Boca Raton 9980 Central Park Blvd Suite 202, N Boca Raton, FL 33428





## VIMIZIM® (elosulfase alfa)

Provider \_

ORDER FORM Date:

| V 11V11Z/11V1 (elosulfase alfa)   |                           |   |                                   |  |
|---|---------------------------|---|-----------------------------------|--|
| PATIENT INFORMATION   |                           |   |                                   |  |
| Name:   | Phone:                    | D   | OOB: SEX: M $\square$ F $\square$ |  |
| □NKDA Allergies:  |                           |   | Weight lbs/kg:                    |  |
| PHYSICIAN INFORMATION   |                           |   |                                   |  |
| Physician Name*: Practice Name:   |                           |   |                                   |  |
| Address:  | Office Contact Name:      |   | Office Contact #:                 |  |
| Phone: Fax:   | Fax: Email (for updates): |   |                                   |  |
| REFERRAL STATUS   |                           |   |                                   |  |
| □New Referral □Referral Renewal □Medication/Order Change □Benefits Verification Only □Discontinuation Order   |                           |   |                                   |  |
| VIMIZIM®:  UMIZIM is indicated for patients with Mucopolysaccharidosis type IVA (MPS IVA; Morquio A syndrome). E76.210  |                           |   |                                   |  |
| 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 ORDERS PATIENT WEIGHTlbs.   | S                                 |  |
| PRE-MEDICATION  |                           | kg  |                                   |  |
| ☐ Tylenol PO 650mg ☐1000 MG ☐ other   |                           | DOCACE  |                                   |  |
| □ Solumedrol 125mg IV □ other   |                           | DOSAGE  300mg IV  |                                   |  |
| □ Benadryl □25mg □50mg □other □ IV □PO  |                           | Other   |                                   |  |
| ☐ Benadryl 50 mg ☐ or PO  |                           |   |                                   |  |
| □ Medication DoseRoute  |                           | FREQUENCY   |                                   |  |
|   |                           | □ 2 mg/kg Weekly  |                                   |  |
| (other)   | (othe                     |   |                                   |  |
| WARNINGS AND PRECAUTIONS https://www.vimizim.com/wp-content/uploads/2018/02/ Prescribing-Information.pdf  WARNINGS AND PRECAUTIONS 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 medical care should symptoms occur. Patients with acute respiratory |                           | REQUIRED DOCUME   | REQUIRED DOCUMENTATION CHECKLIST: |  |
|   |                           | Patient Demograph   | ics                               |  |
|   |                           | Insurance Card/Information  |                                   |  |
|   |                           |   | es addressing VIMIZIM in note     |  |
|   |                           | Recent labs to <b>include CBC, CMP</b> , and please send any other recent labs. |                                   |  |
| illness may be at risk of serious acute exacerbation of thei compromise due to hypersensitivity reactions, and require monitoring.  | r respiratory             | Other   |                                   |  |
|   |                           |   |                                   |  |
| ORDERING PROVIDER   |                           |   |                                   |  |
| Signature <b>X</b>  |                           | Date  | NPI                               |  |

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