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Rabies Vaccine and Immunoglobulin Shortages and Change to Rabies Immunoglobulin (Rablg) Vial Format and Formulation

- Attention: Physicians, Emergency Departments, Nurse Practitioners, Infection Control Practitioners, Occupational Health Professionals, Walk-In Clinics/Urgent Care Clinics, Family Health Teams, Pharmacies, Central LHIN, NSM LHIN
- **Date:** August 2, 2019

We have received notice from the Ministry of Health that there is a provincial supply shortage of publicly funded rabies vaccine and rabies immunoglobulin for post- exposure prophylaxis (PEP) in the management of potential human exposures to the rabies virus. Based on information provided, it is anticipated that the current provincial supply will not be replenished before November 2019 (Memo attached).

At this time, following the recent media reporting of a human rabies death in British Columbia, we have also experienced a substantial increase in reported animal bite exposures. With these challenges, we are requesting a thorough risk assessment be conducted for all reported exposures, and a consultation with public health for any instance where rabies vaccination may be indicated.

When determining the need for post exposure rabies immunization, the current <u>Ministry of Health and Long-Term</u> <u>Care (MOHLTC) Management of Potential Rabies Exposures Guideline, 2019</u> is recommended as a resource. For **rabies vaccine and immunoglobulin**, please contact the Rabies Coordinator at ext. **8894** or after hours at **1-888-225-7851**.

What should health care practitioners do when a patient presents with a bat, wildlife or domestic animal exposure?

1. Please complete the **Rabies and Animal Exposure Incident Report Form** (below) to assist with the collection of required reporting information. All reported incidents will be investigated by a Public Health Inspector (PHI). All animal exposure incidents may be reported by faxing a copy of the completed *Rabies and Animal Exposure Incident Report Form* (page 3 of this document) through our **Designated Rabies Fax line** at 705-725-8132.

Note: Facial, neck or head bites from mammals are higher risk for transmission (with potential for a shorter incubation period) and direct contact with bats is also high risk. Such incidents should also be verbally reported at **705-721-7520** or **1-877-721-7520** ext. **8811** (Monday – Friday 8:30am – 4:30pm) or after hours at **1-888-225-7851** (evenings, weekends or holidays).

2. In instances where there is a bat exposure, please ask the patient if they have the bat available for testing. If so, please provide information about the status and location of the bat in question when reporting to the health unit. If

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possible, bats should be safely captured (or retrieved if dead) and tested for rabies when there has been a human exposure. Such testing can help to avoid or reduce rabies vaccination for exposed patients. Extreme care should always be taken to ensure that there is no further exposure when captured or handled. In the event that a bat is captured, it should be submitted for rabies testing; SMDHU staff are available to facilitate such testing.

Unless exposure from a bat is to the head or neck region, rabies vaccine and immunoglobulin administration can be delayed for up to 48 hours until the rabies test result on the bat is obtained. If vaccination is initiated, it can be discontinued if the bat test is later found to be negative for rabies. Results are usually available within 48 hours, however, result times can occasionally take longer because of collection schedules, courier limitations and delays on weekends.

Change to Rabies Immunoglobulin (Rablg) Vial Format and Formulation

The Ontario Government Pharmaceutical and Medical Supply Service (OGPMSS) has provided notice to all Public Health Units that a new Rabies Immunoglobulin (RabIg) product will be distributed in the near future. The new product will be a new **1mL vial format of HyperRAB**[®] (manufactured by Grifols Therapeutics LLC).

It is important to note that the formulation for the 1mL vial format is different than the 2mL vial format that may be currently in your local stock. The new formulation will be distributed once stocks are depleted:

2 mL vials contain 150 IU/mL formulation 1 mL vials contain 300 IU/mL formulation ** new formulation

Both vial formats contain 300 IU, but the 1 mL format requires less volume to achieve the same dose.

Please use the following formula to determine the dose required: (See attached Dosage Schedule)

For 150 IU/mL Rablg in 2 mL vials:

20 IU/kg x (patient wt in kg) ÷ 150 IU/mL = dose in mL

OR

9.09 IU/lb x (patient wt in lb) ÷ 150 IU/mL = dose in mL

For 300 IU/mL Rablg in 1 mL vials:

20 IU/kg x (patient wt in kg) ÷ 300 IU/mL = dose in mL

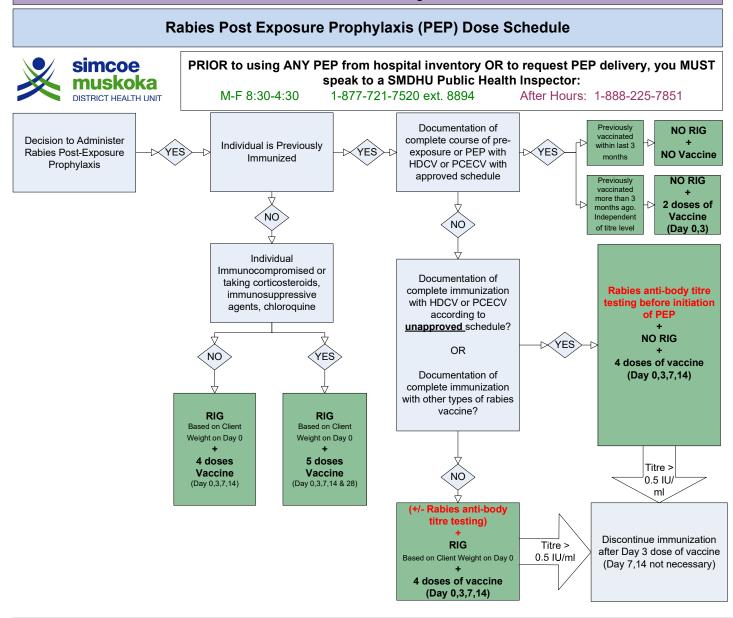
OR

9.09 IU/lb x (patient wt in lb) ÷ 300 IU/mL = dose in mL

If you have any questions, please contact please contact Rabies Coordinator at **705-721-7520** or **1-877-721-7520** ext. **8894** (Monday – Friday 8:30am – 4:30pm) or after hours at **1-888-225-7851** (evenings, weekends or holidays).

STEP 3a. Determine Appropriate Dose/Schedule

If PEP is warranted immediately, and/or physician has decided to proceed with rabies PEP, choose appropriate does based on the following schedule



If PEP is warranted, administration should be initiated within 24 hours of physician decision to administer PEP. Prophylaxis for bites to head and neck region should begin immediately and not be delayed. PEP could be delayed not more than 48 hours if the animal involved is available for testing (does not apply to head/neck bites) or in the case of a captured dog, cat or ferret, may be delayed 10 days while the animal is being observed by public health.

RIG Volume Based on Weight of Client	Weight of Client		IG vials or portion)	Vaccine		
Dose Calculation: 2 different products available:	≤ 15 kg (≤ 33lbs)		1	Vacenie		
	> 15-30 kg (33-66 lbs)		2	Administer 1 ml (1 vial) on each day of		
1ml Vial 300 IU/ml:	> 30-45 kg (66-99 lbs) 3			dosing schedule		
20 IU/kg x (client wt in kg) ÷ 300 IU/ml = dose in ml OR	> 45-60 kg (99-132 lbs)		4	Administer IM into deltoid muscle in older		
2ml Vial 150 lU/ml:	> 60-75 kg (132-165 lbs)		5	children and adults and into vastus lateralis (anterolaterol thigh) in infants		
20 IU/kg x (client wt in kg) ÷ 150 IU/ml = dose in ml	> 75-90 kg (165-19	8 lbs)	6	NEVER administer into gluteal region		
Administered ALL RIG on first day of initiation of therapy (Day 0) Infiltrate into wound and surrounding area (If anatomically feasible) Remaining volume IM at site distant from site of vaccine administration. DO NOT MIX RIG and Vaccine	> 90-105 kg (198-231 lbs) 7		7			
	> 105-120 kg (231-264 lbs)		8	Administer at different anatomical site from RIG.		
	> 120-135 kg (264-2	97 lbs)	9	110.		

 PEP=Post Exposure Prophylaxis (Rabies) may include Rabies Immune Globulin (RIG) and Rabies Vaccine.

 Preparations available for use in Canada: RIG: IMOGAM ® and HYPERRAB ® S/D and Vaccine: IMOVAX ® (HDCV) and RabAvert ® (PCECV)

 HDCV= human diploid cell vaccine (Imovax ®)
 PCECV= purified chick embryo cell culture vaccine (RabAvert ®)

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IMMEDIATELY FAX all animal exposure incidents to the Simcoe Muskoka District Health Unit:										
FAX: 705-725-8132										
For suspicious animal exposures, and head, face, neck exposures: Phone: (705) or (877) 721-7520 ext. 8811 For the physician inquiries and release of rabies post-exposure prophylaxis: Phone: (705) or (877) 721-7520 ext. 8894										
	E									
Date Reported to SM	DHU: <u>YYYY/N</u>	MMM/DD Repor	ting Location (Name of Hospit	al/Office/Police):					
Contact Person and	phone number at	Reporting Locatio	n for addition	al file inform	ation:		Ext:			
(B) PATIENT/VICTIN	M INFORMATIO	N								
					Logal Sov	Male 🗌 F	emale 🗌 X			
Name:			Legal Sex	As found on health card						
Parent Guardian Nam	ne (if patient is und	ler 16yrs of age):								
Date of Birth: <u>YYYY</u>	//MMM/DD		Phone:		(O [.]	ther)				
Permanent Address:	911#	Street Name	Apt	/Unit#	City					
Temporary Address:					Dates Effectiv	ve:				
, ,	911#	Street Name	C	ity	, , , , , , , , , , , , , , , , , , ,					
	ILS									
Date of Incident:	YYYY/MMM/DD)								
Details of Incident:										
Body area affected:		Bite 🗌	Scratch 🗌	Saliva 🗌	Handling 🗌	Other 🗌 :				
Skin broken: Yes	No									
Family Physician:				Phone or o	other contact in	10:				
	R INFORMATIO	N (or person wit	h custody of	animal):						
Owner:		Pł	none:	·	(Oth	er)				
Address of Owner:	911#	Street Name	Apt	/Unit#	City					
Current Location of	Same as Owne	r: 🗌 Other 🗌 :								
Animal:			911#	Street Na	ime	Apt/Unit#	City			
Animal Species:	Dog 🗌 🤅 Cat 🗌	Bat O	ther 🗌 : 🔤							
Breed and Full Descri	-									
		pt alive and availal					of rabies.			
D		animal involved u		-						
1	The Health	Unit does not rem	nove healthy d	omestic anim	nal from their o	wners.				

This personal information is collected under the authority of the Health and Protection and Promotion Act (1990) Reg. 557 Sec. 2. It will be used to conduct investigations and for the purposes of monitoring and surveillance of rabies activity. For further details concerning this collection, contact the SMDHU Privacy Officer at 705-721-7520 or 1-877-721-7520.