

NC Medicaid and NC Health Choice
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 PCSK9 Inhibitors:

1. Is the beneficiary at least 18 years of age? Yes No
2. Is the beneficiary 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 beneficiary's LDL level \geq 70mg/dl after taking atorvastatin (generic for Lipitor) or rosuvastatin (generic for Crestor) for 90 days? Yes No
4. Does the beneficiary 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 beneficiary have a diagnosis of Heterozygous Familial Hypercholesterolemia? Yes No
10. Does the beneficiary have a diagnosis of Homozygous Familial Hypercholesterolemia? Yes No
11. Does the beneficiary 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
12. Does the beneficiary have a diagnosis of Severe Primary Hyperlipidemia (defined as LDL-C \geq 190mg/dL)? Yes No

Clinical Questions for Repatha:

13. Does the beneficiary have a diagnosis of Heterozygous Familial Hypercholesterolemia (HeFH)? Yes No
14. Does the beneficiary have a diagnosis of Homozygous Familial Hypercholesterolemia (HoFH)? Yes No
15. Is the beneficiary 13 years or older? Yes No
16. Does the beneficiary 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
17. Does the beneficiary have a diagnosis of Severe Primary Hyperlipidemia (defined as LDL-C \geq 190mg/dL)? Yes No

Continuation Questions for Praluent and Repatha:

18. Has the provider submitted documentation that indicates a positive clinical response to therapy with this request? Yes No
19. Is the beneficiary continuing to receive other lipid-lowering therapy? Yes No
20. Is the beneficiary currently receiving more than one PCSK9 inhibitor? 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.