Ultomiris

Prior Authorization Request

CVS Caremark administers the prescription benefit plan for the member identified. This patient's benefit plan requires prior authorization for certain medications in order for the drug to be covered. To make an appropriate determination, providing the most accurate diagnosis for the use of the prescribed medication is necessary. **Please respond below and fax this form to CVS Caremark toll-free at 1-855-330-1720**. If you have questions regarding the prior authorization, please contact CVS Caremark at **1-888-877-0518.** For inquiries or questions related to the patient's eligibility, drug copay or medication delivery; please contact the Specialty Customer Care Team: CaremarkConnect® 1-800-237-2767.

The recipient of this fax may make a request to opt-out of receiving telemarketing fax transmissions from CVS Caremark. There are numerous ways you may opt-out: The recipient may call the toll-free number at 877-265-2711, at any time, 24 hours a day/7 days a week. The recipient may also send an opt-out request via email to do_not_call@cvscaremark.com. An opt out request is only valid if it (1) identifies the number to which the request relates, and (2) if the person/entity making the request does not, subsequent to the request, provide express invitation or permission to CVS Caremark to send facsimile advertisements to such person/entity at that particular number. CVS Caremark is required by law to honor an opt-out request within thirty days of receipt.

Patient's Name: _____ **Date**: _____
Patient's ID: _____ **Patient's Date of Birth:** _____
Physician's Name: _____
Specialty: _____ **NPI#:** _____
Physician Office Telephone: _____ **Physician Office Fax:** _____

Referring Provider Info: ❑ **Same as Requesting Provider**
Name: _____ **NPI#:** _____
Fax: _____ **Phone:** _____

Rendering Provider Info: ❑ **Same as Referring Provider** ❑ **Same as Requesting Provider**
Name: _____ **NPI#:** _____
Fax: _____ **Phone:** _____

Approvals may be subject to dosing limits in accordance with FDA-approved labeling, accepted compendia, and/or evidence-based practice guidelines.

Required Demographic Information:

Patient Weight: _____*kg*

*Patient Height:*_____*cm*

Please indicate the place of service for the requested drug:
❑ *Ambulatory Surgical* ❑ *Home* ❑ *Inpatient Hospital* ❑ *Off Campus Outpatient Hospital*
❑ *On Campus Outpatient Hospital* ❑ *Office* ❑ *Pharmacy*

Send completed form to: Case Review Unit CVS Caremark Specialty Programs Fax: 1-855-330-1720

CVS Caremark Specialty Pharmacy ● 2211 Sanders Road NBT-6 ● Northbrook, IL 60062
Phone: 1-888-877-0518 ● Fax: 1-855-330-1720 ● www.caremark.com

Site of Service Questions (SOS):

A. Indicate the site of service requested:
- ❑ On Campus Outpatient Hospital
- ❑ Home infusion, *skip to Clinical Questions*
- ❑ Ambulatory surgical, *skip to Clinical Questions*
- ❑ Off Campus Outpatient Hospital
- ❑ Physician office, *skip to Clinical Questions*
- ❑ Pharmacy, *skip to Clinical Questions*
- ❑ Inpatient hospital, *skip to Clinical Questions*

B. Is the patient less than 21 years old or 65 years of age or older?
- ❑ Yes – less than 21 years old, *skip to Clinical Criteria Questions*
- ❑ Yes – age 65 years or older, *skip to Clinical Criteria Questions*
- ❑ No

C. Is this request to continue previously established treatment with the requested medication?
- ❑ Yes - This is a continuation of an existing treatment.
- ❑ No - This is a new therapy request (patient has not received requested medication in the last 6 months). *skip to Clinical Criteria Questions*

D. Has the patient experienced an adverse event with the requested product that has not responded to conventional interventions (eg acetaminophen, steroids, diphenhydramine, fluids, other pre- medications or slowing of the infusion rate) or a severe adverse event (anaphylaxis, anaphylactoid reactions, myocardial infarction, thromboembolism, or seizures) during or immediately after an infusion? ***ACTION REQUIRED: Attach supporting clinical documentation.*** ❑ Yes, *skip to Clinical Criteria Questions* ❑ No

E. 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? ***ACTION REQUIRED: Attach supporting clinical documentation.*** ❑ Yes, *skip to Clinical Criteria Questions* ❑ No

F. Does the patient have severe venous access issues that require the use of special interventions only available in the outpatient hospital setting? ***ACTION REQUIRED: Attach supporting clinical documentation.***
❑ Yes, *skip to Clinical Criteria Questions* ❑ No

G. 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? ***ACTION REQUIRED: Attach supporting clinical documentation.*** ❑ Yes ❑ No

Criteria Questions:

1. What is the diagnosis?
   - ❑ Paroxysmal nocturnal hemoglobinuria (PNH)
   - ❑ Atypical hemolytic uremic syndrome (aHUS)
   - ❑ Other _____
2. What is the ICD-10 code? _____

Complete the following section based on the patient's diagnosis, if applicable.

Section A: Paroxysmal Nocturnal Hemoglobinuria (PNH)

3. Does the patient have a deficiency of glycosylphosphatidylinositol (GPI)-anchored proteins? ***ACTION REQUIRED: If Yes, attach supporting chart note(s).*** ❑ Yes ❑ No
4. How was the diagnosis established?
   - ❑ Quantification of PNH cells
   - ❑ Quantification of GPI-anchored protein deficient poly-morphonuclear cells, *skip to #6*
   - ❑ None of the above
5. What was the percentage of PNH cells? _____ %
   If percentage of PNH cells is greater than or equal to 5%, skip to #7.
6. What was the percentage of GPI-anchored protein deficient poly-morphonuclear cells? _____ %
7. Has the diagnosis been confirmed by flow cytometry results? ***ACTION REQUIRED: If Yes, attach supporting chart note(s).*** ❑ Yes ❑ No
8. Is the patient currently receiving treatment with the requested medication?
   ❑ Yes ❑ No *If No, no further questions.*
9. Has the patient demonstrated a positive response to therapy (e.g., normalization of lactate dehydrogenase (LDH)

Section B: Atypical Hemolytic Uremic Syndrome (aHUS)

10. Is the disease caused by Shiga toxin? ❑ Yes ❑ No
11. Do tests confirm the absence of Shiga toxin? ❑ Yes ❑ No
12. What is the ADAMTS13 level? ***ACTION REQUIRED: Please attach documentation of ADAMTS13 level.***
    ❑ Yes ❑ No
13. Is the patient currently receiving treatment with the requested medication?
    ❑ Yes ❑ No *If No, no further questions.*
14. Has the patient demonstrated a positive response to therapy (e.g., normalization of lactate dehydrogenase (LDH) levels, platelet counts or improvement in hemoglobin levels)? ❑ Yes ❑ No

I attest that this information is accurate and true, and that documentation supporting this information is available for review if requested by CVS Caremark or the benefit plan sponsor.

X_____
Prescriber or Authorized Signature **Date (mm/dd/yy)**

Send completed form to: Case Review Unit CVS Caremark Specialty Programs Fax: 1-855-330-1720

CVS Caremark Specialty Pharmacy ● 2211 Sanders Road NBT-6 ● Northbrook, IL 60062
Phone: 1-888-877-0518 ● Fax: 1-855-330-1720 ● www.caremark.com