DEPARTMENT OF HEALTH SERVICES

Division of Medicaid Services F-02505 (07/2022)

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. Prescribers 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 and dated by the prescriber before submitting a prior authorization (PA) request on the Portal, by fax, or by mail. Prescribers and pharmacy 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				
Date Prescription Written	7. Refills				
8. Directions for Use					
9. Name – Prescriber					
10 Address Prescriber (Street City State Zin I A Code)					
10. Address – Prescriber (Street, City, State, Zip+4 Code)					
11. Phone Number – Prescriber	12. National Provider Identifier – Prescriber				
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SECTION III – CLINICAL INFORMATION – ALL REQUESTS					
13. Diagnosis Code and Description	· ·				
14. Indicate the member's current low-density lipoprotein	in 15. Date Member's LDL Measured				
(LDL).					
mg/dL	Month Day Year				



SECTION IV - CLINICAL INFORMATION - INITIAL REQUESTS ONLY

Note: Supporting clinical information and a copy of the member's current medical records must be submitted with initial PA requests. The supporting clinical information and 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 PCSK9 inhibitor and statin drug therapies, including the following for each trial:
 - Drug name(s) and dosage
 - Dates taken
 - o 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 drug 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 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	A copy of the memi	ber's current lipid	panel (within	the past 30 day	s) must be subm	itted with renewal PA	
reques	ts.						

requests.				
18. Document the member's PCSK9 inhibitor and statin drug therapies. Include the name, dose, and dosing regimen for each drug, or check "none" if appropriate.				
PCSK9 Inhibitor Drug Name				
Dose Dose Regimen				
□ None (Member is not currently taking a PCSK9 inhibitor drug.)				
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.