



Ultomiris® (ravulizumab-cwvz) Medication Precertification Request

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(All fields must be completed and legible for precertification review.)

Aetna Precertification Notification

Phone: 1-866-752-7021

FAX: 1-888-267-3277

For Medicare Advantage Part B:

Phone: 1-866-503-0857

FAX: 1-844-268-7263

Please indicate: Start of treatment: Start date ____/____/____
 Continuation of therapy, Date of last treatment ____/____/____

Precertification Requested By: _____ Phone: _____ Fax: _____

A. PATIENT INFORMATION

First Name:		Last Name:		DOB:	
Address:			City:		State: ZIP:
Home Phone:		Work Phone:		Cell Phone:	
Patient Current Weight: ____ lbs or ____ kgs		Patient Height: ____ inches or ____ cms		Allergies:	

B. INSURANCE INFORMATION

Aetna Member ID #: _____		Does patient have other coverage? <input type="checkbox"/> Yes <input type="checkbox"/> No	
Group #: _____		If yes, provide ID#: _____ Carrier Name: _____	
Insured: _____		Insured: _____	
Medicare: <input type="checkbox"/> Yes <input type="checkbox"/> No If yes, provide ID #: _____		Medicaid: <input type="checkbox"/> Yes <input type="checkbox"/> No If yes, provide ID #: _____	

C. PRESCRIBER INFORMATION

First Name:		Last Name:		(Check One): <input type="checkbox"/> M.D. <input type="checkbox"/> D.O. <input type="checkbox"/> N.P. <input type="checkbox"/> P.A.	
Address:			City:		State: ZIP:
Phone:		Fax:		St Lic #: NPI #: DEA #: UPIN:	
Provider Email:			Office Contact Name:		Phone:

Specialty (Check one): Hematologist Other: _____

D. DISPENSING PROVIDER/ADMINISTRATION INFORMATION

Place of Administration:		Dispensing Provider/Pharmacy: (Patient selected choice)	
<input type="checkbox"/> Self-administered <input type="checkbox"/> Physician's Office		<input type="checkbox"/> Physician's Office <input type="checkbox"/> Retail Pharmacy	
<input type="checkbox"/> Outpatient Infusion Center Phone: _____		<input type="checkbox"/> Specialty Pharmacy <input type="checkbox"/> Other: _____	
Center Name: _____		Name: _____	
<input type="checkbox"/> Home Infusion Center Phone: _____		Address: _____	
Agency Name: _____		Phone: _____ Fax: _____	
<input type="checkbox"/> Administration code(s) (CPT): _____		TIN: _____ PIN: _____	
Address: _____			

E. PRODUCT INFORMATION

Request is for Ultomiris (ravulizumab-cwvz):

Loading Dose: _____ Frequency: _____

Maintenance Dose: _____ Frequency: _____

F. DIAGNOSIS INFORMATION - Please indicate primary ICD code and specify any other where applicable.

Primary ICD Code: _____ Secondary ICD Code : _____ Other ICD Code: _____

G. CLINICAL INFORMATION - Required clinical information must be completed in its entirety for all precertification requests.

For All Requests (clinical documentation required):

Yes No Is this infusion request in an outpatient hospital setting?

Yes No Has the patient experienced an adverse event with the requested product that has not responded to conventional interventions (e.g., acetaminophen, steroids, diphenhydramine, fluids, other pre-medications or slowing of infusion rate) or a severe adverse event (anaphylaxis, anaphylactoid reactions, myocardial infarction, thromboembolism, or seizures) during or immediately after an infusion?

Yes No Does the patient have severe venous access issues that require the use of special interventions only available in the outpatient hospital setting?

Yes No Does the patient have significant behavioral issues and/or physical or cognitive impairment that would impact the safety of the infusion therapy AND the patient does not have access to a caregiver?

Yes No Is the patient medically unstable which may include respiratory, cardiovascular, or renal conditions that may limit the member's ability to tolerate a large volume or load or predispose the member to a severe adverse event that cannot be managed in an alternate setting without appropriate medical personnel and equipment?

Yes No Please provide a description of the condition: Cardiopulmonary: _____
 Respiratory: _____
 Renal: _____
 Other: _____

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Patient First Name	Patient Last Name	Patient Phone	Patient DOB
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G. CLINICAL INFORMATION (continued) – Required clinical information must be completed in its entirety for all precertification requests.

For Initiation Requests (clinical documentation required):

- Yes No Is the patient switching from eculizumab to the requested drug?
 Yes No Will the loading dose of the requested drug be administered 2 weeks after the last eculizumab infusion?

Paroxysmal nocturnal hemoglobinuria (PNH)

- Yes No Has the diagnosis been confirmed by flow cytometry results?
 Yes No Does the patient have a deficiency of glycosylphosphatidylinositol (GPI)-anchored proteins?
 How was the diagnosis established? Quantification of PNH cells
 Less than 5% Greater than or equal to 5%
 Quantification of GPI-anchored protein deficient poly-morphonuclear cells
 Less than 51% Greater than or equal to 51%
 None of the above

Atypical hemolytic uremic syndrome

- Yes No Is the disease caused by Shiga toxin?
 Yes No Do tests confirm the absence of Shiga toxin?
 What is the ADAMTS13 level? Less than or equal to 5% Greater than 5%

For All Continuation Requests (clinical documentation required):

- Yes No Is there evidence of unacceptable toxicity or disease progression while on the current regimen?
 Yes No Has the patient demonstrated a positive response to therapy (e.g., normalization of lactate dehydrogenase (LDH) levels, platelet counts or improvement in hemoglobin levels)?

H. ACKNOWLEDGEMENT

Request Completed By (Signature Required): _____ **Date:** ____ / ____ / ____

Any person who knowingly files a request for authorization of coverage of a medical procedure or service with the intent to injure, defraud or deceive any insurance company by providing materially false information or conceals material information for the purpose of misleading, commits a fraudulent insurance act, which is a crime and subjects such person to criminal and civil penalties.

The plan may request additional information or clarification, if needed, to evaluate requests.