

Repatha® Coverage Determination Request Form (Page 1 of 2)

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Member Information <small>(required)</small>			Provider Information <small>(required)</small>		
Member Name:			Provider Name:		
Insurance ID#:			NPI#:		Specialty:
Date of Birth:			Office Phone:		
Street Address:			Office Fax:		Office Contact:
City:	State:	Zip:	Office Street Address:		
Phone:			City:	State:	Zip:
Medication Information <small>(required)</small>					
Medication Name: Select one of the following:			Strength:		Dosage Form:
<input type="checkbox"/> Request is for GENERIC					
<input type="checkbox"/> Request is for BRAND (unable to take the generic)					
<input type="checkbox"/> Check if request is for continuation of therapy			Directions for Use:		
Clinical Information <small>(required)</small>					
Select the Type(s) of Coverage Determination Requested:					
<input type="checkbox"/> Prior Authorization - Request is for a drug that requires prior authorization under the plan.					
<input type="checkbox"/> Quantity Limit - Request is for an exception to the plan's quantity limit. Quantity per DAY requested? _____					
Select the diagnosis below:					
<input type="checkbox"/> Atherosclerotic cardiovascular disease (ASCVD)					
<input type="checkbox"/> Homozygous familial hypercholesterolemia (HoFH)					
<input type="checkbox"/> Hyperlipidemia					
<input type="checkbox"/> Other diagnosis: _____ ICD-10 Code(s): _____					
Clinical Information:					
Is there documentation that the patient has an LDL-C 70mg/dL or greater after a minimum 8-week trial of at least moderate-intensity statin therapy? <input type="checkbox"/> Yes <input type="checkbox"/> No					
Select if the patient has an inability to tolerate statin therapy as documented by the following:					
<input type="checkbox"/> Rhabdomyolysis or symptoms with CK (creatinine kinase) exceeding 10 times ULN (upper limit of normal)					
<input type="checkbox"/> Myalgia (no CK elevations) with TWO statins					
<input type="checkbox"/> Myositis (CK less than 10 times ULN) with TWO statins					
<input type="checkbox"/> Hepatotoxicity from statin use (increased AST/ALT exceeding 3 times ULN)					
<input type="checkbox"/> Liver disease documented by Child Pugh A or worse					
<input type="checkbox"/> Liver disease documented by AST/ALT exceeding 3 times ULN for at least 6 weeks					
For atherosclerotic cardiovascular disease (ASCVD), also answer the following:					
Select if the patient's diagnosis is confirmed by of the following:					
<input type="checkbox"/> Stress test					
<input type="checkbox"/> Angiography					
<input type="checkbox"/> Atherosclerotic event (e.g., MI, angina, stroke, claudication, carotid stenosis)					
<input type="checkbox"/> Arterial intervention for atherosclerotic disease (e.g., coronary, peripheral, carotid)					
Reauthorization:					
If this is a reauthorization request, answer the following question:					
Select if there is documentation of sustained reduction in LDL-C from baseline with the following:					
<input type="checkbox"/> 25% reduction of LDL-C from baseline					
<input type="checkbox"/> Sustained LDL-C below 70mg/dL					

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Quantity Limit Requests:

Is there a high risk of significant adverse clinical outcome with medication change or dosage change? Yes No

Is the requested quantity and dose within FDA approved maximum dosing limits or supported by peer-reviewed medical literature, accepted standards of medical practice and/or medical compendia? Yes No

If **yes**, please specify: _____

Are there any other comments, diagnoses, symptoms, medications tried or failed, and/or any other information the physician feels is important to this review?

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