

## Clinical Policy: Tofersen (Qalsody)

Reference Number: LA.PHAR.591

Effective Date: 10.25.23

Last Review Date: 12.11.24

Line of Business: Medicaid

[Coding Implications](#)

[Revision Log](#)

See [Important Reminder](#) at the end of this policy for important regulatory and legal information.

**\*\*Please note: This policy is for medical benefit\*\***

### Description

Tofersen (Qalsody™) is an antisense oligonucleotide.

### FDA Approved Indication(s)

Qalsody is indicated for the treatment of amyotrophic lateral sclerosis (ALS) in adults who have a mutation in the superoxide dismutase 1 (*SOD1*) gene.

This indication is approved under accelerated approval based on reduction in plasma neurofilament light chain observed in patients treated with Qalsody. Continued approval for this indication may be contingent upon verification of clinical benefit in confirmatory trial(s).

### Policy/Criteria

*Provider must submit documentation (such as office chart notes, lab results or other clinical information) supporting that member has met all approval criteria.*

It is the policy of Louisiana Healthcare Connections® that Qalsody is **medically necessary** when the following criteria are met:

#### I. Initial Approval Criteria

##### A. Amyotrophic Lateral Sclerosis (must meet all):

1. Diagnosis of ALS with both of the following (a and b):
  - a. Muscle weakness attributed to ALS;
  - b. Documentation of *SOD1* mutation;
2. Prescribed by or in consultation with a neurologist;
3. Age  $\geq$  18 years;
4. Percent predicted slowed vital capacity (SVC)  $\geq$  50%;
5. Prescribed concurrently with riluzole (at up to maximally indicated doses), unless contraindicated or clinically significant adverse effects are experienced;
6. Member does not have presence of tracheostomy or permanent ventilation;
7. Dose does not exceed 100 mg (1 vial) on days 1, 15, and 29, followed by maintenance dose of 100 mg (1 vial) every 28 days.

**Approval duration: 6 months**

**B. Other diagnoses/indications** (must meet 1 or 2):

1. If this drug has recently (within the last 6 months) undergone a label change (e.g., newly approved indication, age expansion, new dosing regimen) that is not yet reflected in this policy, refer to LA.PMN.255
2. If the requested use (e.g., diagnosis, age, dosing regimen) is NOT specifically listed under section III (Diagnoses/Indications for which coverage is NOT authorized) AND criterion 1 above does not apply, refer to the off-label use policy for the relevant line of business: LA.PMN.53 for Medicaid.

**II. Continued Therapy**

**A. Amyotrophic Lateral Sclerosis** (must meet all):

- a. Currently receiving medication via Louisiana Healthcare Connections benefit or member has previously met initial approval criteria;
1. Member is responding positively to therapy (e.g., no tracheostomy or permanent ventilation);
  2. Prescribed concurrently with riluzole (at up to maximally indicated doses), unless contraindicated or clinically significant adverse effects are experienced;
  3. If request is for a dose increase, new dose does not exceed 100 mg (1 vial) every 28 days.

**Approval duration:** 12 months

**B. Other diagnoses/indications** (must meet 1 or 2):

1. If this drug has recently (within the last 6 months) undergone a label change (e.g., newly approved indication, age expansion, new dosing regimen) that is not yet reflected in this policy, refer to LA.PMN.255
2. If the requested use (e.g., diagnosis, age, dosing regimen) is NOT specifically listed under section III (Diagnoses/Indications for which coverage is NOT authorized) AND criterion 1 above does not apply, refer to the off-label use policy for the relevant line of business: LA.PMN.53 for Medicaid.

**III. Diagnoses/Indications for which coverage is NOT authorized:**

- A. Non-FDA approved indications, which are not addressed in this policy, unless there is sufficient documentation of efficacy and safety according to the off label use policies – LA.PMN.53 for Medicaid or evidence of coverage documents.

**IV. Appendices/General Information**

*Appendix A: Abbreviation/Acronym Key*

ALS: amyotrophic lateral sclerosis

FDA: Food and Drug Administration

LMN: lower motor neuron

*SOD1*: superoxide dismutase 1

*SVC*: slowed vital capacity

*UMN*: upper motor neuron

*Appendix B: Therapeutic Alternatives*

Not applicable

*Appendix C: Contraindications/Boxed Warnings*

None reported

*Appendix D: General Information*

- Revised El Escorial diagnostic criteria for ALS requires the presence of:
  1. Signs of lower motor neuron (LMN) degeneration by clinical, electrophysiological or neuropathologic examination;
  2. Signs of upper motor neuron (UMN) degeneration by clinical examination; and
  3. Progressive spread of signs within a region or to other regions, together with the absence of:
    - a. Electrophysiological evidence of other disease processes that might explain the signs of LMN and/or UMN degenerations; and
    - b. Neuroimaging evidence of other disease processes that might explain the observed clinical and electrophysiological signs.
- Gold Coast consensus diagnostic criteria for ALS requires the presence of:
  1. Progressive motor impairment documented by history or repeated clinical assessment, preceded by normal motor function; and
  2. Presence of upper and lower motor neuron dysfunction in at least 1 body region, (with upper and lower motor neuron dysfunction noted in the same body region if only one body region is involved) or lower motor neuron dysfunction in at least 2 body regions, and
  3. Investigations excluding other disease processes.

*Appendix E: Riluzole Co-administration*

Guidelines support the co-administration of riluzole in ALS:

- The 2009 American Academy of Neurology ALS guideline for the care of the patient with ALS (reaffirmed January 2023) recommends that riluzole should be offered to slow disease progression (Level A).
- The 2020 Canadian best practice recommendations for the management of ALS state the following: riluzole has demonstrated efficacy in improving survival in ALS (level A), there is evidence that riluzole prolongs survival by a median duration of 3 months (level A), and riluzole should be started soon after the diagnosis of ALS (expert consensus).
- The 2024 European Academy of Neurology guideline on the management of ALS recommends offering lifelong riluzole to all people with ALS at diagnosis (strong recommendation in favor).
- Additionally, approximately 62% of patients in the phase 3 VALOR trial were receiving concomitant riluzole.

**V. Dosage and Administration**

Indication	Dosing Regimen	Maximum Dose
<i>SOD1</i> ALS	Initiate recommended dose of 100 mg with 3 loading doses administered intrathecally at 14-day intervals. Maintenance dose of 100 mg should be administered intrathecally once every 28 days thereafter.	100 mg/dose/day

**VI. Product Availability**

Single-dose vial for injection: 100 mg/mL

## VII. References

1. Qalsody Prescribing Information. Cambridge, MA: Biogen; April 2023. Available at: <https://www.biogen.com/us/pdfs/qalsody-prescribing-information.pdf>. Accessed July 19, 2024.
2. Brooks BR, Miller RG, Swash M, et al. El Escorial revisited: revised criteria for the diagnosis of amyotrophic lateral sclerosis. *Amyotroph Lateral Scler Other Motor Neuron Disord*. 2000 Dec;1(5):293-9.
3. Shefner JM, Al-Chalabi A, Baker MR, et al. A proposal for new diagnostic criteria for ALS. *Clin Neurophysiol*. 2020;131(8):1975-1978.
4. Miller RG, Jackson CE, Kasarskis EJ, et al. Practice parameter update: the care of the patient with amyotrophic lateral sclerosis: drug, nutritional, and respiratory therapies (an evidence-based review): report of the Quality Standards Subcommittee of the American Academy of Neurology. *Neurology*. 2009 Oct 13;73(15):1218-26.
5. Shoesmith C, Abrahao A, Benstead T, et al. Canadian best practice recommendations for the management of amyotrophic lateral sclerosis. *CMAJ*. 2020 Nov;192(46):E1453-E1468.
6. Van Damme P, Al-Chalabi A, Andersen PM, et al. European Academy of Neurology (EAN) guideline on the management of amyotrophic lateral sclerosis in collaboration with European Reference Network for Neuromuscular Diseases (ERN EURO-NMD). *Eur J Neurol*. 2024;31(6):e16264.
7. Miller T, Cudkowicz M, Shaw PJ, et al. Phase 1-2 Trial of Antisense Oligonucleotide Tofersen for *SOD1* ALS. *N Engl J Med*. 2020;383(2):109-119.

## Coding Implications

Codes referenced in this clinical policy are for informational purposes only. Inclusion or exclusion of any codes does not guarantee coverage. Providers should reference the most up-to-date sources of professional coding guidance prior to the submission of claims for reimbursement of covered services.

HCPSC Codes	Description
J1304	Injection, tofersen, 1 mg

Reviews, Revisions, and Approvals	Date	LDH Approval Date
Policy created	05.01.23	
Criteria updated per approved FDA labeling: updated SVC $\geq$ 50% to reflect SVC eligibility criteria from VALOR part C trial, added no tracheostomy or permanent assisted ventilation for initial approval criteria and positive response continuation criteria, and updated maximum dose criteria; references reviewed and updated.	07.24.23	09.25.23
Annual Review; Added HCPSC code [J1304].	04.28.24	05.10.24
No significant changes; references reviewed and updated.	12.11.24	

## Important Reminder

This clinical policy has been developed by appropriately experienced and licensed health care professionals based on a review and consideration of currently available generally accepted

standards of medical practice; peer-reviewed medical literature; government agency/program approval status; evidence-based guidelines and positions of leading national health professional organizations; views of physicians practicing in relevant clinical areas affected by this clinical policy; and other available clinical information. LHCC makes no representations and accepts no liability with respect to the content of any external information used or relied upon in developing this clinical policy. This clinical policy is consistent with standards of medical practice current at the time that this clinical policy was approved.

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