

Human Rabies Immune Globulin (RabIg)

KamRAB™

Supplier: Valneva Canada Inc.

INDICATIONS: ^A

- Human Rabies Immune Globulin (RabIg) is used for Rabies Post-Exposure Prophylaxis (RPEP) as determined by the Medical Health Officer. Consultation is available to Medical Health Officers with *Communicable Diseases and Immunization Service (CDIS), BCCDC*.
- RabIg should be given in conjunction with the 1st dose of rabies vaccine. Rabies vaccine and RabIg must be administered with separate needles and syringes at **separate anatomical sites**.
- If a rabies vaccine series has been started or completed elsewhere, and it was not given in accordance with the current WHO standards, administer RabIg (on day 0) in conjunction with the 1st dose of another full series of rabies vaccine. See [Communicable Disease Control Manual, Chapter 1: Rabies, section 4.2.1](#).

DOSES AND SCHEDULE: ^B

- The recommended dosage for children and adults is the same: 20 IU/kg of body weight.
- The dose of KamRAB™ is calculated as:

$$\frac{[20 \text{ IU/kg} \times \text{weight in kg}]}{150 \text{ IU/mL}} = \text{mL}$$

Do not exceed the recommended dose due to interference with active antibody production.

ADMINISTRATION:

- KamRAB™ is supplied in 2 mL vials at a concentration of 150 IU/mL. **Infiltrate as much RabIg as possible deep into and around the wound(s) in order to neutralize the virus.**
 - When more than one wound site exists, each site should be infiltrated with a portion of the RabIg, using a separate syringe and needle for each infiltration.
 - Infiltration of wounds in some anatomical sites (finger tips) must be carried out with care in order to avoid increased pressure in the tissue compartment (compartment syndrome).
 - Any remaining volume should be injected intramuscularly at a site distant from vaccine administration. The deltoid should not be used for RabIg administration. Both deltoid sites should be reserved for administration of rabies vaccine.
- When there is no wound site, the site of administration of RabIg is age specific. See [Immune Globulin Preparations \(HBIG, Ig, TIg, VarIG, RabIG\)](#) for administration information and maximum volume to be administered per site according to age.
- RabIg contains no preservatives. Vials are single dose use. Once entered, discard any unused contents.

BOOSTER DOSES:

None.

^A Since vaccine induced antibodies begin to appear within 1 week, there is no value in administering RabIg more than 7 days after initiation of vaccine.

^B When notification of an exposure is delayed, RPEP may be started as late as 6 or more months after an exposure.

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SEROLOGICAL TESTING:

Serological testing is not recommended before or after immunization.

CONTRAINdications:

None.

PRODUCT COMPONENTS:

Potential allergens: none.

Other components: glycine, sodium hydroxide.

PRECAUTIONS:

- Give RabIg with caution (i.e., in an emergency room setting) if the client has a history of anaphylactic reaction following receipt of any immune globulin product or a history of anaphylactic reaction to any component of KamRAB™ (assess risks versus benefits).
 - Human Ig products are among the safest blood-derived products available. The method of preparation includes one or more steps that exclude or inactivate hepatitis B, C and HIV; therefore the risk of transmission is extremely low. However, it is possible, that unknown infectious agents may be present in such products. The benefits of use of rabies immunoglobulin after exposure to rabies far outweigh the theoretical risk of receipt of a blood product.
 - Persons with IgA deficiency have the potential for developing antibodies to IgA and could have an anaphylactic reaction to subsequent blood products that contain IgA. Therefore, RabIg should only be given to such persons if the expected benefits outweigh the risks, and should be administered in an emergency room setting.
 - Special measures should be considered when administering IM injections to people with bleeding disorders. A smaller gauge needle (23 gauge or smaller) should be used and steady, firm pressure should be applied to the injection site for 5 minutes. If there is concern that the injection may stimulate bleeding, the client should connect with their medical specialist.
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SPECIAL CONSIDERATIONS:

- Document receipt of RabIg in the client's electronic record (e.g., Panorama, PARIS) and/or chart. The following information must be recorded: trade name of product, date, lot number, dosage, route, and site(s).
- Provide a written record to individuals who receive any immune globulin product.
- Regarding RabIg and the administration of live vaccines see [Immune Globulin Preparations or Blood: Timing Intervals for Vaccines Containing Live Measles, Mumps, Rubella or Varicella Virus.](#)

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ADVERSE EVENTS:

Local: pain, bruising.

Systemic: headache, muscle and joint pain, fatigue, abdominal pain, nausea, dizziness, feeling faint.

A potential increased risk of thrombosis (blood clots) has been observed within 24 hours of receipt of immune globulin products, especially when given in large doses (i.e., more than 10 mL). Additional risk factors include: age 45 years and older, history of thrombosis or those with risk factors for thrombosis (e.g., obesity, high blood pressure, diabetes, prolonged periods of immobilization, use of estrogens, a history of heart disease, blood clotting disorders, indwelling central vascular catheters, or diseases that thicken the blood).^{A, B}

^A Daniel GW, Menis M, Sridhar G, et al. Immune globulins and thrombotic events as recorded in a large administrative database in 2008 through 2010. *Transfusion*. 2012; 52:2113-2121.

^B Menis M, Sridhar G, Selvam N, et al. Hyperimmune globulins and same-day thrombotic adverse events as recorded in a large healthcare database during 2008-2011. *Am. J. Hematol.* 2013; 88:1035-1040.