



PRIOR AUTHORIZATION REQUEST
ULTOMIRIS (RAVULIZUMAB)
SOLIRIS (ECULIZUMAB)

PLEASE READ THE INSTRUCTIONS ON THE LAST PAGE OF THIS FORM.

A PATIENT IDENTIFICATION – To be completed by the member.

Patient's last and first name		Relationship with member <input type="checkbox"/> Member <input type="checkbox"/> Spouse <input type="checkbox"/> Dependent child		Patient's date of birth YYYY MM DD	
Member's last and first name			Contract No.		Certificate No.
No., street, apt.			City		Province Postal code
Telephone Nos – Home:		Office:		Extension:	
Email:					

Since the response to this request includes confidential information, please indicate how you would like to be informed of the decision:

By mail (The response to your request will be sent to the address indicated in this section.) By fax:

Coordination of benefits: If the patient has coverage under a private insurance plan or is enrolled in a provincial drug insurance plan, please submit the request to this plan first. Then send us a copy of the decision notice and this form filled out by the physician, so we can analyze the request.

PRIVATE PLAN

Does the patient have drug coverage under a private insurance plan?
 Yes – Please provide a copy of the notice of approval or refusal. → **Copy attached to this form.**
 Specify: Name of the insurer: _____ Contract No.: _____ Certificate No.: _____
 No

PROVINCIAL PLAN

Has a request for reimbursement been submitted under your provincial plan?
 Yes – Please provide a copy of the notice of approval or refusal. → **Copy attached to this form.**
 No – Please explain: _____

PATIENT SUPPORT PROGRAM

Is the patient enrolled in a patient support program? **Yes** **No**
 If so – Program name: _____
 Contact person: _____ Telephone No.: _____ Extension: _____

B1 DECLARATION AND AUTHORIZATION FOR THE COLLECTION AND COMMUNICATION OF PERSONAL INFORMATION

All the information I have provided on the claim form is accurate and complete. I authorize Desjardins Financial Security Life Assurance Company, hereinafter Desjardins Insurance, strictly for the purposes of managing my file and settling this claim to: (a) collect from any person or legal entity, or from any public or parapublic organization, only the information deemed necessary to manage my file. The non-exhaustive list of sources from which information may be collected includes healthcare professionals or facilities, and insurance companies; (b) communicate to the said persons or organizations only the personal information about me that is deemed necessary for the purposes of my file; (c) when necessary use the personal information it may have about me in existing files that are now closed. This authorization is also valid for the collection, use and communication of personal information concerning my dependents, insofar as applicable to the claim. A photocopy of this authorization is as valid as the original.

➤ **Signature of member:** _____ **Date:** _____

Last name and first name of parent/legal guardian (if applicable): _____

Signature of patient or parent/legal guardian (if applicable): _____ **Date:** _____

B2 CONSENT TO THE COMMUNICATION OF PERSONAL INFORMATION TO A THIRD PARTY

To help us process your claim more efficiently, do you authorize Desjardins Insurance to inform the patient support program and the attending physician or the attending physician's medical team of the reasons for the decision on your prior authorization request?

Yes No

➤ **Signature of member:** _____ **Date:** _____

Last name and first name of parent/legal guardian (if applicable): _____

Signature of patient or parent/legal guardian (if applicable): _____ **Date:** _____

C ATTENDING PHYSICIAN SECTION – To be completed by the attending physician.

Physician's last and first name (PLEASE PRINT)		License No.	Specialty		
No., street, suite		City		Province	Postal code
Telephone No.:			Fax No.:		

➤ **Signature of physician:** _____ **Date:** _____

Drug name	Formulation	Strength	Dosage	Patient's weight	Scheduled duration of treatment
-----------	-------------	----------	--------	------------------	---------------------------------

Where is the drug administered? Home Physician's office Private clinic Hospital – Inpatient Hospital – Outpatient
 Other (please specify): _____

- Make sure to fill out all sections so we can process the request faster. If any information is missing, we will send the form back to the member.
- In order to consider any diagnosis not mentioned on this form, we need supporting documents (clinical practice guidelines, clinical studies, etc.) that justify the drug's use in the given context.

DIAGNOSIS

- Paroxysmal nocturnal hemoglobinuria (PNH) Atypical hemolytic uremic syndrome (aHUS)
- Other therapeutic indication(s) – Please specify: _____

INFORMATION RELATING TO PAROXYSMAL NOCTURNAL HEMOGLOBINURIA (PNH)

Please provide serum concentration of lactate dehydrogenase (LDH): _____

Please indicate serum concentration of hemoglobin: _____

Please indicate if one of the following applies to the patient:

- Thrombotic or embolic event which required institution of therapeutic anticoagulant therapy
- Minimum transfusion of 4 units of red blood cells in the previous 12 months
- Anemia defined by a hemoglobin serum concentration measured at least twice, < 100 g/L and accompanied by symptoms of anemia, or ≤ 70 g/L
- Pulmonary insufficiency: debilitating shortness of breath and/or chest pain resulting in limitation of normal activity
- Renal insufficiency demonstrated by an eGFR less than or equal to 60mL/min
- Smooth muscle spasm: recurrent episodes of severe pain requiring hospitalization and/or narcotic analgesia

INFORMATION RELATING TO ATYPICAL HEMOLYTIC UREMIC SYNDROME (aHUS)

Please indicate ADAMTS-13 activity: _____

Please indicate result for the STEC (Shiga toxin-producing E.Coli)-test: _____

Please indicate result for the platelet count: _____

Please indicate what situation apply to the patient:

- Presence of schistocytes Lactate dehydrogenase above normal range
- Low or absent haptoglobin Tissue biopsy confirming TMA

Please provide one of the following: eGFR: _____ Creatinine clearance: _____

Please indicate if one of the following documented clinical features of active organ damage or impairment other than renal impairment (neurological, cardiac, pulmonary, gastro-intestinal): _____

PRIOR MEDICATION OR TREATMENT

Has the patient ever used medication or received treatment for this medical condition? Yes No

If not, please explain: _____

If so, please list any medication already used or any treatment already received for this medical condition:

MEDICATION OR TREATMENT NAME	OUTCOME	TREATMENT PERIOD
Name: _____ Dose: _____	<input type="checkbox"/> Inefficiency <input type="checkbox"/> Intolerance <input type="checkbox"/> Contraindication Specify: _____	From: _____ To: _____
Name: _____ Dose: _____	<input type="checkbox"/> Inefficiency <input type="checkbox"/> Intolerance <input type="checkbox"/> Contraindication Specify: _____	From: _____ To: _____
Name: _____ Dose: _____	<input type="checkbox"/> Inefficiency <input type="checkbox"/> Intolerance <input type="checkbox"/> Contraindication Specify: _____	From: _____ To: _____
Name: _____ Dose: _____	<input type="checkbox"/> Inefficiency <input type="checkbox"/> Intolerance <input type="checkbox"/> Contraindication Specify: _____	From: _____ To: _____

PRESCRIPTION RENEWAL – Please provide an objective evidence of efficacy.

Paroxysmal nocturnal hemoglobinuria

Please indicate serum concentration of lactate dehydrogenase (LDH): _____ LDH concentration before treatment: _____

Atypical hemolytic uremic syndrome

Please provide the following: Platelet count: _____ Haptoglobin count: _____

 Serum concentration of lactate dehydrogenase (LDH): _____

Please provide the following information: eGFR: _____ Baseline eGFR: _____

If one of the following complications was present at initiation of treatment, please indicate which impairment was stabilized with eculizumab:

- Not applicable Cardiac Neurologic Pulmonary Gastro-intestinal

