

OPTIMA HEALTH FAMILY CARE (MEDICAID)

PHARMACY/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, LAB RESULTS **MUST** BE SUBMITTED **WITHIN THE LAST 30 DAYS**. **ALL** criteria **MUST** be met for approval.

- The prescriber is a metabolic geneticist or endocrinologist

AND

- The patient is at least 5 years of age

AND

- The patient has a diagnosis of mucopolysaccharidosis type IVA (MPS IVA) as verified by genetic testing (**labs confirming diagnosis must be submitted**)

AND

- Patient's current height (please note): _____ Patient's current weight (please note): _____

AND

Current FEV1 (**please submit labs**): _____ Current MVV (**please submit lab results within the last 30 days**): _____

AND

(continued on next page)

- ❑ Patient’s current normalized urine keratan sulfate levels (please submit lab results within the last 30 days):

AND

- ❑ 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)

AND

- ❑ Chart notes are attached to document symptoms, prior medical procedures, and prior therapies used in the treatment of MPS IVA

AND

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 of one within the last 30 days is attached (please attach current 6 minute walk time with date noted):

AND

- ❑ Patient’s 6 minute walk time has sustained improvement from baseline

AND

- ❑ Patient’s current height (please note): _____ Patient’s current weight (please note): _____

AND

Current FEV₁ (please submit labs within last 30 days): _____

Current MVV (please submit labs within the last 30 days): _____

AND

- ❑ Patient’s current normalized urine keratan sulfate levels (please submit labs within the last 30 days): _____

AND

- ❑ 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)

Medication being provided by the following:

- ❑ Location/site of administration: _____

NPI or DEA # of administering location: _____

OR

- ❑ Specialty Pharmacy - PropriumRx

(Continued on next page; signature page **MUST** be attached to this request.)

(Signature page **MUST** be included with this request.)

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: 6/30/2018; 8/31/2018; 10/8/2018; (REFORMATTED) 2/5/2019