

Hepatitis B Vaccine Biological Page

Section 7:	Biological Product Information			Standard #: 07.234
Created by:	Provincial Immunization Program Standards and Quality			
Approved by:	Provincial Immunization Program Standards and Quality			
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	Engerix-B Recombivax HB		
Manufacturer	GlaxoSmithKline Inc.	Merck Canada Inc.	
Biological Classification	Non-live: Recombinant		
Indications for Provincially Funded Vaccine	Pre-exposure: Refer to Serology Recommendations and Follow-up for pre-immunization serology recommendations.		
	 basis, generally at 10 years up to and principle should be to offer protection to system Individuals born March 1, 2018 or later who not needed or contraindicated Individuals born in 1981 or later who would hepatitis B vaccine program and who have vaccine. Endemic: Children whose families have immigrated to prevalence of hepatitis B (endemic for hepatigh Endemic Geographic Areas). Non-immune adults who have immigrated prevalence of hepatitis B. See Hepatitis B Areas. 	cine can be provided on a case by case including 18 years of age. The guiding to students prior to them leaving the school en immunization with DTaP-IPV-Hib-HB is d have been eligible for the school universal e not received a series of hepatitis B to Canada from areas where there is a high patitis B). See Hepatitis B Virus Infection - to Canada from areas where there is a high Virus Infection - High Endemic Geographic which hepatitis B is highly endemic, following	
	 Chronic Health Conditions: Individuals with hemophilia and others receiving repeated infusions of blood or bl products (hepatitis B vaccine is not provided for parents providing home infusion their children) Individuals with chronic liver disease from any cause (with the exception of HBsA positive individuals) Individuals with Inflammatory Bowel Disease (IBD), or other chronic inflammatory conditions, who will be on long term immunosuppressive medications including blimited to Imuran or TNF antagonists like Remicade or Humira. Note: Individuals with chronic liver disease with lab confirmation of positive anti-HBs without documentation of any doses of hepatitis B vaccine should be offered a 		

 Individuals with chronic liver disease with lab confirmation of positive anti-HBs with any incomplete series should have their series completed.

Individuals with chronic health conditions that may be HYPORESPONSIVE to hepatitis B vaccine should receive a higher dose of hepatitis B vaccine:

These include:

- Individuals with chronic renal disease or who are undergoing chronic hemodialysis/peritoneal dialysis, including those who are pre-dialysis (progressive renal insufficiency). See Hepatitis B (HBVD) Algorithm for Chronic Renal Disease (alberta.ca) for additional information
- o Individuals with congenital immunodeficiencies
- o Individuals infected with HIV
- o Candidates for and recipients of Solid Organ Transplant (SOT) See:
 - Standard for the Immunization of Transplant Candidates and Recipients
 - Adult SOT Chart
 - Children Expecting SOT Before 18 Months of Age
 - Children Expecting SOT After 18 Months of Age
- Recipients of Hematopoietic Stem Cell Transplant (HSCT) See:
 - Standard for the Immunization of Transplant Candidates and Recipients
 - Adult HSCT Chart
 - Child HSCT Chart

Note:

- Individuals with lab confirmation of positive anti-HBs but without documentation of any doses of hepatitis B vaccine should be offered a series of hepatitis B vaccine to ensure long term immunity.
- Individuals with lab confirmation of positive anti-HBs with any incomplete series should have their series completed.
- Periodic serological testing may be done by the attending physician for hyporesponsive individuals. See Serology section for more information.

Lifestyle Risks:

Individuals with lifestyle risks of infection including:

- Men who have sex with men (MSM)
- o Individuals with more than one sexual partner in the previous 6 months
- o Individuals with a history of a sexually transmitted infection
- Individuals seeking evaluation or treatment for a sexually transmitted infection.
- o Individuals who engage in high risk sexual practices
- o Individuals who have unprotected sex with new partners
- Individuals who use illicit drugs and associated drug-using paraphernalia (for example, needles, tubes used for snorting), resulting in blood exposure.

Occupational/Other Settings:

- Individuals who are workers, volunteers or students (accepted into post-secondary educational programs) and who have a reasonable anticipated risk of exposure to blood/bloody body fluids and/or sharps injuries during the course of their work. See Occupational Considerations for Immunization and Hepatitis B Risk Assessment.
- Children and workers in childcare settings in which there is a hepatitis B infected (acute or chronic) child or worker.
 - o If exceptional circumstances such as biting behavior or special medical conditions exist and Hepatitis B status is unknown, consult with zone MOH/MOH designate.
- Residents and staff of institutions or group homes for the developmentally challenged.
 Inmates in provincial correctional facilities who will be incarcerated for a sufficient length of time to complete a hepatitis B vaccine series.
 - Immunization of inmates in long-term correctional facilities is the responsibility of the Federal Correctional Service. However, vaccine will be provided provincially for completion of immunization of discharged inmates who began their hepatitis B series in federal prisons.

Note:

Hepatitis A and Hepatitis B Combined Vaccine may be considered for individuals 1 year of age and older who are eligible for both pre-exposure hepatitis A and B vaccines if they do not require the double strength hepatitis B vaccine (see <u>HABV Vaccine Biological Page</u>).

Post-exposure:

Refer to <u>Serology Recommendations and Follow-up</u> for post-immunization serology recommendations.

Infants:

Newborns born to hepatitis B surface antigen positive (HBsAg) mothers (acute cases
or carriers) should receive hepatitis B immune globulin (HBIG) and the first dose of
hepatitis B vaccine as soon as possible after birth (within 12 hours) but within 7 days
after birth if HBIG/hepatitis B vaccine is delayed for any reason.

Note:

- o If prenatal screening has not been done prior to delivery, it should be done as soon as possible after admission for delivery. In addition, repeat testing should be considered in uninfected, susceptible women with continuing high risk factors.
- If results can be obtained within 12 hours, the first dose of Hepatitis B vaccine should be administered. HBIG administration should be delayed pending results.
- If results will not be available within 12 hours, administer hepatitis B vaccine and consider administration of HBIG, taking into account maternal risk factors and erring on the side of providing HBIG if there is any question of possible maternal hepatitis B infection.
- Infants (other than newborns) younger than 12 months of age:
 - o Hepatitis B vaccine and HBIG if the mother or primary caregiver is an acute case
 - Hepatitis B vaccine only if the caregiver or significant household contact is a chronic carrier.

Refer to: <u>Public Health Notifiable Disease Management Guidelines – Hepatitis B</u> and <u>Alberta Prenatal Screening Program for Selected Communicable Diseases</u>
Public Health Guidelines – Hepatitis B.

- Susceptible household contacts, sexual partners and needle sharing partners of individuals with acute or chronic hepatitis B infection:
 - Hepatitis B vaccine. HBIG may be recommended for some individuals depending upon the time from exposure and the specific surrounding the exposure.
 Refer to: <u>Public Health Notifiable Disease Management Guidelines – Hepatitis B</u>
 Note:
 - Susceptible household contacts, sexual partners and needle-sharing partners with lab confirmation of positive anti-HBs but without documentation of any doses of hepatitis B vaccine should be offered a series of hepatitis B vaccine to ensure long term immunity.
 - Susceptible household contacts, sexual partners and needle-sharing partners with lab confirmation of positive anti-HBs with any incomplete series should have their series completed.
- Percutaneous (needle stick) or mucosal exposure:
 - Post-exposure follow-up and prophylaxis should be based on the immunization history and antibody status of the exposed person and, if known, the infectious nature of the source.
 - When a susceptible individual sustains a "community needle stick" injury (needle stick in a non-health care setting), the risk of exposure to hepatitis B is increased. If the individual has no history of a hepatitis B vaccine series and the source is HBsAg positive, high risk, unknown or not available for testing, HBIG should be administered (as soon as possible but within seven days of exposure) with the first dose of the hepatitis B vaccine series.

Note:

 Individuals sustaining percutaneous (needle stick) or mucosal exposure with lab confirmation of positive anti-HBs but without documentation of any doses

- of hepatitis B vaccine should be offered a series of hepatitis B vaccine to ensure long term immunity.
- Individuals sustaining percutaneous (needle stick) or mucosal exposure with lab confirmation of positive anti-HBs with any incomplete series should have their series completed.
- Susceptible individuals of sexual assault.
 - HBIG and hepatitis B vaccine should be offered.

Note:

For further guidelines related to post-exposure follow-up refer to the following:

- Public Health Notifiable Disease Management Guidelines Hepatitis B
- Alberta Guidelines for Post-Exposure Management and Prophylaxis: HIV, Hepatitis B, Hepatitis C and Sexually Transmitted Infections
- Canadian Immunization Guide: Hepatitis B Vaccine (Figures 1 & 2)

Serology

See <u>Serology Interpretation</u> and <u>Serology Recommendations and Follow-up</u>.

Schedule and Dose For Healthy Individuals

May consider use of INFANRIX hexa in children 2 months up to and including 23 months of age. See INFANRIX hexa Vaccine Biological page for indications. The schedule and spacing considerations for INFANRIX hexa vaccine varies slightly from those of the individual HBV and DTaP-IPV-Hib vaccines. Ensure the appropriate schedule is followed for the vaccine(s) that are being used.

Individuals being immunized with Engerix-B	
Vaccine	
(20 mcg/1 mL):	

Individuals being immunized with Recombivax HB Vaccine (10 mcg/1 mL):

Newborns born to hepatitis B surface antigen positive (HBsAg) mother (3 doses): Give as 0.5 mL

- Dose 1: at birth, given with HBIG
- o Dose 2: 2 months of age
- Dose 3: 6 months of age

Note:

 Third dose should not be given to infants before 6 months (24 weeks or 168 days) of age.

Newborns with birth weight less than 2,000 grams born to hepatitis B surface antigen positive (HBsAg) mother (4 doses):

Give as 0.5 mL

- o Dose 1: at birth, given with HBIG
- o Dose 2: 1 month of age
- o Dose 3: 2 months of age
- Dose 4: 6 months of age

Note:

- The response to hepatitis B vaccine may be diminished in infants with a birth weight below 2,000 grams.
- Fourth dose should not be given to infants before 6 months (24 weeks or 168 days) of age.
- Serologic testing at minimum 9 months of age and at least 1 month after completion of 4-dose series is recommended.

Other infants from birth up to and including 11 months of age (3 doses): Give as 0.5 mL

- Dose 1: 2 months of age
- Dose 2: 4 months of age
- Dose 3: 12 months of age

Note:

- Third dose should not be given to infants before 6 months (24 weeks or 168 days) of age.
- If the infant is identified as a significant household contact of a hepatitis B carrier, offer hepatitis B vaccine as soon as possible.
- Minimum intervals can be used for children who begin immunization off schedule refer to Spacing Considerations.

Infants with birth weight less than 2,000 grams who receive hepatitis B vaccine at birth (for example, father or other primary caregiver is HBsAg positive) (4 doses): Give as 0.5 mL

- o Dose 1: at birth
- o Dose 2: 1 month of age
- o Dose 3: 2 months of age
- o Dose 4: 6 months of age

Note:

- The response to hepatitis B vaccine may be diminished in infants with a birth weight below 2,000 grams.
- Serologic testing at minimum 9 months of age and at least 1 month after completion of series is recommended.
- Fourth dose should not be given to infants before 6 months (24 weeks or 168 days) of age.

Children 12 months of age up to and including 10 years of age (3 doses): Give as 0.5 mL

- o Dose 1: day 0
- o Dose 2: 1 month after dose 1
- Dose 3: 6 months after dose 1, and 5 months after dose 2

Note:

• If the child is identified as a significant household contact of a hepatitis B carrier, offer hepatitis B vaccine as soon as possible.

Students 11 years of age up to and including 15 years of age (2 doses):

- This includes grade 6 students younger than 11 years of age as eligibility for a two-dose series is determined by grade level.
 - Dose 1: day 0
 - O Dose 2: 6 months after dose 1

Minimal acceptable spacing between the first and second dose is 24 weeks.

Note:

- In the event that a 0.5 mL dose is given, a 3-dose schedule must be followed: Give as 0.5 mL
 - o Dose 1: day 0
 - o Dose 2: 1 month after dose 1
 - Dose 3: 6 months after dose 1, and 5 months after dose 2
- The minimum acceptable interval is 0, 1 and 4 months, with 1 month (28 days) between the first and second dose, at least 2 months (56 days) between the second and third dose and at least 4 months (112 days) between the first and third dose.
- If a student will turn 16 years of age before a 2-dose series can be completed, a 3-dose schedule should be initiated (see below).

Children 16 years of age up to and including 19 years of age (3 doses): Give as 0.5 mL

	Engerix-B	Recombivax HB		
	 Dose 1: day 0 Dose 2: 1 month after dose 1 Dose 3: 6 months after dose 1, and 5 months after dose 2 Note: For individuals who received a 1 mL dose of hepatitis B vaccine as their first dose at 11 to 15 years of age and present at 16 years of age or older for subsequent doses, the series reverts to a 3-dose schedule following appropriate dosing for age. 			
	Adults 20 years of age and older (3 doses): Give as 1 mL			
	 Spacing Considerations: If the recommended schedule cannot be followed, refer to Standard for Recommended Immunization Schedules Section 5: Minimum Age and Minimum Intervals Between Vaccine Doses. Interruption of the immunization schedule does not require any dose(s) be repeat the minimum intervals between doses are respected. For those who may have an alternate immunization history refer to Standard for Recommended Immunization Schedules. If a second hepatitis B immunization series is required, this can be started once the need is identified. Immunization started in another province or territory prior to grade 6 can be completed as they present to public health using the current schedule and dose 			
Schedule and Dose for Hyporesponsive	Hyporesponsive individuals being immunized with Engerix-B Vaccine (20 mcg/1 mL):	Hyporesponsive individuals being immunized with Recombivax HB Vaccine (10 mcg/1 mL):		
Individuals	Birth to 15 years of age (3 doses): Give as 1 mL	Individuals from birth up to and including 15 years of age (3 doses): Give as 1 mL		

Engerix-B	Recombivax HB	
16 years to	19 years of age	
Individuals 16 years of age up to and including 19 years of age (4 doses): Give as 2 mL Dose 1: day 0 Dose 2: 1 month after dose 1 Dose 3: 2 months after dose 1 Dose 4: 6 months after dose 1 Note: Those initiating a 4-dose schedule with Engerix-B should complete the series using the same vaccine whenever possible. Minimum interval between the third and fourth dose is 4 months and at least 6 months between the first and fourth dose.		
If any dose in the series was Engerix-B, a administered for those 16 years of age and		
Hyporesponsive individuals being immunized with Engerix-B Vaccine (20 mcg/1 mL):	Hyporesponsive individuals being immunized with Recombivax HB Adult Dialysis Strength Vaccine (40 mcg/1 mL):	
20 years and older		
Individuals 20 years of age and older	Individuals 20 years of age and older	

Individuals 20 years of age and older (4 doses):

Give as 2 mL

- o Dose 1: day 0
- o Dose 2: 1 month after dose 1
- o Dose 3: 2 months after dose 1
- Dose 4: 6 months after dose 1

Note:

- Those initiating a 4-dose schedule with Engerix-B should complete the series using the same vaccine whenever possible.
- Minimum interval between the third and fourth dose is 4 months and at least 6 months between the first and fourth dose.

Individuals 20 years of age and older (3 doses)

Give as 1 mL

- o Dose 1: day 0
- o Dose 2: 1 month after dose 1
- Dose 3: 6 months after dose 1, and 5 months after dose 2

Note:

- Do not use this formulation for individuals younger than 20 years of age.
- Hyporesponsive persons 20 years of age and older should receive Recombivax HB
 Dialysis Strength Vaccine. If Recombivax HB Dialysis Strength Vaccine is
 unavailable or the person has a medical contraindication to this product, Engerix-B
 may be used following the schedule outlined.
- If any dose in the series was Engerix-B, a total of 4 doses of vaccine should be administered for those 16 years of age and older.

Spacing Considerations:

- Interruption of the immunization schedule does not require any dose(s) be repeated if the minimum intervals between doses are respected.
- For those who may have an alternate immunization history refer to <u>Standard for</u> <u>Recommended Immunization Schedules</u>.

	Engerix-B	Recombivax HB		
	 If a second hepatitis B immunization series is required, this can be started once the need is identified. Immunization started in another province or territory prior to grade 6 can be completed as they present to public health using the current schedule and dose recommended in Alberta. 			
Preferred Use	 There is no preference indicated for the use of Recombivax HB or Engerix-B for those eligible for regular strength vaccine. Both vaccines are safe and immunogenic for all ages Persons with medical contraindications to one product should be offered the alternate product if supply is available 			
Route	 Note: Vaccine should not be administered in the gluteal areas as this may result in lower immune response. If vaccine is inadvertently given in the gluteal area the individual should be tested for immunity and re-immunized if antibody concentrations are inadequate. 			
Contraindications/ Precautions	Contraindications: Known severe hypersensitivity to any component of a hepatitis B containing vaccine. Anaphylactic reactions or other allergic reactions to a previous dose of vaccine containing similar components. For Recombivax HB only: Anaphylactic reactions to latex. Precautions: None identified. 			
Possible Reactions	 Common: Injection site pain, soreness, tenderness, pruritus, erythema, ecchymoses, swelling, induration, warmth and nodule formation. Irritability, headache, fatigue/asthenia, malaise, pharyngitis and fever. Loss of appetite, nausea, vomiting, diarrhea and abdominal pain. Uncommon: Dizziness, myalgia Rare: Lymphadenopathy, paresthesia, rash, urticaria and arthralgia. Anaphylaxis, angioedema, allergic reactions. As with any immunization, unexpected or unusual side effects can occur. Refer to the product monograph for more detailed information. Note: A number of studies have been unable to demonstrate any evidence of a causal association following hepatitis B vaccine and the following chronic illnesses: chronic fatigue syndrome, multiple sclerosis, Guillain-Barré syndrome (GBS) or rheumatoid 			
Pregnancy	 arthritis. Hepatitis B vaccine should be administered to pregnant women when indicated. Data is not available on the effect of hepatitis B vaccine on fetal development; however, the risk is expected to be negligible as the vaccine consists of non-infectious subunits. Eligible pregnant woman should receive provincially funded vaccine. Pregnant women at high risk of hepatitis B infection should be tested for antibody response following receipt of hepatitis B vaccine series. See serology section for more information. 			
Lactation	Can be administered to eligible breastfeeding women. It is not known whether hepatitis B vaccine is excreted in human milk.			
Composition	Each 0.5 mL dose contains: • 10 mcg hepatitis B surface antigen • 0.25 mg aluminum hydroxide. Each 0.5 mL dose contains: • 5 mcg hepatitis B surface antigen • 0.25 mg amorphous aluminum hydroxyphosphate			

Each 1 mL dose contains:		Engerix-B	Recombivax HB	
Blood/Blood Products Contains no human blood/blood products. Contains no bovine or porcine products. Contains no bovine or porcine products. Latex Does not contain latex. Latex in vial stopper. Hepatitis B vaccines produced by different manufacturers can be used interchangeably despite different doses and schedules. The dose administered should be that recommended by the manufacturer for the specific product being used. When possible, series should be completed with the same vaccine, especially with hyporesponsive individuals. If this is not possible, hyporesponsive individuals 16 years and older who have received any doses of Engerix-B vaccine should be completed using the 4-dose schedule. Refer to schedule and dose section for more details. Administration with Other Products Appearance Slightly opaque, white suspension. Storage Storage Storage Contains no human blood/blood products. Latex in vial stopper. When possible, series should be that recommended by the manufacturer for the specially with hyporesponsive individuals. If this is not possible, specially with hyporesponsive individuals. If this is not possible, specially with hyporesponsive individuals. If this is not possible, specially with hyporesponsive individuals. If this is not possible, specially with hyporesponsive individuals. If this is not possible, specially with hyporesponsive individuals. If this is not possible, specially with hyporesponsive individuals. If this is not possible, specially with hyporesponsive individuals. If this is not possible, specially with hyporesponsive individuals. If this is not possible, specially		 20 mcg hepatitis B surface antigen 0.5 mg aluminum hydroxide. Non-medicinal Ingredients: disodium phosphate hydrate sodium chloride sodium dihydrogen phosphate dihydrate water for injection. Single dose presentations are preservative 	 35 mcg sodium borate Water for injection. Each 1 mL dose contains: 10 mcg hepatitis B surface antigen 0.5 mg amorphous aluminum hydroxyphosphate 9 mg sodium chloride 70 mcg sodium borate Water for injection. Each 1 mL dose of Recombivax HB Dialysis Strength contains: 40 mcg hepatitis B surface antigen 0.5 mg amorphous aluminum hydroxyphosphate 9 mg sodium chloride 70 mcg sodium borate Water for injection. The following manufacturing residuals may be found in the above preparations of Recombivax HB vaccine: Less than 1% yeast protein Less than 15 mcg/mL formaldehyde. These presentations are preservative free 	
Bovine/Porcine Products Latex Does not contain latex. Latex in vial stopper. Hepatitis B vaccines produced by different manufacturers can be used interchangeability Hepatitis B vaccines produced by different manufacturers can be used interchangeably despite different doses and schedules. The dose administered should be that recommended by the manufacturer for the specific product being used. When possible, series should be completed with the same vaccine, especially with hyporesponsive individuals. If this is not possible, hyporesponsive individuals 16 years and older who have received any doses of Engerix-B vaccine should be completed using the 4-dose schedule. Refer to schedule and dose section for more details. Administration with Other Products Products Appearance Slightly opaque, white suspension. Storage Storage Storage Storage Storage Contains no bovine or porcine products. Latex in vial stopper. Latex	Dia ad/Dia ad			
Latex Does not contain latex. Latex in vial stopper.		Contains no numan blood/blood products.		
Interchangeability • Hepatitis B vaccines produced by different manufacturers can be used interchangeably despite different doses and schedules. • The dose administered should be that recommended by the manufacturer for the specific product being used. • When possible, series should be completed with the same vaccine, especially with hyporesponsive individuals. If this is not possible, hyporesponsive individuals 16 years and older who have received any doses of Engerix-B vaccine should be completed using the 4-dose schedule. Refer to schedule and dose section for more details. Administration with Other Products • May be given at the same time as other inactivated and live vaccines using a separate needle and syringe for each vaccine. • The same limb may be used if necessary, but different sites must be chosen. Slightly opaque, white suspension. Storage • Store at +2°C to +8°C. • Do not freeze. • Do not use past the expiry date.		Contains no bovine or porcine products.		
interchangeably despite different doses and schedules. The dose administered should be that recommended by the manufacturer for the specific product being used. When possible, series should be completed with the same vaccine, especially with hyporesponsive individuals. If this is not possible, hyporesponsive individuals 16 years and older who have received any doses of Engerix-B vaccine should be completed using the 4-dose schedule. Refer to schedule and dose section for more details. Administration with Other Products May be given at the same time as other inactivated and live vaccines using a separate needle and syringe for each vaccine. The same limb may be used if necessary, but different sites must be chosen. Slightly opaque, white suspension. Storage Storage Store at +2°C to +8°C. Do not freeze. Do not use past the expiry date.	Latex	Does not contain latex.	Latex in vial stopper.	
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Storage Store at +2°C to +8°C. Do not freeze. Do not use past the expiry date.	with Other	separate needle and syringe for each vaccine.		
 Do not freeze. Do not use past the expiry date. 	Appearance	Slightly opaque, white suspension.		
	Storage	Do not freeze.Do not use past the expiry date.		
Vaccine CodeHBVHBV – regular strength productHBVD – dialysis strength vaccine	Vaccine Code	HBV		

Antigen Code HBV Licensed for Individuals of		Recombivax HB regular strength: Licensed for persons of all ages. Vaccine dose for all children from birth up to and including 10 years of age in Alberta is 0.5 mL (5µg). Recombivax HB Dialysis Strength: Licensed for individuals 20 years of age
Licensed for Individuals of		 Licensed for persons of all ages. Vaccine dose for all children from birth up to and including 10 years of age in Alberta is 0.5 mL (5µg). Recombivax HB Dialysis Strength: Licensed for individuals 20 years of age
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for infar 1995 Se for stude 1999 Se students 2011 Au age incl 2016 Ju DTaP-IF 2017 No those w complet 2018 Fe school u hepatitis 2018 Se change 2019 Au dose to 2023 De 2024 Ap 2025 Ja	ats at high risk; Hepatitis B Dialysis eptember: Hepatitis B introduced ents in grade 5. Eptember: Hepatitis B catch-up so so was available from September august: Hepatitis B vaccine change uding students in Grade 5 from 0 aly 1: Infanrix hexa® introduced for PV-Hib and hepatitis B. Every beautified by the properties of the properties	e in dosage for all individuals 0-10 years of .25 mL to 0.5 mL. or children under 2 years of age eligible for recommended documented series for anti-HBs positive and recommend a tive after first series. or later who would have been eligible for the ram and who have not received a series of tis B vaccine. atory Bowel disease (IBD) who will be on a became eligible for hepatitis B vaccine. Ization schedule for hepatitis B vaccine or grade 6. On schedule for hepatitis B changed from 3 prology recommendations updated. The recommendations updated ally funded vaccine updated. Post-
	s B Vaccine Information Sheet	

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Groups	Pre-immunization Serology recommendation	Post- immunization Serology recommendation	Follow-up Special Considerations
Individuals with <u>chronic renal</u> <u>disease</u> including hemodialysis, peritoneal dialysis, and pre-dialysis	Pre-immunization serology is not routinely recommended	Serology (anti- HBs) should be done 1 – 6 months following the primary series of hepatitis B vaccine	Individuals who are hyporesponsive due to renal disease (hemodialysis, peritoneal dialysis and predialysis) often respond suboptimally to hepatitis B vaccine and should receive a higher vaccine dose according to the schedule for hyporesponsive individuals. If protection is achieved and then wanes, subsequent exposure may result in acute disease or carrier state.
			Individuals who are anti-HBs negative after the first series should receive a second series , followed by serology one month later.
			Individuals with chronic renal disease or on dialysis should be evaluated annually for anti-HBs. Should anti-HBs testing show suboptimal protection, a booster dose of vaccine should be given.
			See <u>Hepatitis B (HBVD) Algorithm for Chronic</u> <u>Renal Disease (alberta.ca)</u> for additional information.
			Individuals with lab confirmation of positive anti- HBs but without documentation of any doses of hepatitis B vaccine OR those with incomplete series should be offered a complete series of hepatitis B vaccine to ensure long term immunity.
Individuals with congenital immunodeficiencies Candidates for and recipients of solid organ transplant (SOT)	Pre-immunization serology is not routinely recommended	Serology (anti- HBs) should be done 1 – 6 months following the primary series of hepatitis B	Individuals who are hyporesponsive due to congenital immunodeficiencies, HSCT, SOT and HIV infection often respond suboptimally to hepatitis B vaccine and may need additional antigen to mount a response. If protection is achieved and then wanes, subsequent exposure may result in acute disease or carrier state.
Recipients of hematopoietic stem cell transplant		vaccine	Individuals who are anti-HBs negative after the first series should receive a second series , followed by serology one month later.
(HSCT)			Periodic monitoring (by attending physician) for the presence of anti-HBs should be considered, taking into account the severity of the compromised state and whether or not the risk for hepatitis B infection is still present. Should anti-HBs testing show suboptimal protection, a booster dose of vaccine and retesting should be undertaken.
			Individuals with lab confirmation of positive anti- HBs (and anti-HBc negative) but without documentation of any doses of hepatitis B vaccine OR those with incomplete series should be offered a complete series of hepatitis B vaccine to ensure long term immunity.

Groups	Pre-immunization Serology recommendation	Post- immunization Serology recommendation	Follow-up Special Considerations
 Individuals infected with HIV 	Pre-immunization serology (anti- HBs, HBsAg and anti-HBc) is recommended	Serology (anti- HBs) should be done 1 – 6 months following the primary series of hepatitis B	Individuals who are hyporesponsive due to congenital immunodeficiencies, HSCT, SOT and HIV infection often respond suboptimally to hepatitis B vaccine and may need additional antigen to mount a response. If protection is achieved and then wanes, subsequent exposure may result in acute disease or carrier state.
		vaccine	Individuals who are anti-HBs negative after the first series should receive a second series , followed by serology one month later.
			Periodic monitoring (by attending physician) for the presence of anti-HBs should be considered, taking into account the severity of the compromised state and whether or not the risk for hepatitis B infection is still present. Should anti-HBs testing show suboptimal protection, a booster dose of vaccine and retesting should be undertaken.
			Individuals with lab confirmation of positive anti-HBs (and anti-HBc negative) but without documentation of any doses of hepatitis B vaccine OR those with incomplete series should be offered a complete series of hepatitis B vaccine to ensure long term immunity.
Individuals with <u>chronic liver</u> <u>disease</u>	Pre-immunization serology (anti- HBs, HBsAg and anti-HBc) is recommended	Serology (anti- HBs) should be done 1 – 6 months following the	Individuals who are anti-HBs negative after the first series should receive a second series using a higher dose vaccine schedule for hyporesponsive individuals followed by serology one month later.
		primary series of hepatitis B vaccine	Individuals with lab confirmation of positive anti- HBs (and anti-HBc negative) but without documentation of any doses of hepatitis B vaccine OR those with incomplete series should be offered a complete series of hepatitis B vaccine to ensure long term immunity.
Newborns born to hepatitis B infected mothers Infants (other than newborns) younger than 12 months of	Pre-immunization serology is not recommended	Serology (anti- HBs and HBsAg) is recommended 1 – 6 months following the primary series	If the individual is anti-HBs negative after the first series, a second hepatitis B vaccine series should be administered, with repeat serology testing one month later. Once a positive antibody result is documented no further serology is recommended.
age with hepatitis B infected caregiver or household contact		of hepatitis B vaccine and the infant should be at least 9 months of age	Refer to: Public Health Notifiable Disease Management Guidelines – Hepatitis B and Alberta Prenatal Screening Program for Selected Communicable Diseases Public Health Guidelines – Hepatitis B.

Groups	Pre-immunization Serology recommendation	Post- immunization Serology	Follow-up Special Considerations
Susceptible household contacts, sexual partners and needle-sharing partners of individuals with acute or chronic hepatitis B infection HCWs and Post- Secondary Health Care Students		Serology recommendation with Notifiable ont Guidelines — ement of or High Risk] of B Infection) for	
	partners and needle sharing partners of a hepatitis B case or chronic carrier. • Household		immunity. HCWs who have been previously assessed do not require reassessment or updating at this time. Once a positive anti-HBs result is documented no further serology is recommended.
	contacts of a hepatitis B case or chronic carrier.		
 Individuals who are workers or volunteers and who 	Pre-immunization serology is not	Serology (anti- HBs) should be done 1 – 6	If the individual is anti-HBs negative within 6 months of completion of the first series, a second hepatitis B

Groups	Pre-immunization	Post-	Follow-up
Groups	Serology recommendation	immunization Serology recommendation	Special Considerations
have a reasonable anticipated risk of exposure to blood/bloody body fluids	routinely recommended	months following the primary series of hepatitis B vaccine	vaccine series should be administered, with repeat serology testing one month later.
			If the vaccine series was completed more than 6 months previously and post-immunization serology was not done, testing should be done as part of a routine assessment.
			o If the individual is anti-HBs negative, one booster dose of hepatitis B vaccine should be administered followed by serology one month later. If the individual is still negative after the booster dose, the second series of hepatitis B vaccine should be completed followed by serology one month later.
			Once a positive anti-HBs result is documented no further serology is recommended.
 Individuals who use illicit drugs and associated drug-using 	Pre-immunization serology (anti- HBs, HBsAg and anti-HBc) is recommended	Serology (anti- HBs) should be done 1 – 6 months following the primary series of hepatitis B vaccine	If the individual is anti-HBs negative after the first series, a second hepatitis B vaccine series should be administered, with repeat serology testing one month later.
paraphernalia resulting in blood exposure			If vaccine series was completed more than 6 months previously and post-immunization serology was not done, testing should be done as part of a routine assessment.
			o If the individual is anti-HBs negative, the individual should be given one booster dose of hepatitis B vaccine followed by serology one month later. If the individual is still negative after the booster dose the second series of hepatitis B vaccine should be completed followed by serology one month later.
			Individuals with lab confirmation of positive anti- HBs but without documentation of any doses of hepatitis B vaccine should be offered a series of hepatitis B vaccine to ensure long term immunity.
			Individuals with lab confirmation of positive anti- HBs with any incomplete series should have their series completed.
			Once a positive anti-HBs result is documented no further serology is recommended.
Lifestyle risks • Men who have sex with men (MSM)	Pre-immunization serology (anti- HBs, HBsAg and	Post- immunization serology is not	Reimmunization (that is, booster dose or reimmunization with a complete series) is not generally recommended.
, ,	anti-HBc) is recommended	routinely recommended	For individuals who were immunized as infants, children or adults, testing for anti-HBs years after

Groups	Pre-immunization Serology recommendation	Post- immunization Serology recommendation	Follow-up Special Considerations			
Individuals with more than one sexual partner in the previous 6			immunization might not distinguish vaccine non- responders from responders. Anti-HBs wanes and titres may become non-detectable over time; however immune memory persists.			
months • Individuals with a history of a sexually transmitted infection (STI)			If post-immunization serology was inadvertently done and found to be anti-HBs negative, one booster dose of hepatitis B vaccine should be administered. Additional serology is not required.			
Individuals seeking evaluation or treatment for an STI						
Individuals who engage in high risk sexual practices						
Individuals who have unprotected sex with new partners						
Adults who have immigrated to Canada from endemic areas	Pre-immunization serology (anti- HBs, HBsAg and anti-HBc) is recommended	Post- immunization serology is not recommended	Reimmunization (that is, booster dose or reimmunization with a complete series) is not generally recommended.			
			For individuals who were immunized as infants, children or adults, testing for anti-HBs years after immunization might not distinguish vaccine non-responders from responders. Anti-HBs wanes and titres may become non-detectable over time; however immune memory persists.			
			If post-immunization serology was inadvertently done and found to be anti-HBs negative, these individuals do not quality for additional doses of provincially funded vaccine.			
Individuals with hemophilia and others receiving repeated infusions of blood or blood products	Pre-immunization serology is not routinely recommended	Post- immunization serology is not routinely recommended	Reimmunization (that is, booster dose or reimmunization with a complete series) is not generally recommended. For individuals who were immunized as infants,			
Individuals with Inflammatory Bowel disease (IBD), or other chronic inflammatory conditions, who will be on long term	Pre-immunization serology is not recommended	Post- immunization serology is not recommended	children or adults, testing for anti-HBs years after immunization might not distinguish vaccine non-responders from responders. Anti-HBs wanes and titres may become non-detectable over time; however immune memory persists.			

For individuals who are eligible for more than one reason – follow the most comprehensive serology recommendations for the respective eligibility groups.

Groups	Pre-immunization Serology recommendation	Post- immunization Serology recommendation	Follow-up Special Considerations	
immunosuppressive medications			If post-immunization serology was inadvertently done and found to be anti-HBs negative these individuals do not quality for additional doses of provincially funded vaccine	
Children whose families have immigrated to Canada from an endemic area	Pre-immunization serology is not recommended	Post- immunization serology is not recommended		
Populations or communities in Alberta in which hepatitis B is highly endemic	Pre-immunization serology is not routinely recommended	Post- immunization serology is not recommended		
Children and workers in childcare settings in which there is a hepatitis B infected child or worker Residents and staff of institutions or group homes for the developmentally challenged Inmates in provincial correctional facilities	Pre-immunization serology is not recommended	Post- immunization serology is not recommended	Reimmunization (that is, booster dose or reimmunization with a complete series) is not generally recommended. For individuals who were immunized as infants, children or adults, testing for anti-HBs years after immunization might not distinguish vaccine non-responders from responders. Anti-HBs wanes and titres may become non-detectable over time; however immune memory persists. If post-immunization serology was inadvertently done and round to be anti-HBs negative, these individuals do not qualify for additional doses of provincially funded vaccine. See statements above.	
 Students in Grade 6 Students in Grades 7 through 12 Individuals born in 1981 or later 	Pre-immunization serology is not recommended	Post- immunization serology is not recommended		
Percutaneous (needle stick) or mucosal exposure (blood and body fluid exposures)	Refer to: Alberta Guidelines for Post-Exposure Management and Prophylaxis: for specific serology recommendations and interpretation.			
Susceptible individuals of sexual assault	Refer to: <u>Alberta Guidelines for Post-Exposure Management and Prophylaxis</u> : for specific serology recommendations and interpretation.			

Any individual who fails to respond to the second series of vaccine is unlikely to benefit from further doses. Therefore, if protective levels are not achieved, the individual should be considered a non-responder and susceptible.

SEROLOGY INTERPRETATION				
Serology Result	Interpretation			
anti-HBs <u>positive</u> ** HBsAg negative anti-HBc negative	Considered immune. Refer to Serology Recommendations and Follow-Up Table for those requiring documented doses of hepatitis B vaccine regardless of positive anti-HBs serology.			
 anti-HBs <u>positive</u>** HBsAg negative anti-HBc <u>positive</u> 	Considered immune. No vaccine indicated.			
anti-HBs negativeHBsAg negativeanti-HBc negative	Susceptible. Proceed with immunization as per eligibility criteria.			
anti-HBs negativeHBsAg positiveanti-HBc negative or positive	No vaccine indicated. Refer to: Alberta Public Health Hepatitis B Notifiable Disease Guidelines for interpretation and follow-up.			
anti-HBs negativeHBsAg negativeanti-HBc positive	Refer to: Alberta Public Health Hepatitis B Notifiable Disease Guidelines for interpretation and follow-up.			
**Anti-HBs positive is greater than or equal to 10 IU/L; negative is less than 10 IU/L				