Infant ≥6 months age use only

| Alert | Influenza vaccines can change from year to year with regard to which vaccines are registered by the Therapeutic Goods Administration and the indicated ages for each vaccine. Always check annual seasonal influenza statements published by the Australian Technical Advisory Group on Immunisation on health.gov.au website and consult the product information for each vaccine. All children aged 6 months to less than 5 years are now eligible to receive free annual influenza vaccines under the National Immunisation Program (NIP). | | | | | |
|--------------------------------|---|--|---|--|--|--|
| | The dose of influenza vaccines for all ages is 0.5 mL. The 0.25 mL dose for young children is no longer available. ¹ | | | | | |
| Indication | Infants ≥6 months of age are strongly recommended to receive annual influenza vaccine. ² Preterm infants: Provided they are medically stable and there are no contraindications to vaccination, preterm infants should receive vaccines according to the recommended schedule at their chronological age, without correction for prematurity. ³ | | | | | |
| Action | Quadrivalent inactivated influenza virus vaccine. Active immunisation against influenza A, B virus strains (contained in vaccine). | | | | | |
| Drug type | Vaccine | | | | | |
| Trade name | Vaxigrip Tetra 0.5 mL: All people aged ≥6 months. FluQuadri 0.5 mL: All people aged ≥6 months. Influvac Tetra 0.5 mL. All people aged ≥6 months. (Not funded under NIP in Australia) Flucelvax Quad 0.5mL All people aged ≥6 months. (Not funded under NIP in Australia) | | | | | |
| Presentation | Vaxigrip Tetra 0.5 m monodose pre-filled syringe (All people aged ≥6 months). FluQuadri 0.5 mL monodose pre-filled syringe: (All people aged ≥6 months). Influvac Tetra 0.5 mL monodose pre-filled syringe. Flucelvax Quad 0.5mL monodose pre-filled syringe. | | | | | |
| Dose | 2 doses at least 4 weeks apart are recommended for children aged 6 months to <9 years receiving influenza vaccine for the first time. ² | | | | | |
| | Age | Dose | Number of doses needed in the first year of influenza vaccination | Number of doses needed if one or more doses of influenza vaccination received in the previous season | | |
| | 6 months to 9 years | 0.5ml | 2 (4 weeks apart) | 1 | | |
| Dose adjustment | Not applicable. | | | | | |
| Route | Intramuscular (IM) Note: IM route is preferred to the subcutaneous route because it causes fewer local adverse events. However, if given subcutaneously, the vaccine does not need to be readministered. ² | | | | | |
| | - | | | | | |
| Preparation | However, if given s Not required. | ubcutaneo | usly, the vaccine does not need | d to be readministered. ² | | |
| Administration Administration | However, if given s Not required. For intramuscular in Position the limb to Inject into the ante If only two vaccines If two vaccines nee injections. Pierce the skin at a | njection, us relax the r rolateral th are co-adi d to be adr 90° angle, | usly, the vaccine does not need the a 25 gauge 25 mm long need muscle that the vaccine is bein igh for infants not yet walking ministered, it is recommended ministered in the same muscle, so the needle can be safely ins | d to be readministered. ² dle. g injected into. | | |
| - | However, if given s Not required. For intramuscular in Position the limb to Inject into the ante If only two vaccines If two vaccines nee injections. Pierce the skin at a Inject the vaccine s It is not necessary t done this and a flas injection. Document all vaccine record. The parent professional. All immunisation en | njection, us relax the rolateral the are co-adid to be adrived of blood nes administratives incounters i | is a 25 gauge 25 mm long need muscle that the vaccine is being igh for infants not yet walking ministered, it is recommended ininistered in the same muscle, so the needle can be safely insaccount of 5 seconds. It is not the syringe plunger before appears in the needle hub, wire stered to children in the child's eps this record and presents it including influenza vaccination. | die. g injected into. to give one vaccine into each limb. there should be a distance of 2.5 cm between erted to the hub to reach the muscle layer. ee injecting a vaccine. However, if you have chdraw the needle and select a new site for s clinical file and the individual child health every time the child sees a health | | |
| - | However, if given s Not required. For intramuscular in Position the limb to Inject into the ante If only two vaccines If two vaccines nee injections. Pierce the skin at a Inject the vaccine s It is not necessary t done this and a flas injection. Document all vaccine record. The parent professional. All immunisation en | njection, us relax the relateral the are co-adid to be adrived of the decomposition of the counters in stralian/Ao | is a 25 gauge 25 mm long need muscle that the vaccine is being igh for infants not yet walking ministered, it is recommended ininistered in the same muscle, so the needle can be safely insaccount of 5 seconds. It is not the syringe plunger before a count of the needle hub, wire stered to children in the child's eps this record and presents it including influenza vaccination tearoa (New Zealand) Immuni | die. g injected into. to give one vaccine into each limb. there should be a distance of 2.5 cm between erted to the hub to reach the muscle layer. ee injecting a vaccine. However, if you have chdraw the needle and select a new site for s clinical file and the individual child health every time the child sees a health | | |

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| | Anaphylaxis following any vaccine component. | | |
|-------------------|--|--|--|
| Precautions | Persons with egg allergy, including anaphylaxis, can be safely vaccinated with influenza vaccines that have less than 1 microgram of residual egg ovalbumin per dose. none of the listed influenza vaccines contain >1ug of ovalbumin. If there is significant parental or health professional anxiety, the vaccine may be administered in primary care settings with a longer waiting period of 30 minutes. ^{2,4} Influenza vaccination is generally not recommended for people with a history of Guillain-Barré Syndrome whose first episode occurred within 6 weeks of receiving an influenza vaccine. ² | | |
| Drug interactions | There is a possible small increased risk of fever with co-administration of 13vPCV (13-valent pneumococcal conjugate vaccine). | | |
| Adverse reactions | Drowsiness or tiredness, muscle aches, localised pain, redness and swelling at injection site, occasionally, an injection-site nodule which may last many weeks (no treatment needed), fever and irritability and poor feeding in infants. | | |
| Compatibility | | | |
| Incompatibility | Not applicable. | | |
| Stability | Can remain stable at temperatures up to 12°C for 15 minutes. However, immediate administration is highly recommended. Follow local cold chain guidelines and Department of Health National Vaccine Storage 'Strive for 5' Guidelines for management of vaccines during cold chain breaches. ⁵ | | |
| Storage | Store at 2°C to 8°C (Refrigerate, do not freeze). Protect from light. Discard if vaccine has been frozen. | | |
| Excipients | Vaxigrip Tetra: sodium chloride, potassium chloride, dibasic sodium phosphate dihydrate monobasic potassium phosphate, water for injection FluQuadri: sodium chloride, dibasic sodium phosphate, monobasic sodium phosphate, water for injections Influvac Tetra: potassium chloride, monobasic potassium phosphate, dibasic sodium phosphate dihydrate, sodium chloride, calcium chloride dihydrate, magnesium chloride hexahydrate, water for injections. Flucelvax Quad: sodium chloride, potassium chloride, Magnesium chloride hexahydrate, Dibasic sodium phosphate dihydrate, monobasic potassium phosphate, water for injections. | | |
| Special | Children can receive 13vPCV and inactivated influenza vaccine at the same visit if they need both vaccines. ² | | |
| comments | Doses of intramuscular 1:1000 adrenaline for anaphylaxis 2 <1 year (approx. 5–10 kg) = 0.05 to 0.1 mL 1–2 years (approx. 10 kg) = 0.1 mL | | |
| Evidence | Background Influenza ('the flu') is an infectious disease caused by the influenza virus. Approximately 1 in 5 unvaccinated children and 1 in 10 unvaccinated adults are estimated to be infected by seasonal influenza annually, with rates of symptomatic influenza roughly half of these estimates. ⁶ The symptoms of influenza include sudden fever, headache, muscle aches and pains, fatigue, cough, sore throat, and stuffy or runny nose. The virus can cause a mild or severe illness depending on the type of influenza virus and general health of the affected person. Preterm infants have a high rate of underlying medical conditions – particularly respiratory, cardiac or neurological disease – that increase the risk of complications from influenza. ⁷ The incidence of influenza-associated hospitalization in children, NSW 2001-2011, was markedly increased for infants 0 to 24 months of age with bronchopulmonary dysplasia at 41.6 (95%Cl 15.7-67.5) per 1000 child-years, those with cystic fibrosis 44.5 (6.0-83.0) and other congenital and chronic lung conditions 42.9 (18.1-67.8) compared to all other children without chronic lung disease at 0 to 24 months age 9.3 (4.4- 14.2), 2 to 5 years 0.6 (0.3-1.0) and 5 to 10 years 0.1 (0.0-0.1). [7] The cost/episode (95%Cl) of influenza- associated hospitalisation was AUD\$19704 (95%Cl 11 715-27 693) for children with CLDs compared to \$4557 (95%Cl 4129-4984) for children without. ⁷ It is also important to note that the majority of hospital admissions still occur in infants without any comorbidities (underscoring importance of vaccination for all children) ¹⁵ Efficacy Influenza vaccination of infants and children for prevention of influenza infection in children: Systematic review found inactivated influenza vaccine in children aged 6 months to 16 years reduced influenza (RR 0.41, 95%Cl 0.29 to 0.59; participants = 1628; studies = 7; I ² = 36%; RD -20%, 95% Cl -33 to -7; test for subgroup differences according to age p=0.04) and influenza like illness (ILI) (RR 0.64, 95%Cl 0.54 to 0.76; p | | |

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786; studies = 2; RD -5%, 95%CI -17 to 8; participants = 786; studies = 2). Conclusion: In children aged between 3 and 16 years, live influenza vaccines reduce influenza and ILI over a single influenza season. However live influenza vaccine is not available in Australia. Inactivated vaccines also reduce influenza and ILI. There is limited data on efficacy for infants 6 months to 2 years.^{8,9,10} Influenza vaccination of pregnant women for prevention of influenza infection in infants: A systematic review of maternal influenza vaccination in pregnancy found a 36% reduced risk of infants <6 months having laboratory-confirmed influenza infection (RR 0.64, 95%CI 0.52, 0.78; 4 RCTs, 1099 infants). 11,12 Safety Most influenza vaccines are grown in eggs. Due to changes in influenza vaccine manufacturing, most influenza vaccines currently used contain less than 1 microgram of ovalbumin per dose.^{2,4} Influenza vaccine can be safely given in most patients with egg allergy (including egg prophylaxis) following appropriate guidelines.4, 10, 13 Some studies that included a small number of patients reported that people receiving cancer immunooncology therapies (checkpoint inhibitors) may have a higher risk of immune-related adverse events following immunisation with influenza vaccine, but a more recent study on patients receiving treatment with a single checkpoint inhibitor did not. The clinical importance of this potential interaction is currently inconclusive.2 Children can receive 13vPCV and inactivated influenza vaccine at the same visit if they need both vaccines. One study found a slightly higher risk of fever and febrile convulsions in children aged 6 months to <5 years (especially those aged 12-24 months) when they received inactivated trivalent influenza vaccine and 13vPCV at the same time, compared with receiving the vaccines separately. The risk was about 18 more cases per 100,000 doses in children aged 6 months to <5 years. The highest risk was 45 per 100,000 doses in children aged 16 months. This increased risk is small. A later study did not show the same association between febrile seizures and co-administration of these 2 vaccines.² It is acceptable to administer these vaccines concurrently when both vaccines are indicated. 14 All people ≥6 months of age are strongly recommended to receive annual influenza vaccine.² **Practice points** Two doses at least 4 weeks apart are recommended for children aged 6 months to <9 years receiving influenza vaccine for the first time.² Australian technical advisory group on immunisation (ATAGI) clinical advice. Statement on the References administration of seasonal influenza vaccines in 2025. https://www.health.gov.au/sites/default/files/2025-03/atagi-statement-on-the-administration-ofseasonal-influenza-vaccines-in-2025 0.pdf. Australian Immunisation Handbook. https://immunisationhandbook.health.gov.au/contents. Accessed 30/04/2025. Vaccination for preterm infants. Australian immunisation handbook. https://immunisationhandbook.health.gov.au/contents/vaccination-for-special-riskgroups/vaccination-for-preterm-infants. Accessed on 30/04/2025. 4. ASCIA Guidelines - Vaccination of the egg-allergic individual. https://www.allergy.org.au/hp/papers/vaccination-of-the-egg-allergic-individual. 2017. National Vaccine Storage Guidelines 'Strive for 5'. 3rd edition. https://www.health.gov.au/resources/publications/national-vaccine-storage-guidelines-strive-for-5. Somes MP, Turner RM, Dwyer LJ, Newall AT. Estimating the annual attack rate of seasonal influenza among unvaccinated individuals: A systematic review and meta-analysis. Vaccine. 2018;36:3199-207. Homaira N, Briggs N, Oei JL, Hilder L, Bajuk B, Snelling T, Chambers GM, Jaffe A. Impact of influenza on hospitalization rates in children with a range of chronic lung diseases. Influenza other respi. 2019:233-8. Jefferson T, Rivetti A, Di Pietrantonj C, Demicheli V. Vaccines for preventing influenza in healthy children. Cochrane Database Syst Rev. 2018;2:CD004879. Dhamayanti M, Tarigan R, Fadlyana E, Prasetyo D, Amalia N, Rusmil VK, Sari RM, Bachtiar NS, Rusmil K, Kartasasmita CB. Immunogenicity and safety of Quadrivalent Influenza HA vaccine in Indonesian children: An open-labeled, bridging, clinical study. Vaccine. 2020 Jan 29;38(5):993-1000. 10. Hu Y, Shao M, Hu Y, et al. Immunogenicity and safety of an inactivated quadrivalent influenza vaccine: a randomized, double-blind, controlled phase III clinical trial in children aged 6-35 months in China. Hum Vaccin Immunother. 2020 Jul 2;16(7):1691-1698.

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