



ZAMBIA REPORTING FORM FOR ADVERSE EVENTS FOLLOWING IMMUNIZATION (AEFI)



AEFI Reporting ID:

<p>*Patient name:</p> <p>*Patient's full Address:.....</p> <p>Telephone:</p> <p>Sex: <input type="checkbox"/> M <input type="checkbox"/> F (if Female, <input type="checkbox"/> Pregnant <input type="checkbox"/> Lactating <input type="checkbox"/> N/A)</p> <p>*Date of birth (DD/MM/YYYY):</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"/> < 1 Year <input type="checkbox"/> 1 to 5 Years <input type="checkbox"/> >5 to 18 Years</p> <p><input type="checkbox"/> >18 to 60 Years <input type="checkbox"/> >60 Years</p>	<p>*Reporter's Name:</p> <p>Institution :</p> <p>Designation & Department:</p> <p>Address:</p> <p>Telephone & 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 st , 2 nd , etc.)	*Batch/Lot No.	Expiry date	*Batch/Lot No.	Expiry date	Time of reconstitution

<p>*Adverse event (s):</p> <p><input type="checkbox"/> Severe local reaction <input type="checkbox"/> >3 days <input type="checkbox"/> beyond nearest joint</p> <p><input type="checkbox"/> Seizures <input type="checkbox"/> febrile <input type="checkbox"/> afebrile</p> <p><input type="checkbox"/> Abscess</p> <p><input type="checkbox"/> Sepsis</p> <p><input type="checkbox"/> Encephalopathy</p> <p><input type="checkbox"/> Toxic shock syndrome</p> <p><input type="checkbox"/> Thrombocytopenia</p> <p><input type="checkbox"/> Anaphylaxis</p> <p><input type="checkbox"/> Fever ≥38°C</p> <p><input type="checkbox"/> Other (specify).....</p> <p>Date & Time AEFI started (DD/MM/YYYY):</p> <p>..... <input type="checkbox"/><input type="checkbox"/> Hr <input type="checkbox"/><input type="checkbox"/> Min</p>	<p>Describe AEFI (Signs and symptoms):</p>
<p>*Serious: Yes / No ; ➔ If Yes <input type="checkbox"/> Death <input type="checkbox"/> Life threatening <input type="checkbox"/> Disability <input type="checkbox"/> Hospitalization <input type="checkbox"/> Congenital anomaly</p>	
<p>*Outcome: <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) and other relevant information (e.g. other cases). Use additional sheet if needed :</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:
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Comments:

***Compulsory field**

All completed forms should be emailed to the following email: npvu@zamra.co.zm, aefizambia@gmail.com or

WhatsApp: **+260 956 521094**

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