

## NC 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 #: _____	Provider Fax #: _____
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): <input type="checkbox"/> up to 30 Days <input type="checkbox"/> 60 Days <input type="checkbox"/> 90 Days <input type="checkbox"/> 120 Days <input type="checkbox"/> 180 Days <input type="checkbox"/> 365 Days <input type="checkbox"/> Other _____		

### Clinical Information

<p><b>Clinical Questions for All PSCK9 Inhibitors:</b></p> <p>1. 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? <input type="checkbox"/> <b>Yes</b> <input type="checkbox"/> <b>No</b></p> <p>2. Is the beneficiary's LDL level <math>\geq</math> 70mg/dl after taking atorvastatin (generic for Lipitor) or rosuvastatin (generic for Crestor) for 90 days? <input type="checkbox"/> <b>Yes</b> <input type="checkbox"/> <b>No</b></p> <p>3. 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. <input type="checkbox"/> <b>Yes</b> <input type="checkbox"/> <b>No</b></p> <p>4. Has documentation of clinically significant intolerance or allergic reaction to statin treatment been attached to this prior approval request? <input type="checkbox"/> <b>Yes</b> <input type="checkbox"/> <b>No</b></p> <p>5. Baseline LDL before statin treatment: _____</p> <p>6. LDL after statin treatment: _____</p> <p><b>**LDL lab results before and after statin treatment must be attached to this prior approval request**</b></p> <p>7. Will high dose atorvastatin (generic for Lipitor) or rosuvastatin (generic for Crestor) be continued with the PCSK9 inhibitor? <input type="checkbox"/> <b>Yes</b> <input type="checkbox"/> <b>No</b></p> <p><b>Clinical Questions for Praluent:</b></p> <p>8. Is the beneficiary 18 years of age or older? <input type="checkbox"/> <b>Yes</b> <input type="checkbox"/> <b>No</b></p> <p>9. Does the beneficiary have a diagnosis of Heterozygous Familial Hypercholesterolemia? <input type="checkbox"/> <b>Yes</b> <input type="checkbox"/> <b>No</b></p> <p>10. Does the beneficiary have a diagnosis of Homozygous Familial Hypercholesterolemia? <input type="checkbox"/> <b>Yes</b> <input type="checkbox"/> <b>No</b></p> <p>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? <input type="checkbox"/> <b>Yes</b> <input type="checkbox"/> <b>No</b></p> <p>12. Does the beneficiary have a diagnosis of Severe Primary Hyperlipidemia (defined as LDL -C <math>\geq</math> 190mg/dL)? <input type="checkbox"/> <b>Yes</b> <input type="checkbox"/> <b>No</b></p> <p><b>Clinical Questions for Repatha:</b></p> <p>13. Does the beneficiary have a diagnosis of Heterozygous Familial Hypercholesterolemia (HeFH)? <input type="checkbox"/> <b>Yes</b> <input type="checkbox"/> <b>No</b></p> <p>14. Does the beneficiary have a diagnosis of Homozygous Familial Hypercholesterolemia (HoFH)? <input type="checkbox"/> <b>Yes</b> <input type="checkbox"/> <b>No</b></p> <p>15. Is the beneficiary 10 years or older? <input type="checkbox"/> <b>Yes</b> <input type="checkbox"/> <b>No</b></p> <p>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? <input type="checkbox"/> <b>Yes</b> <input type="checkbox"/> <b>No</b></p> <p>17. Does the beneficiary have a diagnosis of Severe Primary Hyperlipidemia (defined as LDL -C <math>\geq</math> 190mg/dL)? <input type="checkbox"/> <b>Yes</b> <input type="checkbox"/> <b>No</b></p> <p><b>Continuation Questions for Praluent and Repatha:</b></p> <p>18. Has the provider submitted documentation that indicates a positive clinical response to therapy with this request? <input type="checkbox"/> <b>Yes</b> <input type="checkbox"/> <b>No</b></p> <p>19. Is the beneficiary continuing to receive other lipid-lowering therapy? <input type="checkbox"/> <b>Yes</b> <input type="checkbox"/> <b>No</b></p> <p>20. Is the beneficiary currently receiving more than one PCSK9 inhibitor? <input type="checkbox"/> <b>Yes</b> <input type="checkbox"/> <b>No</b></p>
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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.