

Vaccinations in pregnancy

Optimising maternal and infant protection

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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).^{1,2} 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.³

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



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.

(ATAGI), in collaboration with other key stakeholders, develop the handbook's recommendations for the safest and most effective use of vaccines.⁴ 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.⁵⁻⁷

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1. RESOURCES FOR GPs

- **Australian Immunisation Handbook from the Australian Technical Advisory Group on Immunisation:** <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://www.health.gov.au/resources/publications/national-immunisation-program-schedule?language=en>
- **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?language=en>

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-inmates-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>

Recent developments, including the introduction of a new respiratory syncytial virus (RSV) vaccine into the NIP, heightened vaccine hesitancy following the coronavirus disease 2019 (COVID-19) pandemic and the resurgence of pertussis, underscore

the critical need to optimise maternal vaccination uptake.⁸⁻¹⁵ 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.¹ 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.¹⁶

Immunological changes in pregnancy

Pregnancy induces complex immunological adaptations that facilitate fetal tolerance while maintaining maternal immune competence.¹⁷ These adaptations include modulation of innate and adaptive immune responses, increased anti-inflammatory cytokine activity and alterations in cellular immunity.¹⁸ 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.¹⁹⁻²⁷

Benefits of maternal vaccination

Protection of the mother

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

2. PRECONCEPTION VACCINATION ASSESSMENT^{1,49,50}

- Pre-vaccination screening checklist. See: <https://immunisationhandbook.health.gov.au/resources/tables/table-pre-vaccination-screening-checklist>⁵⁰
- 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

Abbreviation: AIR = Australian Immunisation Register.

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.³¹ Immunoglobulin G is actively transported across the placenta via the neonatal Fc receptor, reaching peak concentrations in the third trimester.³² 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 three months old.³³⁻³⁶ Similarly, seasonal influenza vaccination during pregnancy reduces the risk of infection and related complications for mothers and their newborns.³⁷⁻⁴¹

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.^{42,43}

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.^{44,45} In Australia, sustained maternal pertussis immunisation programs have led to a decline in infant pertussis-related hospitalisations.⁴⁶ Moreover, the introduction of maternal RSV vaccination is expected to further decrease RSV-related neonatal morbidity and mortality, and provide additional herd protection beyond this group, while being cost-effective.^{47,48}

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/australian-immunisation-register>), as part of comprehensive preconception care (Box 2).^{1,49,50} 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,^{1,51} 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.¹

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).¹ Should serological testing reveal insufficient immunity, the

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

Pathology request form details should 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.

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.^{1,34,52-59}

Hepatitis B

Vaccination is recommended for all hepatitis B-naïve women to prevent acquisition and mother-to-infant transmission.⁵²

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 conception for at least 28 days after the final dose.⁵³⁻⁵⁵

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.⁵⁶

TABLE 1. RECOMMENDED PRECONCEPTION VACCINES^{1,30,52-59*}

Vaccine-preventable disease	If uncertain history of vaccination or disease, check serology	NIP funded	Vaccine brand (type)	Dosage and timing
Hepatitis B ⁵²	✓	No	<ul style="list-style-type: none"> Engerix-B[†] (recombinant subunit) H-B-Vax II[†] (recombinant subunit) 	<ul style="list-style-type: none"> Three doses at zero, one and six months OR Three doses at zero, one and four months OR Three doses at zero, two and four months
Measles ⁵³	✓	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"> M-M-R II[†] (live attenuated) Priorix[‡] (live attenuated) 	<ul style="list-style-type: none"> Two doses, four weeks apart Avoid pregnancy within 28 days of receiving this live vaccine
Mumps ⁵⁴	✓			
Rubella ⁵⁵	✓			
Varicella (chickenpox) ⁵⁶	✓	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"> Varilrix[‡] (live attenuated) Varivax, refrigerated[‡] (live attenuated) 	<ul style="list-style-type: none"> Two doses, four to eight weeks apart Avoid pregnancy within 28 days of receiving this live vaccine
COVID-19 ⁵⁷	✗	COVID-19 vaccines are free for everyone in Australia regardless of Medicare or visa status	<ul style="list-style-type: none"> Comirnaty JN.1[§] (mRNA) Comirnaty Omicron XBB.1.5[§] (mRNA) 	<ul style="list-style-type: none"> Single dose
Influenza ³⁰	✗	Only for adults aged ≥18 years with specified medical risk conditions ^{#†}	<ul style="list-style-type: none"> Vaxigrip Tetra (inactivated) Flucelvax Quad (inactivated) 	<ul style="list-style-type: none"> Single dose
Pneumococcal ⁵⁸	✗	Only for people with specified medical risk conditions** Valneuvance (15vPCV) and Prevenar 20 (20cPCV) are not currently NIP-funded	<ul style="list-style-type: none"> Prevenar 13 (13vPCV) (conjugate) Vaxneuvance (15vPVC) (conjugate) Prevenar 20 (20vPCV) (conjugate) Pneumovax 23 (23vPPV) (polysaccharide) 	<ul style="list-style-type: none"> Single dose of a pneumococcal conjugate vaccine (13vPCV, 15vPCV, or 20vPCV) followed by: <ul style="list-style-type: none"> – two doses of the pneumococcal polysaccharide vaccine (23vPPV), first dose two to 12 months after the PCV, then second dose five years after first 23vPPV dose

Abbreviations: 13vPCV = 13-valent pneumococcal conjugate vaccine; 15vPCV = 15-valent pneumococcal vaccine; 20vPCV = 20-valent pneumococcal vaccine; COVID-19 = coronavirus disease 2019; MMR = measles, mumps and rubella; MMRV = measles, mumps, rubella, varicella; 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.

[†] Adult formulation. Switching hepatitis B vaccine brands is not recommended.⁵²

[‡] Switching of MMR vaccine brands is permitted. 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.

[§] 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 who are 18 to 64 years of age 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-19.⁵⁹

[#] 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.

[†] See Box 1 for a resource on specified medical conditions associated with increased risk of influenza disease and severe outcomes.

^{||} As of August 2025, the optimal pneumococcal vaccination program for Australia is currently under review.⁵⁸

** See Box 1 for a resource on risk conditions for pneumococcal vaccination.

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.^{1,59} Single-dose mRNA vaccines (Comirnaty JN.1 or

Comirnaty Omicron XBB.1.5) are currently recommended.⁵⁷

Influenza

Women planning pregnancy should receive an influenza vaccine to reduce the risk of influenza-related complications

during pregnancy. Annual vaccination is recommended.³⁰

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.^{1,9,30,57,60-63}

Influenza

Seasonal influenza significantly increases morbidity in pregnant women, with elevated risks of severe respiratory illness, hospitalisation, preterm labour and adverse fetal outcomes.^{37,64,65} Influenza vaccination is therefore strongly recommended for all pregnant women during any trimester, especially those whose pregnancy coincides with the influenza season.⁶⁶ 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.^{1,40}

Pertussis

Pertussis remains a significant public health concern, particularly for infants under six months of age who are at the highest risk for severe disease and death.⁶⁷ Vaccination with reduced antigen diphtheria-tetanus-acellular pertussis vaccine (dTpa) is strongly recommended as a single dose between 20 and 32 weeks' gestation, irrespective of previous vaccination status.^{1,68-71} This timing maximises the

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																	
Influenza Recommended single dose at any stage of pregnancy* <i>Vaxigrip Tetra</i> or <i>Flucelvax Quad</i>																																							
																		Pertussis Recommended single dose in each pregnancy, between 20 and 32 weeks <i>Boostrix</i> or <i>Adacel</i>																					
																												RSV Recommended single dose, from 28 weeks [†] <i>Abrysvo</i> [‡]											
COVID-19 Unvaccinated pregnant women only [§] Recommended single dose at any stage of pregnancy [¶] <i>Comirnaty JN.1</i> or <i>Comirnaty Omicron XBB.1.5</i>																																							

Figure. Recommended vaccines during pregnancy for 2025.^{1,9,30,57,60-63}

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.

* 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 the vaccine before becoming pregnant should be revaccinated during pregnancy to protect the newborn infant.³⁰

† If administered less than 2 weeks before giving birth, the newborn will not be adequately protected.⁹

‡ Abrysvo (RSVpreF) is the only RSV vaccine approved for use in pregnant women.⁹

§ 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.⁵⁷ Unless a woman is otherwise eligible, a dose of COVID-19 vaccine is not routinely recommended in pregnancy.⁵⁷ 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 Omicron XBB.1.5-based vaccines) have not been formally studied in pregnant women, the Australian Technical Advisory Group on Immunisation considers them suitable and safe for use.⁵⁷

¶ People with severe immunocompromise conditions and those who are 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.^{57,63}

Adapted from the National Centre for Immunisation Research and Surveillance. Vaccine recommendations for pregnant women – a guide for health professionals.⁶²

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%.^{33-36,67} Importantly, national and international data confirm no increased risk of adverse pregnancy outcomes associated with pertussis vaccination.⁷²⁻⁷⁴

Respiratory syncytial virus

RSV is a leading cause of severe lower respiratory tract infection among infants, frequently resulting in hospitalisation and intensive care admission.⁷⁵ 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.

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.^{9,76} However, 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.⁷⁷ 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.⁷⁸⁻⁸⁰

In May 2025, the TGA issued an updated warning on the known low risk of people developing Guillain-Barré syndrome following vaccination, noting, however, that the benefit-risk balance continues to remain strongly in favour of vaccination

in the target groups.⁸¹ The TGA further stated they had not received any reports of Guillain-Barré syndrome following vaccination with Abrysvo as of 24 March 2025.

Nirsevimab

The TGA also approved nirsevimab (Beyfortus), a long-acting monoclonal antibody administered as a single intramuscular dose to newborns.⁸² 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.⁸³⁻⁸⁶ In April 2024, Western Australia and Queensland became the first jurisdictions to introduce universal access to nirsevimab for all newborns.^{87,88} As of early 2025, all Australian states and territories offer nirsevimab to all infants through either seasonal or year-round programs.⁸⁹

Nirsevimab is recommended by the 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.⁹

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.

Ongoing data collection will provide important insights into the uptake and integration of these dual RSV prevention strategies. Ultimately, achieving high coverage with at least one of these options for all newborns 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 stillbirth.^{21,27,90,91} The ATAGI recommends that only pregnant women who are unvaccinated receive a single primary dose of COVID-19 vaccine.⁵⁷ Current recommended vaccines include mRNA vaccines Comirnaty JN.1 or Comirnaty Omicron XBB.1.5. COVID-19 vaccination during pregnancy has been shown to be safe and effectively induces maternal antibody transfer, offering early infant protection against severe COVID-19.⁹²⁻⁹⁴

Vaccines not routinely recommended during pregnancy

Most other vaccines are not routinely recommended during pregnancy unless there is a specific high-risk scenario.¹ 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.⁹⁵ Many inactivated bacterial vaccines and inactivated viral vaccines are also not recommended during pregnancy.^{96,97} The *Australian Immunisation Handbook* makes special mention of human papillomavirus and yellow fever vaccines.⁹⁸ 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.¹

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.^{99,100} Extensive research demonstrates that vaccination during lactation does not negatively affect breast milk composition, lactation performance or infant health.¹⁰¹ 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.^{93,102}

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.¹⁰³

Addressing barriers to vaccine uptake

Despite clear recommendations and evidence supporting maternal vaccination, coverage in Australia remains suboptimal.¹⁰⁴ 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.¹⁰⁵⁻¹⁰⁷ 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.¹⁰⁵ Others described wanting more time to learn about vaccines at their own pace, understanding 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.^{108,109} This may be achieved by engaging empathetically, addressing specific concerns and reinforcing the safety and benefits of vaccination.^{110,111} Programs like P3-MumBubVax, now integrated into the NCIRS-led SKAI platform, provide tailored, evidence-based tools to support vaccine discussions in antenatal care.¹¹²

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.¹¹³⁻¹¹⁵ Additionally, healthcare providers should incorporate routine vaccine counselling into all antenatal care visits, supported by clinical decision tools and reminders within electronic medical records.^{112,116}

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 Nation's 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.¹¹⁷⁻¹¹⁹ 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.^{120,121} 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.¹²² Several candidate GBS vaccines are undergoing clinical trials to assess safety, immunogenicity and

efficacy.^{123,124} 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.¹²⁵ Research into effective maternal CMV vaccines is ongoing, focusing on safety, efficacy and the potential for widespread implementation.¹²⁶

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**

References

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Vaccinations in pregnancy

Optimising maternal and infant protection

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