Rabies Post-Exposure Prophylaxis (rPEP) Quick Reference Guide

Rabies Immune Globulin (Rablg) - HyperRAB® / IMOGAM®

- All Rablg is ideally administered on Day 0.
- Dose of 20 IU/kg by body weight for all age groups.

For 2 mL vials: Total dose in mL = 20 IU/kg x (patient weight in kg) 150 IU/mL

For 1 mL vials: Total dose in mL = 20 IU/kg x (patient weight in kg) 300 IU/mL

- Infiltrate each wound/exposure site with portion of Rablg using a separate needle and syringe.
- Do not administer more than the maximum dosage of Rablg per body weight.
- DO NOT administer Rablg in the same anatomical location as rabies vaccine.
- Dilute Rablg (if required) per product monograph.
- Give Rablg to individuals who have not previously received appropriate rabies immunization.*
- Draw up 1 vial at a time.
- If entire calculated Rablg dose cannot anatomically be infiltrated around wound(s)/exposure site(s), inject any remaining Rablg volume IM at site(s) distant from the site of vaccine administration.

TOTAL BODY \	# VIALS	
≤ 33 lbs	≤ 15 Kg	1
>33–66lbs	>15–30 Kg	2
>66–99lbs	>30–45 Kg	3
>99–132lbs	>45–60 Kg	4
>132–165lbs	>60–75 Kg	5
>165–198lbs	>75–90 Kg	6
>198–231lbs	>90–105 Kg	7
>231–264lbs	>105–120 Kg	8
>264–297lbs	>120–135 Kg	9
>297–330lbs	>135–150 Kg	10







Rabies Vaccine - IMOVAX® (HDCV) / RabAvert® (PCECV)

- Administer rabies vaccine dose (1.0 mL) intramuscularly (IM) in the deltoid (children & adults)
 or anterolateral thigh (infants & small children).
- Administer at different anatomical site than Rablg using a separate needle and syringe.
- NEVER administer rabies vaccine in the gluteal region.
- Ensure ALL doses are administered at appropriate intervals. DO NOT administer doses early.
- If a dose is delayed, it should be given ASAP and the schedule resumed with the appropriate intervals from the most recent dose.

Patient Profile	Day 0	Day 3	Day 7	Day 14	Day 28
Patient previously immunized with rabies vaccine (total 2 doses)*					
Immunocompetent patient (Total 4 doses)					
Immunocompromised patient (Total 5 doses)**					

* Completed course of pre- or post-exposure prophylaxis with either HDCV or PCECV -OR- Completed immunization with other types of rabies vaccine (or with HDCV or PCECV according to unapproved schedules) with an acceptable antibody response to be a titre of at least 0.5 IU/mL by rapid fluorescent-focus inhibition test.

** Includes those taking corticosteroids or other immunosuppressive agents, those who have

immunosuppressive illnesses, and those taking chloroquine & other anti-malarials

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COMMON ADVERSE REACTIONS*

*Serious or severe adverse effects are rare and are not listed here. Please consult the product monograph for details.

Rabies Immune Globulin (Rablg) - HyperRAB® / IMOGAM®

Local: pain, erythema, indurationSystemic: headache, low-grade fever

Rabies Vaccine - IMOVAX® (HDCV)

- Local: pain, erythema, swelling, pruritus, induration (60% to 90%)
- Systemic: headache, nausea, abdominal pain, muscle aches, dizziness (6% to 55%)

Rabies Vaccine - RabAvert® (PCECV)

- Local: pain, tenderness, swelling, erythema, induration (10% to 60%)
- Systemic: malaise, myalgia, arthralgia, headache, fever (1% to 10%)

CONTRAINDICATIONS & PRECAUTIONS

- There are no contraindications to the use of rabies vaccine or Rablg after significant exposure to a proven rabid animal. Post-exposure vaccination should never be postponed.
- Pregnancy is not a contraindication to post-exposure prophylaxis with rabies vaccine or Rablg.
- Consultation should be sought regarding administration for patients with a proven history of hypersensitivity to product components or packaging.

Product	Potential Allergens	
IMOGAM®	latex in vial stopper	
IMOVAX®	neomycin, phenol red	
RabAvert®	amphotericin B, chick protein, chlortetracycline, neomycin, polygeline (gelatin)	

DRUG INTERACTIONS

Product	Interactions	Effect
Rabies Vaccine	radiation therapy, chloroquine, corticosteroids, immunosuppressive agents	diminished efficacy of rabies vaccine
Rablg	MMR, MMRV, monovalent varicella vaccines	diminished response to live vaccines; delay vaccination with these vaccines for 4 months after Rablg

VACCINE INTERCHANGEABILITY

Wherever possible, an immunization series should be completed with the same vaccine product. However, if it is not feasible to do so, PCECV and HDCV are considered interchangeable.

REFERENCES:

<u>Management of Potential Rabies Exposures Guideline, 2019</u> – Ministry of Health <u>Canadian Immunization Guide, Part 4</u> – Government of Canada

York Region Public Health

If you have questions regarding a potential animal exposure, rabies risk assessment, or would like to request PEP, call Health Connection at 1-800-361-5653 (weekdays between 8:30 and 4:30pm) or 1-888-335-0111 (after hours, weekends or holidays).

