

Nexletol and Nexlizet

Member Information

1. Last Name: _____ 2. First Name: _____
3. Trillium ID #: _____ 4. Date of Birth: _____ 5. Gender: _____

Prescriber Information

1. Prescriber Name: _____ 2. NPI #: _____
3. Requestor Name (Nurse/Office Staff): _____
4. Mailing Address: _____ City: _____ State: _____ Zip: _____
3. Phone #: _____ Ext. _____ Fax #: _____

Drug Information

1. Drug Name: _____ 2. Strength: _____ 3. Quantity Per 30 Days: _____
4. Length of Therapy (in Days): up to 30 Days 60 Days 90 Days 120 Days 180 Days 365 Days

Clinical Information

Initial Coverage Nexletol questions 1-5) and Nexlizet (questions 1-7)

1. Is the member at least 18 years old or older? Yes No
2. Has the member been diagnosed with heterozygous familial hypercholesterolemia (HeFH) or established atherosclerotic cardiovascular disease (ASCVD) defined 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
3. Has the member failed to achieve a target LDL-C (at least 50% reduction from baseline OR if no baseline is available: <70mg/dL for members with ASCVD and <100mg/dL for members with HeFH, and no history of ASCVD) despite physician attestation that the member is adherent to maximally-tolerated doses of statins for at least 90 days duration prior to the lipid panel demonstrating suboptimal reduction? Yes No
4. Is therapy being used in conjunction with maximally-tolerated doses of a statin? Yes No
5. Will therapy NOT be used with concurrent doses of simvastatin > 20gm or pravastatin > 40mg? Yes No

Initial Coverage Nexlizet (questions 6-7)

6. For Nexlizet- Does the beneficiary have a hypersensitivity to ezetimibe (Zetia®)? Yes No
7. Will Nexlizet be used with concurrent fibrate therapy (excluding fenofibrate)? Yes No

Continuation of Coverage for Nexletol and Nexlizet

8. Does the member continue to meet initial criteria above? Yes No
9. Is the member absent of unacceptable toxicity from therapy. (Examples of unacceptable toxicity include the following: hyperuricemia, tendon rupture)? Yes No
10. Does laboratory analysis demonstrate a reduction in LDL-C when compared to the baseline values (prior to initiating bempedoic acid or bempedoic acid/ezetimibe)? 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.