checkbox"/> &gt; 18 years – 60 years <input type="checkbox"/> &gt; 60 years</p>	<p><b>*Reporter's Name:</b></p> <p>Institution:</p> <p>Designation &amp; Department:</p> <p>Address:</p> <p>Telephone &amp; e-mail:</p> <p>Date patient notified event to health system (DD/MM/YYYY): __/__/____</p> <p>Today's date (DD/MM/YYYY): __/__/____</p>
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Health facility (or vaccination centre) name:									
Vaccine							Diluent		
Name of vaccine (Generic)	*Brand Name incl. Name of Manufacturer	*Date of vaccination	*Time of vaccination	Dose (1 <sup>st</sup> , 2 <sup>nd</sup> , etc.)	*Batch/Lot number	Expiry date	*Batch/ Lot number	Expiry date	Time of reconstitution

<p><b>*Adverse event (s):</b></p> <p> <input type="checkbox"/> Severe local reaction    <input type="checkbox"/> &gt;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  <input type="checkbox"/> Other (specify).....         </p> <p>Date &amp; Time AEFI started (DD/MM/YYYY):          ____ / ____ / ____    <input type="checkbox"/><input type="checkbox"/> Hr <input type="checkbox"/><input type="checkbox"/> Min       </p>	<p>Describe AEFI (Signs and symptoms):</p>          
<p><b>*Serious: Yes / No ; ➔</b> If Yes <input type="checkbox"/> Death <input type="checkbox"/> Life threatening <input type="checkbox"/> Disability <input type="checkbox"/> Hospitalization <input type="checkbox"/> Congenital anomaly <input type="checkbox"/> Other important medical event (Specify _____)</p>	
<p><b>*Outcome:</b> <input type="checkbox"/> Recovering <input type="checkbox"/> Recovered <input type="checkbox"/> Recovered with sequelae <input type="checkbox"/> Not Recovered <input type="checkbox"/> Unknown</p> <p><input type="checkbox"/> Died If died, date of death (DD/MM/YYYY): ____ / ____ / ____ Autopsy done: <input type="checkbox"/> Yes <input type="checkbox"/> No <input type="checkbox"/> Unknown</p>	
<p>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). <i>Use additional sheet if needed :</i></p>	

First Decision making level to complete:

Investigation needed: <input type="checkbox"/> Yes <input type="checkbox"/> No	If yes, date investigation planned (DD/MM/YYYY): ____ / ____ / ____
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National level to complete:

Date report received at national level (DD/MM/YYYY): ____ / ____ / ____	AEFI worldwide unique ID :
Comments:	

*\*Compulsory field*