DEPARTMENT OF HEALTH SERVICES
Division of Medicaid Services
F-02505 (07/2019)

STATE OF WISCONSIN
Wis. Admin. Code § DHS 107.10(2)

FORWARDHEALTH

PRIOR AUTHORIZATION DRUG ATTACHMENT FOR LIPOTROPICS, PROPROTEIN CONVERTASE SUBTILISIN / KEXIN TYPE 9 (PCSK9) INHIBITORS

INSTRUCTIONS: Type or print clearly. Before completing this form, read the Prior Authorization Drug Attachment for Lipotropics, Proprotein Convertase Subtiliskin/Kexin Type 9 (PCSK9) Inhibitors Instructions, F-02505A. Providers may refer to the Forms page of the ForwardHealth Portal at www.forwardhealth.wi.gov/WIPortal/Subsystem/Publications/ForwardHealthCommunications.aspx?panel=Forms for the completion instructions.

Pharmacy providers are required to have a completed Prior Authorization Drug Attachment for Lipotropics, PCSK9 Inhibitors form signed by the prescriber before submitting a PA request on the Portal, by fax, or by mail. Providers may call Provider Services at 800-947-9627 with questions.

SECTION I – MEMBER INFORMATION

1. Name – Member (Last, First, Middle Initial)

2. Member ID Number

3. Date of Birth – Member

SECTION II – PRESCRIPTION INFORMATION

4. Drug Name

5. Drug Strength

6. Date Prescription Written

7. Refills

8. Directions for Use

9. Name – Prescriber

10. National Provider Identifier – Prescriber

11. Address – Prescriber (Street, City, State, ZIP+4 Code)

12. Phone Number – Prescriber

SECTION III – CLINICAL INFORMATION – ALL REQUESTS

13. Diagnosis Code and Description

14. Indicate the member's current low-density lipoprotein (LDL).

_____ mg/dL

15. Date Member's LDL Measured

_____ / _____ / _____
Month Day Year

DT-PA123-123

SECTION IV – CLINICAL INFORMATION – INITIAL REQUESTS ONLY

Note: A copy of the member's medical records must be submitted with initial PA requests. Medical records must include the following:

- Evidence that the member has heterozygous familial hypercholesterolemia (HeFH), homozygous familial hypercholesterolemia (HoFH), or clinical atherosclerotic cardiovascular disease (ASCVD)
- A current lipid panel lab report
- Documentation of the member's current and previous ezetimibe, PCSK9 inhibitor, and statin drug therapies, including:
  - Drug name(s) and dosage
  - Dates taken
  - Lipid panel report prior to and during drug therapy (including dates taken)
  - Reasons for discontinuation if drug therapy was discontinued

16. Indicate which of the following medical conditions the PCSK9 inhibitor is being prescribed to treat.

❑ HeFH
Clinical documentation must support a **definitive** diagnosis of HeFH using either World Health Organization criteria (Dutch Lipid Clinic Network clinical criteria with a score greater than eight) or Simon Broome diagnostic criteria.

❑ HoFH
Genetic testing or clinical confirmation must be submitted.

❑ Clinical ASCVD
Clinical documentation must provide evidence of at least one of the following (check all that apply):

❑ The member has coronary artery disease that is supported by a history of myocardial infarction (heart attack), coronary revascularization, or angina pectoris.

❑ The member has a history of non-hemorrhagic stroke.

❑ The member has symptomatic peripheral arterial disease as evidenced by one of the following (check all that apply):

❑ Intermittent claudication with an ankle-brachial index of less than 0.85

❑ Peripheral arterial revascularization procedure

❑ Amputation due to atherosclerotic disease

❑ Other _____

17. Document the member's current and previous ezetimibe, PCSK9 inhibitor, and statin drug therapies including the following for each trial:
    - Drug name(s) and dosage
    - Dates taken
    - Lipid panel report prior to and during drug therapy (including dates taken)
    - Reasons for discontinuation if drug therapy was discontinued

SECTION V – CLINICAL INFORMATION – RENEWAL REQUESTS ONLY

Note: A copy of the member's current lipid panel (within the past 30 days) must be submitted with renewal PA requests.

18. Document the member's ezetimibe, PCSK9 inhibitor, and statin drug therapies. Include the name, dose, and dosing regimen for each drug, or check "none" if appropriate.

Ezetimibe Name _____

Dose _____ Dose Regimen _____

❑ None (Member is not currently taking ezetimibe.)

PCSK9 Inhibitor Name _____

Dose _____ Dose Regimen _____

❑ None (Member is not currently taking a PCSK9 inhibitor.)

Statin Name _____

Dose _____ Dose Regimen _____

❑ None (Member is not currently taking a statin.)

SECTION VI – AUTHORIZED SIGNATURE – INITIAL AND RENEWAL REQUESTS

19. **SIGNATURE –** Prescriber	20. Date Signed

SECTION VII – ADDITIONAL INFORMATION – INITIAL AND RENEWAL REQUESTS

21. Include any additional information in the space below. Additional diagnostic and clinical information explaining the need for the drug requested may be included here.

Reset Form