BlueCross BlueShield Federal Employee Program.

(ravulizumab-cwvz) ULTOMIRIS
PRIOR APPROVAL REQUEST

Send completed form to:
FAX: 855-895-3504
FOR URGENT FAX: 844-244-0226

Additional information is required to process your claim for prescription drugs. Please complete the patient portion, and have the prescribing physician complete the physician portion and submit this completed form.

Patient Information (required)			**Provider Information (required)**		
Date:			Provider Name:		
Patient Name:			Specialty:	NPI:	
Date of Birth:	Sex: ❑Male ❑Female		Office Phone:	Office Fax:	
Street Address:			Office Street Address:		
City:	State:	Zip:	City:	State:	Zip:
Patient ID: **R**			Physician Signature:		
PHYSICIAN COMPLETES					

Ultomiris

(ravulizumab-cwvz)

****Check www.fepblue.org/formulary to confirm which medication is part of the patient's benefit**

NOTE: Form must be completed in its **entirety** for processing

Is this request for brand or generic? ❑Brand ❑Generic

1. Is the prescriber enrolled in the Ultomiris REMS program? ❑Yes ❑No
2. What is the patient's diagnosis?

❑Atypical Hemolytic Uremic Syndrome (aHUS)

a. Does the patient have Shiga toxin E. coli related hemolytic uremic syndrome (STEC-HUS)? ❑Yes ❑No

b. Will Ultomiris be used in combination with another Prior Authorization (PA) medication for atypical hemolytic uremic syndrome (e.g., Soliris (eculizumab))? ❑Yes* ❑No

If YES, specify the medication: _____

c. Has the patient been on Ultomiris continuously for the last **4 months**, excluding samples? ***Please select answer below:***

❑**NO** – this is **INITIATION** of therapy, please answer the following questions:

i. Does the patient have a documented baseline value for serum lactate dehydrogenase (LDH)? ❑Yes ❑No

ii. Has or will the patient be vaccinated against Neisseria meningitidis at least 2 weeks prior to initiating therapy? ❑Yes ❑No*

***If NO**, is urgent Ultomiris therapy indicated for this patient (e.g., the risks of delaying treatment with Ultomiris outweigh the risk of developing a meningococcal infection)? ❑Yes ❑No

❑**YES** – this is a PA renewal for **CONTINUATION** of therapy, please answer the following questions:

i. Has the patient had a decrease in serum lactate dehydrogenase (LDH) from pretreatment baseline? ❑Yes ❑No

ii. Has the patient experienced unacceptable toxicity while on Ultomiris therapy? ❑Yes ❑No

❑Paroxysmal Nocturnal Hemoglobinuria (PNH)

a. Will Ultomiris be used in combination with another Prior Authorization (PA) medication for paroxysmal nocturnal hemoglobinuria (e.g., Empaveli (pegcetacoplan), Soliris (eculizumab))? ❑Yes* ❑No

If YES, specify the medication: _____

b. Has the patient been on Ultomiris continuously for the last **4 months**, excluding samples? ***Please select answer below:***

❑**NO** – this is **INITIATION** of therapy, please answer the following questions:

i. Does the patient have a documented baseline value for serum lactate dehydrogenase (LDH)? ❑Yes ❑No

ii. Has or will the patient be vaccinated against Neisseria meningitidis at least 2 weeks prior to initiating therapy? ❑Yes ❑No*

***If NO**, is urgent Ultomiris therapy indicated for this patient (e.g., the risks of delaying treatment with Ultomiris outweigh the risk of developing a meningococcal infection)? ❑Yes ❑No

❑**YES** – this is a PA renewal for **CONTINUATION** of therapy, please answer the following questions:

i. Has the patient had a decrease in serum lactate dehydrogenase (LDH) from pretreatment baseline? ❑Yes ❑No

ii. Has the patient experienced unacceptable toxicity while on Ultomiris therapy? ❑Yes ❑No

PLEASE PROCEED TO PAGE 2 FOR ADDITIONAL DIAGNOSES

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PAGE 2 - PHYSICIAN COMPLETES

Patient Name: _____ DOB: _____ Patient ID: R _____

❑Generalized Myasthenia Gravis (gMG)

a. Will Ultomiris be used in combination with another Prior Authorization (PA) C5 complement inhibitor for generalized myasthenia gravis (e.g., Soliris (eculizumab))? ❑Yes***** ❑No

If YES, specify the medication: _____

b. Has the patient been on Ultomiris continuously for the last **4 months**, excluding samples? ***Please select answer below:***

❑ **NO** – this is **INITIATION** of therapy, please answer the following questions:

i. Does the patient have a positive serologic test for anti-AChR antibodies? ❑Yes ❑No

ii. What is the patient's MGFA (Myasthenia Gravis Foundation of America) clinical classification? ***Select answer below:***

❑Class I ❑Class II to IV ❑Class V ❑Unknown

iii. **If Class II to IV**, does the patient have a documented baseline *****MG-Activities of Daily Living (MG-ADL) total score greater than or equal to 6? ❑Yes ❑No

****MG-ADL*: http://c.peerview.com/inReview/programs/150204324/downloads/PVI_practiceaids_RMU.pdf**

iv. Has or will the patient be vaccinated against Neisseria meningitidis at least 2 weeks prior to initiating therapy? ❑Yes ❑No*****

If NO, is urgent Ultomiris therapy indicated for this patient (e.g., the risks of delaying treatment with Ultomiris outweigh the risk of developing a meningococcal infection)? ❑Yes ❑No

v. Does the patient have an intolerance or contraindication or have they had an inadequate treatment response to an acetylcholinesterase inhibitor? ❑Yes ❑No

vi. Does the patient have an intolerance or contraindication or have they had an inadequate treatment response to at least one immunosuppressive therapy either in combination or as monotherapy, such as: azathioprine, cyclosporine, mycophenolate mofetil, tacrolimus, methotrexate, or cyclophosphamide? ❑Yes ❑No

❑ **YES** – this is a PA renewal for **CONTINUATION** of therapy, please answer the following questions:

i. Is there a documented decrease of the *****MG-Activities of Daily Living (MG-ADL) total score from baseline of greater than or equal to 2 points? ❑Yes ❑No

****MG-ADL*: http://c.peerview.com/inReview/programs/150204324/downloads/PVI_practiceaids_RMU.pdf**

ii. Has the patient experienced unacceptable toxicity while on Ultomiris therapy? ❑Yes ❑No

❑None of the above