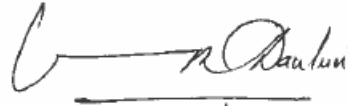


Prior Authorization Review Panel

Prior Authorization Review Panel

CHC-MCO Policy Submission

A separate copy of this form must accompany each policy submitted for review.
Policies submitted without this form will not be considered for review.

Plan: PA Health & Wellness	Submission Date: 05/01/2022
Policy Number: PA.CP.PHAR.161	Effective Date: 01/2018 Revision Date: 04/2022
Policy Name: Galsulfase (Naglazyme)	
<p>Type of Submission – <u>Check all that apply</u>:</p> <p> <input type="checkbox"/> New Policy <input checked="" type="checkbox"/> Revised Policy* <input type="checkbox"/> Annual Review - No Revisions <input type="checkbox"/> Statewide PDL - <i>Select this box when submitting policies for Statewide PDL implementation and when submitting policies for drug classes included on the Statewide PDL.</i> </p>	
<p>*All revisions to the policy <u>must</u> be highlighted using track changes throughout the document.</p> <p>Please provide any changes or clarifying information for the policy below:</p> <p>2Q 2022 annual review: added requirement for documentation of member's current weight for dose calculation purposes; references reviewed and updated.</p>	
Name of Authorized Individual (Please type or print): Venkateswara R. Davuluri, MD	Signature of Authorized Individual: 

Clinical Policy: Galsulfase (Naglazyme)

Reference Number: PA.CP.PHAR.161

Effective Date: 01/18

Last Review Date: 04/2022

[Coding Implications](#)

[Revision Log](#)

Description

Galsulfase (Naglazyme®) is a hydrolytic lysosomal glycosaminoglycan-specific enzyme.

FDA approved indication

Naglazyme is indicated for the treatment of patients with mucopolysaccharidosis VI (MPS VI; Maroteaux-Lamy syndrome). Naglazyme has been shown to improve walking and stair-climbing capacity.

Policy/Criteria

Provider must submit documentation (such as office chart notes, lab results or other clinical information) supporting that member has met all approval criteria.

It is the policy of PA Health & Wellness that Naglazyme is **medically necessary** when the following criteria are met:

I. Initial Approval Criteria

A. Maroteaux-Lamy Syndrome (Mucopolysaccharidosis VI [MPS VI]): (must meet all):

1. Diagnosis of Maroteaux-Lamy syndrome (MPS VI) confirmed by one of the following:
 - a. Enzyme assay demonstrating a deficiency in N-acetylgalactosamine 4-sulfatase (arylsulfatase B) activity;
 - b. DNA testing;
2. Age \geq 3 months;
3. Documentation of member's current weight (in kg);
4. Dose does not exceed 1 mg/kg/week.

Approval duration: 6 months

B. Other diagnoses/indications: Refer to PA.CP.PMN.53

II. Continued Approval

A. Maroteaux-Lamy Syndrome (MPS VI): (must meet all):

1. Currently receiving medication via PA Health & Wellness benefit or member has previously met all initial approval criteria; or the Continuity of Care policy (PA.LTSS.PHAR.01) applies;
2. Member is responding positively to therapy as evidenced by improvement in the individual member's MPS VI (Maroteaux-Lamy syndrome) manifestation profile (*see Appendix D for examples*);
3. Documentation of member's current weight (in kg);
4. If request is for a dose increase, new dose does not exceed 1 mg/kg/week.

Approval duration: 12 months

B. Other diagnoses/indications (must meet 1 or 2):

1. Currently receiving medication via PA Health & Wellness benefit and documentation supports positive response to therapy; or the Continuity of Care policy (PA.LTSS.PHAR.01) applies; or
2. Refer to PA.CP.PMN.53

III. Appendices/General Information

Appendix A: Abbreviation/Acronym Key

FDA: Food and Drug Administration

MPS VI: mucopolysaccharidosis VI

Appendix B: Therapeutic Alternatives

Not applicable

Appendix C: Contraindications/Boxed Warnings

- Contraindication(s): none reported.
- Boxed warning(s): none reported.

Appendix D: General Information

The presenting symptoms and clinical course of MPS VI can vary from one individual to another. Some examples, however, of improvement in MPS VI disease as a result of Naglazyme therapy may include improvement in:

- 12-minute walking test distance;
- 3-minute stair climb rate;
- Poor endurance;
- Vision problems;
- Respiratory infections;
- Breathing problems, sleep apnea;
- High blood pressure;
- Joint stiffness;
- Hepatomegaly, splenomegaly.

IV. Dosage and Administration

Indication	Dosing Regimen	Maximum Dose
MPS VI	1 mg/kg IV once weekly	1 mg/kg/week

V. Product Availability

Vial: 5 mg/5 mL

VI. References

1. Naglazyme Prescribing Information. Novato, CA: BioMarin Pharmaceutical, Inc.; December 2019. Available at <http://www.naglazyme.com>. Accessed February 26, 2022.
2. Muenzer J. The mucopolysaccharidoses: a heterogeneous group of disorders with variable pediatric presentations. J Pediatr. 2004; 144(5 Suppl): S27-S34.

3. Akyol MU, Alden TD, Amartino H, et al. Recommendations for the management of MPS VI: systematic evidence- and consensus-based guidance. Orphanet J of Rare Dis, 2019;12(118)1-21.

Coding Implications

Codes referenced in this clinical policy are for informational purposes only. Inclusion or exclusion of any codes does not guarantee coverage. Providers should reference the most up-to-date sources of professional coding guidance prior to the submission of claims for reimbursement of covered services.

HCPCS Codes	Description
J1458	Injection, galsulfase, 1 mg

Reviews, Revisions, and Approvals	Date	Approval Date
2Q 2018 annual review: Modified age restriction to 3 months per PI. Added prescriber requirement. Added max dose criteria. Added requirement for positive response to therapy. References reviewed and updated.	02.13.18	04.18
2Q 2019 annual review: references reviewed and updated.	04/2019	
2Q 2020 annual review: references reviewed and updated.	04/2020	
2Q 2021 annual review: references reviewed and updated.	04/2021	
2Q 2022 annual review: added requirement for documentation of member's current weight for dose calculation purposes; references reviewed and updated.	04/2022	