Philadelphia/Center City 1528 Walnut Street Suite 1205 Philadelphia, PA 19102

Provider \_





Philadelphia/King Of Prussia 216 Mall Blvd Suite#1 King Of Prussia, PA, 19046

## VIMIZIM® (elosulfase alfa)

ORDER FORM Date: \_\_\_\_\_

	PATIEN	IT INFORMA	ATION						
Name:	Phone:		DOB:	SEX: M □ F □					
□NKDA Allergies:			We	eight lbs/kg:					
	PHYSICI	AN INFORM	MATION						
Physician Name*:	Prac	tice Name:	<u>,                                      </u>						
Address:	Offic	e Contact Name: Office Contact #:							
hone: Fax: Emai		il (for updates):							
	REFEI	RRAL STATUS							
□New Referral □Referral Renewal □N	1edication/Orde	r Change □Be	nefits Verification Only	☐ Discontinuation Order					
VIMIZIM <sup>®</sup> :									
□ VIMIZIIVI. □ VIMIZIM is indicated for patients with Muco	nolveaccharidos	is type IVA (MPS I	VA: Morquio A syndron	ne) F76 210					
- VIVIIZIIVI IS INdicated for patients with Mideo	рогузасспаниоз	is type TV/T (IVII 5 T	v/t, /viorquio/t syndron	ne). L7 0.2 10					
DOSAGE AND ADMINISTRATION: Recommended Dose Pre-treatment with antihistamines with or without antipyretics is recommended 30 to 60 minutes prior to the start of the infusion.		VIMIZIM ( PATIENT W	EIGHT _ lbs.						
PRE-MEDICATION			_ kg						
□ Tylenol PO 650mg □1000 MG □other		DOSAGE							
□ Solumedrol 125mg IV □ other		□ 300mg IV							
□ Benadryl □25mg □50mg □other □	IV □PO	□ Other	<del></del>						
□ Benadryl 50 mg □ or PO		FREQUENC	Y						
□ Medication DoseRoute		□ 2 mg/kg Weekly							
(other)	(other)		X weeks						
	(00101)			weeks					
WARNINGS AND PRECAUTIONS https://www.vimizim.com/wp-content/uploads/2018/02/ Prescribing-Information.pdf		Other							
WARNING: RISK OF ANAPHYLAXI									
Life-threatening anaphylactic reactions have occurred in some patients during VIMIZIM (elosulfase alfa) infusions.  Anaphylaxis, presenting as cough, erythema, throat tightness, urticaria, flushing, cyanosis, hypotension, rash, dyspnea, chest discomfort, and gastrointestinal symptoms (e.g., nausea, abdominal pain, retching, and vomiting) in conjunction with urticaria, have been reported to occur during VIMIZIM (elosulfase alfa) infusions, regardless of duration of the course of treatment.  Closely observe patients during and after VIMIZIM (elosulfase alfa) administration and be prepared to manage anaphylaxis. Inform patients of the signs and symptoms of anaphylaxis and have them seek immediate		REQUIRED DOCUMENTATION CHECKLIST:							
		Patient Demographics Insurance Card/Information Recent Progres notes addressing VIMIZIM in note Recent labs to include CBC, CMP, and please send any other recent labs.							
					medical care should symptoms occur. Patients with acut		Other		
					illness may be at risk of serious acute exacerbation of the compromise due to hypersensitivity reactions, and requimonitoring.		Other		
ORDERING PROVIDER									
Signature X		Date		NPI					

\_\_\_\_\_\_ Phone \_\_\_\_\_\_ Fax \_\_\_