

Wegovy[®] (semaglutide) Prior Authorization Form

Member Name: _____ Date of Birth: _____ Member ID#: _____

Drug Information

Pharmacy Billing (NDC: _____) Start Date (or date of next dose): _____

Dose: _____ Regimen: _____

Pharmacy Information

Pharmacy NPI: _____ Pharmacy Name: _____

Pharmacy Phone: _____ Pharmacy Fax: _____

Prescriber Information

Prescriber NPI: _____ Prescriber Name: _____

Prescriber Phone: _____ Prescriber Fax: _____ Specialty: _____

Criteria

For Initial Authorization: (Page 1 of 2)

1. Please indicate the diagnosis and information:

- To reduce the risk of major adverse cardiovascular (CV) events
- Other _____

2. Does the member have established cardiovascular disease (CVD)? Yes ___ No ___

3. Is the member obese or overweight? Yes ___ No ___

4. Does the member have a history of any of the following? (check all that apply and clinical documentation must be submitted with the request)

- Previous myocardial infarction
- Previous stroke
- Symptomatic peripheral arterial disease confirmed by 1 of the following:
 - Intermittent claudication with ankle-brachial index <0.85 at rest
 - Peripheral arterial revascularization procedure
 - Amputation due to atherosclerotic disease

5. Member's body mass index (BMI): _____ Date Taken: _____

6. Member's hemoglobin A1C (HbA1c): _____ Date Taken: _____

7. Does the member have type 1 diabetes mellitus (T1DM) or type 2 diabetes mellitus (T2DM)? Yes ___ No ___

8. Will member use Wegovy[®] in combination with other semaglutide-containing products or any other glucagon-like peptide-1 (GLP-1) receptor agonist? Yes ___ No ___

9. Is the member currently receiving guideline-directed management and therapy (GDMT) for CVD (e.g., antihypertensives, lipid-lowering agents, antiplatelets), as documented in the member's pharmacy claims history, unless contraindicated? Yes ___ No ___

a. Please provide the member's current GDMT for CVD: _____

b. If contraindicated, please provide details: _____

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PLEASE PROVIDE THE INFORMATION REQUESTED AND RETURN TO:

University of Oklahoma College of Pharmacy
Pharmacy Management Consultants
Product Based Prior Authorization Unit

Fax: 1-800-224-4014
Phone: 1-800-522-0114 Option 4

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Criteria

For Initial Authorization: (Page 2 of 2)

10. Will Wegovy[®] be used in conjunction with diet and exercise? Yes ___ No ___ (clinical documentation of member's diet and exercise program must be included with the request)
11. Request is for:
- Titration dosing
 - Maintenance dosing
12. Initial approvals will be for the titration period to allow initial and escalation dosing. A separate request must be submitted for each dose. Approvals will be for 4 weeks at a time to allow for proper dose escalation. An additional 4 weeks for each dose may be approved for those who experience intolerable adverse effects during dose escalation with proper documentation.

Additional information: _____

For Authorization of Maintenance Dosing: (approvals will be for 1 year)

1. Date of last dose: _____
2. Is the member tolerating maintenance dosing? Yes ___ No ___
3. Has the member developed T1DM or T2DM? Yes ___ No ___
4. Is the member continuing all of the following?
- Reduced calorie diet
 - Increased physical activity
 - GDMT for CVD where applicable

Additional Information: _____

(Page 2 of 2)

Prescriber Signature: _____ Date: _____

I certify that the indicated treatment is medically necessary and all information is true and correct to the best of my knowledge. Please do not send in chart notes. Specific information will be requested if necessary. Failure to complete this form in full will result in processing delays.

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