

AEFI CASE INVESTIGATION REPORT

To: Dept. of Communicable Disease Surveillance & Control Fax No. 24601832		Institution:	
Date of Notification: / / (dd/mm/yy)			
1st name:	2nd name:	3rd name:	Tribes:
Sex: M / F DOB/Age:	Nationality:	MR2 No.:	OPD/IPD No.:
Village:	Wilayat:	Region:	Tel.No.
House No., Landmark:			
Name of the suspect vaccine:		No. of doses per vial:	Storage temp:
Batch No. of vaccine	Manufactured by:	Manufactured date: / / (dd/mm/yy)	Expiry date: / / (dd/mm/yy)
Diluent batch No. (if relevant)	Manufactured by:	Manufactured date: / / (dd/mm/yy)	Expiry date: / / (dd/mm/yy)
Syringe used (company name)	Manufactured by:	Manufactured date: / / (dd/mm/yy)	Expiry date: / / (dd/mm/yy)
Date of Injection: / / (dd/mm/yy)	Time of injection:	Institution:	Site of Injection:
Injection given by (name/designation)	Time of reactions: (onset of symptoms)	Time of recovery:	Admission: Yes / No if yes, Date of admission: / /
Outcome:			
II. Immunization History <i>(attach photocopy of child health card)</i>	III. Laboratory Findings (if relevant)	IV. Management:	
V. Medical History: H/o reactions to previous doses, drug allergies etc.		VI. Event summary:	
<input type="checkbox"/> Injection-site abscess (requiring drainage): <input type="checkbox"/> Yes <input type="checkbox"/> No		<input type="checkbox"/> Lymphadenitis: <input type="checkbox"/> Suppurative lymphadenitis <input type="checkbox"/> BCG-adenitis \geq 1.5 cm	
<input type="checkbox"/> Local Adverse Reactions <input type="checkbox"/> Pain, swelling and redness at the site of injection <input type="checkbox"/> Redness and/or swelling centered at the site of injection AND one of the following <input type="checkbox"/> Swelling beyond the nearest joint <input type="checkbox"/> Pain, redness and swelling of more than 3 days duration <input type="checkbox"/> Requires hospitalization <input type="checkbox"/> Injection site hypersensitivity reactions like rash/hives (urticaria) or itching at the injection site; hypotonic hypo-responsive episode			
<input type="checkbox"/> Systemic Adverse Reactions <input type="checkbox"/> Non-specific symptoms (eg. Fever above 39°C, Malaise, Headache, persistent screaming.... etc) Specify:			
<input type="checkbox"/> CNS adverse events: <input type="checkbox"/> Acute paralysis <input type="checkbox"/> Meningitis <input type="checkbox"/> Encephalopathy <input type="checkbox"/> Seizures (febrile or afebrile) <input type="checkbox"/> Encephalitis <input type="checkbox"/> Others (specify)		<input type="checkbox"/> Allergic Reactions: <input type="checkbox"/> Generalized urticaria <input type="checkbox"/> Breathing difficulty <input type="checkbox"/> Anaphylactic shock <input type="checkbox"/> Acute hypersensitivity reaction (anaphylactoid) <input type="checkbox"/> Toxic-shock syndrome <input type="checkbox"/> Disseminated BCG-adenitis <input type="checkbox"/> Hypotension <input type="checkbox"/> Other (specify)	
<input type="checkbox"/> Other Adverse Reactions (specify): eg. Osteitis, Osteomyelitis, Arthralgia etc.			

➤ Immunization History – attach photocopy of child health card

➤ If the child admitted – attach the discharge summary sheet