Diphtheria-Tetanus-Acellular Pertussis-Polio Adsorbed (DTaP-IPV)

QUADRACEL® **Supplier: Sanofi Pasteur Limited**

INFANRIX®-IPV **Supplier: GlaxoSmithKline Inc.**

INDICATIONS:

- School entry booster given at 4-6 years of age. This is the 5th dose in a routinely immunized child.
- Used to complete the primary series and booster for children under 7 years of age in whom Hib is not indicated (see Part 4 – Biological Products, Haemophilus b Conjugate Vaccine).

Not approved for use in those 7 years of age and older.

DOSES AND SCHEDULE:

4-6 years of age: 1 dose given as 0.5 mL **IM**. [A]

Primary series completion: See Part 1 – Immunization Schedules for appropriate schedule.

BOOSTER DOSES:

There are no routine additional doses of these vaccines required outside of the above indications. [B]

SEROLOGICAL TESTING:

Serological testing is not recommended before or after immunization.

CONTRAINDICATIONS:

1. History of anaphylactic reaction to a previous dose of diphtheria or tetanus toxoids or to pertussis or polio vaccines or any component of the product.
2. History of Guillain-Barré syndrome (GBS) within 8 weeks of receipt of a tetanus toxoid without any other cause being identified.

PRODUCT COMPONENTS:

QUADRACEL®:
Potential allergens: neomycin, polymyxin B, bovine serum albumin, polysorbate 80.
Other components: aluminum phosphate, 2-phenoxyethanol, formaldehyde, glutaraldehyde.

INFANRIX®-IPV:
Potential allergens: neomycin, polymyxin.
Other components: aluminum hydroxide, Medium 199.

PRECAUTIONS:

Not applicable.

[A] Not necessary if the 4th dose of DTaP or Tdap-containing vaccine was given after the 4th birthday.
[B] Refer to Part 1 – Immunization Schedules for information regarding booster doses using the age appropriate tetanus and diphtheria-containing vaccines.

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SPECIAL CONSIDERATIONS:

- A hypotonic-hyporesponsive episode (HHE) following a prior dose of DTaP-containing vaccine is not a contraindication to future immunization.
- Any dose(s) of oral polio vaccine (OPV) received on or after April 1, 2016 will not be considered as a valid dose within the routine BC immunization schedule. For more information, see Part 4 – Biological Products, Polio Vaccine, Special Considerations.

ADVERSE EVENTS:

Local: redness, swelling, pain.

Systemic: fever, decreased appetite, vomiting, diarrhea, decreased activity, irritability, drowsiness, increased crying.