

**Policy:** Publicly Funded Vaccine/Immunoglobulin Eligibility Policy

**Originating Branch:** Office of the Chief Medical Officer of Health

**Original Approval Date:** July 6, 2015      **Effective Date:** July 6, 2015

**Revised Approval Date:** February 1, 2019      **Effective Date:** February 1, 2019

**Approved By:** 

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**Dr. Robert Strang, Chief Medical Officer of Health, Health and Wellness**

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**1. POLICY STATEMENT**

- 1.1. Providing immunization to residents of Nova Scotia is a responsibility shared between the Department of Health and Wellness (DHW), the Nova Scotia Health Authority (NSHA), the Izaak Walton Killam Health Centre (IWK), primary care providers and health care organizations.
- 1.2. DHW provides policies, standards and guidelines for each of the vaccine programs and procures the vaccines/immunoglobulins to be included in the publicly funded program.
- 1.3. NSHA, the IWK, primary care providers and health care organizations implement the programs to Nova Scotians in adherence with those policies, standards and guidelines.

**2. DEFINITIONS**

- 2.1. N/A

**3. POLICY OBJECTIVES**

- 3.1. To protect residents of Nova Scotia and others as identified in the policy from vaccine preventable diseases.
- 3.2. To provide guidance for public health providers and other immunization providers to identify which vaccines/immunoglobulins are publicly funded in Nova Scotia and who is eligible to receive them.

**4. APPLICATION**

- 4.1. This policy applies to all public health and other immunization providers who provide publicly funded vaccine.

## 5. POLICY DIRECTIVES

### Eligibility

#### 5.1. Residents of Nova Scotia

- 5.1.1. All residents of Nova Scotia with a valid Nova Scotia health card are eligible to receive publicly funded vaccines/immunoglobulins as described in Appendix A.
- 5.1.2. Individuals who have become residents of Nova Scotia and started an immunization series out of province:
  - Will finish the series as appropriate based on the Nova Scotia schedule.
  - Will follow the same eligibility as residents of Nova Scotia, regardless of eligibility out of province.
- 5.1.3. Individuals who have started a series of immunizations as part of post exposure prophylaxis out of province will be able to have the series completed in Nova Scotia.

#### 5.2. Visitors and/or Temporary Residents

- 5.2.1. Vaccines/immunoglobulins, with the exception of the influenza vaccine, are not routinely provided through the publicly funded immunization program to visitors or temporary residents of Nova Scotia.
- 5.2.2. The eligibility criteria for these individuals may change based on their circumstances.
- 5.2.3. A risk assessment approach in consultation with the Medical Officer of Health is to be used when making a decision regarding immunization of non-residents with publicly funded vaccines.
- 5.2.4. Individuals who have started a series of immunizations as part of post exposure prophylaxis out of province will be able to have the series completed in Nova Scotia.

#### 5.3. School Based Program

- 5.3.1. Youth who have moved to Nova Scotia are eligible for each of the vaccines included in the school based program:
  - 1) if they would have been in grade 7 (regardless of where they lived) at the time each of the vaccines were added to the school based program. (For example, HPV for males in grade 7 was implemented in 2015); and
  - 2) if they are up to and including 18 years of age.
- 5.3.2. Youth who have missed or refused immunizations included in the school based program are eligible for each of the vaccines:
  - 1) if they would have been in grade 7 at the time the missed or refused vaccine was added to the school based program (For example, HPV for males in grade 7 was implemented in 2015); and
  - 2) if they are up to and including 18 years of age.

## 6. POLICY GUIDELINES

- 6.1. Publicly funded vaccines may be provided through the publicly funded program to residents/non-residents of Nova Scotia as part of outbreak/pandemic management, contact management or prevention of communicable diseases in high risk populations more susceptible regardless of residency status: immigrants, refugees and other individuals establishing residency in Nova Scotia.

- 6.2. Products included in Appendix A may vary based on national contracts and availability.

## 7. ACCOUNTABILITY

- 7.1. DHW is responsible to ensure the policy is current, evidence informed and reviewed every two years.
- 7.2. The NSHA and the IWK are accountable to ensure this policy is communicated to public health staff and other immunization providers within NSHA and the IWK respectively.
- 7.3. Public Health staff and other immunization providers who provide publicly funded immunizations are responsible for adhering to this policy.

## 8. MONITORING / OUTCOME MEASUREMENT

- 8.1. DHW is responsible for defining and monitoring strategic outcomes associated with this policy.
- 8.2. The NSHA and the IWK are responsible for monitoring the implementation of this policy.

## 9. REPORTS

- 9.1. N/A

## 10. REFERENCES

- 10.1. Government of New Brunswick (2013). *Policy 2.2: Eligibility Criteria for Publicly Funded Vaccine and Biologics*.
- 10.2. Public Health Agency of Canada. *Canadian Immunization Guide*. Retrieved from <https://www.canada.ca/en/public-health/services/canadian-immunization-guide.html>
- 10.3. Public Health Agency of Canada. *National Advisory Committee On Immunization Recommendations, Statements and Updates*. Retrieved from <http://www.phac-aspc.gc.ca/naci-ccni/index-eng.php>

## 11. APPENDICES

- 11.1. Appendix A: Nova Scotia Publicly Funded Vaccine/Immunoglobulin Eligibility

## 12. INQUIRIES

Communicable Disease Prevention and Control  
Nova Scotia Department of Health and Wellness  
Tel: (902) 424-8160  
Email: [cdpc@novascotia.ca](mailto:cdpc@novascotia.ca)

## Appendix A: Nova Scotia Publicly Funded Vaccine/Immunoglobulin Eligibility

Abbreviation	National Active Immunizing Agent (Type)	Vaccine Products e.g.	Eligibility
<a href="#">DTaP-IPV-Hib</a>	Diphtheria, tetanus toxoid, acellular pertussis, inactivated polio, haemophilus influenzae type b	Pediacel	<ul style="list-style-type: none"> <li>Routine immunization of children 2 months to 6 years of age</li> <li>*Re-immunization of individuals 7 years of age and older post Hematopoietic Stem Cell Transplant (HSCT)</li> </ul>
<a href="#">Tdap IPV</a>	Tetanus toxoid, diphtheria, acellular pertussis, inactivated polio	Adacel Polio Boostrix Polio	<ul style="list-style-type: none"> <li>Routine immunization booster for children 4 to 6 years of age</li> <li>Immunization of individuals 7 to 17 years of age who are unimmunized or have incomplete immunization</li> <li>Immunization of adults who are unimmunized</li> </ul>
<a href="#">Tdap</a>	Tetanus toxoid, diphtheria, acellular pertussis	Adacel Boostrix	<ul style="list-style-type: none"> <li>Grade 7 students (school based immunization program)</li> <li>Youth who have missed or refused Tdap vaccine as part of the school based program, up to and including 18 years of age</li> <li>Pregnant women, in every pregnancy, irrespective of previous Tdap history</li> <li>Immunization of individuals 18 years of age and older who are unimmunized or have incomplete immunization</li> <li>Adults who require a tetanus or pertussis vaccine and have not received a pertussis containing vaccine in adulthood should receive a single dose of Tdap</li> </ul>
<a href="#">Td</a>	Tetanus toxoid, diphtheria	Td adsorbed	<ul style="list-style-type: none"> <li>Adult booster every 10 years following one dose of Tdap as an adult</li> <li><a href="#">Post-exposure/wound management</a></li> </ul>

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<a href="#">Hib</a>	Haemophilus influenzae type b	Act-Hib	<ul style="list-style-type: none"> <li>• *Pre-exposure prophylaxis for individuals 5 years of age and older with the following high risk conditions:                             <ul style="list-style-type: none"> <li>◦ Cancer: Malignant hematologic disorders only e.g. Leukemia or Lymphoma</li> <li>◦ Cochlear implant</li> <li>◦ Congenital immunodeficiency</li> <li>◦ Hematopoietic stem cell transplant (HSCT) if not receiving <a href="#">DTaP-IPV-Hib</a></li> <li>◦ Lung transplants only</li> <li>◦ <a href="#">Splenic disorders</a> including sickle cell disease or other hemoglobinopathies</li> </ul> </li> </ul>
<a href="#">HA</a>	Hepatitis A	Havrix Vaqta	<ul style="list-style-type: none"> <li>• Post-exposure prophylaxis</li> <li>• Outbreak control</li> <li>• *Pre-exposure prophylaxis for those at increased risk of infection or severe Hepatitis A:                             <ul style="list-style-type: none"> <li>◦ Chronic liver disease</li> <li>◦ Men who have sex with men</li> <li>◦ High risk sexual practices</li> <li>◦ HIV</li> <li>◦ Illicit drug use or alcoholism</li> <li>◦ Individuals receiving repeated replacement of plasma derived clotting factors</li> <li>◦ Children 6 months to 2 years of age who are living in a household with an individual who is at increased risk of infection or severe Hepatitis A</li> </ul> </li> </ul>

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<a href="#">HB</a>	Hepatitis B	Engerix Recombivax	<ul style="list-style-type: none"> <li>• Grade 7 students (school based immunization program)</li> <li>• Youth who have missed or refused HB vaccine as part of the school based program, up to and including 18 years of age</li> <li>• <a href="#">Post-exposure prophylaxis</a></li> <li>• Outbreak control</li> <li>• *Pre-exposure prophylaxis for those at increased risk of Hepatitis B infection or severe Hepatitis B:                             <ul style="list-style-type: none"> <li>◦ Chronic liver disease</li> <li>◦ Men who have sex with men</li> <li>◦ Chronic renal disease</li> <li>◦ Congenital immunodeficiency</li> <li>◦ Hematopoietic stem cell transplant (HSCT)</li> <li>◦ Hemophilia and other bleeding disorders</li> <li>◦ High risk sexual practices</li> <li>◦ HIV</li> <li>◦ Illicit drug use or alcoholism</li> <li>◦ Solid organ transplant</li> <li>◦ Splenic disorders including sickle cell disease or other hemoglobinopathies</li> </ul> </li> </ul>
<a href="#">HAHB</a>	Hepatitis A and B	Twinrix	<ul style="list-style-type: none"> <li>• *Pre-exposure prophylaxis for those at increased risk of Hepatitis A and Hepatitis B infection or severe Hepatitis A and Hepatitis B:                             <ul style="list-style-type: none"> <li>◦ Chronic liver disease</li> <li>◦ Men who have sex with men</li> <li>◦ High risk sexual practices</li> <li>◦ HIV</li> <li>◦ Illicit drug use or alcoholism</li> </ul> </li> </ul>

6 \*Refer to the [publicly funded vaccine eligibility for individuals at high risk of acquiring vaccine preventable disease policy](#) for eligibility criteria.

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<a href="#">HPV</a>	Human papillomavirus	Gardasil	<ul style="list-style-type: none"> <li>• Grade 7 students (school based immunization program)</li> <li>• Females: Youth who have missed or refused HPV vaccine as part of the school based program up to and including 18 years of age</li> <li>• Males: Youth who have missed or refused HPV vaccine as part of the school based program (beginning September 2015) up to and including 18 years of age</li> <li>• Men who have sex with men – for those up to and including 45 years of age</li> <li>• *Pre-exposure prophylaxis for the following high-risk condition: <ul style="list-style-type: none"> <li>◦ HIV – for those up to and including 45 years of age</li> </ul> </li> </ul>
<a href="#">Inf</a>	Influenza - inactivated	Fluzone FluLaval Tetra	<ul style="list-style-type: none"> <li>• Quadrivalent standard dose products: Residents and non-residents of NS, 6 months of age and older</li> </ul>
		Fluzone High-Dose	<ul style="list-style-type: none"> <li>• Trivalent high-dose product: Long-term Care Facility (Nursing Homes and Residential Care Facilities) residents of NS 65 years of age and older.</li> </ul>
<a href="#">IPV</a>	Inactivated polio	Imovax Polio	<ul style="list-style-type: none"> <li>• Immunization of adults who are unimmunized or have incomplete immunization with polio vaccine or combination vaccines such as Tdap-IPV.</li> </ul>
<a href="#">Men-B</a>	Meningococcal B	Bexsero	<ul style="list-style-type: none"> <li>• <a href="#">Post-exposure prophylaxis</a> for Serotype B</li> <li>• Outbreak control</li> <li>• *Pre-exposure prophylaxis for the following high risk conditions: <ul style="list-style-type: none"> <li>◦ Congenital immunodeficiency</li> <li>◦ Hematopoietic stem cell transplant (HSCT)</li> <li>◦ HIV</li> <li>◦ Immunosuppressive therapy using eculizumab (Solaris)</li> <li>◦ Solid organ transplant</li> <li>◦ <a href="#">Splenic disorders</a> including sickle cell disease or other hemoglobinopathies</li> </ul> </li> </ul>
<a href="#">Men-C-C</a>	Meningococcal - Conjugate	NeisVac-C Menjugate	<ul style="list-style-type: none"> <li>• Routine immunization of children &lt; 5 years of age</li> <li>• <a href="#">Post-exposure prophylaxis</a> for Serotype C</li> <li>• Outbreak control</li> </ul>

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<a href="#">Men-C-ACYW-135</a>	Meningococcal - Conjugate	Menveo Menactra Nimenrix	<ul style="list-style-type: none"> <li>• Grade 7 students (school based immunization program)</li> <li>• Youth who have missed or refused meningococcal vaccine as part of the school based program up to and including 18 years of age</li> <li>• <a href="#">Post-exposure prophylaxis</a> for Serotypes A, C, Y, W-135</li> <li>• Outbreak control</li> <li>• *Pre-exposure prophylaxis for the following high risk conditions:                             <ul style="list-style-type: none"> <li>◦ Congenital immunodeficiency</li> <li>◦ <a href="#">Hematopoietic stem cell transplant (HSCT)</a></li> <li>◦ HIV</li> <li>◦ Immunosuppressive therapy using eculizumab (Solaris)</li> <li>◦ Solid organ transplant</li> <li>◦ Splenic disorders including sickle cell disease or other hemoglobinopathies</li> </ul> </li> </ul>
<a href="#">MMR</a>	Measles, mumps, rubella	MMR 11 Priorix	<ul style="list-style-type: none"> <li>• Routine immunization of children if not receiving MMRV.</li> <li>• Immunization of children 6 to 11 months of age travelling to regions where measles is a concern (<a href="https://travel.gc.ca/travelling/health-safety/travel-health-notice">https://travel.gc.ca/travelling/health-safety/travel-health-notice</a>)</li> <li>• Adults born in 1970 or later who have not had measles disease or mumps disease or received two doses of measles or mumps containing vaccine</li> <li>• Post-partum women who are found to be non-immune to rubella</li> <li>• Post-exposure prophylaxis</li> <li>• Outbreak control</li> <li>• *Pre-exposure prophylaxis for the following high risk conditions <b>once immunocompetent:</b> <ul style="list-style-type: none"> <li>◦ <a href="#">Hematopoietic stem cell transplant (HSCT)</a></li> <li>◦ HIV</li> <li>◦ Immunosuppressive therapy</li> <li>◦ Solid organ transplant</li> </ul> </li> </ul>
<a href="#">MMRV</a>	Measles, mumps, rubella & varicella	Priorix Tetra	<p>Routine immunization of children, up to and including 12 years of age, born 2006 and later and not previously immunized with MMR and Varicella are eligible for 2 doses</p> <p>*Pre-exposure for the following high risk conditions up to and including 12 years of age, <b>once immunocompetent:</b></p> <ul style="list-style-type: none"> <li>◦ <a href="#">Hematopoietic stem cell transplant (HSCT)</a></li> <li>◦ HIV</li> <li>◦ Immunosuppressive therapy</li> <li>◦ Solid organ transplant</li> </ul>

8 \*Refer to the [publicly funded vaccine eligibility for individuals at high risk of acquiring vaccine preventable disease policy](#) for eligibility criteria.



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<a href="#">Var</a>	Varicella	Varilrix Varivax	<ul style="list-style-type: none"> <li>• Routine immunization of children not receiving MMRV</li> </ul> <p>Individuals born 1996 -2005 are eligible for one dose (the first dose) of varicella vaccine Individuals born in 2006 and later are eligible for 2 doses of varicella vaccine if not receiving MMRV</p> <p>Post-exposure prophylaxis</p> <p>*Pre-exposure prophylaxis for the following high risk conditions <b>once immunocompetent</b> (if not receiving MMRV):</p> <ul style="list-style-type: none"> <li>◦ <a href="#">Hematopoietic stem cell transplant (HSCT)</a></li> <li>◦ HIV</li> <li>◦ <a href="#">Immunosuppressive therapy</a></li> <li>◦ <a href="#">Solid organ transplant</a></li> </ul> <p>*Pre-exposure prophylaxis for the following high risk conditions (if not receiving MMRV)</p> <ul style="list-style-type: none"> <li>◦ <a href="#">Chronic renal disease</a></li> <li>◦ <a href="#">Chronic salicylate therapy</a></li> <li>◦ <a href="#">Cystic fibrosis</a></li> <li>◦ <a href="#">Splenic disorders</a></li> </ul> <p>Pre-exposure prophylaxis for others (if not receiving MMRV):</p> <ul style="list-style-type: none"> <li>◦ Non-immune health care workers</li> <li>◦ Post-partum women who are found to be non-immune to varicella</li> <li>◦ Non-immune individuals <i>who live with or care for</i> anyone in the following categories:                         <ul style="list-style-type: none"> <li>✓ blood dyscrasias</li> <li>✓ leukemia (except Acute Lymphoblastic Leukemia)</li> <li>✓ lymphoma</li> <li>✓ other malignancies affecting the bone marrow or lymphatic system</li> <li>✓ other defects of cell-mediated immunity</li> </ul> </li> </ul> <p>receiving treatment associated with T-cell abnormalities (e.g. intensive chemotherapy)</p>
<a href="#">Pneu-C-13</a>	Pneumococcal-Conjugate	Pevnar 13	<ul style="list-style-type: none"> <li>• Routine immunization of children</li> <li>• *Pre-exposure prophylaxis for the following high risk conditions:                         <ul style="list-style-type: none"> <li>◦ Cancer</li> <li>◦ Congenital immunodeficiency</li> <li>◦ <a href="#">Hematopoietic stem cell transplant (HSCT)</a></li> <li>◦ HIV</li> <li>◦ <a href="#">Immunosuppressive therapy</a></li> <li>◦ <a href="#">Solid organ transplant</a></li> <li>◦ <a href="#">Splenic disorders</a> including sickle cell disease or other hemoglobinopathies</li> </ul> </li> </ul>

9 \*Refer to the [publicly funded vaccine eligibility for individuals at high risk of acquiring vaccine preventable disease policy](#) for eligibility criteria.

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<a href="#">Pneu-P-23</a>	Pneumococcal - Polysaccharide	Pneumovax 23	<ul style="list-style-type: none"> <li>• Adults 65 years and older</li> <li>• *Pre-exposure prophylaxis for Individuals 2 years and older with the following high risk conditions:                             <ul style="list-style-type: none"> <li>◦ Cancer</li> <li>◦ Chronic cerebral spinal fluid (CSF) leak</li> <li>◦ Chronic liver disease</li> <li>◦ Chronic lung disease (not asthma)</li> <li>◦ Chronic neurological conditions that may impair clearance of oral secretions</li> <li>◦ Chronic renal disease</li> <li>◦ Cochlear implant</li> <li>◦ Congenital immunodeficiency</li> <li>◦ Cystic fibrosis</li> <li>◦ Diabetes</li> <li>◦ Heart disease</li> <li>◦ Hematopoietic stem cell transplant (HSCT)</li> <li>◦ HIV</li> <li>◦ Homelessness</li> <li>◦ Illicit drug use or alcoholism</li> <li>◦ Immunosuppressive therapy</li> <li>◦ Residing in long term care facilities</li> <li>◦ Solid organ transplant</li> <li>◦ <a href="#">Splenic disorders</a> including sickle cell disease or other hemoglobinopathies</li> </ul> </li> </ul>
<a href="#">Rab</a>	Rabies	Imovax Rabies Rabavert	<ul style="list-style-type: none"> <li>• Post-exposure prophylaxis</li> </ul>

## Other Biological Products [\(Canadian Immunization Guide\)](#)

Abbreviation	National Agent (Type)	Trade Name E.g.	Eligibility
BAtx	Botulism antitoxin		<ul style="list-style-type: none"> <li>• People with established or suspected botulism (therapeutic)</li> <li>• Asymptomatic people strongly suspected of having eaten food contaminated with botulism toxin (prophylaxis)</li> </ul>
DAtx	Diphtheria antitoxin		<ul style="list-style-type: none"> <li>• Clinical suspicion of diphtheria regardless of bacteriological confirmation</li> </ul>
Ig	Immunoglobulin	GamaSTAN	<p><b>Hepatitis A</b></p> <ul style="list-style-type: none"> <li>• Post exposure prophylaxis for the following: <ul style="list-style-type: none"> <li>◦ Infants &lt; 6 months of age</li> <li>◦ Immunocompromised people who may not respond to the vaccine</li> <li>◦ Immunocompetent individuals <math>\geq</math> 60 years of age</li> <li>◦ Individuals with chronic liver disease</li> <li>◦ People for whom Hepatitis A vaccine is contraindicated</li> </ul> </li> </ul> <p><b>Measles (<i>Rubeola</i>)</b></p> <ul style="list-style-type: none"> <li>• Post exposure prophylaxis for the following susceptible contacts of measles: <ul style="list-style-type: none"> <li>◦ Infants &lt; 6 months of age</li> <li>◦ Immunologically compromised individuals for whom measles vaccine is contraindicated</li> <li>◦ Susceptible immunocompetent people who present more than 72 hours but less than 1 week after exposure, i.e., too late for vaccine</li> </ul> </li> </ul>
HBIG	Hepatitis B immunoglobulin	HepaGamB HyperHEPB	<ul style="list-style-type: none"> <li>• Post exposure prophylaxis for the following high risk situations: <ul style="list-style-type: none"> <li>◦ Acute percutaneous or mucosal exposure to blood containing Hepatitis B virus</li> <li>◦ Perinatal exposure of infants born to mothers with acute or chronic Hepatitis B virus</li> <li>◦ Sexual contacts of individuals with acute or chronic Hepatitis B</li> </ul> </li> </ul>
RabIg	Rabies immunoglobulin	HyperRAB	<ul style="list-style-type: none"> <li>• Post exposure prophylaxis</li> </ul>
TIG	Tetanus immunoglobulin	HyperTET	<ul style="list-style-type: none"> <li>• <a href="#">Post exposure/wound management</a></li> </ul>

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Varlg	Varicella immunoglobulin	VariZIG	<ul style="list-style-type: none"> <li>• Post exposure prophylaxis for the following high-risk conditions:                             <ul style="list-style-type: none"> <li>◦ Pregnant women</li> <li>◦ Immunocompromised patients, such as those with congenital or acquired immunodeficiency</li> <li>◦ Newborn infants of mothers who have varicella that began during the 5 days before to 48 hours after delivery</li> <li>◦ For the management of significant varicella exposure in a neonatal or pediatric intensive care setting, consultation with the infectious diseases/infection control specialist regarding the potential use of VariZIG™ is advised</li> </ul> </li> </ul>

