Influenza Vaccine Program Summary, 2018 – 2019 Influenza Season (FLU)

Biological	Indications	Initial Series	Specific Contraindications	Expected Reactions
Quadrivalent Influenza vaccine (QIV) (Split virion, inactivated) A/Michigan/45/20 15)(H1N1) pdm09-like strain A/Singapore/INFI MH-16-0019/2016 (H3N2)-like strain B/Phuket/3073/20 13–like strain B/Colorado/6/201 7- like strain B/Colorado/6/201 80% 6 – 23 months of age 80% of health care workers 100% of immunizers	 Universal Program for anyone 6 months of age and older who lives, works or studies in Alberta. People at high risk of influenza-related complications or hospitalization: Anyone 65 years of age and older. Children 6 – 59 months of age Pregnant women at any stage of pregnancy and lactating women. Indigenous peoples Adults and children with the following health conditions: Chronic cardiac Chronic respiratory (including asthma) Immunosuppression/ immunodeficiency/HIV Diabetics/other metabolic diseases Cancer Renal disease Anemia Cognitive dysfunction, spinal cord injury, seizure disorder, neuromuscular disorder Children 6 months to 18 years of age on long term ASA therapy Neurologic or neurodevelopment conditions Morbid obesity (BMI ≥ 40) People capable of transmitting influenza to those at risk: Health care workers All household contacts of high risk individuals Members of household expecting a newborn Regular child caregivers for children ≤ 59 months of age. Others: People in direct contact during culling operations with poultry infected with avian influenza Health individuals aged 5 – 64 years Travelers 	Individuals 6 months up to and including 8 years of age who HAVE NOT received influenza vaccine in a previous season: • 2 doses, 4 weeks apart (document: 1 of 2, 2 of 2) Individuals 6 months up to and including 8 years of age who HAVE received influenza vaccine in a previous season: • 1 dose (document: Annual) Individuals 9 years of age and older: • 1 dose Dose: 0.5 ml IM Minimum age: 6 months of age Maximum age: no limit Notes: • A reason code must be documented for each dose administered; do NOT leave blank or use "unknown" • Influenza vaccine status/eligibility should be assessed during routine contact: • Well-child clinics (everyone with child) • Seniors clinics • Diabetes events • Prenatal/postnatal classes • Etc. • May be given at the same time as other inactivated and live vaccines using a separate needle and syringe for each vaccine.	 Infants under 6 months of age Anaphylactic reaction to a previous dose of influenza vaccine or to any constituent of the vaccine severe ORS following previous doses of influenza vaccine (See ORS Algorithm) Individuals who developed GBS within 6 weeks of previous influenza immunization. Individuals with acute, febrile illness should be deferred until better. Note: Egg allergy is NOT a contraindication to inactivated influenza immunization. Individuals severely allergic to eggs should be monitored for 30 minutes following immunization. 	 Common: Injection site pain, tenderness, redness, swelling Irritability, abnormal crying, malaise, fatigue, anorexia, myalgia, headache, fever, dizziness, gastrointestinal symptoms, arthralgia Rare: Anaphylaxis, allergic reaction Oculo-respiratory syndrome Guillain-Barré Syndrome As with any immunization, unexpected or unusual side effects can occur.

Note:

Fluzone® and FLULVAL TETRA multidose vials: once entered can use for 28 days if sterile technique and cold chain maintained. .

Protect from light. Do not inject intradermally, subcutaneously or intravenously. •

Cold Chain must be strictly observed. •

• QIV: Quadrivalent influenza vaccine.