

# Vaccinations in pregnancy

## Optimising maternal and infant protection in 2026

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In an era marked by rising vaccine hesitancy and resurgent infectious threats, maternal vaccination stands as a powerful safeguard for the mother and for the infant in their first six months of life. This article unpacks the latest recommendations, demystifies safety concerns and explores bold strategies to improve vaccine uptake across the preconception, pregnancy and breastfeeding continuum, providing clinicians with practical tools to champion vaccination and protect Australia's next generation.

Vaccination plays a crucial role in safeguarding both expectant mothers and their newborns from various vaccine preventable diseases (VPDs).<sup>1,2</sup> By administering specific vaccines preconception, during pregnancy and while breastfeeding, immunisation confers direct protection to the pregnant woman and subsequent passive immunity to their child, thereby reducing morbidity and mortality rates among these vulnerable populations.<sup>3</sup>

For healthcare professionals, a key indispensable resource is the Australian Immunisation Handbook, the national clinical guideline for vaccination recommendations. A list of resources is provided in Box 1. Using the best scientific evidence available, the Australian Technical Advisory Group on Immunisation (ATAGI), in



### KEY POINTS

- Maternal vaccination protects both the pregnant woman and her infant in their first six months of life, with antibody transfer in utero and through breast milk offering vital early life protection against severe infections.
- Preconception vaccination offers a critical opportunity to address immunity gaps, particularly for rubella, varicella and hepatitis B, reducing the risk of serious congenital infections.
- Vaccines recommended during pregnancy in Australia include influenza, pertussis (diphtheria-tetanus-acellular pertussis) and, most recently, respiratory syncytial virus, with the timing optimised to maximise passive immunity for the newborn.
- Vaccination during breastfeeding is safe and beneficial, with most vaccines posing no risk to the infant and potentially enhancing passive immune protection.
- Barriers to vaccine uptake, such as misinformation, access issues and variable healthcare provider engagement, must be addressed through co-ordinated public health capacity building and communication, integrated antenatal vaccine delivery and digital health solutions.
- Future maternal vaccines, including those for group B streptococcus and cytomegalovirus, are on the horizon and sustained investment in research, surveillance and trust-building will be key to success.

collaboration with other key stakeholders, develop the handbook's recommendations for the safest and most effective use of vaccines.<sup>4</sup> Augmenting the ongoing development of the handbook is the National Centre for Immunisation Research and Surveillance (NCIRS), with many of its collaborative works including the Sharing Knowledge About Immunisation (SKAI) and AusVaxSafety programs, providing reliable information to support healthcare professionals in discussing vaccination and VPDs with their patients.

Most, but not all, recommended vaccines are funded and delivered through the National Immunisation Program (NIP), a joint Commonwealth, State and Territory government initiative, acknowledged as one of the world's most comprehensive immunisation programs.<sup>5-7</sup>

Recent developments underscore the critical need to optimise maternal vaccination uptake. These include the national launch of the Respiratory Syncytial Virus Mother and Infant Protection

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Program (RSV-MIPP) on 3 February 2025, providing free maternal RSV vaccination under the NIP, alongside state- and territory-funded infant nirsevimab programs,

heightened vaccine hesitancy following the coronavirus disease 2019 (COVID-19) pandemic, and a profound resurgence of pertussis.<sup>8-13</sup> Australia recorded 57,146

pertussis notifications in 2024; the highest annual total since national monitoring began in 1991, with more than 82,000 cases cumulatively across 2024–25.<sup>14-16</sup> Encouragingly, early 2026 data indicate pertussis activity has returned to inter-seasonal levels, although vaccination during pregnancy remains the most effective strategy to protect infants during their highest-risk period.<sup>16</sup> In addition, 2026 marks Australia's transition from quadrivalent to trivalent seasonal influenza vaccines and the introduction of an intranasal live attenuated influenza vaccine (LAIV) for children; developments with implications for maternal counselling.<sup>17</sup>

Pregnant women may also be recommended to receive additional vaccines if they are at increased risk of other specific VPDs because of factors including occupation, personal behaviour or medical conditions.<sup>1</sup> This article offers healthcare professionals an in-depth review of vaccination during preconception, pregnancy and breastfeeding, focusing on current recommendations, safety profiles, barriers to uptake and strategies to improve immunisation rates among pregnant women.

### Rationale for maternal vaccination

Maternal vaccination is a crucial public health strategy designed to prevent or reduce the severity of maternal infection, fetal or congenital infection and infant infection during their first six months of life. This is achieved through a co-ordinated immunisation approach for every pregnancy. The scheduling of vaccine administration varies based on the at-risk individual (mother, fetus and/or infant), the period when the risk of infection is greatest or becomes significant, and the length of protective immunity following vaccination.<sup>18</sup>

### Immunological changes in pregnancy

Pregnancy induces complex immunological adaptations that facilitate fetal tolerance while maintaining maternal immune competence.<sup>19</sup> These adaptations include modulation of innate and adaptive immune responses, increased anti-inflammatory

## 1. RESOURCES FOR GPs

- **Australian Immunisation Handbook from the Australian Technical Advisory Group on Immunisation (ATAGI):** <https://immunisationhandbook.health.gov.au>
- **National Centre for Immunisation Research and Surveillance (NCIRS):** <https://ncirs.org.au>
- **Sharing Knowledge About Immunisation from the NCIRS:** <https://skai.org.au>
- **AusVaxSafety from the NCIRS:** <https://www.ausvaxsafety.org.au>
- **National Immunisation Program Schedule:** <https://health.gov.au/resources/publications/national-immunisation-program-schedule>
- **ATAGI General Practitioner Bulletin (commenced November 2025):** <https://www.health.gov.au/committees-and-groups/atagi>
- **COVID-19 vaccination shared decision making guide for women who are pregnant, breastfeeding or planning pregnancy:** <https://www.health.gov.au/resources/publications/covid-19-vaccination-shared-decision-making-guide-for-women-who-are-pregnant-breastfeeding-or-planning-pregnancy>

### Resources from the Australian Immunisation Handbook on vaccine considerations in special risk groups planning pregnancy

- **Specified medical conditions associated with increased risk of influenza disease and severe outcomes:** <https://immunisationhandbook.health.gov.au/resources/tables/table-specified-medical-conditions-associated-with-increased-risk-of-influenza-disease-and-severe-outcomes>
- **Risk conditions for pneumococcal vaccination and eligibility for National Immunisation Program funding:** <https://immunisationhandbook.health.gov.au/resources/tables/table-risk-conditions-for-pneumococcal-vaccination-and-eligibility-for-nip-funding>
- **Specified immunocompromised conditions:** <https://immunisationhandbook.health.gov.au/contents/vaccination-for-special-risk-groups/vaccination-for-people-who-are-immunocompromised>
- **Aboriginal and Torres Strait Islander people:** <https://immunisationhandbook.health.gov.au/contents/vaccination-for-special-risk-groups/vaccination-for-aboriginal-and-torres-strait-islander-people>
- **Occupational risks:** <https://immunisationhandbook.health.gov.au/contents/vaccination-for-special-risk-groups/vaccination-for-people-at-occupational-risk>
- **International travellers:** <https://immunisationhandbook.health.gov.au/contents/vaccination-for-special-risk-groups/vaccination-for-international-travellers>
- **Individuals who have had an adverse event following immunisation:** <https://immunisationhandbook.health.gov.au/contents/vaccination-for-special-risk-groups/vaccination-for-people-who-have-had-an-adverse-event-following-immunisation>
- **Individuals who have recently received normal human immunoglobulin and other blood products:** <https://immunisationhandbook.health.gov.au/contents/vaccination-for-special-risk-groups/vaccination-for-people-who-have-recently-received-normal-human-immunoglobulin-and-other-blood-products>
- **High-risk groups:** <https://immunisationhandbook.health.gov.au/contents/vaccination-for-special-risk-groups/vaccination-for-other-groups-in-mates-men-who-have-sex-with-men-people-who-inject-drugs-and-sex-workers>
- **Migrants, refugees and people seeking asylum:** <https://immunisationhandbook.health.gov.au/contents/vaccination-for-special-risk-groups/vaccination-for-migrants-refugees-and-people-seeking-asylum-in-australia>
- **Bleeding disorders:** <https://immunisationhandbook.health.gov.au/contents/vaccination-for-special-risk-groups/vaccination-for-people-with-bleeding-disorders>

## 2. PRECONCEPTION VACCINATION ASSESSMENT<sup>1,50,51</sup>

- Pre-vaccination screening checklist. See: <https://immunisationhandbook.health.gov.au/resources/tables/table-pre-vaccination-screening-checklist><sup>51</sup>
- Check prior immunisation history on the AIR to identify potentially missing or incomplete vaccinations
- Consider occupational and lifestyle factors (including travel) that may increase the risk of specific infections
- Evaluate existing medical conditions that may influence vaccine recommendations and necessity
- Seek further expert advice, when necessary, from a specialist immunisation clinic, a medical practitioner experienced in vaccination, the immunisation section in state or territory health authorities, or a local public health unit.

Abbreviations: AIR = Australian Immunisation Register.

cytokine activity and alterations in cellular immunity.<sup>20</sup> Although there is little robust evidence demonstrating that pregnant women are more susceptible to VPDs including influenza and COVID-19, multiple studies confirm they are at increased risk of severe complications, with higher rates of hospitalisation, intensive care unit admission and maternal mortality.<sup>21-27</sup>

### Benefits of maternal vaccination

#### Protection of the mother

Maternal vaccination reduces the risk of severe illness, hospitalisation and adverse pregnancy outcomes.<sup>28</sup> Strong, robust evidence demonstrates that influenza vaccination during pregnancy decreases the likelihood of influenza-related complications, including pneumonia and preterm birth.<sup>29,30</sup>

#### Passive immunity for the infant

One of the primary advantages of maternal immunisation is the transplacental transfer of antibodies, which provides passive immunity to the infant that is sustained during the vulnerable neonatal period.<sup>31</sup> Immunoglobulin G is actively transported across the placenta via the neonatal Fc receptor,

reaching peak concentrations in the third trimester.<sup>32</sup> Studies have demonstrated that maternal pertussis vaccination results in a 70 to 90% reduction in pertussis cases and an up to 90.5% reduction in related hospitalisations among infants younger than 3 months old.<sup>33-36</sup> Similarly, seasonal influenza vaccination during pregnancy reduces the risk of infection and related complications for mothers and their newborns.<sup>37-41</sup>

#### Reduction in disease burden

Widespread maternal vaccination programs have led to significant reductions in neonatal morbidity and mortality. For instance, global implementation of maternal tetanus immunisation programs has contributed to the near elimination of neonatal tetanus in many countries.<sup>42,43</sup>

#### Public health impact

Maternal vaccination not only benefits individuals but also reduces the overall burden of VPDs within the broader community, as herd immunity extends to unvaccinated individuals, including premature infants and immunocompromised neonates.<sup>44,45</sup> In Australia, sustained maternal pertussis immunisation programs have led to a decline in infant pertussis-related hospitalisations.<sup>46</sup> With the national rollout of maternal RSV vaccination through the RSV-MIPP since February 2025, early program data indicate an estimated 140,000 infants had been protected in the program's first year through either maternal vaccination or infant nirsevimab, delivering meaningful reductions in RSV-related neonatal morbidity and mortality while being cost effective, though uptake of the complementary infant nirsevimab program has varied markedly across jurisdictions.<sup>47-49</sup>

### Preconception vaccination: ensuring immunity before pregnancy

Women planning pregnancy should have their vaccination history thoroughly assessed, including using the Australian Immunisation Register (AIR; available at: <https://www.servicesaustralia.gov.au/>

## 3. SEROLOGICAL TESTING IN CASES OF UNCERTAIN VACCINATION OR INFECTION HISTORY

Pathology request form details to include the following (where appropriate)

- Hepatitis B serology (HBsAg, HBsAb, HBcAb)
- Measles serology (IgG antibodies)
- Mumps serology (IgG antibodies)
- Rubella serology (IgG antibodies)
- VZV serology (IgG antibodies)

Clinical notes should specify the purpose

- 'To assess preconception immune status'

Abbreviations: HBcAb = hepatitis B core antibody; HBsAb = hepatitis B surface antibody; HBsAg = hepatitis B surface antigen; IgG = immunoglobulin G; VZV = varicella zoster virus.

australian-immunisation-register) as part of comprehensive preconception care (Box 2).<sup>1,50,51</sup> This process is crucial to ensure optimal protection against VPDs that could result in significant pregnancy complications or congenital infections. Furthermore, given that live vaccines are generally contraindicated during pregnancy because of a theoretical risk of transmission,<sup>1,52</sup> the preconception period provides an important opportunity for these vaccines to be administered without undue concern for adverse fetal effects.

Additionally, individuals residing in the same household should also have their vaccination status reviewed and updated to minimise the risk of transmitting VPDs to the prospective mother and the newborn.<sup>1</sup>

### Managing uncertain vaccination or infection history

If a woman's vaccination history or infection status is unclear, serological testing should be performed to verify immunity against key pathogens (Box 3).<sup>1</sup> Should serological testing reveal insufficient immunity, the necessary vaccines should be administered, taking into consideration recommended waiting periods before conception.

### Recommended preconception vaccines

Women planning pregnancy should be up to date with the following vaccines, as summarised in Table 1.<sup>1,34,53-62</sup>

**Hepatitis B**

Vaccination is recommended for all hepatitis B-naïve women to prevent acquisition and mother-to-infant transmission.<sup>53</sup>

**Measles, mumps and rubella**

Infection with any of these viruses during pregnancy can have severe consequences such as congenital rubella syndrome.

Women lacking confirmed immunity should receive two doses of measles, mumps and rubella (MMR) live vaccine, spaced four weeks apart, and should avoid

**TABLE 1. RECOMMENDED PRECONCEPTION VACCINES<sup>1,30,53-59\*</sup>**

Vaccine-preventable disease	If uncertain history of vaccination or disease, check serology	NIP funded	Vaccine brand (type)	Dosage and timing
Hepatitis B <sup>53</sup>	✓	No	<ul style="list-style-type: none"> <li>Engerix-B<sup>†</sup></li> <li>(recombinant subunit)</li> <li>H-B-Vax II<sup>†</sup></li> <li>(recombinant subunit)</li> </ul>	<ul style="list-style-type: none"> <li>Three doses at zero, one and six months OR</li> <li>Three doses at zero, one and four months OR</li> <li>Three doses at zero, two and four months</li> </ul>
Measles <sup>54</sup>	✓	Only for adolescents and adults aged <20 years who have been incompletely vaccinated or do not have evidence of immunity to either measles, mumps or rubella	<ul style="list-style-type: none"> <li>M-M-R II<sup>‡</sup></li> <li>(live attenuated)</li> <li>Priorix<sup>‡</sup></li> <li>(live attenuated)</li> </ul>	<ul style="list-style-type: none"> <li>Two doses, four weeks apart</li> <li><b>Avoid pregnancy within 28 days of receiving this live vaccine</b></li> </ul>
Mumps <sup>55</sup>	✓			
Rubella <sup>56</sup>	✓			
Varicella – chickenpox <sup>57</sup>	✓	Only for adolescents and adults aged <20 years who have been incompletely vaccinated or do not have evidence of immunity to varicella	<ul style="list-style-type: none"> <li>Varilrix<sup>‡</sup></li> <li>(live attenuated)</li> <li>Varivax Refrigerated<sup>‡</sup></li> <li>(live attenuated)</li> </ul>	<ul style="list-style-type: none"> <li>Two doses, four to eight weeks apart</li> <li><b>Avoid pregnancy within 28 days of receiving this live vaccine</b></li> </ul>
COVID-19 <sup>58</sup>	✗	COVID-19 vaccines are free for everyone in Australia regardless of Medicare or visa status	<ul style="list-style-type: none"> <li>Comirnaty JN.1<sup>§</sup> (mRNA)</li> <li>Comirnaty Omicron LP8.1<sup>§</sup> (mRNA)</li> </ul>	<ul style="list-style-type: none"> <li>Single dose</li> </ul>
Influenza <sup>30</sup>	✗	Only for adults aged ≥18 years with specified medical risk conditions <sup>#1</sup>	<ul style="list-style-type: none"> <li>Vaxigrip (inactivated, trivalent)</li> <li>Flucelvax (inactivated, trivalent, cell-based)</li> </ul>	<ul style="list-style-type: none"> <li>Single dose</li> </ul>
Pneumococcal <sup>II</sup> <sup>59</sup>	✗	Only for people with specified medical risk conditions <sup>**</sup> Vaxneuvance (15vPCV) and Prevenar 20 (20vPCV) are not currently NIP-funded	<ul style="list-style-type: none"> <li>Prevenar 13 (conjugate)</li> <li>Vaxneuvance (15vPCV) (conjugate)</li> <li>Prevenar 20 (20vPCV) (conjugate)</li> <li>Pneumovax 23 (23vPPV)</li> <li>(polysaccharide)</li> </ul>	<ul style="list-style-type: none"> <li>Single dose of a pneumococcal conjugate vaccine (13vPCV, 15vPCV, or 20vPCV) followed by – Two doses of the pneumococcal polysaccharide vaccine (23vPPV), first dose two to 12 months after the PCV, then second dose five years after the first 23vPPV dose</li> </ul>

Abbreviations: 13vPCV = 13-valent pneumococcal conjugate vaccine; 15vPCV = 15-valent pneumococcal conjugate vaccine; 20vPCV = 20-valent pneumococcal conjugate vaccine; 23vPPV = 23-valent pneumococcal polysaccharide vaccine; COVID-19 = coronavirus disease 2019; MMR = measles, mumps and rubella; MMRV = measles, mumps, rubella and varicella; mRNA = messenger ribonucleic acid; NIP = National Immunisation Program; PCV = pneumococcal conjugate vaccine.

\* Also ensure all age-related and relevant specified medical risk group recommended vaccinations are up to date.

<sup>†</sup> Adult formulation. Switching hepatitis B vaccine brands is not recommended.<sup>53</sup>

<sup>‡</sup> Switching of MMR vaccine brands is permitted. Measles-mumps-rubella-varicella (MMRV) vaccine is not recommended for use in people ≥14 years of age as no data are available on safety, immunogenicity or efficacy in this age group. If a person ≥14 years of age is inadvertently given a dose of MMRV vaccine, this dose does not need to be repeated.

<sup>§</sup> Formulation for individuals ≥12 years of age. Adults ≥18 years of age without severe immunocompromise who have not previously received a COVID-19 vaccine are recommended a single primary dose. Adults aged 18–64 years without severe immunocompromise can consider further doses every 12 months based on a risk-benefit assessment, such as the presence of other medical conditions that may increase the risk of severe COVID-19.<sup>58,63</sup>

<sup>#</sup> In some states and territories, influenza vaccines for all persons ≥6 months of age may also be provided free of charge to residents, regardless of NIP eligibility. From 2026, all seasonal influenza vaccines in Australia are trivalent formulations (TIVs); the previous quadrivalent brand suffixes “Tetra” and “Quad” have been discontinued for 2026.<sup>17</sup> The live attenuated vaccine (FluMist) is contraindicated in pregnancy.

<sup>1</sup> See Box 1 for a resource on specified medical conditions associated with increased risk of influenza disease and severe outcomes.

<sup>II</sup> As of April 2026, the optimal adult pneumococcal vaccination program for Australia remains under review: the Australian Immunisation Handbook pneumococcal chapter (updated 19 January 2026) now includes interim recommendations for the use of extended-valency vaccines (Vaxneuvance [15vPCV] and Prevenar 20 [20vPCV] in adults, although these are not currently NIP-funded for adults at risk. Note that from 1 September 2025, Prevenar 20 replaced Prevenar 13 and Pneumovax 23 on the NIP childhood schedule.<sup>59</sup>

<sup>\*\*</sup> See Box 1 for a resource on risk conditions for pneumococcal vaccination.

Trimester 1												Trimester 2												Trimester 3															
1	2	3	4	5	6	7	8	9	10	11	12	13	14	15	16	17	18	19	20	21	22	23	24	25	26	27	28	29	30	31	32	33	34	35	36	37	38	39	40
												Antibody transfer												Peak antibody transfer															
<b>Influenza</b> Recommended single dose at any stage of pregnancy* <i>Vaxigrip or Flucelvax</i>																																							
												<b>Pertussis</b> Recommended single dose in each pregnancy, ideally between 20 and 32 weeks (may be given up until delivery) <i>Boostrix or Adacel</i>																											
																								<b>RSV</b> Recommended single dose, from 28 weeks† <i>Abrysvo‡</i>															
<b>COVID-19</b> Unvaccinated pregnant women only§ Recommended single dose at any stage of pregnancy¶ <i>Comirnaty JN.1 or Comirnaty LP8.1</i>																																							

**Figure.** Recommended vaccines during pregnancy for 2026.<sup>1,9,17,30,58,64,65</sup>

Abbreviations: COVID-19 = coronavirus disease 2019; RSV = respiratory syncytial virus.

The National Immunisation Program funded vaccine brands are provided above, in addition to the COVID-19 vaccines, which are free for everyone in Australia regardless of Medicare or visa status. Additional brands of influenza vaccines are available but may incur a cost to the patient. From 2026, all Australian seasonal influenza vaccines are trivalent formulations.<sup>17</sup> The intranasal live attenuated influenza vaccine (FluMist) is contraindicated in pregnancy.

\* Women who received the previous year's seasonal influenza vaccine early in their pregnancy are advised to receive the current seasonal influenza vaccine (when it becomes available) later in the same pregnancy. Women who received vaccine before becoming pregnant should be revaccinated during pregnancy to protect the unborn infant.<sup>30</sup>

† If administered less than 2 weeks before giving birth, the newborn will not be adequately protected.<sup>9</sup> If Abrysvo is inadvertently administered before 28 weeks' gestation, it does not need to be re-administered.

‡ Abrysvo (RSVpreF) is the only RSV vaccine approved for use in pregnant women; Arexvy (RSVpreF3) is not approved for pregnancy.<sup>9</sup>

§ Unvaccinated pregnant women are recommended to receive a primary course of COVID-19 vaccine. Dosing and vaccine choice recommendations are the same as for non-pregnant people of the same age.<sup>58</sup> Unless a woman is otherwise eligible, a dose of COVID-19 vaccine is not routinely recommended in pregnancy.<sup>58</sup> Pregnant women who have previously been vaccinated can discuss with their healthcare provider whether to have a further dose during their pregnancy, based on an individual risk-benefit assessment. Although the latest mRNA COVID-19 vaccines (Comirnaty JN.1 and LP.8.1-based vaccines) have not been formally studied in pregnant women, the Australian Technical Advisory Group on Immunisation considers them suitable and safe for use.<sup>58</sup>

¶ People with severe immunocompromise conditions, who are over six months of age or older are recommended two primary doses and are eligible for a third primary dose based on an individual risk-benefit assessment.<sup>58</sup>

Adapted from the National Centre for Immunisation Research and Surveillance. Vaccine recommendations for pregnant women – a guide for health professionals (updated 27 February 2026).<sup>65</sup>

conception for at least 28 days after the final dose.<sup>54,56</sup>

### Varicella (chickenpox)

Vaccination is recommended for seronegative women to prevent congenital varicella syndrome or neonatal varicella. Two live vaccine doses should be administered four to eight weeks apart, with conception avoided for 28 days following the final dose.<sup>57</sup>

### COVID-19

COVID-19 vaccination is recommended as a primary course for women who are not previously vaccinated, because of increased risk of severe disease with COVID-19 in pregnant women.<sup>1,63</sup> Single-dose mRNA vaccines (Comirnaty JN.1 or Comirnaty LP8.1) are currently recommended.<sup>58</sup>

### Influenza

Women planning pregnancy should receive an influenza vaccine to reduce the risk of influenza-related complications during pregnancy. Annual vaccination is recommended.<sup>30</sup>

### Special risk groups considerations

Some women may require additional vaccines or vaccine schedule alterations (Box 1). This includes women who are immunocompromised because of a congenital or medical condition, or following immunosuppressive treatment, those who have previously experienced an adverse event following vaccination, and those planning overseas travel where relevant exposures are anticipated. Expert consultation may be necessary for

individualised risk assessment and vaccine planning.

### Vaccination during pregnancy: protecting mother and baby

Vaccination during pregnancy provides dual protection by directly safeguarding the mother against severe illness and indirectly protecting the infant via transplacental transfer of maternal antibodies.

### Vaccines recommended during pregnancy

The Australian Immunisation Handbook recommended vaccines during pregnancy are detailed in the Figure.<sup>1,9,17,30,58,64,65</sup>

### Influenza

Seasonal influenza significantly increases

morbidity in pregnant women, with elevated risks of severe respiratory illness, hospitalisation, preterm labour and adverse fetal outcomes.<sup>37,66,67</sup> Influenza vaccination is therefore strongly recommended for all pregnant women during any trimester, especially those whose pregnancy coincides with the influenza season.<sup>68</sup> Clinical studies have consistently shown influenza vaccination during pregnancy to be safe, with no increased risk of congenital malformations or adverse maternal or fetal outcomes.<sup>1,40</sup>

Two developments in the 2026 influenza program warrant mention. First, Australia has completed the transition from quadrivalent to trivalent seasonal influenza vaccines, following the WHO and Australian Influenza Vaccine Committee recommendation that inclusion of the B/Yamagata lineage virus (which has not circulated for several years and is considered potentially extinct) is no longer warranted.<sup>17</sup> All 2026 Australian vaccines are therefore trivalent formulations. Second, an intranasally administered LAIV (FluMist) is available for the first time in Australia for children aged 2 years to less than 18 years, through the private market and selected state-based programs. Importantly, LAIV is contraindicated in pregnancy, and pregnant women should receive the inactivated influenza vaccine only.<sup>17</sup>

### **Pertussis**

Pertussis remains a significant public health concern, particularly for infants under 6 months of age who are at the highest risk for severe disease and death.<sup>69</sup> Australia's recent pertussis resurgence with 57,146 notifications in 2024 (the highest annual total since 1991) and more than 82,000 cumulative notifications across 2024–25, has underscored the critical role of maternal pertussis immunisation.<sup>14–16</sup> Vaccination with reduced antigen diphtheria-tetanus-acellular pertussis vaccine (dTPa) is strongly recommended as a single dose, ideally between 20 and 32 weeks' gestation, although it may be given up until delivery, irrespective of previous vaccination

status.<sup>1,70–73</sup> This timing maximises the transfer of protective antibodies to the fetus, offering protection during the critical early life period. Multiple studies confirm maternal pertussis immunisation as highly effective, reducing the risk of pertussis infection in infants younger than three months by up to 90%.<sup>33–36,69</sup> Importantly, national and international data confirm no increased risk of adverse pregnancy outcomes associated with pertussis vaccination.<sup>74–76</sup> Moreover, a 2026 Australian population-based cohort study of 13,918 singleton pregnancies found no association between concomitant (same-day) administration of influenza and pertussis vaccines during pregnancy and adverse pregnancy, birth, or neonatal outcomes, supporting co-administration as a safe strategy to optimise vaccine uptake.<sup>77</sup>

### **Respiratory syncytial virus**

RSV is a leading cause of severe lower respiratory tract infection among infants, frequently resulting in hospitalisation and intensive care admission.<sup>78</sup> Until recently, preventive options were limited; however, 2023 to 2025 marked a significant shift in RSV prevention in Australia with the approval and rollout of two complementary strategies: maternal vaccination and the monoclonal antibody nirsevimab for newborns and infants, both now delivered through the national RSV-MIPP.<sup>8,9,49</sup>

### **Maternal vaccination**

The maternal RSV vaccine (Abrysvo, RSVpreF) was approved by the TGA in early 2024 and is recommended for use in pregnant women from 28 weeks' gestation.<sup>9,79</sup> Infants are unlikely to receive adequate protection if they are born within 14 days of their mother receiving the vaccine, so administration by 36 weeks' gestation is strongly recommended.<sup>80</sup> If Abrysvo is inadvertently administered before 28 weeks' gestation, it does not need to be re-administered.<sup>9</sup> Maternal immunisation enables transplacental transfer of RSV-specific antibodies, providing passive protection to infants during their highest risk period. Clinical trial data demonstrated robust

safety profiles and high vaccine efficacy, significantly reducing infant RSV-related lower respiratory tract infections and hospitalisations.<sup>81–83</sup>

In May 2025, the TGA updated the Product Information for Abrysvo (and the related older-adult vaccine Arexvy) to reflect a known low risk of Guillain–Barré syndrome following vaccination.<sup>84</sup> It is important to note that this safety signal is derived from postmarketing observational data in adults aged 65 years and older, where an estimated nine excess cases of Guillain–Barré syndrome per million doses of Abrysvo were identified within 42 days of vaccination; no equivalent signal has been demonstrated in the pregnant population to date. The TGA has reiterated that the benefit–risk balance continues to remain strongly in favour of vaccination in approved populations.

Since the RSV-MIPP commenced on 3 February 2025, Abrysvo has been available free of charge under the NIP for all Medicare-eligible pregnant women.<sup>8</sup> First-year program data are encouraging: an estimated 140,000 infants had been protected through the program (via maternal vaccination or infant nirsevimab) by late 2025.<sup>49</sup>

### **Nirsevimab**

The TGA also approved nirsevimab (Beyfortus), a long-acting monoclonal antibody administered as a single intramuscular dose to newborns.<sup>85</sup> Nirsevimab provides direct passive immunity and has demonstrated efficacy in reducing medically attended RSV illness and hospitalisations, including among infants with heart or lung disease, premature infants, and those under the age of two years who are immunocompromised.<sup>86–89</sup> In April 2024, Western Australia and Queensland became the first jurisdictions to introduce universal access to nirsevimab for all newborns.<sup>90,91</sup> As of 2026, all Australian states and territories offer nirsevimab to at least high-risk infants through either seasonal or year-round programs, under the RSV-MIPP umbrella.<sup>92</sup>

Nirsevimab is recommended by ATAGI for infants whose mothers did not receive RSV vaccine in pregnancy or were vaccinated less than two weeks before delivery, as well as infants at increased risk of severe RSV disease, regardless of maternal vaccination status.<sup>9</sup>

The availability of both maternal vaccination and nirsevimab raises important considerations for clinical practice. While maternal immunisation is preferred for healthy term pregnancies because of the broader benefits of protecting both mother and infant, nirsevimab offers an effective alternative or adjunct, particularly in cases of late or missed maternal vaccination, early delivery or increased infant risk. Clinicians should counsel women on both options, balancing time, infant risk and vaccine availability.

First-year RSV-MIPP data have shown substantial variation in nirsevimab uptake between jurisdictions, with reported early uptake ranging from about 13% in New South Wales to about 41% in Western Australia.<sup>49</sup> Ongoing data collection will provide important insights into the uptake and integration of these dual RSV prevention strategies, and achieving high coverage with at least one option for all newborns, particularly for First Nations and regional or remote populations, will be crucial to reducing the national burden of RSV-related morbidity.

### COVID-19

Unvaccinated pregnant women face heightened risks of severe COVID-19 associated with increased rates of hospitalisation, intensive care admission, preterm birth and still-birth.<sup>27,92-94</sup> The ATAGI recommends that only pregnant women who are unvaccinated receive a single primary dose of COVID-19 vaccine.<sup>58</sup> Current recommended mRNA vaccines are Comirnaty JN.1 and Comirnaty LP.8.1; although these most recent variant-containing formulations have not been formally studied in pregnant women, ATAGI considers them suitable and safe for use in pregnancy. COVID-19 vaccination during pregnancy has been shown to be safe

and effectively induces maternal antibody transfer, offering early infant protection against severe COVID-19.<sup>96-99</sup>

Previously vaccinated pregnant women with no medical risk conditions are not routinely recommended to have a further dose during pregnancy; however, a further dose may be considered based on underlying risk conditions or personal preference.<sup>58</sup>

### Vaccines not routinely recommended during pregnancy

Most other vaccines are not routinely recommended during pregnancy unless there is a specific high-risk scenario.<sup>1</sup> Notably, live virus vaccines such as MMR and varicella vaccines are contraindicated during pregnancy because of theoretical risks of inducing infection and subsequent complications, including adverse fetal outcomes.<sup>99</sup> As noted above, the intranasal LAIV, newly available in Australia in 2026 for children aged 2 years to less than 18 years, is also contraindicated in pregnancy.<sup>17</sup> Many inactivated bacterial vaccines and inactivated viral vaccines are similarly not recommended during pregnancy.<sup>101,102</sup> The Australian Immunisation Handbook makes special mention of human papillomavirus and yellow fever vaccines.<sup>103</sup> Nonetheless, there are scenarios in which the potential benefit of vaccination during pregnancy may outweigh theoretical risks, particularly among women with significant medical or exposure risks. These cases should be individually assessed by a specialist immunisation service or healthcare provider experienced in maternal immunisation, as absolute contraindications for vaccination during pregnancy are rare.<sup>1</sup>

### Vaccination during breastfeeding: ensuring ongoing protection

Breastfeeding provides essential immunological protection to infants through the transfer of maternal antibodies, particularly immunoglobulin A. Vaccination during lactation is a vital component of postpartum care, ensuring continued maternal immunity and enhancing passive immunity for the infant.

### Safety of vaccination while breastfeeding

Most vaccines, including both inactivated and live-attenuated vaccines, are safe during breastfeeding.<sup>1,103</sup> Extensive research demonstrates that vaccination during lactation does not negatively affect breast milk composition, lactation performance or infant health.<sup>104</sup> Vaccines administered to breastfeeding women do not pose risks of vaccine-associated infections to the infant, making postpartum vaccination a practical and safe preventive health measure.<sup>97,106</sup>

### Recommended vaccines for breastfeeding mothers

Breastfeeding mothers who did not receive recommended vaccines during pregnancy, especially for pertussis, should consider postpartum vaccination to ensure ongoing protection.<sup>107</sup>

### Addressing barriers to vaccine uptake

Despite clear recommendations and evidence supporting maternal vaccination, coverage in Australia remains suboptimal.<sup>108</sup> Uptake is shaped by a complex interplay of psychological, structural and systemic factors. The COVID-19 pandemic highlighted several challenges, such as the amplification of misinformation and the erosion of trust; however, it also introduced opportunities, including the accelerated innovation in health communication and engagement.<sup>109-111</sup> To improve maternal vaccination rates, it is essential to understand and address these barriers comprehensively.

### Addressing and overcoming vaccine hesitancy

Pregnancy is a time of heightened caution. Many women delay or avoid vaccination because of concerns about vaccine safety for their unborn child, unfamiliarity with newer vaccines, or confusing or inconsistent messaging. The COVID-19 pandemic intensified these hesitations. Rapid vaccine development, the absence of initial trial data for pregnant women, and experiences of coercive public health measures (such as

mandates) led some people to feel pressured or mistrustful of authorities.<sup>108</sup> Others described wanting more time to learn about vaccines at their own pace, underscoring the importance of early, consistent and evidence-based communication.

Clinicians remain the most trusted source of information and can improve transparency and mitigate hesitancy by clearly recommending vaccines while sharing the robust, real-world, ongoing vaccine safety and surveillance data available for maternal immunisations.<sup>112,113</sup> This may be achieved by engaging empathetically, addressing specific concerns and reinforcing the safety and benefits of vaccination.<sup>114,115</sup> Programs like P3-MumBubVax, now integrated into the NCIRS-led SKAI platform, provide tailored, evidence-based tools to support vaccine discussions in antenatal care.<sup>116</sup>

### Clinician knowledge and recommendations

Healthcare providers significantly influence maternal vaccine uptake. Regular training and updates should be provided to GPs, practice and immunisation nurses, and obstetricians and midwives on the latest immunisation recommendations and communication strategies.<sup>117-119</sup> The ATAGI's new *General Practitioner Bulletin*, the first edition of which was published in November 2025, provides a concise, practice-focused summary of current immunisation priorities for GPs and is a useful complement to the Australian Immunisation Handbook. Additionally, healthcare providers should incorporate routine vaccine counselling into all antenatal care visits, supported by clinical decision tools and reminders within electronic medical records.<sup>116,120</sup>

### Reducing systemic and logistical barriers

Barriers to vaccine access during pregnancy include limited availability in antenatal settings, logistical issues such as transport and storage needs, and fragmented care models that can lead to vaccination being seen as another healthcare provider's responsibility. This is particularly problematic for

Australia's culturally and linguistically diverse (CALD) communities, including First Nations and Pacific peoples, and those in rural and remote regions. Integrating vaccines into routine antenatal appointments, and providing maternal immunisation through community pharmacies and mobile clinics, offer flexibility and ease of access to help bridge the gap.<sup>121-123</sup> Utilising digital health solutions – such as automated reminders, nudge notifications, and educational tools – can facilitate timely vaccine administration.

### Public health initiatives and strategies

Tailored public awareness campaigns addressing vaccine safety, benefits and misconceptions should be specifically designed to include CALD communities. The Vaccine Champions initiative leverages credible healthcare influencers and community healthcare workers to disseminate evidence-based vaccine information and advocate for immunisation within their own networks.<sup>124,125</sup> These champions can address vaccine concerns and shift social norms towards vaccine acceptance.

Effective public health communication must recognise these contextual influences. Strategies that combine respectful dialogue, consistent messaging and local engagement are essential to rebuilding trust and improving maternal vaccine confidence post-COVID-19 pandemic.

### Future directions and research priorities

To sustain and enhance maternal and infant health protection through vaccination, continued research and development is essential.

### Emerging vaccines for pregnant women

#### Group B streptococcus vaccine

Group B streptococcus (GBS) remains a leading cause of neonatal sepsis and meningitis, resulting in significant morbidity and mortality.<sup>126</sup> Several candidate GBS vaccines are undergoing clinical trials to

assess safety, immunogenicity and efficacy, with the hexavalent capsular polysaccharide conjugate candidate GBS6 now in Phase 3 evaluation.<sup>127,128</sup> Successful development and integration of a maternal GBS vaccine could substantially reduce neonatal infections.

#### Cytomegalovirus vaccine

Congenital cytomegalovirus (CMV) infection is a significant cause of birth defects and neurological impairment.<sup>129</sup> Research into effective maternal CMV vaccines is ongoing, focusing on safety, efficacy and the potential for widespread implementation.<sup>130</sup>

### Conclusion

Vaccination preconception, during pregnancy and while breastfeeding remains a cornerstone of maternal and neonatal healthcare, significantly reducing the risks posed by VPDs. Despite compelling evidence supporting the safety, efficacy and necessity of maternal vaccination, barriers such as vaccine hesitancy, limited healthcare provider engagement and systemic accessibility challenges persist. Addressing these challenges requires co-ordinated efforts involving healthcare professionals, public health authorities, policymakers and communities.

Continued research into emerging vaccines, such as those targeting GBS and CMV, alongside initiatives aimed at improving vaccine confidence, will be crucial to advancing maternal and neonatal health outcomes. Through enhanced awareness, strengthened clinical recommendations, improved accessibility and supportive public health policies, Australia can ensure comprehensive protection and optimal health outcomes for mothers and their infants. **MT**

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