



Rivfloza® (nedosiran) (Subcutaneous)

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I. Length of Authorization

Coverage will be provided for 6 months initially and may be renewed annually thereafter.

II. Dosing Limits

Max Units (per dose and over time) [HCPCS Unit]:

• 160 mg every month

III. Initial Approval Criteria ¹

Coverage is provided in the following conditions:

Patient is at least 9 years of age; AND

Universal Criteria 1-4

- Patient has not had a liver transplant; AND
- Must be prescribed by, or in consultation with, a specialist in genetics, nephrology or urology;
 AND
- Patient does not have severe renal impairment defined as an eGFR <30 mL/min/1.73 m²; AND
- Will not be used in combination with other urinary oxalate reducing agents (i.e., lumasiran, etc.);
 AND

Primary Hyperoxaluria Type 1 (PH1) † Φ 1-4

- Patient has a definitive diagnosis of primary hyperoxaluria type 1 as evidenced by one of the following:
 - Patient has a biallelic pathogenic mutation in the alanine: glyoxylate aminotransferase
 (AGXT) gene as identified on molecular genetic testing; OR
 - Identification of alanine: glyoxylate aminotransferase (AGT) enzyme deficiency on liver biopsy; AND
- Patient has a baseline for one or more of the following:
 - Urinary oxalate excretion level (corrected for BSA)
 - Spot urinary oxalate: creatinine ratio
 - Estimated glomerular filtration rate (eGFR)

Plasma oxalate level

† FDA Approved Indication(s); ‡ Compendium Recommended Indication(s); • Orphan Drug

IV. Renewal Criteria 1-4

Coverage can be renewed based upon the following criteria:

- Patient continues to meet the universal and other indication-specific relevant criteria identified in section III; AND
- Absence of unacceptable toxicity from the drug. Examples of unacceptable toxicity include: severe injection site reactions, etc.; AND
- Disease response as evidenced by at least one of the following:
 - o Decrease in urinary oxalate excretion level (corrected for BSA) from baseline
 - Reduction in spot urinary oxalate: creatinine ratio from baseline
 - Stabilization of estimated glomerular filtration rate (eGFR)
 - Decrease in plasma oxalate level from baseline

V. Dosage/Administration ¹

Indication	Dose
Primary Hyperoxaluria	For administration by a healthcare professional, caregiver or patient as a subcutaneous injection.
Type 1 (PH1)	Children 9 to 11 years of age
	- ≥ 50 kg: 160 mg pre-filled syringe once monthly
	- < 50 kg: 3.3 mg/kg once monthly, not to exceed 128 mg (vial, dose volume rounded to the nearest 0.1 mL)
	Adults and adolescents 12 years of age and older
	- ≥ 50 kg: 160 mg pre-filled syringe once monthly
	- < 50 kg: 128 mg pre-filled syringe once monthly

VI. Billing Code/Availability Information

HCPCS:

- J3490 Unclassified drugs
- C9399 Unclassified drugs or biologicals (Hospital outpatient use ONLY)

NDC:

- Rivfloza 80 mg/0.5 mL in a single-dose vial: 00169-5308-xx
- Rivfloza 128 mg/0.8 mL in a single-dose pre-filled syringe: 00169-5307-xx
- Rivfloza 160 mg/1 mL in a single-dose pre-filled syringe: 00169-5306-xx



VII. References

- Rivfloza [package insert]. Plainsboro, NJ; Novo Nordisk, Inc., September 2023. Accessed December 2024.
- 2. Milliner DS, Harris PC, Sas DJ, et al. Primary Hyperoxaluria Type 1. Initial Posting: June 19, 2002; Last Update: August 15, 2024. In: Adam MP, Feldman J, Mirzaa GM, et al., editors. GeneReviews® [Internet]. Seattle (WA): University of Washington, Seattle; 1993-2024. Available from: https://www.ncbi.nlm.nih.gov/books/NBK1283/. Accessed January 8, 2025.
- 3. Baum MA, Langman C, Cochat P, et al. PHYOX2: a pivotal randomized study of nedosiran in primary hyperoxaluria type 1 or 2. Kidney International, Volume 103, Issue 1, 2023, Pages 207-217, ISSN 0085-2538, https://doi.org/10.1016/j.kint.2022.07.025.
- 4. Groothoff J, Sellier-Leclerc AL, Deesker L, et al. Nedosiran Safety and Efficacy in PH1: Interim Analysis of PHYOX3. Kidney Int Rep. 2024 Mar 4;9(5):1387-1396. doi: 10.1016/j.ekir.2024.02.1439. PMID: 38707801; PMCID: PMC11068990.

Appendix 1 – Centers for Medicare and Medicaid Services (CMS)

The preceding information is intended for non-Medicare coverage determinations. Medicare coverage for outpatient (Part B) drugs is outlined in the Medicare Benefit Policy Manual (Pub. 100-2), Chapter 15, §50 Drugs and Biologicals. In addition, National Coverage Determinations (NCDs) and/or Local Coverage Determinations (LCDs) may exist and compliance with these policies is required where applicable. Local Coverage Articles (LCAs) may also exist for claims payment purposes or to clarify benefit eligibility under Part B for drugs which may be self-administered. The following link may be used to search for NCD, LCD, or LCA documents: https://www.cms.gov/medicare-coverage-database/search.aspx. Additional indications, including any preceding information, may be applied at the discretion of the health plan.

Medicare Part B Covered Diagnosis Codes (applicable to existing NCD/LCA/LCD): N/A

Medicare Part B Administrative Contractor (MAC) Jurisdictions			
Jurisdiction	Applicable State/US Territory	Contractor	
E (1)	CA, HI, NV, AS, GU, CNMI	Noridian Healthcare Solutions, LLC	
F (2 & 3)	AK, WA, OR, ID, ND, SD, MT, WY, UT, AZ	Noridian Healthcare Solutions, LLC	
5	KS, NE, IA, MO	Wisconsin Physicians Service Insurance Corp (WPS)	
6	MN, WI, IL	National Government Services, Inc. (NGS)	
H (4 & 7)	LA, AR, MS, TX, OK, CO, NM	Novitas Solutions, Inc.	
8	MI, IN	Wisconsin Physicians Service Insurance Corp (WPS)	
N (9)	FL, PR, VI	First Coast Service Options, Inc.	
J (10)	TN, GA, AL	Palmetto GBA	
M (11)	NC, SC, WV, VA (excluding below)	Palmetto GBA	
L (12)	DE, MD, PA, NJ, DC (includes Arlington & Fairfax counties and the city of Alexandria in VA)	Novitas Solutions, Inc.	



Medicare Part B Administrative Contractor (MAC) Jurisdictions				
Jurisdiction	Applicable State/US Territory	Contractor		
K (13 & 14)	NY, CT, MA, RI, VT, ME, NH	National Government Services, Inc. (NGS)		
15	KY, OH	CGS Administrators, LLC		

