

## IMMUNISATION ADVERSE EVENT REPORTING FORM

| OFFICE USE ONLY |
|-----------------|
| ACT CASE NO:    |
| TGA CASE NO:    |

| . DETAILS OF PERSON WHO EXPERIENCED THE ADVERSE EVENT  |
|--|
| ameDOB//   |
| ender M/F/ unknown (circle) Address  |
| tate Mobile Mobile   |
| a child, Parent/Guardian Name  |
| boriginal Torres Strait Islander Aboriginal & Torres Strait Islander Neither Not stated  |
|  |
| . PAST MEDICAL HISTORY   |
| ny known allergies?  |
| ny medical conditions?   |
| oes the person take any routine medications?   |
| ny prior reactions following immunisation? <b>Yes/No/Unknown</b> : If Yes, provide details   |
| eneral PractitionerPhone   |
|  |
| as the person ill before the vaccine was given? Yes/No If Yes, provide details   |
|  |
|  |
| . VACCINATION DETAILS  |
| chool program Yes/No   |
| accine Provider Name   |
| rovider Address Post code  |
| honeFaxEmail   |
|  |
| Vaccine Dose No Date Time Batch No Route/Site/Side (left or right)   |
| Brand/Type Administered Administered   |
|  |
|  |
|  |
|  |
|  |
|  |
| Vere any other vaccines given within 4 weeks prior to the adverse event? <b>NO/YFS:</b> If Yes, specify details:                   |
| /ere any other vaccines given within 4 weeks prior to the adverse event? <b>NO/YES:</b> If Yes, specify details:                   |
| /ere any other vaccines given within 4 weeks prior to the adverse event? NO/YES: If Yes, specify details:                          |
|  |
| . ADVERSE EVENT DETAILS  |
| . ADVERSE EVENT DETAILS Onset of event   |
| ADVERSE EVENT DETAILS Onset of event rate and time reaction occurred   |
| ADVERSE EVENT DETAILS  Onset of event  ate and time reaction occurred  unknown, time elapsed between vaccination and adverse event |
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| Management of event  |   |
|--|---|
| None/Nurse/GP/Hospital ED Was hospitalization required? yes/no   |   |
| Date of admission / / Date of discharge / /  |   |
| Detailed description of any treatment including medications  | _ |
|  |   |
|  |   |
|  |   |
|  |   |
| Outcome  |   |
| Have the symptoms resolved? yes/no/unknown If yes, time and date   | _ |
| If no, symptoms ongoing as of (time and date):   | - |
| Please describe ongoing symptoms   |   |
|  |   |
|  |   |
| 5. DETAILS OF PERSON REPORTING THIS ADVERSE EVENT  |   |
| Name Phone Date://   |   |
| AddressSuburbPost code   |   |
| Reporter type GP/ Medical Specialist/Medial Practitioner/Nurse RN/EN/Vaccinated person/parent/guardian/<br>Other   |   |
| <b>Consent statement</b> Please advise the parent/patient that they may be contacted if additional information is required. The contact details will be used for this purpose.         |   |
| If the parent/patient does not wish to be contacted, please fill out the following:  |   |
| I, the parent/patient do not agree to be contacted. Parent/patient signature The person has advised that they do not wish to be contacted (reporter to sign) Date                      |   |
| For verbal reports, indicate how consent was obtained:   |   |
| Please circle most relevant answer where appropriate.  On completion, fax this form to 02 5124 9307, or email to: <a href="mailto:immunisation@act.gov.au">immunisation@act.gov.au</a> |   |
| Office use only  |   |
| Is this considered a serious AEFI? Yes/No If yes, please specify   |   |
| Is follow up on patient required? Yes/No Immediately/next day/next 30 days/next 60 days  |   |
| Data report received Data scanned to TCA   |   |

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