

# UnitedHealthcare Pharmacy Clinical Pharmacy Programs

Program Number	2025 P 2282-4
Program	Prior Authorization/Medical Necessity
Medication	Radicava ORS® (edaravone)
P&T Approval Date	7/2022, 7/2023, 7/2024, 3/2025
Effective Date	6/1/2025

### 1. Background:

Radicava ORS® is indicated for the treatment of amyotrophic lateral sclerosis (ALS).<sup>1</sup>

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

## A. Initial Authorization

- 1. Radicava ORS will be approved based on <u>one</u> of the following criteria:
  - a. **Both** of the following criteria:
    - (1) Patient has been established on therapy with edaravone for amyotrophic lateral sclerosis under an active UnitedHealthcare medical benefit prior authorization

#### -AND-

- (2) <u>All</u> of the following:
  - (a) Diagnosis of ALS
  - (b) Prescribed by, or in consultation with, a neurologist with expertise in the diagnosis of ALS
  - (c) Patient is currently receiving edaravone therapy
  - (d) Patient is not dependent on invasive ventilation

### -OR-

- b. All of the following criteria:
  - (1) Submission of medical records (e.g., chart notes, previous medical history, diagnostic testing including: imaging, nerve conduction studies, laboratory values) to support the diagnosis of ALS

### -AND-

(2) Prescribed by, or in consultation with, a neurologist with expertise in the diagnosis of ALS

#### -AND-

(3) Submission of the most recent ALS Functional Rating Scale-Revised



(ALSFRS-R) score confirming that the patient has scores  $\geq 2$  in all items of the ALSFRS-R criteria at the start of treatment

#### -AND-

(4) Submission of medical records (e.g., chart notes, laboratory values) confirming that the patient has a % forced vital capacity (%FVC) ≥ 80% at the start of treatment

Authorization will be issued for 12 months.

# B. Reauthorization

- 1. Radicava ORS will be approved based on <u>all</u> of the following criteria:
  - a. Diagnosis of ALS

### -AND-

b. Prescribed by, or in consultation with, a neurologist with expertise in the diagnosis of ALS

### -AND-

c. Patient is currently receiving **Radicava ORS** therapy

#### -AND-

d. Patient is **not** dependent on invasive ventilation

# 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 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. Radicava ORS [package insert]. Jersey City, NJ: Mitsubishi Tanabe Pharma Corporation. November 2022.
- 2. Subcommittee on Motor Neuron Diseases of World Federation of Neurology Research Group on Neuromuscular Diseases, El Escorial "Clinical Limits of ALS" Workshop Contributors. El



- Escorial World Federation of Neurology criteria for the diagnosis of amyotrophic lateral sclerosis. J Neurol Sci 1994; 124: 96–107.
- 3. Takahashi F, Takei K, Tsuda K, Palumbo J. Post-hoc analysis of MCI186-17, the extension study to MCI186-16, the confirmatory double-blind, parallel-group, placebo-control006Ced study of edaravone in amyotrophic lateral sclerosis. Amyotrophic Lateral Sclerosis and Frontotemporal Degeneration. 2017;18(sup1):32-39.

Program	Prior Authorization/Medical Necessity – Radicava ORS® (edaravone)	
Change Control		
7/2022	New program.	
7/2023	Annual review with no changes to the coverage criteria.	
7/2024	Annual review. Clarified criteria for existing prior authorization for under the medical benefit. Updated initial authorization and	
	reauthorization to 12 months.	
3/2025	Updated reference to Radicava IV to reflect that edaravone IV is available generically. Simplified diagnosis requirement. Updated invasive ventilation requirement with no change to clinical intent. Updated references.	