

UnitedHealthcare Pharmacy  
Clinical Pharmacy Programs

Program Number	2024 P 2177-7
Program	Prior Authorization/Medical Necessity
Medication	Firdapse® (amifampridine)
P&T Approval Date	11/2019, 11/2020, 11/2021, 3/2022, 11/2022, 11/2023, 11/2024
Effective Date	2/1/2025

**1. Background:**

Firdapse (amifampridine) is a potassium channel blocker indicated for the treatment of Lambert-Eaton myasthenic syndrome (LEMS) in adults and pediatric patients 6 years of age and older.<sup>1</sup>

**2. Coverage Criteria<sup>a</sup>:****A. Initial Authorization**

1. **Firdapse** will be approved based on **all** of the following criteria:

a. Diagnosis of Lambert-Eaton myasthenic syndrome (LEMS)

**-AND-**

b. Prescribed by or in consultation with a specialist in the treatment of LEMS (e.g., neurologist or oncologist)

**-AND-**

c. Patient is not receiving Firdapse in combination with similar potassium channel blockers [e.g., Ampyra (dalfampridine)]

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

**B. Reauthorization**

1. **Firdapse** will be approved based on **both** the following criteria:

a. Documentation of positive clinical response to Firdapse therapy

**-AND-**

b. Patient is not receiving Firdapse in combination with similar potassium channel blockers [e.g., Ampyra (dalfampridine)]

**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 and/or therapeutic class.
- Supply limits may be in place.

**4. References:**

1. Firdapse [package insert]. Coral Gables, FL: Catalyst Pharmaceuticals, Inc.; May 2024.

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<b>Change Control</b>	
11/2019	New program
11/2020	Annual review with no changes to coverage criteria. Updated references.
11/2021	Annual review with no changes to coverage criteria. Updated references.
3/2022	Removed step through Ruzurgi due to FDAs conversion of Ruzurgi from full approval to tentative approval.
11/2022	Updated background to reflect new pediatric indication for patients 6 years of age and older.
11/2023	Added “Diagnosis of” to initial criteria with no change to clinical intent.
11/2024	Annual review with no changes to coverage criteria. Updated reference.