

### UnitedHealthcare Pharmacy Clinical Pharmacy Programs

| Program Number    | 2023 P 1148-10   |
|-------------------|--|
| Program           | Prior Authorization/Notification                                   |
| Medications       | Cerdelga <sup>®</sup> (eliglustat)                                 |
| P&T Approval Date | 12/2014, 11/2015, 9/2016, 9/2017, 9/2018, 9/2019, 9/2020, 10/2021, |
|                   | 10/2022, 10/2023   |
| Effective Date    | 1/1/2024   |

# 1. Background:

Cerdelga<sup>®</sup> (eliglustat) is a glucosylceramide synthase inhibitor indicated for the long-term treatment of adult patients with Gaucher disease type 1 who are CYP2D6 extensive metabolizers (EMs), intermediate metabolizers (IMs), or poor metabolizers (PMs) as detected by an FDA-cleared test. CYP2D6 ultra-rapid metabolizers may not achieve adequate concentrations of Cerdelga to achieve a therapeutic effect. A specific dosage cannot be recommended for CYP2D6 indeterminate metabolizers.<sup>1</sup>

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

## A. Initial Authorization

- 1. Cerdelga will be approved based on both of the following criteria:
  - a. Diagnosis of Gaucher disease type 1

#### -AND-

- b. Patient is <u>one</u> of the following as detected by an FDA-cleared test:
  - (1) CYP2D6 extensive metabolizer
  - (2) CYP2D6 intermediate metabolizer
  - (3) CYP2D6 poor metabolizer

#### Authorization will be issued for 12 months.

## B. <u>Reauthorization</u>

- 1. Cerdelga will be approved based on the following criterion:
  - a. Documentation of positive clinical response to Cerdelga 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.



### **3.** Additional Clinical Programs:

- Notwithstanding Coverage Criteria, UnitedHealthcare may approve initial and re-authorization 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.
- Supply limits may be in place.

### 4. References:

1. Cerdelga [package insert]. Genzyme Ireland, Ltd. Waterford, Ireland. December 2022.

| Program        | Prior Authorization/Notification - Cerdelga (eliglustat)              |
|----------------|---|
| Change Control |   |
| 12/2014        | New program.  |
| 11/2015        | Annual review. Updated to align with Indication Section of FDA label. |
| 9/2016         | Annual review. No changes to coverage criteria.                       |
| 9/2017         | Annual review. No changes.  |
| 9/2018         | Annual review. No changes.  |
| 9/2019         | Annual review. No changes to coverage criteria. Updated background    |
|                | and references.   |
| 9/2020         | Annual review. No changes to coverage criteria.                       |
| 10/2021        | Annual review. Updated reauthorization duration. Updated reference.   |
| 10/2022        | Annual review. No changes to coverage criteria. Added state mandate.  |
|                | Updated reference.  |
| 10/2023        | Annual review. No changes to coverage criteria. Updated reference.    |