

## UnitedHealthcare Pharmacy Clinical Pharmacy Programs

Program Number	2024 P 1401-3
Program	Prior Authorization/Notification
Medication	Cuvrior <sup>™</sup> (trientine tetrahydrochloride)*
P&T Approval Date	1/2023, 1/2024, 2/2024
Effective Date	5/1/2024

### 1. Background:

Cuvrior\* (trientine tetrahydrochloride) is a copper chelator indicated for the treatment of adult patients with Wilson's disease who are de-coppered and tolerant to penicillamine.

## 2. Coverage Criteria<sup>a</sup>:

## A. Initial Authorization

- 1. Cuvrior\* will be approved based on all of the following criteria:
  - a. Diagnosis of Wilson's disease

#### -AND-

b. Patient is de-coppered [i.e., serum non-ceruloplasmin copper (NCC) level  $\geq$  25 and  $\leq$  150 mcg/L]

#### -AND-

c. Patient is tolerant to penicillamine

#### -AND-

d. Patient will discontinue penicillamine before starting therapy with Cuvrior

Authorization will be issued for 12 months.

## B. Reauthorization

- 1. **Cuvrior** will be approved based upon the following criterion:
  - a. Documentation of positive clinical response to Cuvrior therapy

## Authorization will be issued for 12 months.

<sup>a</sup> State mandates may apply. Any federal regulatory requirements and the member specific benefit plan coverage may also impact coverage criteria. Other policies and utilization management programs may apply.

<sup>\*</sup>Cuvrior is typically excluded from coverage. Tried/Failed criteria may be in place. Please refer to plan specifics to determine exclusion status.



# 3. Additional Clinical Rules:

- Notwithstanding Coverage Criteria, UnitedHealthcare may approve initial and reauthorization based solely on previous claim/medication history, diagnosis codes (ICD-10) and/or claim logic. Use of automated approval and re-approval processes varies by program and/or therapeutic class.
- Supply limits may be in place.

## 4. References:

1. Cuvrior [package insert]. Chicago, IL: Orphalan; April 2022.

Program	Prior Authorization/Notification – Cuvrior (trientine tetrahydrochloride)
Change Control	
1/2023	New program.
1/2024	Annual review with no changes to coverage criteria.
2/2024	Annual review. Added footnote indicating Cuvrior is typically
	excluded from coverage. Updated authorization durations to 12
	months.