

**Tiglutik (riluzole) Oral Suspension**  
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**Effective 01/01/2026**

<b>Plan</b>	<input type="checkbox"/> MassHealth UPPL <input checked="" type="checkbox"/> Commercial/Exchange	<b>Program Type</b>	<input checked="" type="checkbox"/> Prior Authorization <input type="checkbox"/> Quantity Limit <input type="checkbox"/> Step Therapy
<b>Benefit</b>	<input checked="" type="checkbox"/> Pharmacy Benefit <input type="checkbox"/> Medical Benefit		
<b>Specialty Limitations</b>	N/A		
<b>Contact Information</b>	<b>Medical Benefit</b> <b>Pharmacy Benefit</b>	Phone: 833-895-2611 Phone: 800-711-4555	Fax: 888-656-6671 Fax: 844-403-1029
<b>Exceptions</b>	N/A		

### Overview

Tiglutik (riluzole) is indicated for the treatment of amyotrophic lateral sclerosis (ALS)

### Coverage Guidelines

Authorization may be granted for members new to the plan within the past 90 days who are currently receiving treatment with the requested medication excluding when the product is obtained as samples or via manufacturer's patient assistance programs

#### OR

Authorization may be granted for members when all the following criteria have been met:

1. Member is diagnosed with amyotrophic lateral sclerosis (ALS)
2. Member is at least 18 years of age
3. Documentation member is unable to swallow oral riluzole tablets

### Limitations

1. Initial and reauthorizations will be granted for 24 months.
2. Requests for reauthorization will be reviewed against initial criteria.

### References

1. Bensimon G, Lacomblez L, Meininger V. A controlled trial of riluzole in amyotrophic lateral sclerosis. ALS/Riluzole Study Group. *N Engl J Med* 1994; 330:585.
2. Kennel P, Revah F, Bohme GA, et al. Riluzole prolongs survival and delays muscle strength deterioration in mice with progressive motor neuronopathy (pmn). *J Neurol Sci* 2000; 180:55.
3. Miller RG, Mitchell JD, Moore DH. Riluzole for amyotrophic lateral sclerosis (ALS)/motor neuron disease (MND). *Cochrane Database Syst Rev* 2012; CD001447.
4. Tiglutik (riluzole) [prescribing information]. Paoli, PA: EDW Pharma, Inc; September 2025.

### Review History

02/20/19 – Reviewed

07/21/2021 – Reviewed July P&T; update document to include started and stabilized statement, added new formulation Exserval; references updated

09/21/2022 - Reviewed at Sept P&T; references updated; no clinical changes.

10/08/2025 – Reviewed and updated at September P&T. Removed Exservan from the policy due to product discontinuation and updated policy title to reflect this change. Updated language for members who are new to the Plan. Effective 1/1/2026.

