

OPTIMA FAMILY CARE MEDALLION 4.0

MEDICAL PRIOR AUTHORIZATION/STEP-EDIT REQUEST

Directions: The prescribing physician must sign and clearly print name (preprinted stamps not valid) on this request. All other information may be filled in by office staff; fax to 1-844-723-2094. No additional phone calls will be necessary if all information (including phone and fax #s) on this form is correct. Incomplete form will delay the authorization process.

Drug Requested: Vimizim® IV (elosulfase alfa) (J1322) (Medical)

****IV INFUSION PERFORMED AT SENTARA INFUSION CENTERS ONLY****

DRUG INFORMATION: Complete all information below or authorization process will be delayed.

Drug Form/Strength/Month: _____

Dosing Schedule: _____ Length of Therapy: _____

Diagnosis: _____ ICD Code: _____

****If approved, max dose allowed is 2mg/kg to be administered once weekly.**

CLINICAL CRITERIA: Check **ALL** boxes that apply to ensure authorization process will **NOT** be delayed.

- Due to high risk of anaphylaxis and infusion reactions, Vimizim® infusion should be administered **ONLY** by trained medical professionals and will **NOT** be approved for self-administration or for administration by home healthcare providers.

For Initial Approval: Initial approval will be for **24 weeks**. CHART NOTES **MUST** BE SUBMITTED. Boxes below **MUST** be checked to ensure authorization is **NOT** delayed.

- The prescriber is a metabolic geneticist or endocrinologist
- The patient is at least 5 years of age
- The patient has a diagnosis of mucopolysaccharidosis type IVA (MPS IVA) as verified by genetic testing (*labs confirming diagnosis must be submitted*)
- Patient's current height (please note): _____ Patient's current weight (please note): _____
Current FEV₁ (*please submit labs*): _____ Current MVV (*please submit labs*): _____
- Baseline 6 minute walk time of a distance of **at least** 30 meters is attached (*please attach baseline 6 minute walk time with date noted*)
- Chart notes are attached to document symptoms, prior medical procedures, and prior therapies used in the treatment of MPS IVA

For Continued Approval: Continued approval will be based on patient maintaining sustained improved walk time above baseline walk time and evidence of clinical improvement. Continued approval is for **12 months**.

- Current 6 minute walk time is attached (*please attach current 6 minute walk time with date noted*): _____
- Patient's 6 minute walk time has sustained improvement from baseline
- Patient's current height (*please note*): _____ Patient's current weight (*please note*): _____
Current FEV₁ (*please submit labs*): _____ Current MVV (*please submit labs*): _____
- Patient's current normalized urine keratan sulfate levels (*please submit labs*): _____
- Chart notes are attached to document current disease status, any medical procedures performed since last approval of this medication, and evidence of clinical improvement from baseline (*please attach chart notes*)

(continued on next page)

Medication being provided by Sentara Infusion Centers. Complete all information below:

- Location/site of administration: _____
NPI or DEA # of administering location: _____

****Use of samples to initiate therapy does not meet step edit/ preauthorization criteria.****

Previous therapies will be verified through pharmacy paid claims or submitted chart notes.

Patient Name: _____
Member Optima #: _____ Date of Birth: _____
Prescriber Name: _____
Prescriber Signature: _____ Date: _____
Office Contact Name: _____
Phone Number: _____ Fax Number: _____
Prescriber's DEA OR NPI #: _____

*Approved by Pharmacy and Therapeutics Committee: 4/19/2018
REVISED/UPDATED: 7/1/2018