



Vaccines Adverse Event Reporting Form (VAER)

| 1. Patient Details: | | | | | | |
|--|---|---|-------------------------|--|-------------------|-------------|
| Name: | Civil ID: | Nationality: | | | | |
| Date of birth: | Gender: <input type="checkbox"/> Male <input type="checkbox"/> Female | Age at vaccination:YearsMonths.....Days | | | | |
| Address: | Phone number : | E-mail : | | | | |
| Pregnant at time of vaccination?: <input type="checkbox"/> Yes <input type="checkbox"/> No <input type="checkbox"/> Unknown (If yes, describe the event, any pregnancy complications, and estimated due date if known) | | | | | | |
| 2. Health Facility (place or vaccination center) : | | | | | | |
| Name: | Type (Government , Private) | Address: | Phone Number : | | | |
| 3. Vaccine / Vaccination Program : | | | | | | |
| Name of vaccine | Type of vaccination program (Mandatory , Voluntary) | Date and time of vaccination | Route of administration | Dose (1st, 2nd, etc.) | Batch /Lot number | Expiry date |
| | | | | | | |
| 4. Description of adverse reaction: | | | | | | |
| Describe the adverse event(s), treatment, and outcome(s), if any: (symptoms, signs, time course, etc.) | | | | Result or outcome of adverse event(s): | | |
| Past medical history (including history of similar reaction or other allergies), concomitant medication and other relevant information (e.g. other cases). | | | | <input type="checkbox"/> Doctor or other healthcare professional office/clinic visit <input type="checkbox"/> Emergency room/department or urgent care <input type="checkbox"/> Hospitalization: Number of days (if known) Hospital name: <input type="checkbox"/> Prolongation of existing hospitalization (vaccine received during existing hospitalization) <input type="checkbox"/> Life threatening illness (immediate risk of death from the event) <input type="checkbox"/> Disability or permanent damage <input type="checkbox"/> Patient died Date of death: (...../...../.....) <input type="checkbox"/> Congenital anomaly or birth defect <input type="checkbox"/> Other | | |
| Medical tests and laboratory results related to the adverse event(s): (include dates) | | | | | | |
| Has the patient recovered from the adverse event(s)? <input type="checkbox"/> Yes <input type="checkbox"/> No <input type="checkbox"/> Unknown | | | | | | |
| Date reaction started: | | Date reaction stopped: | | | | |
| 5. Reporter data: (All data will be kept confidential) | | | | | | |
| Name: | Profession(Specialty): | Date: | | | | |
| Email: | Phone: | Signature: | | | | |

Please send this form by fax to: 24837245

It's easy to report online to adr_reporting@moh.gov.kw

Note: identities of reporter, patient and institution will remain confidential