



PRIOR AUTHORIZATION REQUEST

Repatha

PATIENT: Name _____
Address: _____
City, State, Zip _____
D.O.B. _____
Member ID: _____

Prescriber: Name _____
Address _____
City, State, Zip _____
Phone _____
Fax _____
NPI _____

Medication Requested: _____ **Qty Requested:** _____

Your patient's prescription benefit requires that we review certain requests for coverage with the prescriber. You have prescribed a medication for your patient that requires Prior Authorization before benefit coverage or coverage of additional quantities can be provided. Please complete the following questions then fax this form to the toll free number listed below. Upon receipt of the completed form, prescription benefit coverage will be determined based on the plan's rules.

SECTION A: Please answer the following questions

1. Yes No Is the requested medication being used concurrently with Praluent or Juxtapid?
2. Yes No Is the requested medication being prescribed by, or in consultation with, a cardiologist; an endocrinologist; or a physician who focuses in the treatment of cardiovascular (CV) risk management and/or lipid disorders?
3. What is the indication or diagnosis?
 - Atherosclerotic cardiovascular disease (ASCVD) [clinical] – **Please answer questions 16 – 26**
 - Heterozygous familial hypercholesterolemia (HeFH) – **Please answer questions 4 – 11 & 21 – 26**
 - Homozygous familial hypercholesterolemia (HoFH) - **Please answer questions 12 – 15 & 21 – 26**
 - Primary Hyperlipidemia – **Please answer questions 23 – 30**
 - All other indications or diagnoses – Please specify _____
4. Yes No Does the patient have an untreated low-density lipoprotein cholesterol (LDL-C) level greater than or equal to 190 mg/dL (that is, prior to treatment with antihyperlipidemic agents)?
5. Yes No Does the patient have genetic confirmation of HeFH by mutations in the low-density lipoprotein receptor (LDLR), apolipoprotein B (APOB), proprotein convertase subtilisin kexin type 9 (PCSK9) or low-density lipoprotein receptor adaptor protein 1 (LDLRAP1) gene?
6. Yes No Has the patient been diagnosed with HeFH by the prescriber using the Dutch Lipid Network criteria?
7. Yes No Does the patient have a score of greater than 5?
8. Yes No Has the patient been diagnosed with HeFH by the prescriber using the Simon Broome criteria?
9. Yes No Has the patient met the threshold for "definite" or "possible" familial hypercholesterolemia?
10. Yes No Does the patient have clinical manifestations of HeFH?

**If you have any questions, call:
800-753-2851**

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11. Yes No Does the patient have a treated low-density lipoprotein cholesterol (LDL-C) level greater than or equal to 100 mg/dL (that is, after treatment with antihyperlipidemic agents but prior to treatment with PCSK9 inhibitor therapy such as Praluent or Repatha)?
12. Yes No Does the patient have genetic confirmation of two mutant alleles at the low-density lipoprotein receptor (LDLR), apolipoprotein B (APOB), proprotein convertase subtilisin/kexin type 9 (PCSK9) or low-density lipoprotein receptor adaptor protein 1 (LDLRAP1) gene locus?
13. Yes No Does the patient have an untreated LDL-C level greater than 500 mg/dL (that is, prior to treatment with antihyperlipidemic agents)?
14. Yes No Does the patient have a treated LDL-C level of 300 mg/dL or greater (that is, after treatment with antihyperlipidemic agents but prior to agents such as Repatha, or Juxtapid)?
15. Yes No Does the patient have patient has clinical manifestations of HoFH?
16. Yes No Has the patient had a previous myocardial infarction (MI) or history of an acute coronary syndrome (ACS)?
17. Yes No Does the patient have a diagnosis of angina (stable or unstable)?
18. Yes No Does the patient have a past history of stroke or transient ischemic attack (TIA)?
19. Yes No Does the patient have peripheral arterial disease (PAD)?
20. Yes No Has the patient undergone a coronary or other arterial revascularization procedure in the past?
21. Yes No Has the patient tried one high-intensity statin therapy (that is, atorvastatin 40 mg or greater daily; rosuvastatin 20 mg or greater daily [as a single-entity or as a combination product]) for at least 8 weeks continuously?
22. Yes No Does the patient's low-density lipoprotein cholesterol (LDL-C) level after this treatment remain greater than or equal to 70 mg/dL?
23. Yes No Has the patient been determined to be statin intolerant by experiencing statin-related rhabdomyolysis?
24. Yes No Has the patient been determined to be statin intolerant by experiencing skeletal-related muscle symptoms?
25. Yes No Did the skeletal-related muscle symptoms occur while receiving separate trials of both atorvastatin and rosuvastatin (as single-entity or as combination products)?
26. Yes No When receiving separate trials of both atorvastatin and rosuvastatin (as single-entity or as combination products) did the skeletal-related muscle symptoms resolve upon discontinuation of each respective statin therapy (atorvastatin AND rosuvastatin)?
27. Yes No Is the patient's coronary artery calcium or calcification (CAC) score greater than or equal to 300 Agatston units?
28. Yes No Has the patient tried one high-intensity statin therapy (that is, atorvastatin 40 mg or greater daily; rosuvastatin 20 mg or greater daily [as a single-entity or as a

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- 29. Was the high-intensity statin therapy... combination product]?
30. Does the patient's low-density lipoprotein cholesterol (LDL-C) level after this treatment regimen remain greater than or equal to 100 mg/dL?

Please document the diagnoses, symptoms, and/or any other information important to this review:

Two horizontal lines for documentation.

SECTION B Physician Signature

PHYSICIAN SIGNATURE DATE
FAX COMPLETED FORM TO: 877-251-5896

Disclaimer: An authorization is not a guarantee of payment. Member must be eligible at the time services are rendered. Services must be a covered Health Plan Benefit and medically necessary with prior

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