

PRODUCT MONOGRAPH

INFANRIX hexa **Adsorbed Hib reconstituted with PEDIARIX**

Combined diphtheria and tetanus toxoids, acellular pertussis, hepatitis B (recombinant), inactivated poliomyelitis and adsorbed conjugated *Haemophilus influenzae* type b vaccine

Sterile suspension for injection

Single dose pre-filled syringe PEDIARIX (suspension for injection)
and
Single dose vial adsorbed hib (lyophilized powder for injection)

Active immunizing agent

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Table of Contents

PART I: HEALTH PROFESSIONAL INFORMATION.....	3
SUMMARY PRODUCT INFORMATION	3
DESCRIPTION.....	3
INDICATIONS AND CLINICAL USE.....	4
CONTRAINDICATIONS	5
WARNINGS AND PRECAUTIONS.....	6
ADVERSE REACTIONS.....	9
DRUG INTERACTIONS	13
DOSAGE AND ADMINISTRATION	14
OVERDOSAGE	17
ACTION AND CLINICAL PHARMACOLOGY	18
STORAGE AND STABILITY.....	22
DOSAGE FORMS, COMPOSITION AND PACKAGING	22
PART II: SCIENTIFIC INFORMATION	24
PHARMACEUTICAL INFORMATION.....	24
CLINICAL TRIALS.....	24
DETAILED PHARMACOLOGY	30
MICROBIOLOGY	30
TOXICOLOGY	30
REFERENCES	31
PART III: CONSUMER INFORMATION.....	35

INFANRIX hexa

Combined diphtheria and tetanus toxoids, acellular pertussis, hepatitis B (recombinant), inactivated poliomyelitis, and adsorbed conjugated *Haemophilus influenzae* type b vaccine

PART I: HEALTH PROFESSIONAL INFORMATION

SUMMARY PRODUCT INFORMATION

Route of Administration	Dosage Form/Strength	Nonmedicinal Ingredients
Intramuscular injection	Sterile suspension for injection/ After reconstitution, 1 dose (0.5 ml) contains 25 limit of flocculation (Lf) [30 International Units (IU)] diphtheria toxoid; 25 mcg filamentous haemagglutinin (FHA); 10 mcg hepatitis B surface antigen (HBsAg); 8 mcg pertactin (PRN); 25 mcg pertussis toxoid (PT); 40 D-antigen units (DU) of type 1 poliovirus, 8 DU type 2 poliovirus, and 32 DU type 3 poliovirus ; 10 Lf (40 IU) tetanus toxoid; 10 mcg of adsorbed purified capsular polysaccharide of <i>Haemophilus influenzae</i> type b (Hib) (PRP) covalently bound to approximately 25 mcg of tetanus toxoid per 0.5 mL dose.	Aluminum (as aluminum salts), lactose, medium 199, sodium chloride, water for injection. Residues*: disodium phosphate, formaldehyde, glutaraldehyde, glycine, monopotassium phosphate, neomycin sulphate, polymyxin B sulphate, polysorbate 20 and 80, and potassium chloride.

*From the manufacturing process.

DESCRIPTION

INFANRIX hexa (combined diphtheria and tetanus toxoids, acellular pertussis, hepatitis B (recombinant), inactivated poliomyelitis, and adsorbed conjugated *Haemophilus influenzae* type b vaccine) contains diphtheria toxoid, tetanus toxoid, three purified pertussis antigens [pertussis toxoid (PT), filamentous haemagglutinin (FHA) and pertactin (PRN) (69 kiloDalton outer membrane protein)], hepatitis B virus surface

antigen recombinant, adsorbed onto aluminum salts, purified, inactivated poliovirus types 1, 2 and 3, *Haemophilus influenzae* type b polysaccharide conjugated to tetanus toxoid.

INDICATIONS AND CLINICAL USE

Pediatrics:

Primary Immunization

INFANRIX hexa (combined diphtheria and tetanus toxoids, acellular pertussis, hepatitis B (recombinant), inactivated poliomyelitis, and adsorbed conjugated *Haemophilus influenzae* type b vaccine) is indicated for:

- active primary immunization against diphtheria, tetanus, pertussis, hepatitis B, poliomyelitis, and disease caused by *Haemophilus influenzae* type b in infants and children 6 weeks to 2 years.

INFANRIX hexa will not prevent hepatitis caused by other agents, such as hepatitis A, C and E viruses, or other pathogens known to infect the liver. As hepatitis D (caused by the delta virus) does not occur in the absence of hepatitis B infection, it can be expected that hepatitis D will also be prevented by INFANRIX hexa vaccination.

Where a dose of hepatitis B vaccine is given at birth, INFANRIX hexa can be used for the second dose from the age of six weeks. If a second dose of hepatitis B vaccine is required before this age, monovalent Hepatitis B vaccine should be used.

Booster Vaccination

The administration of the booster dose should be given at 12 to 23 months as stated in the Canadian Immunization Guide.

INFANRIX hexa can be used for the booster dose provided that the infant has received a full primary vaccination course of each of the antigens contained in INFANRIX hexa, regardless of whether these were administered as monovalent or combination vaccines.

Other combinations of antigens have been studied in clinical trials following primary vaccination with INFANRIX hexa and may be used for a booster dose, these include diphtheria, tetanus, acellular pertussis (DTaP) and DTaP-Hib.

CONTRAINDICATIONS

INFANRIX hexa (combined diphtheria and tetanus toxoids, acellular pertussis, hepatitis B (recombinant), inactivated poliomyelitis, and adsorbed conjugated *Haemophilus influenzae* type b vaccine):

- should not be administered to subjects with known hypersensitivity to any component of this vaccine (see DOSAGE FORMS, COMPOSITION AND PACKAGING) or to subjects having shown signs of hypersensitivity after a previous dose of this vaccine or any injection containing diphtheria, tetanus, pertussis, hepatitis B, poliovirus or *Haemophilus influenzae* type b (see WARNINGS AND PRECAUTIONS, General section for information on treatment of immediate allergic reactions).
- should be used with caution in subjects with known hypersensitivity to the antibiotics neomycin and polymyxin, as INFANRIX hexa contains traces of these antibiotics.
- is contraindicated for use after an immediate anaphylactic reaction temporally associated with a previous dose of this vaccine or any injection containing diphtheria, tetanus, pertussis, hepatitis B, poliovirus, or *Haemophilus influenzae* type b. Because of the uncertainty as to which component of the vaccine might be responsible, no further vaccination with any of these components should be given. Alternatively, because of the importance of tetanus vaccination, such individuals may be referred to an allergist for evaluation.
- should not be administered to persons 7 years of age or older because diphtheria toxoid may cause severe but transient local and febrile reactions in children and adults, the frequency increasing with age, the dose of toxoid and the number of doses given.
- is contraindicated if the infant has experienced an encephalopathy of unknown etiology, occurring within 7 days following previous vaccination with a pertussis containing vaccine. In these circumstances, pertussis vaccination should be discontinued and the vaccination should be continued with diphtheria-tetanus, Hepatitis B, polio, and Hib vaccines.

Immunization should be deferred during the course of a moderate or severe acute febrile illness or acute infection (see WARNINGS AND PRECAUTIONS). The presence of a minor infection, however is not a contraindication.

Elective immunization of individuals over 6 months should be deferred during an outbreak of poliomyelitis.

WARNINGS AND PRECAUTIONS

General

Do not administer INFANRIX hexa (combined diphtheria and tetanus toxoids, acellular pertussis, hepatitis B (recombinant), inactivated poliomyelitis, and adsorbed conjugated *Haemophilus influenzae* type b vaccine) intravascularly or intradermally.

As with any other vaccine, a protective immune response may not be elicited in all vaccinees for all component antigens in the vaccine. This product is not recommended for treatment of actual infections.

INFANRIX hexa will not prevent disease caused by pathogens other than *Corynebacterium diphtheria*, *Clostridium tetani*, *Bordetella pertussis*, hepatitis B virus, poliovirus or *Haemophilus influenzae* type b.

As for all diphtheria, tetanus and pertussis vaccines, each injection should be given deep intramuscularly and each injection of the immunization series should be made at a different site.

As with other injectable vaccines, epinephrine injection (1:1000) and other appropriate agents used for the control of immediate allergic reactions must be immediately available should an acute anaphylactic reaction occur. For this reason, the vaccinee should remain under medical supervision for 30 minutes after immunization.

It is good clinical practice that vaccination should be preceded by a review of the medical history (especially with regard to previous vaccination and possible occurrence of undesirable events) and a clinical examination.

Where passive protection is required, Tetanus Immune Globulin and/or Diphtheria Antitoxin may also be administered at separate sites. Because of the substantial risks of complications from pertussis disease, completion of a primary series of vaccine early in life is strongly recommended.

If any of the following events occur in temporal relation to administration of whole-cell DTP or acellular DTP vaccine, the decision to give subsequent doses of vaccine containing the pertussis component should be carefully considered:

- Temperature of $> 40.5^{\circ}\text{C}$ within 48 hours of vaccination not due to another identifiable cause.
- Collapse or shock-like state (hypotonic hyporesponsive episode) within 48 hours of vaccination.
- Persistent, inconsolable crying lasting ≥ 3 hours, occurring within 48 hours of vaccination.
- Convulsions with or without fever occurring within 3 days of vaccination.

There may be circumstances, such as high incidence of pertussis, in which the potential benefits outweigh possible risks, particularly since these events have not been proven to cause permanent sequelae.

Data from clinical studies indicate that, when INFANRIX hexa is co-administered with pneumococcal conjugate vaccine (PREVNAR, PREVNAR 13 or SYNFLORIX), the rate of febrile reactions is higher compared to that occurring following the administration of INFANRIX hexa alone.

Increased reporting rates of convulsions (with or without fever) and hypotonic hyporesponsive episode were observed with concomitant administration of INFANRIX hexa and PREVNAR 13 (see ADVERSE REACTIONS).

Antipyretic treatment should be initiated according to local treatment guidelines.

Syncope (fainting) can occur following, or even before, any vaccination as a psychogenic response to the needle injection. It is important that procedures are in place to avoid injury from faints.

Hematologic

INFANRIX hexa should be administered with caution to subjects with thrombocytopenia or a bleeding disorder since bleeding may occur following intramuscular administration to these subjects.

Immune

Hepatitis B has a long incubation period. Hepatitis B vaccination may not prevent hepatitis B infection in individuals who had an unrecognized hepatitis B infection at the time of vaccine administration.

INFANRIX hexa is not contraindicated for use in individuals with HIV infection. The expected immunological response may not be obtained after vaccination of immunosuppressed patients.

Immunosuppressive therapies, including irradiation, antimetabolites, alkylating agents, cytotoxic drugs and corticosteroids (used in greater than physiologic doses), may reduce the immune response to vaccines (See DRUG INTERACTIONS).

Hepatitis B

Infants born of HBsAg-positive mothers should receive hepatitis B immune globulin (HBIG) and Hepatitis B vaccine at birth and should complete the Hepatitis B vaccination series given according to a particular schedule. Infants born of mothers of unknown HBsAg status should receive Hepatitis B vaccine at birth and should complete the Hepatitis B vaccination series given according to a particular schedule (see Manufacturer's package insert for Hepatitis B vaccine).

The subsequent administration of INFANRIX hexa for completion of the Hepatitis B vaccination series in infants who were born of HBsAg-positive mothers and received HBIG, or infants born of mothers of unknown status has not been studied.

Neurologic

Experience with INFANRIX (DTaP) and other INFANRIX-based combinations has not revealed any cases of encephalopathy or permanent neurologic damage causally linked to vaccination. While acute encephalopathy and permanent neurologic damage have not been reported to be causally linked nor in a temporal association with administration of INFANRIX hexa data is limited at this time.

In children with progressive neurological disorders, including infantile spasms, uncontrolled epilepsy or progressive encephalopathy, it is better to defer pertussis (Pa or Pw) immunization until the condition is corrected or stable. However, the decision to give pertussis vaccine must be made on an individual basis after careful consideration of the risks and benefits.

A history of convulsions or other central nervous system disorders in parents or siblings is not a contraindication for INFANRIX hexa, an acellular DTP vaccine. Vaccinees with a history of febrile convulsions should be closely followed up as such adverse events may occur within 2 to 3 days post vaccination.

Studies suggest that when given whole-cell DTP vaccine, infants and children with a history of convulsions in first-degree family members (i.e., siblings and parents) have a 2.4-fold increased risk for neurologic events compared to those without such histories.

Respiratory

Although a moderate or severe illness with or without fever is a reason to defer vaccination, minor illnesses such as mild upper respiratory infections with or without low-grade fever are not a contraindication.

Special Populations

Pregnant Women: As INFANRIX hexa is not intended for use in adults, adequate human data on use during pregnancy and adequate animal reproduction studies are not available.

Nursing Women: As INFANRIX hexa is not intended for use in adults, adequate human data on use during lactation and adequate animal reproduction studies are not available.

Pediatrics: Clinical data indicate that INFANRIX hexa can be given to preterm infants, however, as expected in this population, a lower immune response has been observed for some antigens (see ADVERSE REACTIONS and CLINICAL TRIALS). The potential risk of apnea and the need for respiratory monitoring for 48-72h should be considered when administering the primary immunization series to very preterm infants (born after at

least 24 weeks of gestational age) and particularly for those with a previous history of respiratory immaturity. As the benefit of vaccination is high in these infants, vaccination should not be withheld or delayed. Safety and effectiveness of INFANRIX hexa have not been established in infants below the age of 6 weeks and children over 2 years of age.

Interference with laboratory testing

The Hib component of the vaccine does not protect against diseases due to capsular serotypes other than type b of *Haemophilus influenzae* or against meningitis caused by other organisms. Excretion of capsular polysaccharide antigen in the urine has been described following administration of Hib vaccines, and therefore antigen detection may not have a diagnostic value in suspected Hib disease within 1-2 weeks of vaccination. Other tests should be performed in order to confirm Hib infection during this period.

ADVERSE REACTIONS

Adverse Drug Reaction Overview

INFANRIX hexa (combined diphtheria and tetanus toxoids, acellular pertussis, hepatitis B (recombinant), inactivated poliomyelitis, and adsorbed conjugated *Haemophilus influenzae* type b vaccine) is generally well tolerated.

Clinical Trial Adverse Drug Reactions

Because clinical trials are conducted under very specific conditions the adverse reaction rates observed in the clinical trials may not reflect the rates observed in practice and should not be compared to the rates in the clinical trials of another drug. Adverse drug reaction information from clinical trials is useful for identifying drug-related adverse events and for approximating rates.

During a study conducted in the United States, a total of 785 documented doses of study vaccines were given to 267 subjects included in the According To Protocol (ATP) reactogenicity analysis. Solicited and unsolicited symptoms occurring during the 8-day follow-up period after vaccination were reported. Most reported solicited local symptoms and solicited general symptoms were mild to moderate in intensity. There were no statistically significant differences between the two groups in the incidence of soreness, redness or swelling at the injection site (regardless of side/site/dose) or fever. The percentage of subjects per group experiencing symptoms (both solicited and unsolicited) during the 8 days after vaccination is outlined in **Table 1**.

Table 1 Percentage of U.S. Infants with Local or Systemic Reactions within 8 Days of Primary Vaccination with either INFANRIX hexa or Commercially Available INFANRIX, ENGERIX-B, and OPV Administered Simultaneously with Hib at Separate Sites (Per subject analysis)

Event	INFANRIX hexa (N=134)	INFANRIX, ENGERIX-B, Hib vaccine, OPV (N=133)
Local	%	%
Pain, any	42.54	52.63
Pain, severe	1.49	2.26
Redness, any	48.51	47.37
Redness, > 20 mm	2.24	3.01
Swelling, any	35.82	40.60
Swelling, > 20 mm	3.73	4.51
Systemic	%	%
Temperature ≥ 38°C	55.97	51.88
> 39.5°C	0.75	2.26
Diarrhea, any	35.82	33.08
Grade 3	0.75	2.26
Eating/drinking less than usual, any	49.25	57.14
Grade 3	2.24	2.26
Irritability/fussiness, any	82.84	86.47
Grade 3	6.72	6.02
Sleeping less than usual, any	50.75	56.39
Grade 3	2.24	3.76
Sleeping more than usual, any	62.69	67.67
Grade 3	3.73	1.50
Unusual crying for more than one hour, any	42.54	41.35
Grade 3	3.73	2.26
Vomiting, any	25.37	20.30
Grade 3	0.75	0.75

N= number of infants

The safety profile presented below is based on data from more than 16,000 subjects.

As has been observed for DTaP and DTaP-containing combinations, an increase in local reactogenicity and fever was reported after booster vaccination with INFANRIX hexa with respect to the primary course.

Frequencies per dose as defined by CIOMS:

Very common: ≥ 10%

Appetite lost, irritability, crying abnormal, restlessness, pain, redness, local swelling at the injection site (≤ 50 mm), fever $\geq 38^\circ\text{C}$, and fatigue.

Common: ≥ 1% and < 10%

Nervousness, vomiting, diarrhea, local swelling at the injection site (> 50 mm)*, fever >39.5°C, pruritis** and injection site reactions, including induration.

Uncommon: ≥ 0.1% and < 1%

Upper respiratory tract infection, somnolence, cough** and diffuse swelling of the injected limb, sometimes involving the adjacent joint*.

Rare: ≥ 0.01% and < 0.1%

Bronchitis and rash.

Very rare: < 0.01%

Convulsions (with or without fever)***, dermatitis, bronchospasm, and urticaria**.

* Children primed with acellular pertussis vaccines are more likely to experience swelling reactions after booster administration in comparison with children primed with whole cell vaccines. These reactions resolve over an average of 4 days.

**Observed only with other GSK DTaP-containing vaccines.

*** Analysis of post-marketing reporting rates suggests a potential increased risk of convulsions (with or without fever) and hypotonic hyporesponsive episode when comparing groups which reported use of INFANRIX hexa with PREVNAR 13 to those which reported use of INFANRIX hexa alone.

Local reactions after immunization usually consist of swelling or induration, tenderness, and redness or erythema at the injection site. More severe local reactions occasionally occur, such as inflammatory cellulitis without bacterial infection after DTP-containing vaccines.

Post-Marketing Adverse Drug Reaction

Over 12 million doses of INFANRIX hexa have been distributed overall for primary and booster vaccinations. Extremely rare cases of Sudden Unexpected Death (SUD) in close temporal association to vaccination with INFANRIX hexa have been reported in the first year of life. However, a causal relationship has not been established. The observed number of SUD cases following INFANRIX hexa is below the number of cases expected to occur by chance.

Blood and lymphatic system disorders

Lymphadenopathy, thrombocytopenia.

Immune system disorders

Allergic reactions (including anaphylactic and anaphylactoid reactions).

Nervous system disorders

Collapse or shock-like state (hypotonic hyporesponsive episode)***.

Respiratory, thoracic and mediastinal disorders

Apnea** [see section “WARNINGS AND PRECAUTIONS” for apnea in very premature infants (≤ 28 weeks of gestation)].

Skin and subcutaneous tissue disorders

Angioneurotic oedema**.

General disorders and administration site conditions

Extensive swelling reactions, swelling of the entire injected limb*, vesicles at the injection site.

*Children primed with acellular pertussis vaccines are more likely to experience swelling reactions after booster administration in comparison with children primed with whole cell vaccines. These reactions resolve over an average of 4 days.

**Observed only with other GSK DTaP-containing vaccines.

*** Analysis of post-marketing reporting rates suggests a potential increased risk of convulsions (with or without fever) and hypotonic hyporesponsive episode when comparing groups which reported use of INFANRIX hexa with PREVNAR 13 to those which reported use of INFANRIX hexa alone.

Safety in preterm infants

INFANRIX hexa has been administered to more than 1,000 preterm infants (born after a gestation period of 24 to 36 weeks) in primary vaccination studies and in more than 200 preterm infants as a booster dose in the second year of life. In comparative studies, similar rates of symptoms were observed in preterm and full-term infants.

Experience with hepatitis B vaccine

Paralysis, neuropathy, Guillain-Barré syndrome, encephalopathy, encephalitis, meningitis, allergic reactions mimicking serum sickness, neuritis, hypotension, vasculitis, lichen planus, erythema multiforme, arthritis and muscular weakness have been reported extremely rarely during post-marketing surveillance following vaccination with

ENGERIX-B (Hepatitis B vaccine) in infants < 2 years old. The causal relationship to the vaccine has not been established.

DRUG INTERACTIONS

Overview

INFANRIX hexa (Combined diphtheria and tetanus toxoids, acellular pertussis, hepatitis B (recombinant), inactivated poliomyelitis, and adsorbed conjugated *Haemophilus influenzae* type b vaccine) should not be mixed with any other vaccine in the same syringe or vial.

Drug-Drug Interactions

Tetanus Immune Globulin or Diphtheria Antitoxin, if used, should be given at a separate site, with a separate needle and syringe.

Anticoagulants

As with other intramuscular injections, INFANRIX hexa should not be given to infants or children on anticoagulant therapy unless the potential benefit clearly outweighs the risk of administration (see WARNINGS AND PRECAUTIONS).

Immunosuppressive Therapies

Immunosuppressive therapies, including irradiation, antimetabolites, alkylating agents, cytotoxic drugs and corticosteroids (used in greater than physiologic doses), may reduce the immune response to vaccines. Although no specific data are available from studies with INFANRIX hexa under these conditions, if immunosuppressive therapy will be discontinued shortly, it would be reasonable to defer immunization until the patient has been off therapy for 3 months; otherwise, the patient should be vaccinated while still on therapy. If INFANRIX hexa is administered to a person receiving immunosuppressive therapy, or a recent injection of immune globulin, an adequate immunologic response may not be obtained.

Use with Other Vaccines

INFANRIX hexa can be given concomitantly with pneumococcal conjugate, MenC conjugate, MenACWY conjugate, rotavirus, measles, mumps, rubella and varicella vaccines. Data have shown no clinically relevant interference in the antibody response to each of the individual antigens in INFANRIX hexa.

Data from clinical studies indicate that, when INFANRIX hexa is co-administered with pneumococcal conjugate vaccine, the rate of febrile reactions is higher compared to that occurring following the administration of INFANRIX hexa (see WARNINGS AND PRECAUTIONS, General). The incidence of fever following administration of the two vaccines in the primary series was lower than that observed after the booster vaccination.

Drug-Food Interactions

Interactions with food have not been established.

Drug-Herb Interactions

Interactions with herbal products have not been established.

Drug-Laboratory Interactions

Interactions with laboratory tests have not been established.

DOSAGE AND ADMINISTRATION

The use of reduced volume (fractional doses) is not recommended. The effect of such practices on the frequency of serious adverse events and on protection against disease has not been determined.

Pre-term infants (born after at least 24 weeks of gestational age) should be vaccinated according to their chronological age from birth.

Recommended Dose

Primary Immunization

The primary immunization course for infants born of HBsAg-negative mothers is 2 or 3 doses of INFANRIX hexa 0.5 mL, given intramuscularly, which should be administered according to official recommendations (See CLINICAL TRIALS for schedules evaluated in clinical trials). INFANRIX hexa can be considered for the booster if the antigen composition is in accordance with the official recommendations. INFANRIX hexa should not be administered to any infant before the age of 6 weeks.

Primary vaccination	Booster vaccination	General considerations
<i>Full-term infants (≥ 6 weeks of age)</i>		
3-dose	A booster dose must be given.	<ul style="list-style-type: none">• There should be an interval of at least 1 month between primary doses.• When giving a booster dose, this should be at least 6 months after the last priming dose and given at 12 to 23 months as stated in the Canadian Immunization Guide.
2-dose	A booster dose must be given.	<ul style="list-style-type: none">• There should be an interval of at least 1 month between primary doses.• When giving a booster dose, this should be at least 6 months after the last priming dose and preferably between 11 and 13 months of age.
<i>Preterm infants born after at least 24 weeks of gestational age (≥ 6 weeks of age)</i>		
3-dose	A booster dose must be given.	<ul style="list-style-type: none">• There should be an interval of at least 1 month between primary doses.• When giving a booster dose, this should be at least 6 months after the last priming dose and

		given at 12 to 23 months as stated in the Canadian Immunization Guide.
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Children Previously Vaccinated with One or More Doses of Hepatitis B Vaccine

Children who receive one dose of Hepatitis B vaccine at or shortly after birth may be administered a 3 dose series of INFANRIX hexa vaccine starting as early as 6 weeks of age. There is no data to support the use of a 3 dose series of INFANRIX hexa in infants who have previously received more than one dose of Hepatitis B vaccine. INFANRIX hexa may be administered to infants otherwise scheduled to receive concurrent INFANRIX (diphtheria, tetanus and acellular pertussis vaccine) and Hepatitis B vaccine and in whom vaccination against poliovirus is also desired.

Missed Dose

Interruption of the recommended schedule with a delay between doses should not interfere with the final immunity achieved with INFANRIX hexa. There is no need to start the series over again regardless of the time elapsed between doses.

Additional Dosing Information

If any recommended dose of pertussis vaccine cannot be given, diphtheria and tetanus toxoids (DT) for pediatric use should be given as needed to complete the series.

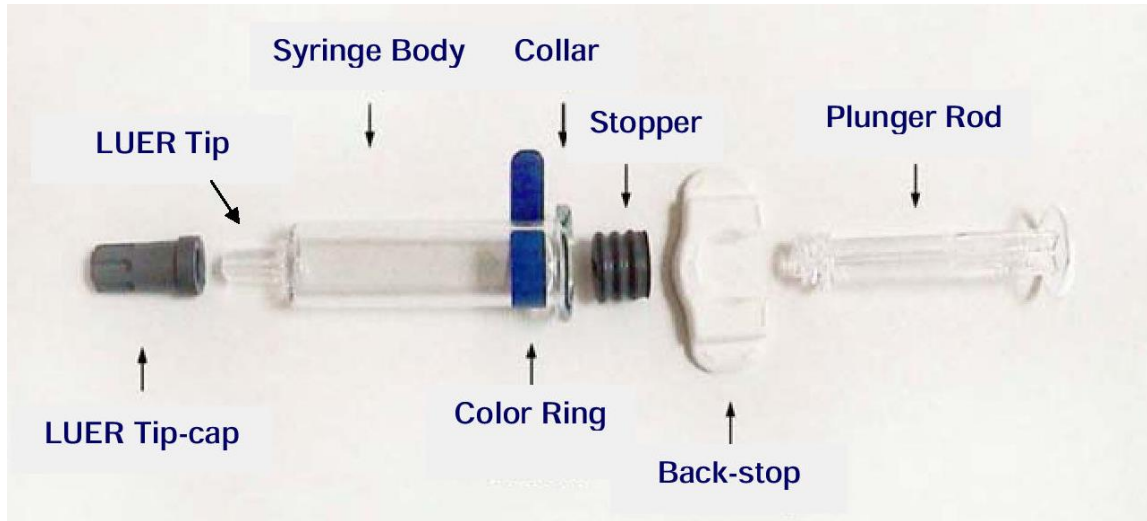
For persons 7 years of age or older, Tetanus and Diphtheria Toxoids (Td) for adult use should be given for routine booster immunization against tetanus and diphtheria.

Administration

Preparation for Administration

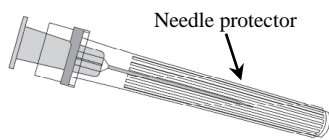
The vaccine is reconstituted by adding the entire contents of the syringe (PEDIARIX) to the vial containing the Hib pellet.

Do not remove the white back-stop from the syringe. Prior to administration, ensure that the plunger rod is firmly attached to the rubber stopper by turning the plunger clockwise until slight resistance is felt. **Do not** over tighten. Remove syringe LUER Tip-cap and needle cap. Attach needle by pressing and twisting in a clockwise rotation until secured to the syringe.

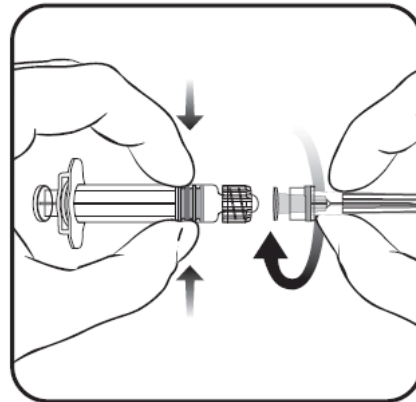
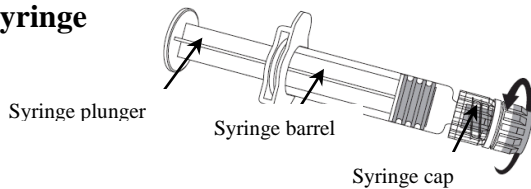


Specific instructions for the pre-filled syringe with a LUER lock adaptor

Needle



Syringe



Holding the syringe **barrel** in one hand (avoid holding the syringe plunger), unscrew the syringe cap by twisting it anticlockwise. To attach the needle to the syringe, twist the needle clockwise into the syringe until you feel it lock (see picture). Remove the needle protector, which on occasion can be a little stiff.

Reconstitution

Upon storage, a white deposit and clear supernatant may be observed in the syringe. This is a normal observation and does not constitute a sign of deterioration. Shake the syringe well before use. With thorough agitation, DTaP-HB-IPV (combined diphtheria and tetanus toxoids, acellular pertussis, hepatitis B (recombinant) and inactivated poliomyelitis vaccine, tradename PEDIARIX) is a homogeneous white turbid suspension. The syringe and the vial containing the Hib pellet should be inspected visually for any foreign particulate matter and/or variation of physical aspect. In the event of either being observed, discard the vaccine. The vaccine is reconstituted by adding the entire contents of the syringe (PEDIARIX) to the vial containing the Hib pellet. After the addition of the PEDIARIX vaccine to the pellet, the mixture should be well shaken until the pellet is completely dissolved. The vaccine should not be mixed with other vaccines.

It is good clinical practice to only inject a vaccine when it has reached room temperature. In addition, a vial at room temperature ensures sufficient elasticity of the rubber closure to minimise any coring of rubber particles. To achieve this, the vial should be kept at room temperature (25 ± 3 °C) for at least five minutes before connecting the syringe and reconstituting the vaccine.

The reconstituted vaccine presents as a slightly more cloudy suspension than the liquid component alone. This is a normal observation. The reconstituted vaccine should be inspected visually for any foreign particulate matter and/or abnormal physical appearance. In the event of either being observed, discard the vaccine.

Since this product is a suspension containing an adjuvant, shake vigorously to obtain a uniform suspension prior to withdrawal from the vial. **Do not use if resuspension does not occur with vigorous shaking.** Withdraw the entire contents of the vial.

INFANRIX hexa should be administered by intramuscular injection. The preferred sites are the anterolateral aspects of the thigh or the deltoid muscle of the upper arm. The vaccine should not be injected in the gluteal area or areas where there may be a major nerve trunk. Before injection, the skin at the injection site should be cleaned and prepared with a suitable germicide. After insertion of the needle, aspirate to ensure that the needle has not entered a blood vessel.

Do not administer this product subcutaneously or intravenously.

After reconstitution, the vaccine should be injected promptly. However the vaccine may be kept for up to 8 hours at room temperature (21°C).

OVERDOSAGE

For management of a suspected drug overdose, contact your regional Poison Control Centre.

ACTION AND CLINICAL PHARMACOLOGY

Diphtheria

Diphtheria is a serious communicable disease, primarily a localized and generalized intoxication caused by diphtheria toxin, an extracellular protein metabolite of toxigenic strains of *Corynebacterium diphtheriae*. The disease occurs most frequently in unimmunized or partially immunized individuals. The incidence of diphtheria in Canada has decreased from 9,000 cases reported in 1924 to extremely low levels. Only 1 or 2 cases have been reported annually in recent years. The case fatality rate remains 5% to 10 %, with the highest death rates in the very young and elderly. If immunization levels are allowed to fall and adults do not receive booster doses, disease re-emergence may appear as demonstrated in the Commonwealth of Independent States (former Soviet Union), where tens of thousands of cases with substantial mortality have been reported. Protection against disease is due to the development of neutralizing antibodies to the diphtheria toxin. Following adequate immunization with diphtheria toxoid, it is thought that protection persists for at least 10 years. Serum antitoxin levels of at least 0.01 antitoxin units per mL are generally regarded as protective.

This significantly reduces both the risk of developing diphtheria and the severity of clinical illness. Immunization with diphtheria toxoid does not, however, eliminate carriage of *C. diphtheriae* in the pharynx, nose or on the skin.

Tetanus

Tetanus is an intoxication manifested primarily by neuromuscular dysfunction caused by a potent exotoxin released by *Clostridium tetani*. Immunization is highly effective, provides long-lasting protection and is recommended for the whole population. Only 1 to 7 with an average of 5 cases of tetanus are now reported annually in Canada, while no deaths have been recorded since 1995. The disease continues to occur almost exclusively among persons who are unvaccinated, inadequately vaccinated or whose vaccination histories are unknown or uncertain.

Spores of *C. tetani* are ubiquitous. Naturally acquired immunity to tetanus toxin does not occur. Thus, universal primary immunization and timed booster doses to maintain adequate tetanus antitoxin levels are necessary to protect all age groups. Protection against disease is due to the development of neutralizing antibodies to the tetanus toxin. Tetanus toxoid is a highly effective antigen and a completed primary series generally induces serum antitoxin levels of at least 0.01 antitoxin units per mL, a level which has been reported to be protective. It is thought that protection persists for at least 10 years.

Pertussis

Pertussis (whooping cough) is a disease of the respiratory tract caused by *Bordetella pertussis*. Pertussis is highly communicable (attack rates in unimmunized household contacts of up to 90% have been reported) and can affect individuals of any age; however, severity is greatest among young infants. Precise epidemiologic data do not exist, since bacteriological confirmation of pertussis can be obtained in less than half of the suspected cases. Most reported illness from *B. pertussis* occurred in infants and young children in whom complications can be severe. Older children, adolescents and adults, in whom classic signs are often absent, may go undiagnosed and may serve as reservoirs of disease. Pertussis epidemics are cyclic and occur every 3 to 4 years. Pertussis has been controlled in Canada through immunization. During the last 40 years, the incidence of pertussis has decreased by > 90% although outbreaks continue to occur.

A recent study was conducted in Germany to assess the efficacy of pertussis vaccine after partial and completed primary vaccination series for preventing hospitalizations due to pertussis under field conditions. Data was acquired by a nationwide, hospital based, active surveillance system. After one dose of the vaccine, vaccine effectiveness was as high as 68%, increasing to 91.8% after receipt of the second dose. Vaccine effectiveness of 3 and 4 doses of acellular vaccine were estimated to be 99.8% and 98.6%, respectively.

Antigenic components of *B. pertussis* believed to contribute to protective immunity include: pertussis toxin (PT); filamentous hemagglutinin (FHA); and pertactin. Although the role of these antigens in providing protective immunity in humans is not well understood, clinical trials which evaluated candidate acellular DTP vaccines manufactured by GlaxoSmithKline supported the efficacy of 3 component INFANRIX (DTaP). Recently published data suggests a higher importance of the PT and pertactin components in providing protection against pertussis.

INFANRIX contains 3 pertussis antigens (PT, FHA and pertactin), and has been shown to be effective in preventing World Health Organization (WHO)-defined pertussis as well as clinically milder disease in two published clinical trials when administered as a primary series.

A double-blind, randomized, placebo (DT)-controlled trial conducted in Italy, sponsored by the U.S. National Institutes of Health (NIH), assessed the absolute protective efficacy of INFANRIX when administered at 2, 4 and 6 months of age. A total of 15,601 infants were immunized with 1 of 2 tri-component acellular DTP vaccines (containing inactivated PT, FHA and pertactin), or with a whole-cell DTP vaccine manufactured by Sanofi Pasteur, or with DT vaccine alone. The mean length of follow-up was 17 months, beginning 30 days after the third dose of vaccine. The population used in the primary analysis of vaccine efficacy included 4,481 INFANRIX vaccinees, 4,348 whole-cell DTP vaccinees and 1,470 DT vaccinees. After 3 doses, the protective efficacy of INFANRIX against WHO-defined typical pertussis (21 days or more of paroxysmal cough with infection confirmed by culture and/or serologic testing) was 84% (95% CI: 76% to 89%) while the efficacy of the whole-cell DTP vaccine was 36% (95% CI: 14% to 52%).

When the definition of pertussis was expanded to include clinically milder disease with respect to type and duration of cough, with infection confirmed by culture and/or serologic testing, the efficacy of INFANRIX was calculated to be 71% (95% CI: 60% to 78%) against > 7 days of any cough and 73% (95% CI: 63% to 80%) against ≥ 14 days of any cough. A longer follow-up of the Italian trial showed that after 3 doses, the absolute efficacy of INFANRIX against WHO-defined pertussis remained high at 84% among children up to 4 years of age.

A prospective, blinded efficacy trial was also conducted in Germany employing a household contact study design. In preparation for this study, 3 doses of INFANRIX were administered at 3, 4 and 5 months of age to more than 22,000 children living in 6 areas of Germany in a large safety and immunogenicity trial. Infants who did not participate in this trial could have received whole-cell DTP vaccine (manufactured by Chiron Behring, Germany) or DT vaccine. Calculation of vaccine efficacy was based on attack rates of pertussis in household contacts classified by vaccination status. Of the 173 unvaccinated household contacts, 96 developed WHO-defined pertussis (21 days or more of paroxysmal cough with infection confirmed by culture and/or serologic testing), as compared to 7 of 112 contacts vaccinated with INFANRIX and 1 of 75 contacts vaccinated with whole-cell DTP vaccine. The protective efficacy of INFANRIX was calculated to be 89% (95% CI: 77% to 95%), with no indication of waning of immunity up until the time of the booster. The protective efficacy of the whole-cell DTP vaccine was calculated to be 98% (95% CI: 83% to 100%). When the definition of pertussis was expanded to include clinically milder disease, with infection confirmed by culture and/or serologic testing, the efficacy of INFANRIX against ≥ 7 days of any cough was 67% (95% CI: 52% to 78%) and against ≥ 7 days of paroxysmal cough was 81% (95% CI: 68% to 89%). The corresponding efficacy rates of INFANRIX against ≥ 14 days of any cough or paroxysmal cough were 73% (95% CI: 59% to 82%) and 84% (95% CI: 71% to 91%), respectively.

Hepatitis B

Several hepatitis viruses are known to cause a systemic infection resulting in major pathologic changes in the liver (e.g., A, B, C, D, E). Hepatitis B infection can have serious consequences including acute massive hepatic necrosis, chronic active hepatitis and cirrhosis of the liver. It has been estimated that more than 350 million people in the world are persistently infected with hepatitis B virus.

Among infected infants, very few (5 -10%) recover completely; the majority (up to 90%) become chronic carriers with the risk of becoming a chronic carrier decreasing with age (children < 5 years 25% to 50%, adults 6% to 10%). Those patients who become chronic carriers can infect others and are at increased risk of developing either cirrhosis or primary hepatocellular carcinoma.

Among other factors, infection with hepatitis B may be the single most important factor for development of this carcinoma. Considering the serious consequences of infection, immunization should be considered for all persons.

Mothers infected with hepatitis B virus can infect their infants at, or shortly after, birth if they are carriers of the hepatitis B surface antigen (HBsAg) or develop an active infection during the third trimester of pregnancy. Infected infants usually become chronic carriers. Therefore, screening of pregnant women for hepatitis B is recommended. According to the Canadian Immunization Guide, hepatitis B prevention should include programs for universal immunization of children, pre-exposure vaccination of high-risk groups, universal HBsAg screening of all pregnant women and post-exposure intervention for those exposed to disease, particularly infants born to hepatitis B-infected mothers. There is no specific treatment for acute hepatitis B infection. However, those who develop anti-HBs antibodies after active infection are usually protected against subsequent infection. Antibody titers ≥ 10 mIU/mL against HBsAg are recognized as conferring protection against hepatitis B. Seroconversion is defined as antibody titers ≥ 1 mIU/mL.

Poliomyelitis

Poliovirus is an enterovirus that belongs to the picornavirus family. Three serotypes of poliovirus have been identified (types 1, 2 and 3). Poliovirus is highly contagious with the predominant mode of transmission being person-to-person via the fecal-oral route. Infection may be spread indirectly through contact with infectious saliva or feces or by contaminated water or sewage.

Replication of poliovirus in the pharynx and intestine is followed by a viremic phase where involvement of the central nervous system can occur. While poliovirus infections are asymptomatic or cause nonspecific symptoms (low-grade fever, malaise, anorexia and sore throat) in 90% to 95% of individuals, 1% to 2% of infected persons will develop paralytic disease.

Following the introduction of inactivated poliovirus vaccines (IPV) in Canada in 1955, the indigenous disease has been eliminated. Since 1980, 12 paralytic cases have been reported in Canada, 11 of which were determined to be vaccine-associated paralytic poliomyelitis (VAPP), with Oral Polio Vaccine (OPV). The last reported case of VAPP occurred in 1995.

Forty seven studies involving over 19,000 infants and children have been conducted in developed and developing countries with GlaxoSmithKline's enhanced inactivated poliovirus vaccine, as trivalent IPV vaccine or as a part of DTaP-IPV based combinations.

Haemophilus Influenzae type b

Haemophilus influenzae type b (Hib) was the most common cause of bacterial meningitis and a leading cause of other serious invasive infections in young children prior to the introduction of other Hib vaccines. About 55% to 65% of affected children had meningitis while the remainder had epiglottitis, bacteremia, cellulitis, pneumonia or septic arthritis. The case fatality rate of meningitis is about 5%. Severe neurologic sequelae occur in 10% to 15% of survivors and deafness in 15% to 20% (severe in 3% to 7%).

Before the introduction of Hib conjugate vaccines in Canada in 1988, there were approximately 2,000 cases of Hib disease annually. Since then the overall incidence has fallen by more than 99%. The majority of cases occur now in children too old to have received primary vaccination. In 1998, only 15 cases were reported in children < 5 years of age.

STORAGE AND STABILITY

Store INFANRIX hexa (combined diphtheria and tetanus toxoids, acellular pertussis, hepatitis B (recombinant), inactivated poliomyelitis, and adsorbed conjugated *Haemophilus influenzae* type b vaccine) at 2° to 8°C. Do not use after the expiration date shown on the label. After reconstitution, immediate use is recommended. However, stability of the vaccine has been demonstrated for 8 hours at 21°C after reconstitution.

Do not freeze. Discard if the vaccine has been frozen.

Protect from light.

During transport, recommended conditions of storage must be respected.

Stability data indicate that the vaccine components are stable at temperatures up to 25°C for 72 hours. These data are intended to guide healthcare professionals in case of temporary temperature excursion only.

DOSAGE FORMS, COMPOSITION AND PACKAGING

Dosage Forms

Syringe and Vial

PEDIARIX (combined diphtheria and tetanus toxoids, acellular pertussis, hepatitis B (recombinant) and inactivated poliomyelitis vaccine) is supplied as a turbid suspension in a pre-filled syringe (Type I glass) (0.5 mL) with plunger stoppers (butyl).

Haemophilus influenzae type b vaccine is supplied as a pellet in a 3.0 mL vial (Type I glass) with stopper (butyl).

Composition

INFANRIX hexa (single 0.5mL dose) contains 25 Lf (30 IU) diphtheria toxoid, 25 mcg filamentous haemagglutinin, 10 mcg HBsAg, 8 mcg pertactin, 25 mcg pertussis toxoid, 40 D-antigen Units (DU) of type 1 poliovirus, 8 DU type 2 poliovirus, 32 DU type 3 poliovirus, 10 Lf (40 IU) tetanus toxoid, and 10 mcg of adsorbed purified capsular polysaccharide of Hib (PRP) covalently bound to approximately 25 mcg of tetanus toxoid, 10 Lf (40 IU) tetanus toxoid. It also contains 0.82 mg aluminum (as aluminum salts), 12.6 mg lactose, 1.15 mg medium 199, 4.5 mg sodium chloride, water for

injection. Residues from the manufacturing process: disodium phosphate, formaldehyde, glutaraldehyde, glycine, monopotassium phosphate, neomycin sulphate, polymyxin B sulphate, polysorbate 20 and 80, and potassium chloride.

Packaging

Pack sizes of:

Syringe and Vial:

Supplied as a kit in pack sizes of 10 with or without needles.

PART II: SCIENTIFIC INFORMATION

PHARMACEUTICAL INFORMATION

Drug Substance

Proper name: combined diphtheria and tetanus toxoids, acellular pertussis, hepatitis B (recombinant), inactivated poliomyelitis, and adsorbed conjugated *Haemophilus influenzae* type b vaccine

Product Characteristics

INFANRIX hexa (combined diphtheria and tetanus toxoids, acellular pertussis, hepatitis B (recombinant), inactivated poliomyelitis, and adsorbed conjugated *Haemophilus influenzae* type b vaccine) contains combined diphtheria and tetanus toxoids, hepatitis B virus surface antigen recombinant, three purified pertussis antigens [pertussis toxoid (PT), filamentous haemagglutinin (FHA), pertactin (69 kiloDalton outer membrane protein)], adsorbed onto aluminum salts, and purified, inactivated poliovirus types 1, 2 and 3, and *Haemophilus influenzae* type b polysaccharide conjugated to tetanus toxoid.

CLINICAL TRIALS

Study results

The immunogenicity of INFANRIX hexa has been evaluated in clinical studies from 6 weeks of age. The vaccine was assessed in 2-dose and 3-dose priming schedules, including the schedule for the Expanded Program on Immunization, and as a booster dose. The results of these clinical studies are summarized below.

Immune Response to INFANRIX hexa Administered as a 3 Dose Primary Series

A total of 14,600 doses of INFANRIX hexa (combined diphtheria and tetanus toxoids, acellular pertussis, hepatitis B (recombinant), inactivated poliomyelitis, and adsorbed conjugated *Haemophilus influenzae* type b vaccine) have been administered to 4,890 infants from 6 weeks of age and up as a primary series in clinical studies.

The immune responses to each of the antigens contained in INFANRIX hexa were evaluated in sera obtained 1 month after the third dose of vaccine as compared to that following administration of commercially available vaccines (INFANRIX (diphtheria, tetanus, and acellular pertussis vaccine), ENGERIX-B (hepatitis B vaccine (recombinant)), Hib vaccine, and Oral Polio Virus vaccine) simultaneously at separate sites, in a study conducted in the U.S. The schedule of administration was 2, 4, and 6 months of age. One month after the third dose of INFANRIX hexa, immune response rates to each antigen were comparable to rates seen following separately administered vaccines (see Table 2).

The immunogenicity of INFANRIX hexa was evaluated across three studies including approximately 300 preterm infants (born after a gestation period of 24 to 36 weeks) following a 3-dose primary vaccination course at 2, 4 and 6 months of age. The immunogenicity of a booster dose at 18 to 24 months of age was evaluated in approximately 200 preterm infants (see Immunogenicity in preterm infants below).

Table 2 Antibody Responses to Each Antigen Following INFANRIX hexa as Compared to INFANRIX, ENGERIX-B, Hib vaccine, and OPV (One Month After Administration of Dose 3)

	INFANRIX hexa (N=78-106)	INFANRIX, ENGERIX-B, Hib vaccine, OPV (N=71-98)
Anti-Diphtheria % ≥ 0.1 IU/mL GMT	100.0 1.431	99.0 1.009
Anti-Tetanus % ≥ 0.1 IU/mL GMT	100.0 1.979	100.0 1.486
Anti-PT (V.R.) % R GMT	99.0 67.4	97.9 41.8
Anti-FHA (V.R.) % R GMT	100.0 288.0	98.7 302.8
Anti-Pertactin (V.R.) % R GMT	96.2 168.2	95.8 136.9
Anti-HBs % ≥ 10 mIU/mL GMT	99.1 1239.5	100.0 934.3
Anti-Polio 1 % ≥ 8 GMT	100.0 494.8	98.6 1278.2
Anti-Polio 2 % ≥ 8 GMT	98.8 507.4	100.0 1350.4
Anti-Polio 3 % ≥ 8 GMT	98.8 1275.1	98.6 367.5
Anti-PRP % ≥ 0.15 mcg/mL	100.0	96.9
Anti-PRP % ≥ 1.0 mcg/mL GMT	84.0 2.648	91.8 5.527

OPV manufactured by Wyeth

OmniHib manufactured by Sanofi Pasteur

% R = in initially seronegative subjects, appearance of antibodies (titre ³⁵ EL.U./mL); in initially seropositive subjects, at least maintenance of prevaccination titre

GMT = Geometric mean antibody titre

PT = Pertussis Toxoid

FHA = Filamentous Haemagglutinin

HBs = Hepatitis B surface (antigen)

V.R. = Vaccine Response (Vaccine response is defined as appearance of antibodies in initially seronegative subjects or as at least maintenance of pre-vaccination antibody titres in initially seropositive subjects).

Polio = Poliovirus
PRP = Polyribosyl-ribitol-phosphate

Clinical trials have investigated the tolerability and immunogenicity of the vaccine in various schedules (i.e. 2, 3, 4 months; 3, 4, 5 months; 2, 4, 6 months; 3, 5, 11 months; 1.5, 2.5, 3.5 months). Results obtained in all of the clinical studies for each of the components are summarized below:

DTaP component:

Immunological data:

One month after the 3 dose primary vaccination course, 98.5 to 100% of infants vaccinated with INFANRIX hexa had antibody titres of ≥ 0.1 IU/mL for both tetanus and diphtheria.

Following administration of a 4th dose of INFANRIX hexa in the second year of life, 100% of infants had antibody titres of ≥ 0.1 IU/mL for both tetanus and diphtheria.

One month after the 3 dose primary vaccination course, the overall response rate for each of the 3 individual pertussis antigens (PT, FHA, and pertactin) was between 97.2-99.3%, 95.2-100% and 95.9-99.3%, respectively.

Following administration of a 4th dose of INFANRIX hexa in the second year of life, a booster response was seen in at least 97.2%, 94.1%, and 100% of vaccinated infants against the respective pertussis antigens. Since a serological correlation for protection against pertussis disease does not exist, the efficacy of the pertussis component presently relies on efficacy trials described below.

Protective efficacy data:

The protective efficacy of the DTaP component, against WHO-defined typical pertussis (≥ 21 days of paroxysmal cough) was demonstrated after 3 dose primary immunization in the studies tabulated below:

Study	Country	Schedule	Vaccine efficacy	Considerations
Household contact study (prospective blinded)	Germany	3,4,5 months	88.7%	Based on data collected from secondary contacts in households where there was an index case with typical pertussis.
Efficacy study (NIH sponsored)	Italy	2,4,6 months	84%	In a follow-up of the same cohort, the efficacy was confirmed up to 60 months after completion of primary vaccination without administration of a booster dose of pertussis.

Hepatitis B component:

After the primary vaccination course with INFANRIX hexa, 98.5 to 100% of infants developed protective antibody titres of ≥ 10 mIU/mL.

At one month after the booster dose, administered 18 months after primary vaccination, 97 to 100% of these subjects had protective titres of ≥ 10 mIU/mL.

IPV component:

One month after the primary vaccination, the seroprotection rates for each of the three serotypes (types 1, 2, and 3) were 99.2 to 100%, 94.5 to 99.0%, and 98.8 to 100%, respectively.

Following administration of the booster dose, at least 98.5%, 98.5%, and 100% of infants were seroprotected for the three serotypes, respectively.

Hib component:

One month after completion of the primary vaccination course, the Geometric Mean Concentration (GMC) of antibodies ranged from 1.52 to 3.53 mcg/mL, with between 93.5 and 100% of the subjects reaching antibody titres ≥ 0.15 mcg/mL.

One month after the booster dose given in the second year of life, the GMC ranged from 19.1 to 94.0 mcg/mL, with 99.5 to 100% of the subjects reaching antibody titres ≥ 0.15 mcg/mL.

These GMCs are numerically lower when compared to GMCs resulting from separate administration of the Hib component, however they are not different from those elicited by comparator vaccines DTaP-Hib and DTaP-IPV-Hib vaccines.

Induction of immunological memory has been shown to be an important and intrinsic part of the protective immune response following administration of Hib conjugate vaccines. Children primed with INFANRIX hexa had an anamnestic response (defined as a rapid and substantial increase in antibody level) on subsequent exposure to the antigen.

Immunogenicity in preterm infants (3 Dose Primary Series)

One month after 3-dose primary vaccination at least 98.7% of subjects were seroprotected against diphtheria, tetanus and poliovirus types 1 and 2; at least 90.9% had seroprotective antibody levels against the hepatitis B, PRP and poliovirus type 3 antigens; and all subjects were seropositive for antibodies against FHA and PRN while 94.9% were seropositive for anti-PT antibodies.

One month after the booster dose at least 98.4% of subjects had seroprotective or seropositive antibody levels against each of the antigens except against PT (at least 96.8%) and hepatitis B (at least 88.7%). The response to the booster dose in terms of fold increases in antibody concentrations (15- to 235-fold), indicate that preterm infants were adequately primed for all the antigens of INFANRIX hexa.

In a follow-up study involving 74 preterm infants, approximately 2.5 to 3 years after the booster dose, 85.3% of the children were still seroprotected against hepatitis B and at least 95.7% were seroprotected against the three poliovirus types and PRP.

Post-marketing experience:

The effectiveness of the Hib component of INFANRIX hexa was investigated via an extensive post-marketing surveillance study conducted in Germany. Over a seven year follow-up period, the effectiveness of the Hib components of two hexavalent vaccines, of which one was INFANRIX hexa, was 89.6% for a 3-dose full primary series and 100% for a 3-dose full primary series plus booster dose (irrespective of the Hib vaccine used for priming)

Immune Response to INFANRIX hexa Administered as a 2 Dose Primary Series

After a complete vaccination according to a 2-dose primary and booster schedule with INFANRIX hexa, at least 97.9% of the subjects had developed seroprotective or seropositive antibody levels against each of the vaccine antigens (see Table 3).

Table 3 Percentage of subjects with antibody titres \geq assay cut-off and GMC/GMT one month after 2-dose primary (Post-dose 2) and one month after booster (Post-dose 3) vaccination with INFANRIX hexa

Antibody (cut-off)	Vaccination at 2-4-12 months	
	Post-dose 2 N=223 (1 study)	Post-dose 3 N=196 (1 study)
	%	%
Anti-diphtheria (0.1 IU/ml) † GMC	99.6 1.197	100.0 5.307
Anti-tetanus (0.1 IU/ml) † GMC	100 1.380	100.0 4.720
Anti-PT (5 EL.U/ml) GMC	100 49.3	99.5 86.1
Anti-FHA (5 EL.U/ml) GMC	100 172.3	100.0 451.2
Anti-PRN (5 EL.U/ml) GMC	99.6 74.5	100.0 242.9
Anti-HBs (10 mIU/ml) † GMC	99.5 -	99.8 -
Anti-Polio type 1 (1/8 dilution) † GMT	89.6 28.3	98.4 313.7
Anti-Polio type 2 (1/8 dilution) † GMT	85.6 25.1	98.4 429.2
Anti-Polio type 3 (1/8 dilution) † GMT	92.8 62.2	97.9 541.9
Anti-PRP (0.15 μ g/ml) † GMC	84.3 0.671	100.0* 13.737

N=number of subjects

* Post booster, 94.4% of subjects in the 2-4-12 months schedule had anti-PRP concentration \geq 1 μ g/ml indicative of long-term protection.

† cut-off accepted as indicative of protection

Persistence of the immune response:

The persistence of the immune response to a 3-dose primary and booster schedule with INFANRIX hexa was evaluated in children 4-8 years of age. Protective immunity against the three poliovirus types and PRP was observed in at least 91.0% of children and against diphtheria and tetanus in at least 64.7% of children. At least 25.4% (anti-PT), 97.5% (anti-FHA) and 87.0% (anti-PRN) of children were seropositive against the pertussis components.

With regards to hepatitis B, protective immunity following a 3-dose primary and booster schedule with INFANRIX hexa has been shown to persist in \geq 85% of subjects 4-5 years

of age and in $\geq 72\%$ of subjects 7-8 years of age. Additionally, following a 2-dose primary and booster schedule, protective immunity against hepatitis B persisted in $\geq 48\%$ of subjects 11-12 years of age.

Hepatitis B immunological memory was confirmed in children 4 to 12 years of age. These children had received INFANRIX hexa as primary and booster vaccination in infancy, and when an additional dose of monovalent HBV vaccine was administered, protective immunity was induced in at least 96.8% of subjects.

DETAILED PHARMACOLOGY

Not applicable.

MICROBIOLOGY

Not applicable.

TOXICOLOGY

Not applicable.

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PART III: CONSUMER INFORMATION

INFANRIX hexa

Combined diphtheria and tetanus toxoids, acellular pertussis, hepatitis B (recombinant), inactivated poliomyelitis, and adsorbed conjugated *Haemophilus influenzae* type b vaccine

This leaflet is Part III of a three-part "Product Monograph" published for INFANRIX hexa (Combined diphtheria and tetanus toxoids, acellular pertussis, hepatitis B (recombinant), inactivated poliomyelitis, and adsorbed conjugated *Haemophilus influenzae* type b vaccine) approved for sale in Canada, and is designed specifically for Consumers. This leaflet is a summary and will not tell you everything about INFANRIX hexa. Contact your doctor or pharmacist if you have any questions about the drug.

ABOUT THIS VACCINE

What the vaccine is used for:

INFANRIX hexa is a vaccine used in children for protection against diphtheria, tetanus (lockjaw), pertussis (whooping cough), hepatitis B, poliomyelitis (Polio) and *Haemophilus influenzae* type b diseases.

Vaccination is the best way to protect against these diseases.

What it does:

INFANRIX hexa works by helping the body to make its own protection (antibodies) which protect your child against these diseases.

When it should not be used:

INFANRIX hexa should not be used:

- in children who have a known allergy to any component of the vaccine (see "What the important nonmedicinal ingredients are" section) or children having shown signs of an allergic reaction after a previous dose of this vaccine or any injection containing diphtheria, tetanus, pertussis, hepatitis B, poliovirus, or *Haemophilus influenzae* type b. Signs of an allergic reaction may include skin rash, shortness of breath and swelling of the face or tongue.
- in persons 7 years of age or older.
- in infants who experienced problems of the nervous system within 7 days following previous vaccination with a pertussis (whooping cough) vaccine.

- if your child has an infection with a high temperature (over 38°C). A minor infection such as a cold should not be a problem, but talk to your doctor first.
- if your child has breathing difficulties, please contact your doctor. This may be more common in the first three days following vaccination if your child is born prematurely (before or at 24 weeks of pregnancy).

What the medicinal ingredient is:

INFANRIX hexa contains the following medicinal ingredients: combined diphtheria and tetanus toxoids, hepatitis B (recombinant), three purified pertussis antigens [pertussis toxoid, filamentous haemagglutinin and pertactin (69 kiloDalton outer membrane protein)], inactivated polio virus types 1, 2 and 3, and conjugated *Haemophilus influenzae* type b.

None of the components in the vaccine are infectious. You cannot get the diseases from the INFANRIX hexa vaccine.

What the nonmedicinal ingredients are:

Aluminum (as aluminum salts), lactose, medium 199, sodium chloride, and water for injection. Residues from the manufacturing process: disodium phosphate, formaldehyde, glutaraldehyde, glycine, monopotassium phosphate, neomycin sulphate, polymyxin B sulphate, polysorbate 20 and 80, and potassium chloride.

What dosage forms it comes in:

INFANRIX hexa is a sterile suspension for injection, with the following components:

- PEDIARIX, supplied as a sterile, cloudy suspension for injection in a pre-filled glass syringe.
- *Haemophilus influenzae* type b vaccine, supplied as a pellet in a glass vial.

The 2 components are mixed together before they are given to your child.

WARNINGS AND PRECAUTIONS

Before you use INFANRIX hexa talk to your doctor or pharmacist if:

- your child had any problems (such as high fever, collapse or shock-like state or persistent crying lasting 3 hours or more) within 48 hours or fits (with or without a fever) within 3 days of vaccination with INFANRIX hexa or another vaccine against pertussis (whooping cough).

- your child has a weakened immune system, for example due to HIV infection or due to medicines that suppress the immune system, as your child may not get the full benefit from INFANRIX hexa.
- you have a family history of convulsions.
- your child is suffering from neurological disorders, including infantile spasms, uncontrolled epilepsy or progressive encephalopathy (disease of brain).
- your child has a bleeding problem or bruises easily. INFANRIX hexa should be given with caution since bleeding may occur following vaccination.
- your child has a high temperature (over 38°C).
- your child has any known allergies.
- your child is taking any other medicine or has recently received any other vaccine.
- your child has any serious health problem.
- your child is younger than 6 weeks of age.

Fainting can occur following, or even before, any needle injection; therefore, tell the doctor or nurse if your child fainted with a previous injection.

High incidence of fever (> 39.5°C) was reported in infants receiving INFANRIX hexa and pneumococcal conjugate vaccine (PREVNAR, PREVNAR 13 or SYNFLORIX) compared to infants receiving INFANRIX hexa alone.

Increased reporting rates of fits (with or without fever) and collapse or shock-like state were observed with concomitant administration of INFANRIX hexa and PREVNAR 13.

INTERACTIONS WITH THIS VACCINE

As with other vaccines, INFANRIX hexa should not be given to children on anticoagulant (medicine that prevents blood from clotting) therapy unless the benefits clearly outweigh the risks. Talk to your doctor.

Patients receiving immunosuppressive therapy (medicine that lowers the body's normal immune system response) should delay receiving INFANRIX hexa vaccination until they have been off therapy for 3 months; otherwise you may not be fully protected against the diseases.

PROPER USE OF THIS VACCINE

In case of drug overdose, contact a health care practitioner, hospital emergency department or regional Poison Control Centre immediately, even if there are no symptoms.

Usual dose:

Your child will receive 2 or 3 doses given intramuscularly (into a muscle) with an interval of at least 1 month between each one. Your doctor or nurse will let you know when to come back for the next dose and if an additional booster dose is necessary. INFANRIX hexa should not be administered to your child before the age of 6 weeks.

Missed Dose:

If your child misses a scheduled injection, talk to your doctor and arrange another visit.

Make sure your child finishes the complete vaccination course. If not, your child may not be fully protected against the diseases.

SIDE EFFECTS AND WHAT TO DO ABOUT THEM

Like all vaccines, INFANRIX hexa may occasionally cause unwanted effects.

As with other vaccines in any age group, allergic reactions may occur very rarely (in less than 1 in 10,000 doses of the vaccine). This can be recognised by symptoms such as itchy rash of the hands and feet, swelling of the eyes and face, difficulty in breathing or swallowing and a sudden drop in blood pressure and loss of consciousness. Such reactions will usually occur before leaving the doctor's office. However, you should seek immediate treatment in any event.

See your doctor straight away if your child has any of the following serious side effects:

- collapse
- times when they lose consciousness or have a lack of awareness
- fits – this may be when they have a fever

These side effects have happened very rarely with other vaccines against whooping cough. They usually happen within 2 to 3 days after vaccination.

Other side effects:

Very common side effects (in more than 1 in 10 doses of the vaccine) after having INFANRIX hexa are loss of appetite, irritability, unusual crying, restlessness, pain, redness and swelling at injection site, fever more than 38°C and feeling tired.

Common side effects (in more than 1 in 100 doses of the vaccine) after having INFANRIX hexa are nervousness, vomiting, diarrhea, swelling larger than 5 cm at injection site, fever higher than 39.5°C, itching and a hard lump at injection site.

Uncommon side effects (in more than 1 in 1,000 doses of the vaccine) after having INFANRIX hexa are upper respiratory tract infection, sleepiness, cough and swelling occurring over a large area of the injected limb.

Rare side effects (in more than 1 in 10,000 doses of the vaccine) after having INFANRIX hexa are bronchitis and rash.

Very rare side effects (in less than 1 in 10,000 doses of the vaccine) after having INFANRIX hexa are skin rash, hives, wheeziness or cough, swollen glands in the neck, armpit or groin, bleeding or bruising more easily than normal, temporarily stopping breathing, in babies born very prematurely (at or before 28 weeks of gestation) longer gaps than normal between breaths may occur for 2-3 days after vaccination, swelling of the face, lips, mouth, tongue or throat which may cause difficulty in swallowing or breathing, swelling of the entire injected limb, and blisters at the injection site.

If these symptoms continue or become severe, tell the doctor or nurse.

If your child develops any other symptom within days following the vaccination, tell your doctor as soon as possible.

Do not be alarmed by this list of possible side effects. It is possible that your child will have no side effects from vaccination.

This is not a complete list of side effects. For any unexpected effects while taking INFANRIX hexa, contact your doctor or pharmacist.

HOW TO STORE IT

Store INFANRIX hexa in a refrigerator at 2° to 8°C. **Do not freeze.** Discard if the vaccine has been frozen. Store in the original package in order to protect from light.

After reconstitution immediate use is recommended.

Do not use after expiration date shown on the label. The date for last use corresponds to the last day of the month mentioned.

Store all vaccines out of the reach and sight of children.

REPORTING SUSPECTED SIDE EFFECTS

To monitor vaccine safety, the Public Health Agency of Canada collects case reports on adverse events following immunization.

For health care professionals:

If a patient experiences an adverse event following immunization, please complete the appropriate Adverse Events following Immunization (AEFI) Form and send it to your local Health Unit in [your province/territory](#).

For the General Public:

Should you experience an adverse event following immunization, please ask your doctor, nurse, or pharmacist to complete the Adverse Events following Immunization (AEFI) Form.

If you have any questions or have difficulties contacting your local health unit, please contact Vaccine Safety Section at Public Health Agency of Canada:

By toll-free telephone: 1-866-844-0018

By toll-free fax: 1-866-844-5931

By email: caefi@phac-aspc.gc.ca

At the following website:

<http://www.phac-aspc.gc.ca/im/vs-sv/index-eng.php>

By regular mail:

The Public Health Agency of Canada

Vaccine Safety Section

130 Colonnade Road

Ottawa, Ontario

K1A 0K9 Address Locator 6502A

NOTE: Should you require information related to the management of the side effect, please contact your health care provider before notifying the Public Health Agency of Canada. The Public Health Agency of Canada does not provide medical advice.

MORE INFORMATION

This document plus the full product monograph, prepared for health professionals can be found at:

<http://www.gsk.ca> or by contacting the sponsor,

GlaxoSmithKline Inc.,

7333 Mississauga Road

Mississauga, Ontario

L5N 6L4

1-800-387-7374

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