

## PRIOR AUTHORIZATION POLICY

**POLICY:** Amifampridine Products Prior Authorization Policy

- Firdapse® (amifampridine tablets – Catalyst)
- Ruzurgi® (amifampridine tablets – Jacobus [approval withdrawn])

**REVIEW DATE:** 07/31/2024

---

### OVERVIEW

Amifampridine, a broad spectrum potassium channel blocker, is indicated for the **treatment of Lambert-Eaton myasthenic syndrome (LEMS)**.<sup>1,2</sup>

- Firdapse is indicated in **adults and pediatric patients ≥ 6 years of age**.<sup>1</sup>
- Ruzurgi was indicated in **patients 6 years to < 17 years of age** (prior to withdrawal of FDA approval).<sup>2</sup>

As of February 01, 2022, the FDA has withdrawn approval for Ruzurgi. Firdapse was approved by the FDA on November 28, 2018, for the treatment of LEMS in adults, with 7 years of orphan-drug exclusivity (ODE). On May 6, 2019, Ruzurgi was approved by the FDA for the treatment of LEMS in patients 6 to < 17 years of age. On June 12, 2019, Catalyst (manufacturer of Firdapse) brought suit against the FDA, challenging the FDA's approval of Ruzurgi stating that it violated the ODE for Firdapse. In 2022, the Court of Appeals for the Eleventh Circuit sided with Catalyst; therefore, the FDA had to withdraw approval for Ruzurgi. Due to the 7-year ODE for Firdapse, Ruzurgi may not be approved for marketing until ODE has expired on November 28, 2025.

### Disease Overview

LEMS is a rare autoimmune disorder affecting the connection between nerves and muscles and causing proximal muscle weakness, autonomic dysfunction, and areflexia.<sup>3,4</sup> The characteristic weakness is thought to be caused by antibodies generated against the P/Q-type voltage-gated calcium channels present on presynaptic nerve terminals and by diminished release of acetylcholine.<sup>4</sup> The diagnosis of LEMS is confirmed by electrodiagnostic studies, including repetitive nerve stimulation, or anti-P/Q-type voltage-gated calcium channels antibody testing.

### Clinical Efficacy

Firdapse was approved based on two pivotal trials.<sup>1,5</sup> One pivotal trial enrolled both amifampridine-naïve and treatment-experienced patients; patients were initially entered into an open-label run-in phase lasting 90 days.<sup>5</sup> During the open-label run-in phase, Firdapse was titrated for each individual patient to a dose that produced optimal neuromuscular benefit and tolerability in the opinion of the investigator. In order to continue in the study, treatment-naïve patients were required to have an improvement of at least three points in the quantitative myasthenia gravis score from the initial evaluation. For its pediatric indication, use is supported by evidence from studies of Firdapse in adults with LEMS, pharmacokinetic data in adults, pharmacokinetic modeling and simulation to identify the dosing regimen in pediatric patients, and safety data from pediatric patients ≥ 6 years of age.

### Safety

Firdapse and Ruzurgi are contraindicated in patients with a history of seizures.<sup>1,2</sup> There is also a Warning/Precaution in the prescribing information for these medications because seizures have been observed in patients with and without a history of seizures taking amifampridine at the recommended doses.

07/31/2024

© 2024. All Rights Reserved.

This document is confidential and proprietary. Unauthorized use and distribution are prohibited.

Many of these patients were taking medications or had comorbidities that may have lowered their seizure threshold. Seizures may be dose-dependent.

### POLICY STATEMENT

Prior Authorization is recommended for prescription benefit coverage of amifampridine. All approvals are provided for the duration noted below. In cases where the approval is authorized in months, 1 month is equal to 30 days. Because of the specialized skills required for evaluation and diagnosis of patients treated with amifampridine as well as the monitoring required for adverse events and long-term efficacy, initial approval requires amifampridine to be prescribed by or in consultation with a physician who specializes in the condition being treated.

**Automation:** None.

### RECOMMENDED AUTHORIZATION CRITERIA

Coverage of amifampridine is recommended in those who meet the following criteria:

#### FDA-Approved Indications

1. **Lambert-Eaton Myasthenic Syndrome (LEMS).** Approve amifampridine for the duration noted if the patient meets ONE of the following (A or B):
  - A) Initial therapy. Approve amifampridine for 3 months if the patient meets ALL of the following (i, ii, iii, and iv):
    - i. Patient is  $\geq 6$  years of age; AND
    - ii. Patient has confirmed LEMS based on at least ONE of the following, according to the prescriber:
      - a) Electrodiagnostic study (e.g., repetitive nerve stimulation); OR
      - b) Anti-P/Q-type voltage-gated calcium channels antibody testing; AND
    - iii. Patient does not have a of seizures; AND
    - iv. Amifampridine is being prescribed by or in consultation with a neurologist or a neuromuscular specialist; OR
  - B) Patient is Currently Receiving amifampridine. Approve amifampridine for 1 year if the patient is continuing to derive benefit from amifampridine, according to the prescriber.

Note: Examples of continued benefit include improved muscle strength and improvements in mobility.

### CONDITIONS NOT RECOMMENDED FOR APPROVAL

Coverage of amifampridine is not recommended in the following situations:

1. Coverage is not recommended for circumstances not listed in the Recommended Authorization Criteria. Criteria will be updated as new published data are available.

### REFERENCES

1. Firdapse® tablets [prescribing information]. Coral Gables, FL: Catalyst; May 2024.
2. Ruzurgi® tablets [prescribing information]. Princeton, NJ: Jacobus; April 2020.
3. FDA news release. FDA approves first treatment for children with Lambert-Eaton myasthenic syndrome, a rare autoimmune disorder. Issued on: May 6, 2019. Available at: <https://www.fda.gov/news-events/press-announcements/fda-approves-first-treatment-children-lambert-eaton-myasthenic-syndrome-rare-autoimmune-disorder>. Accessed on July 29, 2024.

4. Kesner VG, Oh SJ, Dimachkie MM, et al. Lambert-Eaton Myasthenic Syndrome. *Neurol Clin.* 2018;36(2):379-394.
5. Oh S, Shcherbakova N, Kostera-Pruszczyk A, et al. Amifampridine phosphate (Firdapse®) is effective and safe in a phase 3 clinical trial in LEMS. *Muscle Nerve.* 2016;53(5):717-725.