

SmartPA Criteria Proposal

Drug/Drug Class:	Lambert-Eaton Myasthenic Syndrome (LEMS) Clinical Edit
First Implementation Date:	January 30, 2020
Proposed Date:	October 17, 2023
Prepared for:	MO HealthNet
Prepared by:	MO HealthNet/Conduent
Criteria Status:	<input type="checkbox"/> Existing Criteria <input checked="" type="checkbox"/> Revision of Existing Criteria <input type="checkbox"/> New Criteria

Executive Summary

Purpose: Ensure appropriate utilization and control of agents for Lambert-Eaton myasthenic syndrome (LEMS)

Why Issue Selected: Lambert-Eaton myasthenic syndrome (LEMS) is a rare autoimmune disorder that affects the connection between nerves and muscles and causes weakness and other symptoms in affected patients. Current treatment strategies for LEMS include initial therapy to increase the amount of acetylcholine available at the post-synaptic membrane with agents such as pyridostigmine, amifampridine, and guanidine; since pyridostigmine is readily available and well-tolerated, it is usually the first step in therapy. Amifampridine is a broad spectrum potassium channel blocker; the exact mechanism in which it exerts its therapeutic effect in LEMS is unknown. On November 28, 2018, Firdapse® (amifampridine) was FDA approved for the treatment of LEMS in adults. It was the first FDA-approved drug for treatment of LEMS in adults, which comprise the majority of LEMS patients.

Due to high cost and specific approved indication, MO HealthNet will impose clinical criteria to ensure appropriate utilization of agents for LEMS Disease.

Program-Specific Information:	Date Range FFS 7-1-22 to 6-30-23			
	Drug	Claims	Cost per tab	Cost per month (based on 30mg/day)
	FIRDAPSE 10 MG TABLET	0	\$218.66 MAC	\$19,679.40 MAC

Type of Criteria: ☐ Increased risk of ADE ☐ Preferred Drug List
☒ Appropriate Indications ☒ Clinical Edit

Data Sources: ☐ Only Administrative Databases ☒ Databases + Prescriber-Supplied

Setting & Population

- Drug class for review: agents for the treatment of Lambert-Eaton myasthenic syndrome (LEMS)
- Age range: All appropriate MO HealthNet participants aged 6 years or older

Approval Criteria

- Participant aged 6 years or older **AND**
- Diagnosis of LEMS consistent with 1 of the following:
 - Repetitive Nerve Stimulation (RNS) showing reproducible post-exercise increase in compound muscle action potential (CMAP) amplitude of at least 60% compared with pre-exercise baseline value or a similar increment on high-frequency repetitive nerve stimulation without exercise **OR**
 - Positive anti-P/Q type voltage-gated calcium channel antibody test **AND**
- Documentation of clinical symptoms suggestive of LEMS (proximal weakness affecting legs, eyes, face, throat) **AND**
- For first claim only:
 - Documented trial of pyridostigmine defined as 15 days of therapy in the past 30 days **AND**
 - Documentation of N-acetyltransferase 2 (NAT2) testing prior to initiation of therapy for dose determination **AND**
- Renewal Criteria:
 - Initial approval of prior authorization is 3 months
 - Renewal of prior authorization may be up to 12 months following documentation of the following:
 - All approval criteria listed above
 - Lack of ADE/ADR to therapy
 - Documentation of clinical benefit of therapy (less than expected decline in functional ability and/or symptoms of disease)

Denial Criteria

- Therapy will be denied if all approval criteria are not met
- Documented history of a seizure disorder

Required Documentation

Laboratory Results:
MedWatch Form:

X

Progress Notes:
Other:

X

Disposition of Edit

Denial: Exception code "0682" (Clinical Edit)
Ryle Type: CE

Default Approval Period

3 months

References

- FIRDAPSE® (amifampridine) [package insert]. Coral Gables, FL: Catalyst Pharmaceuticals, Inc; May 2023.
- Muscular Dystrophy Association. Lambert-Eaton Myasthenic Syndrome (LEMS). <https://www.mda.org/disease/lambert-eaton-myasthenic-syndrome>. Accessed August 28, 2023
- IPD Analytics. New Drug Approval: Firdapse (amifampridine). December 2018.
- IPD Analytics. Patients with LEMS Will Transition to Catalyst's Firdapse as Ruzurgi Approval Overturned. Trending Industry & Strategy Topics. February 7, 2022.

SmartPA Clinical Proposal Form

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