

## Prior Authorization Form



**Note:** Please provide as much information as possible on this form. Missing data may cause processing delays for requested authorization(s). Attach additional sheets to this form if necessary.

Please fax the completed PA form and any additional informational sheets to Nirvanahealth at the following fax number:  
+1(866) 871-8565

Patient Information		Prescriber Information	
Patient Name: _____		Prescriber Name: _____	
Health Plan Name: _____		Prescriber Address: _____	
Patient Insurance Id: _____		_____	
Patient Date of Birth: _____		Prescriber Phone: (     ) _____	
Patient Phone: _____		Prescriber Fax: (     ) _____	
		Prescriber Specialty: _____	
		Prescriber DEA: _____	
		Prescriber NPI: _____	

  

Medication & Medical Information	
Requested Drug(s) & Strength(s):	[ ] Empaveli 1,080 mg/20 mL subcutaneous solution
Requested Daily Quantity Limit – Amount:	
Requested Daily Quantity Limit – Days:	
Requested Quantity Limit Over Time – Amount:	
Requested Quantity Limit Over Time – Days:	
Requested Quantity Per Rx – Amount:	
Expected Length of Therapy:	
Directions:	
Diagnosis and Diagnosis Codes (ICD-10 Standard Codes):	
List drugs used previously to treat the same condition:	
Additional clinical information or history. Please include any relevant test results and/or medical record notes:	

## Questionnaire

Q1: I, as the provider or designated representative of the provider, certify and attest that the information provided is complete and accurate and that, upon request, I shall provide any information to RxAdvance that RxAdvance determines is reasonably necessary to verify my responses. (Check only one that apply)

[ ] Yes

[ ] No

Q2: Is the member currently treated with this medication? (Check only one that apply)

[ ] Yes (please list start date of therapy (month/day/year)) \_\_\_\_\_  
(\*Required)

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☐ No

Q3: What is the member's diagnosis? (Check only one that apply)

☐ Paroxysmal Nocturnal Hemoglobinuria (PNH)

☐ Other (please specify the member's diagnosis and provide clinical rationale for the request)  
\_\_\_\_\_\*Required)

Q4: Does the member have documentation supporting positive clinical response to therapy (e.g., improvement in hemoglobin level, hemoglobin stabilization, decrease in the number of red blood cell transfusions)? (Check only one that apply)

☐ Yes (please provide documentation(s) supporting the positive response of the therapy)  
\_\_\_\_\_\*Required)

☐ No (please provide medical justification for continuation of therapy)  
\_\_\_\_\_\*Required)

Q5: What is the member's diagnosis? (Check only one that apply)

☐ Paroxysmal Nocturnal Hemoglobinuria (PNH)

☐ Other (please specify the member's diagnosis and provide clinical rationale for the request)  
\_\_\_\_\_\*Required)

Q6: Has the member had an inadequate response, intolerance or experienced contraindication(s) to Ultomiris (ravulizumab)? (Check only one that apply)

☐ Yes (please specify corresponding contraindication(s) or intolerance(s) experienced and the start and end date(s) of therapy (month/year)) \_\_\_\_\_\*Required)

☐ No (please provide clinical rationale for the request) \_\_\_\_\_  
(\*Required)

**Attestation:** I attest the information provided is true and accurate to the best of my knowledge. I understand that the Health Plan, Insurer, Medical Group or its designated representatives may perform a routine audit and request the medical information necessary to verify the accuracy of the information reported on this form.

Signature of Prescriber or Authorized Representative:

Date:

Print Authorized Representative Name: