



AETNA BETTER HEALTH®
Coverage Policy/Guideline

Name:	Wegovy (semaglutide injection) Cardiovascular	Page:	1 of 3
Effective Date:	7/24/2024	Last Review Date:	6/13/2024
Applies to:	<input checked="" type="checkbox"/> Illinois <input checked="" type="checkbox"/> Maryland	<input type="checkbox"/> Virginia <input checked="" type="checkbox"/> Florida Kids	<input checked="" type="checkbox"/> New Jersey <input checked="" type="checkbox"/> Pennsylvania Kids

Intent:

The intent of this policy/guideline is to provide information to the prescribing practitioner outlining the coverage criteria for Wegovy under the patient's prescription drug benefit.

Description:

FDA-Approved Indication

Wegovy is indicated in combination with a reduced calorie diet and increased physical activity to reduce the risk of major adverse cardiovascular events (cardiovascular death, non-fatal myocardial infarction, or non-fatal stroke) in adults with established cardiovascular disease and either obesity or overweight.

Limitation of Use

Wegovy contains semaglutide. Coadministration with other semaglutide-containing products or with any other GLP-1 receptor agonist is not recommended.

Use of Wegovy for the indication of weight loss only is an excluded benefit and will not be covered.

Applicable Drug List:

Wegovy

Policy/Guideline:

Criteria for Initial Approval:

The requested drug will be covered with prior authorization when the following criteria are met:

- The requested drug will be used to reduce the risk of major adverse cardiovascular events (cardiovascular death, non-fatal myocardial infarction, or non-fatal stroke) in an adult with established cardiovascular disease and either obesity or overweight

AND

- The requested drug is being used with a reduced calorie diet and increased physical activity

AND

- The request is NOT for continuation of therapy

AND

- The patient has established cardiovascular disease with a history of ONE of the following: [Documentation is required for approval.]
 - A. Previous myocardial infarction (MI)
 - B. Previous stroke



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- C. Symptomatic peripheral arterial disease (PAD), as evidenced by intermittent claudication with ankle-brachial index (ABI) less than 0.85 (at rest), peripheral arterial revascularization procedure, or amputation due to atherosclerotic disease
- D. Prior history of revascularization (coronary artery bypass grafting (CABG), percutaneous coronary intervention (PCI), or angioplasty)
- E. Positive nuclear stress test
- F. Ischemic cardiomyopathy.

AND

- The patient has a baseline body mass index (BMI) greater than or equal to 27 kg/m². [Documentation is required for approval.]

AND

- The patient does NOT have type 2 diabetes
[NOTE: Ozempic is indicated to reduce the risk of major cardiovascular events in adults with type 2 diabetes mellitus and established cardiovascular disease. Patients with type 2 diabetes may be treated for risk reduction of cardiovascular events with Ozempic.]

AND

- The patient is currently receiving guideline-directed management and therapy (GDMT) for cardiovascular disease OR the patient has clinical reason not to be treated with GDMT for cardiovascular disease (e.g., lipid-lowering agent, antiplatelet, beta-blocker, renin-angiotensin inhibitor, etc.)
[Documentation is required for approval.]

OR

- The request is for continuation of therapy

AND

- The patient has established cardiovascular disease with a history of ONE of the following:
 - A. Previous myocardial infarction (MI)
 - B. Previous stroke
 - C. Symptomatic peripheral arterial disease (PAD), as evidenced by intermittent claudication with ankle-brachial index (ABI) less than 0.85 (at rest), peripheral arterial revascularization procedure, or amputation due to atherosclerotic disease



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- D. Prior history of revascularization (coronary artery bypass grafting (CABG), percutaneous coronary intervention (PCI), or angioplasty)
- E. Positive nuclear stress test
- F. Ischemic cardiomyopathy

AND

- The patient is being treated with a maintenance dosage of the requested drug

Approval Duration and Quantity Restrictions:

Initial Approval: 7 months

Renewal Approval: 12 months

Quantity Level Limit:

Drug	Dosage	1 Month Limit
Wegovy (semaglutide)	0.25 mg/0.5 mL	2 mL (1 package of 4 pens each) / 30 days
	0.5 mg/0.5 mL	
	1 mg/0.5 mL	
	1.7 mg/0.75 mL	3 mL (1 package of 4 pens each) / 30 days
	2.4 mg/0.75 mL	

References:

1. Wegovy [package insert]. Plainsboro, NJ: Novo Nordisk Inc.; March 2024.
2. Lexicomp Online, AHFS DI (Adult and Pediatric) Online. Waltham, MA: UpToDate, Inc.; 2024. <https://online.lexi.com>. Accessed March 18, 2024.
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4. Lincoff AM, Brown-Frandsen K, Colhoun HM, et al. Semaglutide and Cardiovascular Outcomes in Obesity without Diabetes. *N Engl J Med*. 2023;389:2221-2232.
5. Virani SS, Newby LK, Arnold SV, et al. 2023 AHA/ACC/ACCP/ASPC/NLA/PCNA Guideline for the Management of Patients With Chronic Coronary Disease: A Report of the American Heart Association/American College of Cardiology Joint Committee on Clinical Practice Guidelines. *Circulation*. 2023;148:e9-e119.
6. Kleindorfer DO, Towfighi A, Chaturvedi S, et al. 2021 Guideline for the Prevention of Stroke in Patients With Stroke and Transient Ischemic Attack: A Guideline From the American Heart Association/American Stroke Association. *Stroke*. 2021;52(7):e364-e467.
7. Gerhard-Herman MD, Gornik HL, Barrett C, et al. 2016 AHA/ACC Guideline on the Management of Patients with Lower Extremity Peripheral Artery Disease: A Report of the American College of Cardiology/American Heart Association Task Force on Clinical Practice Guidelines. *Circulation*. 2017;135(12):e726-e779.