Significant events in COVID-19 vaccination practice in Australia

Year	Month	Intervention
2020	October	Vaxzevria (adenovirus viral vector) COVID-19 vaccine granted provisional determination by the Therapeutic Goods Administration (TGA), making it eligible for provisional registration
	October	Comirnaty (mRNA) COVID-19 vaccine granted provisional determination by the TGA, making it eligible for provisional registration
	November	COVID-19 vaccine Janssen (adenovirus viral vector) granted provisional determination by the TGA, making it eligible for provisional registration
2021	January	Nuvaxovid (protein-based) COVID-19 vaccine granted provisional determination by the TGA, making it eligible for provisional registration Comirnaty provisionally registered for use in individuals aged ≥16 years
	February	Vaxzevria provisionally registered for use in individuals aged ≥18 years Nationally funded COVID-19 vaccination program began. The rollout was carried out in phases, with population groups prioritised according to ATAGI advice *Phase 1a* • Quarantine and border workers • Frontline healthcare workers • Aged care and disability care staff • Aged care and disability care residents *Phase 1b* • Healthcare workers currently employed and not included in Phase 1a • Household contacts of quarantine and border workers • Critical and high-risk workers who are currently employed, including defence, police, fire, emergency services and meat processing • Essential outbound travellers with a travel exemption • Elderly people aged ≥80 years • Elderly people aged ≥70 years • Aboriginal and Torres Strait Islander people aged ≥50 years • Adults with an underlying medical condition or significant disability *Phase 2a* • People aged ≥50 years • Aboriginal and Torres Strait Islander people aged 16–49 years • Other critical and high-risk workers *Phase 2b* • People aged 16–49 years *Phase 3* • People aged less than 16 years
	March	Phase 1b began
	April	Recommendations for use of Vaxzevria changed due to an association with thrombosis with thrombocytopenia syndrome (TTS): • Comirnaty is preferred over Vaxzevria in individuals aged <50 years • Those who have received their first dose of Vaxzevria with no TTS could still receive their second dose of Vaxzevria
	May	Phase 2a began in people aged ≥50 years

2021 (cont.)	June	Recommendations for use of Vaxzevria changed due to further emerging data on the association with TTS: • Comirnaty is preferred over Vaxzevria in individuals aged <60 years
	June	COVID-19 vaccine Janssen provisionally registered for use in individuals aged ≥18 years
	June	Spikevax (mRNA) COVID-19 vaccine granted provisional determination by the TGA, making it eligible for provisional registration
	July	Comirnaty indication age extended to include use in individuals aged ≥12 years
	August	Spikevax provisionally registered for use in individuals aged ≥18 years
	September	Spikevax indication age extended to include use in individuals aged ≥12 years
	October	A 3rd primary dose of COVID-19 vaccine recommended in severely immunocompromised populations 2 to 6 months after the 2nd dose of vaccine. An mRNA vaccine is preferred to Vaxzevria for this 3rd dose Comirnaty provisionally registered for a booster dose 6 months after the 2nd dose in immunocompetent individuals aged ≥18 years A booster dose was recommended for immunocompetent individuals aged ≥18 years who had their primary COVID-19 vaccine course ≥6 months ago. The highest-priority groups recommended to receive booster doses are those with risk factors for severe COVID-19 and/or those at increased occupational risk of COVID-19. Comirnaty is preferred irrespective of the primary COVID-19 vaccine used
	December	Comirnaty indication age extended to include use in individuals aged ≥5 years Spikevax provisionally registered for a booster dose 6 months post-dose 2 in immunocompetent individuals aged ≥18 years *12 December:* Due to ongoing transmission of Omicron and Delta variants, the recommended minimum interval between the primary course and the booster dose was shortened from 6 months to 5 months *24 December:* The recommended minimum interval between the primary course and the booster dose was shortened from 5 months to 4 months and, when capacity permits (in late January 2022), 3 months
2022	January	Nuvaxovid provisionally registered for use in individuals aged ≥18 years for the primary course Comirnaty provisionally registered for a booster dose 6 months post-dose 2 in individuals aged 16–17 years Funded vaccination roll-out began in children aged 5–11 years Severely immunocompromised children aged 5–11 years recommended to receive a 3rd primary dose of COVID-19 vaccine 2 to 6 months after their 2nd dose, in line with other severely immunocompromised age cohorts People aged ≥18 years who received a 3-dose primary course due to severe immunocompromise recommended to receive a booster (4th) dose ≥4 months after their 3rd dose
	February	A booster dose was recommended for adolescents aged 16–17 years who had their primary COVID-19 vaccine course ≥3 months ago Spikevax indication age extended to include use in individuals aged ≥6 years Vaxzevria provisionally registered for a booster dose 6 months post dose 2, in individuals aged ≥18 years

2022 (cont.)	March	A winter booster dose was recommended for the following groups ≥4 months after the person has received their first booster dose, or ≥4 months after a confirmed SARS-CoV-2 infection, if infection occurred since the person's first COVID-19 booster dose: • adults aged ≥65 years • residents of aged care or disability care facilities • individuals aged ≥16 years with severe immunocompromise (as defined in the ATAGI statement on the use of a 3rd primary dose of COVID-19 vaccine in individuals who are severely immunocompromised) • Aboriginal and Torres Strait Islander people aged ≥50 years
	April	Comirnaty vaccine provisionally registered for use as a booster dose for individuals aged 12–15 years
	May	The winter booster recommendation was expanded to include people aged 16–64 years with medical risk conditions or disability
	June	Nuvaxovid (Novavax) vaccine provisionally registered for use as a booster in individuals aged ≥18 years Recommendations for boosters in high-risk adolescents aged 12–15 years
	July	Nuvaxovid (Novavax) vaccine indication age extended to include use in individuals aged ≥12 years Spikevax vaccine indication age extended to include use in individuals aged ≥6 months A winter booster dose was recommended for individuals aged >50 years, and individuals aged 30–49 years ,could receive a winter booster after discussion with their regular medical provider to review their individual health needs and the benefits and risks of a second booster dose A 3-month interval between a recent SARS-CoV-2 infection or the first booster dose and a winter booster dose was recommended
	August	Nuvaxovid (Novavax) vaccine provisionally registered for use in individuals aged 12–17 years for the primary course Spikevax bivalent (original/omicron BA.1) vaccine provisionally registered for use as a booster dose in individuals aged ≥18 years
	September	Comirnaty vaccine provisionally registered for use as a booster dose in individuals aged 5–11 years Comirnaty vaccine indication age extended to include use in individuals aged ≥6 months
	October	Spikevax vaccine provisionally registered for use as a booster dose in individuals aged ≥12 years Spikevax vaccine was recommended as a paediatric booster dose in children aged 5–11 years with high-risk medical conditions or disability Comirnaty bivalent (original/omicron BA.1) vaccine provisionally registered for use as a booster dose in individuals aged ≥18 years

2023	February	A booster dose was recommended for the following groups if their last vaccine dose or confirmed infection was ≥6 months:
		• all adults aged ≥65 years
		• adults aged 18–64 years who have medical comorbidities or disability
		• the following groups, after a risk–benefit assessment:
		  ○ all adults aged 18–64 years without risk factors for severe COVID-19
		  ○ children aged 5–17 years with high-risk medical conditions or disability
		Comirnaty bivalent (original/omicron BA.4/5) vaccine provisionally registered for use as a booster dose in adolescents and individuals aged ≥12 years
		Moderna bivalent (original/omicron BA.4/5) vaccine provisionally registered for use as a booster dose in adolescents and individuals aged ≥12 years