

## Clinical Policy: Nitisinone (Nityr, Orfadin)

Reference Number: PA.CP.PHAR.132

Effective Date: 10.17.18

Last Review Date: 10/2023

[Revision Log](#)

### Description

Nitisinone (Nityr™, Orfadin®) is a hydroxy-phenylpyruvate dioxygenase inhibitor.

### FDA Approved Indication(s)

Nityr and Orfadin are indicated for the treatment of adult and pediatric patients with hereditary tyrosinemia type 1 (HT-1) in combination with dietary restriction of tyrosine and phenylalanine.

### 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 Nityr and Orfadin are **medically necessary** when the following criteria are met:

### I. Initial Approval Criteria

#### A. Hereditary Tyrosinemia Type 1 (must meet all):

1. Diagnosis of HT-1 as confirmed by one of the following (a or b);
  - a. Genetic testing confirms a mutation of the *FAH* gene;
  - b. Biochemical testing confirms elevated levels of succinylacetone in blood or urine;\*

*\* The lower limit of normal for succinylacetone is laboratory- and/or treatment center-specific; refer to laboratory- or clinic-specific reference ranges to determine elevated levels.*
2. Prescribed by or in consultation with an endocrinologist or a metabolic or genetic disease specialist;
3. Request is for use as an adjunct to dietary restriction of tyrosine and phenylalanine;
4. Member is not using two different nitisinone products concurrently;
5. For requests for Nityr and Orfadin 2 mg, 5 mg, or 10 mg strengths, member must use generic nitisinone, unless contraindicated or clinically significant adverse effects are experienced;
6. Dose does not exceed 2 mg/kg per day.

**Approval duration:** 6 months

#### B. Other diagnoses/indications

1. Refer to the off-label use policy if diagnosis is NOT specifically listed under section III (Diagnoses/Indications for which coverage is NOT authorized): PA.CP.PMN.53.

### II. Continued Therapy

#### A. Hereditary Tyrosinemia Type 1 (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;
  3. Request is for use as an adjunct to dietary restriction of tyrosine and phenylalanine;
  4. Member is not using two different nitisinone products concurrently;
  5. For requests for Nityr and Orfadin 2 mg, 5 mg, or 10 mg strengths, member must use generic nitisinone, unless contraindicated or clinically significant adverse effects are experienced;
  6. If request is for a dose increase, new dose does not exceed 2 mg/kg per day.
- Approval duration:** 12 months

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

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.  
**Approval duration: Duration of request or 6 months (whichever is less);** or
2. Refer to the off-label use policy if diagnosis is NOT specifically listed under section III (Diagnoses/Indications for which coverage is NOT authorized): PA.CP.PMN.53.

**III. Diagnoses/Indications for which coverage is NOT authorized:**

- A. Non-FDA approved indications, which are not addressed in this policy, unless there is sufficient documentation of efficacy and safety according to the off label use policy – PA.CP.PMN.53 or evidence of coverage documents.

**IV. Appendices/General Information**

*Appendix A: Abbreviation/Acronym Key*

FDA: Food and Drug Administration

HT-1: hereditary tyrosinemia type 1

*Appendix B: Therapeutic Alternatives*

*This table provides a listing of preferred alternative therapy recommended in the approval criteria. The drugs listed here may not be a formulary agent for all relevant lines of business and may require prior authorization.*

Drug Name	Dosing Regimen	Dose Limit/ Maximum Dose
Generic nitisinone (Orfadin, Nityr)	0.5 mg/kg PO BID	2 mg/kg

*Therapeutic alternatives are listed as Brand name<sup>®</sup> (generic) when the drug is available by brand name only and generic (Brand name<sup>®</sup>) when the drug is available by both brand and generic.*

*Appendix C: Contraindications/Boxed Warnings*

- Contraindication(s): none reported
- Box warning(s): none reported

**V. Dosage and Administration**

Drug Name	Dosing Regimen	Maximum Dose
Nitisinone (Nityr)	0.5 mg/kg PO BID	2 mg/kg
Nitisinone (Orfadin)	0.5 mg/kg PO BID	2 mg/kg

**VI. Product Availability**

Drug Name	Availability
Nitisinone (Nityr)	Tablets: 2 mg, 5 mg, 10 mg
Nitisinone (Orfadin)	Capsules: 2 mg, 5 mg, 10 mg, 20 mg Oral suspension: 4 mg/mL

**VII. References**

1. Orfadin Prescribing Information. Waltham, MA: Sobi, Inc.; November 2021. Available at: <http://www.orfadin.com/>. Accessed June 28, 2023.
2. Nityr Prescribing Information. Centro Insema, Manno Switzerland: Rivopharm; June 2021. Available at: [www.nityr.us](http://www.nityr.us). Accessed August 11, 2023.
3. Chinsky JM, Singh R, Ficicioglu C, et al. Diagnosis and treatment of tyrosinemia type I: a US and Canadian consensus group review and recommendations. *Genetics in Medicine*. Dec 2017;19(12).
4. Van Ginkel WG, Rodenburg IL, Harding CO, et al. Long-term outcomes and practical considerations in the pharmacological management of tyrosinemia type 1. *Pediatr Drugs*. 2019;21:413–26. <https://doi.org/10.1007/s40272-019-00364-4>.

Reviews, Revisions, and Approvals	Date	P&T Approval Date
Policy created	10/2018	
4Q 2019 annual review: No changes per Statewide PDL implementation 01-01-2020	10/2019	
4Q 2020 annual review: added requirement for adjunctive dietary restriction of tyrosine and phenylalanine, in line with the FDA-approved indication; references reviewed and updated.	10/2020	
4Q 2021 annual review: added requirement for diagnosis confirmation by either genetic or biochemical testing; references reviewed and updated.	10/2021	
4Q 2022 annual review: no significant changes; references reviewed and updated.	10/2022	
4Q 2023 annual review: no significant changes; added exclusion against concomitant use of multiple different nitisinone products; added generic redirection for 2 mg, 5 mg, 10 mg strengths (generic nitisinone 20 mg strength is either NF or same tier level as brand Orfadin 20 mg); references reviewed and updated.	10/2023	