

# COVID-19 vaccine: NSW Health adverse event following immunisation case notification form

## Instructions

- This form has been designed to collect initial clinical information regarding an Adverse Event Following Immunisation (AEFI) related to COVID-19 vaccination.
- The information provided will be used to investigate the reported adverse event following immunisation and will be reported to the Therapeutics Goods Administration to support vaccine safety surveillance.
- The form should be completed by a health professional and submitted by email on [MOH-covidaefi@health.nsw.gov.au](mailto:MOH-covidaefi@health.nsw.gov.au). Alternatively, cases can be notified by phone to the local Public Health Unit on **1300 066 055**.
- The Public Health Unit may contact you during business hours for further information regarding this case notification.

Vaccinated person's details		
Surname		
First name		
Date of birth		
Age		
Gender	Male	<input type="checkbox"/>
	Female	<input type="checkbox"/>
	Other	<input type="checkbox"/>
Street address		
Postcode		
Suburb		
State		
Phone number		
Aboriginal status: Is the person of Aboriginal or Torres Strait Islander origin?	<input type="checkbox"/> No <input type="checkbox"/> Yes, Aboriginal <input type="checkbox"/> Yes, Torres Strait Islander <input type="checkbox"/> Yes, both Aboriginal and Torres Strait Islander	

Reporter details	
Surname	
First name	

Practice name (if relevant)		
Street Address		
Suburb		
Postcode		
Phone – Landline (incl. area code)		
Mobile phone		
Email address		
Fax		
Date of report		
Reporter type	Medical practitioner	<input type="checkbox"/>
	Register nurse	<input type="checkbox"/>
	Vaccinated person	<input type="checkbox"/>
	Parent/guardian	<input type="checkbox"/>
	Other (specify)	
Is the reporter the vaccination provider	No	<input type="checkbox"/>
	Yes	<input type="checkbox"/>

Vaccine details (if known)							
Vaccine (brand name)	Dose no.	Batch number	Date given	Time given	Route of administration	Injection site	
					IM <input type="checkbox"/> SC <input type="checkbox"/> ID <input type="checkbox"/>	RL <input type="checkbox"/> RA <input type="checkbox"/>	LL <input type="checkbox"/> LA <input type="checkbox"/>
					IM <input type="checkbox"/> SC <input type="checkbox"/> ID <input type="checkbox"/>	RL <input type="checkbox"/> RA <input type="checkbox"/>	LL <input type="checkbox"/> LA <input type="checkbox"/>

Adverse Event Details	
Date of onset	
Time of onset	
Description of events, including timeline	
Management of event (tick as many as apply)	None <input type="checkbox"/>
	Nurse assessment <input type="checkbox"/>
	GP assessment <input type="checkbox"/>
	Hospital emergency department <input type="checkbox"/>

	Hospital admission (specify number of days and date of discharge)	
	Unknown	<input type="checkbox"/>
	Other (describe)	

Please specify the treatment/care provided (e.g. antibiotics, adrenaline, advice, counselling, etc.):	
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**Outcome**

Have the symptoms resolved?	Yes (specify date and time resolved)	Date	
		Time	
	No (Symptoms are ongoing as of)	Date	
		Time	
		Describe	
	Unknown	<input type="checkbox"/>	