

**GRIFOLS** 

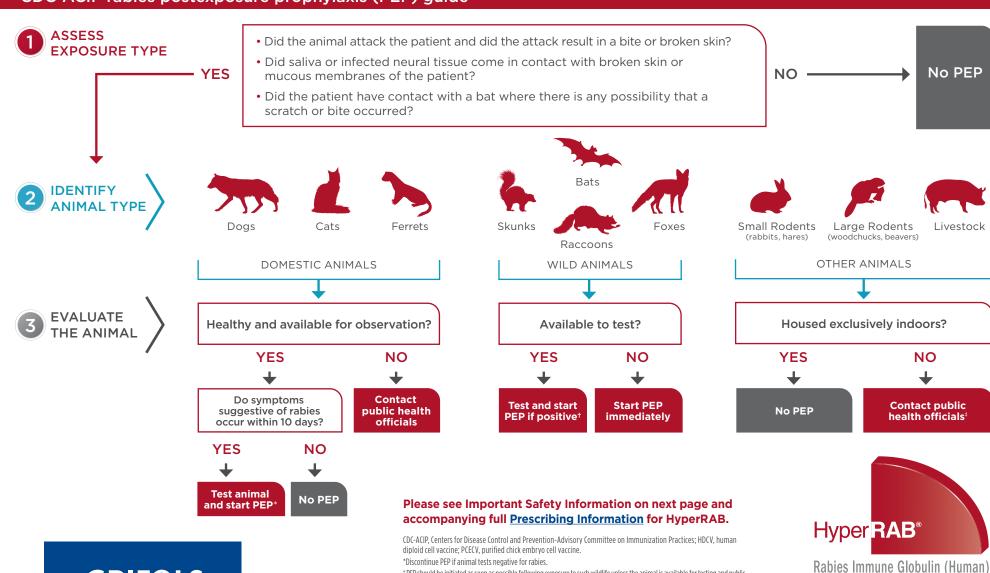
Efficiency and flexibility matter in your choice of human rabies immune globulin (HRIG)

# **BE RABIES READY**with HyperRAB®

(rabies immune globulin [human]) 300 IU/mL

A comprehensive HRIG solution designed with your patients, healthcare team, and institution in mind.

## CDC-ACIP rabies postexposure prophylaxis (PEP) guide<sup>1</sup>



<sup>1</sup>Typically, exposure to these animals does not require PEP.

† PEP should be initiated as soon as possible following exposure to such wildlife unless the animal is available for testing and public health authorities are facilitating expeditious laboratory testing or it is already known that the brain material from the animal has

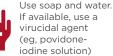
tested negative. The animal should be euthanized and tested as soon as possible. Holding for observation is not recommended.

300 IU/mL

#1 Prescribed HRIG in the US2

# Follow the CDC guidelines<sup>3</sup>

# **CLEANSE ALL WOUNDS**





\*Inject full dose around and into the wound (if possible). For remaining volume (or if there is no wound), inject HRIG in the deltoid area opposite of the vaccine injection,<sup>3</sup>

†Inject full dose in the deltoid area opposite of the HRIG injection or wound.<sup>3</sup>

#### PREVIOUSLY VACCINATED

#### **VACCINE (NO HRIG)**

Vaccinate with HDCV or PCECV

NOT PREVIOUSLY VACCINATED

Administer HRIG around

Vaccinate with HDCV or

PCECV using a new syringe<sup>†</sup>

and into the wound'

**HRIG + VACCINE** 





# Dosing recommendations for previously unvaccinated patients<sup>4</sup>

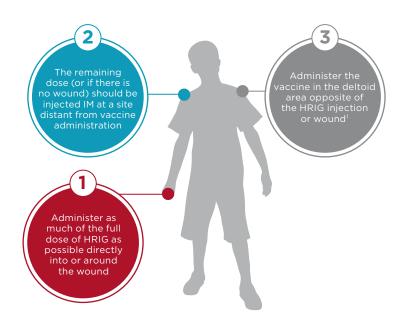
Dosing for HyperRAB® (rabies immune globulin [human]) is based on weight

• The recommended dose is 20 IU/kg (0.0665 mL/kg) of actual body weight

## When a patient is exposed to rabies<sup>3,5,6</sup>:

DAY 0	DAY 3	DAY 7	DAY 14
Administer HRIG and the FIRST rabies vaccine dose (1 mL IM). Antibodies start working immediately at the site of infection	Administer rabies vaccine (1 mL IM)	Administer rabies vaccine (1 mL IM), antibody production begins	Administer rabies vaccine (1 mL IM)

## Administer HRIG and the vaccine correctly<sup>3</sup>



<sup>‡</sup>For children, the anterolateral aspect of the thigh is also acceptable.

HRIG should never be administered in the same syringe or needle or in the same anatomical site as the first vaccine dose.<sup>3</sup> Do not inject rabjes vaccine or HRIG in the gluteal area due to risk of diminished immunologic response and injury to the sciatic nerve (unless the exposure site is in the gluteal region).7

#### **HRIG Dosing Recommendation** lb 44 110 132 154 176 198 220 242 264 22 66 88 287 309 Patient Weight kg 10 20 30 40 50 60 70 80 90 100 110 120 130 140 200 400 600 800 1000 1200 1400 1600 1800 2000 2200 2400 2600 2800 Recommended Dosing (IU) 1x5-ml HyperRAB (300 IU/mL) 1x3-mL 1x3-mL 2x3-mL 1x5-mL 1x5-mL 1x1-mL 2x1-mL 2x1-mL 1x3-mL 1x5-mL 2x3-mL 2x3-mL 1x3-mL 2x5-mL 1x1-mL 1x1-mL 1x1-mL 1x3-mL 1x3-mL Vials Required 1x1-mL

#### IMPORTANT SAFETY INFORMATION

#### **Indication and Usage**

HYPERRAB® (rabies immune globulin [human]) is indicated for postexposure prophylaxis, along with rabies vaccine, for all persons suspected of exposure to rabies.

#### **Limitations of Use**

Persons who have been previously immunized with rabies vaccine and have a confirmed adequate rabies antibody titer should receive only vaccine.

For unvaccinated persons, the combination of HYPERRAB and vaccine is recommended for both bite and nonbite exposures regardless of the time interval between exposure and initiation of postexposure prophylaxis.

Beyond 7 days (after the first vaccine dose), HYPERRAB is not indicated since an antibody response to vaccine is presumed to have occurred.

#### **Important Safety Information**

#### For infiltration and intramuscular use only.

Severe hypersensitivity reactions may occur with HYPERRAB. Patients with a history of prior

systemic allergic reactions to human immunoglobulin preparations are at a greater risk of developing severe hypersensitivity and anaphylactic reactions. Have epinephrine available for treatment of acute allergic symptoms, should they occur.

HYPERRAB is made from human blood and may carry a risk of transmitting infectious agents, eg, viruses, the variant Creutzfeldt-Jakob disease (vCJD) agent, and, theoretically, the Creutzfeldt-Jakob disease (CJD) agent.

The most common adverse reactions in >5% of subjects during clinical trials were injectionsite pain, headache, injection-site nodule, abdominal pain, diarrhea, flatulence, nasal congestion, and oropharyngeal pain.

Do not administer repeated doses of HYPERRAB once vaccine treatment has been initiated as this could prevent the full expression of active immunity expected from the rabies vaccine.

Other antibodies in the HYPERRAB preparation may interfere with the response to live vaccines such as measles, mumps, polio, or rubella. Defer immunization with live vaccines for 4 months after HYPERRAB administration.

### Please see accompanying full Prescribing Information for HYPERRAB.

References: 1. Manning SE, Rupprecht CE, Fishbein D, et al; Advisory Committee on Immunization Practices. Human rabies prevention—United States. 2008. MMWR Recomm Rep. 2008:57(RR-3):1-28. 2. Data on file, Grifols. 3. Rupprecht CE. Briags D. Brown CM. et al: Advisory Committee on Immunization Practices. Use of a reduced (4-dose) vaccine schedule for postexposure prophylaxis to prevent human rabies. MMWR Recomm Rep. 2010:59(RR-2):1-9. 4. HyperRAB\* (rabies immune globulin [human]) Prescribing Information. Grifols. 5. Baxter D. Active and passive immunity, vaccine types, excipients and licensing. Occup Med (Lond). 2007;57(8):552-556. 6. Siegrist CA. Vaccine immunology. In: Plotkin SA, Orenstein WA, Offit PA, eds. Vaccines. 6th ed. Elsevier-Saunders; 2013:17-36. 7. Atkinson WL, Pickering LK, Schwartz B, Weniger BG, Iskander JK, Watson JC; Centers for Disease Control and Prevention. General recommendations on immunization. Recommendations of the Advisory Committee on Immunization Practices (ACIP) and the American Academy of Family Physicians (AAFP). MMWR Recomm Rep. 2002;51(RR-2):1-35.

