



Please complete ALL information below and fax your request to 1-888-671-5285

### Praluent® & Repatha® Prior Authorization Request Form

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Member Information (required)			Provider Information (required)		
Member Name:			Provider Name:		
Insurance ID#:			NPI#:	Specialty:	
Date of Birth:			Office Phone:		
Street Address:			Office Fax:		
City:	State:	Zip:	Office Street Address:		
Phone:			City:	State:	Zip:

Medication Information (required)		
Medication Name:	Strength:	Dosage Form:
<input type="checkbox"/> Check if <b>generic substitution</b> is acceptable	Directions for Use:	
<input type="checkbox"/> Check if request is for <b>continuation of therapy</b>		

Clinical Information (required)	
<b>Select the diagnosis below:</b>	
<input type="checkbox"/> Atherosclerotic cardiovascular disease	
<input type="checkbox"/> Homozygous familial hypercholesterolemia (Repatha only)	
<input type="checkbox"/> Hyperlipidemia	
<input type="checkbox"/> Other diagnosis: _____ ICD-10 Code(s): _____	

<p><b>Clinical Information:</b></p> <p>Does the patient have LDL-C 70 mg/dL or greater after a minimum 8-week trial of at least moderate-intensity statin therapy?* <input type="checkbox"/> <b>Yes</b> <input type="checkbox"/> <b>No</b></p> <p><i>*Please note: Moderate intensity statins are defined as lowering LDL-C by approximately 30% to less than 50% (Atorvastatin 10 or 20 mg daily, fluvastatin XL 80 mg daily, fluvastatin 40 mg twice daily, lovastatin 40mg daily, pitavastatin 2 to 4 mg daily, pravastatin 40 mg or 80 mg daily, rosuvastatin 5 or 10 mg daily, simvastatin 20 to 40 mg daily)</i></p> <p>Does the patient have an inability to tolerate statin therapy as documented by having had rhabdomyolysis or symptoms with creatine kinase (CK) exceeding 10 times the upper limit of normal (ULN) on <b>any</b> statin? <input type="checkbox"/> <b>Yes</b> <input type="checkbox"/> <b>No</b></p> <p>Select if the patient has had an inability to tolerate statin therapy with <b>TWO</b> statins as documented by <b>ONE</b> of the following:</p> <ul style="list-style-type: none"> <li><input type="checkbox"/> Myalgia (no CK elevation)</li> <li><input type="checkbox"/> Myositis (CK less than 10 times ULN)</li> <li><input type="checkbox"/> Hepatotoxicity from statin use (increased AST/ALT exceeding 3 times ULN)</li> <li><input type="checkbox"/> Liver disease documented by Child Pugh A or worse OR AST/ALT exceeding 3 times ULN for at least 6 weeks</li> </ul>
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<p><b>For atherosclerotic cardiovascular disease, also answer the following:</b></p> <p>Is the patient diagnosed by either stress test, angiography, atherosclerotic event (e.g., MI, angina, stroke, claudication, carotid stenosis) or arterial intervention for atherosclerotic disease (e.g. coronary, peripheral, carotid)? <input type="checkbox"/> <b>Yes</b> <input type="checkbox"/> <b>No</b></p>
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<p><b>Continuation:</b></p> <p>Has the patient had a sustained reduction in LDL-C of at least 25% since initiation of the requested medication? <input type="checkbox"/> <b>Yes</b> <input type="checkbox"/> <b>No</b></p>
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Are there any other comments, diagnoses, symptoms, medications tried or failed, and/or any other information the physician feels is important to this review?

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Please note: This request may be denied unless all required information is received.

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