



PRIOR AUTHORIZATION REQUEST

**JUXTAPID (LOMITAPIDE)
LEQVIO (INCLISIRAN)
PRALUENT (ALIROCUMAB)
REPATHA (EVOLOCUMAB)**

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? <input type="checkbox"/> Yes – Please provide a copy of the notice of approval or refusal. → <input type="checkbox"/> Copy attached to this form. Specify: Name of the insurer: _____ Contract No.: _____ Certificate No.: _____ <input type="checkbox"/> No
PROVINCIAL PLAN	Has a request for reimbursement been submitted under your provincial plan? <input type="checkbox"/> Yes – Please provide a copy of the notice of approval or refusal. → <input type="checkbox"/> Copy attached to this form. <input type="checkbox"/> No – Please explain: _____
PATIENT SUPPORT PROGRAM	Is the patient enrolled in a patient support program? <input type="checkbox"/> Yes <input type="checkbox"/> 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:** _____

CONTINUED ON THE BACK

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
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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.**

Clinical atherosclerotic cardiovascular disease : We are not accepting requests for Juxtapid, Leqvio, Praluent and Repatha for this diagnosis.
Heterozygous familial hypercholesterolemia (HeFH) : We are not accepting requests for Juxtapid for this diagnosis.
Homozygous familial hypercholesterolemia (HoFH) : We are not accepting requests for Praluent for this diagnosis.
We will consider requests for Juxtapid for this diagnosis only if there has already been an adequate trial with Repatha, along with a diet and other lipid-lowering treatments (statins, ezetimibe and LDL apheresis).

DIAGNOSTIC

Homozygous familial hypercholesterolemia (HoFH) Heterozygous familial hypercholesterolemia (HeFH)
 Other therapeutic indication(s) – Please specify:

INFORMATION RELATING TO HOMOZYGOUS FAMILIAL HYPERCHOLESTEROLEMIA (HoFH)

The treatment will be administered in combination with other lipid-lowering treatments: Yes No
Until now, the patient has been on a low-cholesterol diet: Yes No
For **at least three months** before the start of treatment, the patient's LDL cholesterol was above 2 mmol/L despite taking two or more statins: Yes No
Will the treatment be administered in combination with a low-density lipoproteins (LDL) apheresis treatment? Yes No
If not, please specify the reason:
The patient has one or more functional mutations in both LDL receptor alleles or alleles known to affect LDL receptor functionality: Yes No
The patient's LDL cholesterol was above 13 mmol/L before treatment: Yes No
The patient had xanthomas before age 10: Yes No
Both biological parents have been diagnosed with heterozygous familial hypercholesterolemia (HeFH): Yes No

INFORMATION RELATING TO HETEROZYGOUS FAMILIAL HYPERCHOLESTEROLEMIA (HeFH)

Does the patient have atherosclerotic cardiovascular disease? Yes – specify: _____ No
LDL-C at the time of diagnosis: _____ mmol/L Date: _____
LDL-C before the start of the requested treatment: _____ mmol/L Date: _____

Please check any element that apply to the patient:

- DNA-based evidence of an LDL receptor mutation or other FH-related gene mutation
- Family history of HeFH, confirmed by genotyping, in a first degree relative
- Presence of a mutation causing familial hypercholesterolemia of LDLR, ApoB or PCSK9 genes in a first-degree relative
- Presence of xanthomas in the person or in one of the parents of the first or second degree
- Presence of an arcus cornealis before the age of 45 in a first degree relative
- Family history of LDL-C > 4.9 mmol/L in an adult first degree relative or ≥ 4 mmol/L in a first degree relative younger than 18 years of age
- Family history of a total cholesterol concentration > 7.5 mmol/L in a first- or second-degree adult parent or > 6.7 mmol/L in a first-degree parent under 16 years of age

