

Vaccine Clinic Resource for Immunizers

Disclaimer: this Quick Reference is not intended to replace other product specific vaccine references. The document is intended as a quick reference for frequently referred to information. Please refer to the product monograph and vaccine specific resources for all current and complete information.

Title:	Mpox Vaccine (IMVAMUNE [®]) Quick Reference Guide		
	IMVAMUNE [®] vaccine is indicated for active immunization against smallpox, mpox and related orthopoxvirus infection for those 18 years of age older at high risk of infection		
Effective Date:	September 11, 2024		
Approver:	Final		

Mpox Vaccine Resources:

Fact sheet

Mpox vaccine factsheet

Product Monograph https://pdf.hres.ca/dpd_pm/00071931.PDF

Eligibility Criteria

For the most up to date information on eligibility criteria refer to www.gov.mb.ca/health/publichealth/diseases/mpox.html

NACI – Interim Guidance on the use of Imvamune®

https://www.canada.ca/content/dam/phac-aspc/documents/services/immunization/national-advisory-committee-on-immunization-naci/guidance-imvamune-monkeypox/guidance-imvamune-monkeypox-en.pdf

Canadian Immunization Guide:

For additional guidance on contraindications, precautions and special populations refer to the vaccine specific section: https://www.canada.ca/en/public-health/services/publications/healthy-living/canadian-immunization-guide-part-4-active-vaccines/page-21-smallpox-vaccine.html



Mpox Vaccine

Product	Storage and Handling	Mpox Eligibility Criteria	Recommendations for Use
IMVAMUNE® Modified Vaccinia Ankara-Bavarian Nordic® (live-attenuated, non-replicating) * Format: Single or 10/per box 0.5 ml vials Product is latex, preservative and adjuvant free. Potential Allergens: Benzonase, gentamicin and ciprofloxacin Other Ingredients: Tris-hydroxymethyl- amino methane, sodium chloride, hydrochloric acid, trometamol *Imvamune®, 3rd generation orthopoxvirus vaccine, is labelled as JYNNEOS® and Imvanex® in other jurisdictions. These vaccines are also considered valid when assessing previous vaccination status.	 Storage: Store frozen at -20°C ± 5°C or -50°C ± 10°C or -80°C ± 10°C. Expiry date depends on storage temperature Thaw at room temperature. After thawing, the vaccine can be stored at 2°C – 8°C for up to 2 months prior to use. Do not refreeze a vial once it has been thawed. Protect from light. Handling: After thawing, the drug product should appear as a pale milky colored homogeneous suspension. The liquid vaccine should be visually inspected for any foreign particulate matter prior to administration. 	 Pre-exposure prophylaxis (PrEP): cisgender, transgender, or two-spirit people who self- identify as belonging to the gay, bisexual and other men who have sex with men (gbMSM) community and who meet at least one of the following criteria: have received a diagnosis of a sexually transmitted infection in the past year; have had two or more sexual partners in the past 90 days; have attended locations for sexual contact (e.g. bath houses or sex clubs) or are planning to; have had anonymous sex in the past 90 days (i.e. using apps, online sites, formal/informal gatherings) or are planning to; engaged in sex work or plan to, as a worker or client; or any sexual contacts of the individuals described above. individuals who self-identify as sex workers, regardless of self-identified sex/gender staff or volunteers in sex-on-premises venues where workers may have contact with objects or materials that may be contaminated with the mpox virus without the use of personal protective equipment individuals who engage in sex tourism regardless of gender, sex assigned at birth, or sexual orientation individuals who anticipate experiencing any of the above scenarios For the most current eligibility criteria refer to: www.gov.mb.ca/health/publichealth/diseases/mpox.html 	 Pre-exposure prophylaxis (PrEP): 2 doses Interval: 28 days (4 weeks) apart* Dosage: 0.5 ml Route: SC (5/8" 25g needle) *28 days is the minimum/preferred interval. 2nd dose can be given later than 28 days to complete the series > Imvamune can be administered concurrently (i.e., same day) or at any time before or after live or non-live vaccines. > Imvamune® may be offered to the following populations if vaccination is recommended based on high-risk criteria: Individuals who are immunocompromised due to disease or treatment Individuals who are lactating/breastfeeding Children and youth (< 18 years of age)



Mpox Vaccine

Product	Storage and Handling	Mpox Eligibility Criteria	Recommendations for Use
IMVAMUNE [®] Modified Vaccinia Ankara-Bavarian Nordic [®]	Storage: Store frozen at -20°C ± 5°C or -50°C ± 10°C	Post-exposure prophylaxis (PEP):	Post-exposure prophylaxis (PEP): 1 or 2 doses
(live-attenuated, non-replicating) *	or -80°C ± 10°C. Expiry date depends on storage temperature	High risk close contacts of a confirmed or probable mpox case. *	 Interval: 1st dose preferably to be given 0-4 days after last exposure or up to 14 days after last exposure*
Format: Single or 10/per box 0.5 ml vials	Thaw at room temperature. After thawing, the vaccine can be stored at 2°C – 8°C for up to 2 months prior to use.	• Previously unimmunized with Imvamune [®] Since we do not know the effectiveness of the previous generations of smallpox vaccination against current mpox infection, all individuals who received previous	 2nd dose given 28 days (4 weeks) apart** Dosage: 0.5 ml
Product is latex, preservative and adjuvant free. Potential Allergens: Benzonase, gentamicin and ciprofloxacin Other Ingredients: Tris-hydroxymethyl- amino methane, sodium chloride,	Do not refreeze a vial once it has been thawed. Protect from light. Handling: After thawing, the drug product should appear as a pale milky colored	generations of smallpox vaccine, as well as unimmunized individuals, should receive 1 dose of Imvamune [®] for PEP if they have been exposed to a probable or confirmed case of mpox. If the person also meets eligibility criteria for PrEP, a second dose of Imvamune [®] should be offered for administration 28 days later.	Route: SC (5/8" 25g needle) * Vaccine should be given within 4 days from the date of exposure to prevent onset of the disease. However, vaccine can be given up to 14 days after the date of exposure, and may reduce the symptoms of disease, but may not prevent the disease.
aydrochloric acid, trometamol "Imvamune®, 3rd generation orthopoxvirus vaccine, is labelled as JYNNEOS® and mvanex® in other jurisdictions. These vaccines are also considered valid when assessing previous vaccination status.	appear as a pale milky colored homogeneous suspension. The liquid vaccine should be visually inspected for any foreign particulate matter prior to	 1 previous dose of Imvamune® For contacts that previously received one dose of Imvamune® prior to exposure to a confirmed or probable case, 1 dose of Imvamune® should be administered for PEP if it has been at least 28 days since the pre-exposure dose was received. 2 previous doses of Imvamune® Consider contact as fully immunized and no PEP required. *For further information on contacts and PEP refer to the Mpox (Orthopoxvirus) Protocol 	 **28 days is the minimum/preferred interval. 2nd dose can be given later than 28 days to complete the series Imvamune[®] can be administered concurrently (i.e., same day) or at any time before or after live or non-live vaccines. Imvamune[®] may be offered to the following populations if vaccination is recommended based on high-risk criteria: Individuals who are immunocompromised due to disease or treatment Individuals who are pregnant Individuals who are lactating/breastfeeding Children and youth (< 18 years of age)