

ADVERSE EVENT FOLLOWING IMMUNISATION REPORTING FORM

MCA-F-305/02

Off Bertil Harding Highway, Kotu East, P.O. BOX 3162, Serekunda, email / website: info@mca.gm / www.mca.gm

Sex:	tials / number (first, I	years	Reporter's Name: Institution: Designation & Department: Region: Telephone & e-mail:						
Health facility (or vaccination centre) name:					Today's date (day/month/year)://				
Vaccine Diluent									
vaccine	Name of	Date of	Time of vaccination	Dose (1 st , 2 nd etc.)	Batch/Lot d _, number	Expiry date	Batch/Lot number	Expiry date	Time of reconstitution (24hrs)
Adverse event(s): ☐ Severe local reaction ☐ >3 days ☐ beyond nearest joint ☐ Seizures ☐ febrile ☐ afebrile ☐ Abscess ☐ Sepsis ☐ Encephalopathy ☐ Toxic shock syndrome ☐ Thrombocytopenia ☐ Anaphylaxis ☐ Fever≥38°C						EFI (Signs aı	nd symptor	ms):	
Other (specify)									
Date (day/month/year) & Time (24 Hr/Min) AEFI started://									
Serious: ☐ Yes ☐ No If Yes: ☐ Death* ☐ Life threatening ☐ Disability ☐ Hospitalisation ☐ Congenital anomaly ☐ Important medical event ☐ Other (Specify) ☐ Recovered ☐ Recovered with sequelae ☐ Not Recovered ☐ Unknown									
*Date of death (day/month/year):/ Autopsy done:									
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). <i>Use additional sheet if needed</i> :									
Date report received by MCA:					MCA AEFI number:				
Comments:									