



Pharmacy Prior Approval Request for PCSK9 Inhibitors

Beneficiary Information

1. Beneficiary Last Name: \_\_\_\_\_ 2. First Name: \_\_\_\_\_
3. Beneficiary ID #: \_\_\_\_\_ 4. Beneficiary Date of Birth: \_\_\_\_\_ 5. Beneficiary Gender: \_\_\_\_\_

Prescriber Information

6. Prescribing Provider NPI #: \_\_\_\_\_
7. Requester Contact Information - Name: \_\_\_\_\_ Phone #: \_\_\_\_\_ Ext. \_\_\_\_\_

Drug Information

8. Drug Name: \_\_\_\_\_ 9. Strength: \_\_\_\_\_ 10. Quantity Per 30 Days: \_\_\_\_\_
11. Length of Therapy (In days):  up to 30 Days  60 Days  90 Days  120 Days  180 Days  365 Days  Other \_\_\_\_\_

Clinical Information

Clinical Questions for All PSCK9 Inhibitors:
1. Is the member at least 18 years of age?  Yes  No
2. Is the member currently taking the maximum dose, for his/her age, of atorvastatin (generic for Lipitor) or rosuvastatin (generic for Crestor) AND has completed 90 days of treatment?  Yes  No
3. Is the member's LDL level >= 70mg/dl after taking atorvastatin (generic for Lipitor) or rosuvastatin (generic for Crestor) for 90 days?  Yes  No
4. Does the member have a significant intolerance or allergic reaction to atorvastatin (generic for Lipitor) or rosuvastatin (generic for Crestor)? Examples of significant intolerance include severe muscle pain, significant liver abnormalities, and rhabdomyolysis. Intolerance does not include fatigue, cognitive impairment, or mild aches.  Yes  No
5. Has documentation of clinically significant intolerance or allergic reaction to statin treatment been attached to this prior approval request?  Yes  No
6. Baseline LDL before statin treatment: \_\_\_\_\_
7. LDL after statin treatment: \_\_\_\_\_
\*\*LDL lab results before and after statin treatment must be attached to this prior approval request\*\*
8. Will high dose atorvastatin (generic for Lipitor) or rosuvastatin (generic for Crestor) be continued with the PCSK9 inhibitor?  Yes  No
Clinical Questions for Praluent:
9. Does the member have a diagnosis of Heterozygous Familial Hypercholesterolemia?  Yes  No
10. Does the member have clinical atherosclerotic cardiovascular disease such as acute coronary syndromes, or a history of myocardial infarction, stable or unstable angina, coronary or other arterial revascularization, stroke, transient ischemic attack, or peripheral arterial disease of atherosclerotic origin?  Yes  No
11. Does the member have a diagnosis of Severe Primary Hyperlipidemia (defined as LDL-C >= 190mg/dL)?  Yes  No
Clinical Questions for Repatha:
12. Does the member have a diagnosis of Heterozygous Familial Hypercholesterolemia (HeFH)?  Yes  No
13. Does the member have a diagnosis of Homozygous Familial Hypercholesterolemia (HoFH)?  Yes  No
14. Is the member 13 years or older?  Yes  No
15. Does the member have clinical atherosclerotic cardiovascular disease such as acute coronary syndromes, or a history of myocardial infarction, stable or unstable angina, coronary or other arterial revascularization, stroke, transient ischemic attack, or peripheral arterial disease of atherosclerotic origin?  Yes  No
16. Does the member have a diagnosis of Severe Primary Hyperlipidemia (defined as LDL-C >= 190mg/dL)?  Yes  No

Signature of Prescriber: \_\_\_\_\_ Date: \_\_\_\_\_
(Prescriber Signature Mandatory)

I certify that the information provided is accurate and complete to the best of my knowledge, and I understand that any falsification, omission, or concealment of material fact may subject me to civil or criminal liability.