

Nulibry [®] (fosdenopterin) Effective 11/01/2021

Plan	☑ MassHealth□Commercial/Exchange			\boxtimes Prior Authorization	
Benefit	□ Pharmacy Benefit ⊠ Medical Benefit (NLX)		Program Type	□ Quantity Limit □ Step Therapy	
Specialty Limitations	N/A				
	Specialty Medications				
	All Plans	Pł	Phone: 866-814-5506 Fax: 866-249-6155		
	Non-Specialty Medications				
Contact	MassHealth	Pł	none: 877-433-7643	Fax: 866-255-7569	
Information	Commercial	Pł	Phone: 800-294-5979 Fax: 888-836-0730		
	Exchange	Pl	none: 855-582-2022	Fax: 855-245-2134	
	Medical Specialty Medications (NLX)				
	All Plans	Pł	none: 844-345-2803	Fax: 844-851-0882	
Exceptions	N/A				

Overview

Nulibry is cyclic pyranopterin monophosphate (cPMP) indicated to reduce the risk of mortality in patients with molybdenum cofactor deficiency (MoCD) Type A.

Coverage Guidelines

Authorization may be reviewed for members new to AllWays Health Partners who are currently receiving treatment with Nulibry excluding when the product is obtained as samples or via manufacturer's patient assistance programs.

OR

Authorization may be granted for members when ALL the following criteria are met, and documentation is provided:

- 1. Appropriate diagnosis of molybdenum cofactor deficiency (MoCD) Type A confirmed by genetic testing
- 2. Prescriber is a specialist in genetic or metabolic diseases or consult is provided
- 3. Appropriate dosing
- 4. Member's current weight.

Continuation of Therapy

Reauthorization will be granted if member meets all of the following criteria:

- 1. Improved change in molybdenum cofactor deficiency biomarkers
- 2. Improved growth parameters

Limitations

1. Initial approvals will be granted for 3 months

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2. Reauthorizations will be granted for 12 months

References

- 1. Nulibry [package insert]. Boston, MA: Origin Biosciences, Inc.; February 2021.
- 2. Atwal PS, Scaglia F. Molybdenum cofactor deficiency. Mol Genet Metab. 2016;117(1):1-4.
- 3. Schwahn BC, Van Spronsen FJ, Belaidi AA, et al. Efficacy and safety of cyclic pyranopterin monophosphate substitution in severe molybdenum cofactor deficiency type A: a prospective cohort study. Lancet. 2015; 386: 1955-1963.
- 4. ClinicalTrials.gov. Study of ORGN001 (formerly ALXN1101) in neonates with molybdenum cofactor deficiency (MOCD) type A. Available at: <u>https://clinicaltrials.gov/ct2/show/NCT02629393.</u>
- ClinicalTrials.gov. Safety & efficacy study of ORGN001 (formerly ALXN1101) in pediatric patients with MoCD type A currently treated with rcPMP. Available at: https://clinicaltrials.gov/ct2/show/NCT02047461.

Review History

09/22/2021 - Reviewed and Created for Sept P&T. Effective 11/01/2021

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