

## THE PRESCRIBER **C-LETTER**



## Upcoming Changes to MassHealth Coverage of Weight Loss Drugs

MassHealth began covering anti-obesity medications in January 2024. At that time, Wegovy<sup>®</sup> (semaglutide) and Saxenda<sup>®</sup> (liraglutide) were designated as preferred glucagon-like peptide-1 (GLP-1) receptor agonists for the treatment of overweight and obesity.

### Effective October 1, 2024, Zepbound® (tirzepatide) Is a Preferred Drug

Effective October 1, 2024, Zepbound<sup>®</sup> (tirzepatide) is a preferred GLP-1 for the treatment of overweight and obesity. With this update, Zepbound<sup>®</sup> (tirzepatide) no longer requires a trial with Wegovy<sup>®</sup> (semaglutide) or Saxenda<sup>®</sup> (liraglutide) for prior authorization (PA) approval.

# Effective January 1, 2025, Wegovy<sup>®</sup> (semaglutide) and Saxenda<sup>®</sup> (liraglutide) Will Be Designated as Non-Covered Agents

Effective January 1, 2025, Wegovy<sup>®</sup> (semaglutide) and Saxenda<sup>®</sup> (liraglutide) will be non-covered agents for the treatment of overweight and obesity in adults.

- MassHealth members ≥18 years of age receiving Wegovy<sup>®</sup> (semaglutide) or Saxenda<sup>®</sup> (liraglutide) for the treatment of overweight or obesity will be required to switch to Zepbound<sup>®</sup> (tirzepatide) on January 1, 2025.
- MassHealth members ≥12 and <18 years of age may continue to use Wegovy<sup>®</sup> (semaglutide) or Saxenda<sup>®</sup> (liraglutide).
- MassHealth will continue to cover Wegovy<sup>®</sup> (semaglutide) for the indication of reduction of the risk of major adverse cardiovascular events in adults with established cardiovascular disease and either obesity or overweight. A new PA needs to be submitted if not initially approved for this indication.

To aid in transitioning, all MassHealth members ≥18 years of age approved for Wegovy<sup>®</sup> (semaglutide) or Saxenda<sup>®</sup> (liraglutide) for the treatment of overweight or obesity that would have had an approval duration beyond December 31, 2024, will automatically have an approval put in place to allow Zepbound<sup>®</sup> (tirzepatide) to pay at the pharmacy. The Zepbound<sup>®</sup> (tirzepatide) approval will be effective January 1, 2025, and expire six-months after the initial date the current Wegovy<sup>®</sup> (semaglutide) or Saxenda<sup>®</sup> (liraglutide) authorization was approved. At that time, a recertification PA for Zepbound<sup>®</sup> (tirzepatide) will be required and will be reviewed using baseline weight. Please see **Table 1** for guidance on the dosing equivalencies when switching members to Zepbound<sup>®</sup> (tirzepatide). Prescribers should work with their patients to determine which doses are appropriate. There are no restrictions on doses or durations needed. Example:

- Member started on Wegovy<sup>®</sup> (semaglutide) on September 1, 2024, and was issued an initial PA approval with an expiration date of March 1, 2025.
- The Wegovy<sup>®</sup> (semaglutide) PA will be end-dated effective December 31, 2024, and the Zepbound<sup>®</sup> (tirzepatide) approval will automatically be entered for January 1, 2025, through March 1, 2025.

• After March 1, 2025, a new PA needs to be submitted and reviewed for recertification based on weight loss from the baseline weight (prior to initiation of ANY GLP-1).

## Effective January 6, 2025, Phentermine Trial Will Be Required before Initiating Weight Loss GLP-1

Effective January 6, 2025:

- Generic phentermine will be available without PA for all MassHealth members ≥12 years of age. Lomaira<sup>®</sup> (phentermine) will also be available without PA for members ≥12 to <17 years of age. For members ≥18 years of age, Lomaira<sup>®</sup> (phentermine) will be available with PA.
- Requests for new starts for GLP-1 medication for the treatment of overweight or obesity will require a step through phentermine, with or without topiramate. Members already stable on GLP-1 therapy will not require this step through. For all new starts on Zepbound<sup>®</sup> (tirzepatide), PAs will need to document the following:
  - Diagnosis of obesity or overweight
  - Member age is  $\geq 18$  years
  - Baseline BMI of ≥30 kg/m2 or ≥27 kg/m2 with at least one weight-related comorbid condition
  - Baseline weight
  - Member has been counseled to continue reduced-calorie diet and increased physical activity
  - Inadequate response, adverse reaction, or contraindication to phentermine with or without topiramate

### Summary of PA Status of Anti-Obesity Agents Effective January 6, 2025

Available without PA

- Generic phentermine (≥12 years of age)
- Lomaira<sup>®</sup> (phentermine) (≥12 to <17 years of age)
- PA Required
  - benzphetamine
  - diethylpropion
  - diethylpropion ER
  - Lomaira<sup>®</sup> (phentermine) (≥18 years of age)
  - phendimetrazine
  - phendimetrazine ER
  - Xenical<sup>®</sup> (orlistat)
  - Zepbound<sup>®</sup> (tirzepatide)

Non-Covered

- Saxenda<sup>®</sup> (liraglutide)
- Wegovy<sup>®</sup> (semaglutide)

Agent	Comparative Doses (mg)								
Liraglutide once daily	0.6	1.2	1.8-3						
Semaglutide once weekly		0.25	0.5	1	2-2.4				
Tirzepatide once weekly			2.5		5	7.5	10	12.5	15

#### Table 1. Dosing Equivalencies for Injectable Anti-Obesity Agents1

Additional information can be found on the MassHealth Drug List.

#### References

 Whitley HP, Trujillo JM, Neumiller JJ; Special Report: Potential Strategies for Addressing GLP-1 and Dual GLP-1/GIP Receptor Agonist Shortages. Clin Diabetes 1 July 2023; 41 (3): 467–473.

The Prescriber e-Letter is an update designed to enhance the transparency and efficiency of the MassHealth drug prior-authorization (PA) process and the MassHealth Drug List. Each issue highlights key clinical information and updates to the MassHealth Drug List. The Prescriber E-Letter was prepared by the MassHealth Drug Utilization Review Program and the MassHealth Pharmacy Program.