

**PHARMACY PRIOR AUTHORIZATION REQUEST FORM**
**MYASTHENIA GRAVIS**

Rystiggo®, Soliris®, Ultomiris®, Vyvgart®, Vyvgart® Hytrulo, Zilbrysq®

**For authorization, please answer each question and fax this form PLUS chart notes back to the U of U Health Plans Prior Authorization Department at 801-213-1547.**

**Failure to submit clinical documentation to support this request will result in a dismissal of the request.**

If you have medical pharmacy prior authorization questions, please call for assistance: Individual Exchange: 833-981-0214, Commercial Groups: 833-981-0213, MHC: 844-262-1500

Disclaimer: Prior authorization request forms are subject to change in accordance with Federal and State notice requirements.

Date:	Member Name:	ID#:
DOB:	Gender:	Physician:
Office Phone:	Office Fax:	Office Contact:
Height/Weight:	HCPCS Code:	

**Member must try formulary preferred drugs before a request for a non-preferred drug may be considered. If treatment with preferred products has not been successful, you must submit which preferred products have been tried, dates of treatment, and reason for failure. Reasons for failure must meet the Health Plan medical necessity criteria.**

Preferred/Non-preferred

1. 1<sup>st</sup> line preferred agents:
  - A. Rystiggo® (rozanolixizumab-noli) subcutaneous infusion, Vyvgart® (efgartigimod alfa-fcab) intravenous infusion
2. 2<sup>nd</sup> line non-preferred agents; after trial and failure of the preferred first-line agents:
  - A. Ultomiris® (ravulizumab) intravenous infusion
3. Excluded/Not covered unless failure or contraindication to all other agents:
  - A. Soliris® (eculizumab) intravenous infusion; Vyvgart® Hytrulo (efgartigimod alfa/hyaluronidase) subcutaneous infusion, Zilbrysq® (zilucoplan)

Product being requested: \_\_\_\_\_

Dosing/Frequency: \_\_\_\_\_

**If the request is for reauthorization, proceed to reauthorization section**

Questions	Yes	No	Comments/Notes
<b>MYASTHENIA GRAVIS (gMG)</b>			
1. Is the request being made by or in consultation with a neurologist or other specialist in the treatment of gMG?	<input type="checkbox"/>	<input type="checkbox"/>	
2. Does the member have a diagnosis of gMG?	<input type="checkbox"/>	<input type="checkbox"/>	
3. Does the member have a positive serologic test for anti-acetylcholine receptor (anti-AchR) antibodies?	<input type="checkbox"/>	<input type="checkbox"/>	<b>Please provide documentation</b>
4. If the request is for Rystiggo®, does the member have a positive serologic test for anti-acetylcholine receptor (anti-AchR) antibodies OR anti-muscle-specific kinase (anti-MuSK) antibodies?	<input type="checkbox"/>	<input type="checkbox"/>	<b>Please provide documentation</b>
5. Has the member been diagnosed with class II to IV gMG according to the Myasthenia Gravis Foundation of America?	<input type="checkbox"/>	<input type="checkbox"/>	<b>Please provide documentation</b>

6. Has the member tried and failed pyridostigmine AND at least two immunosuppressive therapies (e.g. rituximab, methotrexate, mycophenolate mofetil, azathioprine, cyclosporine) for a total duration of at least 12 months?	<input type="checkbox"/>	<input type="checkbox"/>	<b>Please provide documentation</b>
7. Has the member tried and failed intravenous immunoglobulin (IVIG)?	<input type="checkbox"/>	<input type="checkbox"/>	<b>Please provide documentation</b>
8. Will the requested therapy be used in combination with IVIG or other biologic agents for gMG treatment?	<input type="checkbox"/>	<input type="checkbox"/>	
9. If the request is for Rystiggo®, is the member's Myasthenia Gravis Activities of Daily Living (MG-ADL) score ≥ 3?	<input type="checkbox"/>	<input type="checkbox"/>	<b>Please provide documentation</b>
10. If the request is for Vyvgart®, is the member's MG-ADL score ≥ 5?	<input type="checkbox"/>	<input type="checkbox"/>	<b>Please provide documentation</b>
11. If the request is for Soliris® or Ultomiris®, is the member's MG-ADL score ≥ 6?	<input type="checkbox"/>	<input type="checkbox"/>	<b>Please provide documentation</b>
12. If the request is for Soliris® or Ultomiris®, is the prescribing physician enrolled in Soliris® or Ultomiris® REMS program?	<input type="checkbox"/>	<input type="checkbox"/>	
<b>REAUTHORIZATION</b>			
1. Is the request for reauthorization of therapy?	<input type="checkbox"/>	<input type="checkbox"/>	
2. If the request is for reauthorization of Vyvgart® or Rystiggo®, has the member had a positive clinical response to treatment shown by a ≥ 2 points reduction in MG-ADL score?	<input type="checkbox"/>	<input type="checkbox"/>	<b>Please provide documentation</b>
3. If the request is for reauthorization of Soliris® or Ultomiris®, has the member had a positive clinical response to treatment shown by a ≥ 2 points reduction in MG-ADL score or a ≥ 3 points reduction in quantitative myasthenia gravis (QMG) score?	<input type="checkbox"/>	<input type="checkbox"/>	<b>Please provide documentation</b>
<b>What medications and/or treatment modalities have been tried in the past for this condition? Please document name of treatment, reason for failure, treatment dates, etc.</b>			
Additional information:			
Physician's Signature:			

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Policy: PHARM-M046  
 Origination Date: 08/03/2023  
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