SENTARA COMMUNITY PLAN
(MEDICAID)

PHARMACY PRIOR AUTHORIZATION/STEP-EDIT REQUEST*

Directions: **The prescribing physician must sign and clearly print name (preprinted stamps not valid) on this request**. All other information may be filled in by office staff; **fax to 1-800-750-9692**. No additional phone calls will be necessary if all information (including phone and fax #s) on this form is correct. **If the information provided is not complete, correct, or legible, the authorization process can be delayed.**

Weight Loss Management

Drug Requested: (check box below that applies)

PREFERRED MEDICATIONS ***Requires prior authorization**	
❑ **Adipex-P®/Suprenza™** (phentermine HCl)	❑ **Xenical®** (orlistat)
❑ **Bontril®/Bontril PDM®** (phendimetrazine)	❑ **Radtue®** (diethylpropion)
❑ **Didrex®/Regimex®** (benzphetamine)	
NON-PREFERRED MEDICATIONS ***Saxenda, Wegovy, Zepbound requires trial and failure of one (1) oral preferred agent** **For Wegovy and Zepbound**: the member has tried and failed the selected product, **Saxenda®**, as indicated on the PDL.	
❑ **Saxenda®** (liraglutide) *ages 12 and older	❑ **Wegovy®** (semaglutide) *ages 12 and older
❑ **Imcivree™**(setmelanotide) *ages 6 and older	❑ **Zepbound™** (tirzepatide) *ages 18 and older

MEMBER & PRESCRIBER INFORMATION: Authorization may be delayed if incomplete.

Member Name: _____

Member Sentara #: _____ **Date of Birth:** _____

Prescriber Name: _____

Prescriber Signature: _____ **Date:** _____

Office Contact Name: _____

Phone Number: _____ **Fax Number:** _____

NPI #: _____

DRUG INFORMATION: Authorization may be delayed if incomplete.

Drug Name/Form/Strength: _____

Dosing Schedule: _____ **Length of Therapy**: _____

Diagnosis: _____ **ICD Code, if applicable:** _____

Weight (if applicable): _____ **Date weight obtained:** _____

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CLINICAL CRITERIA: Check below all that apply. All criteria must be met for approval. To support each line checked, all documentation, including lab results, diagnostics, and/or chart notes, must be provided or request may be denied.

Initial Request Requirements:

- ❒ Provider must submit current height and weight measurements **(verified by chart notes)**

  Height: _____ **Current Weight:** _____ **BMI:** _____ **Date:** _____

Coverage for all medications will be limited to the following:

Absence of medical contraindications:

- ❑ No contraindications to use; (i.e. uncontrolled hypertension, hyperthyroidism etc. for stimulant based products); **AND**
- ❑ No malabsorption syndromes, cholestasis, pregnancy and/or lactation (for orlistat); **AND**
- ❑ No history of an eating disorder (e.g., anorexia, bulimia); **AND**
- ❑ No acute pancreatitis, acute suicidal behavior/ideation, personal or family history of medullary thyroid cancer or multiple endocrine neoplasia 2 syndrome (if requesting a GLP-1 Receptor Agonists); **AND**

Additional qualifying criteria to include the following: (excludes Imcivree™)

- ❑ Participation in nutritional counseling; **AND**
- ❑ Participation in physical activity program, unless medically contraindicated; **AND**
- ❑ Commitment to continue the above weight-loss treatment plan; **AND**

The provider attests that the patient's obesity is disabling and life threatening (i.e., puts the patient at risk for high-morbidity conditions):

- ❑ Yes
- ❑ No

Written documentation must include the following:

- ❑ Current medical status and weight loss plan. An individualized weight loss program should include a specific reduced calorie meal plan, recommended routine physical activity, and behavioral intervention including lifestyle modification as needed to improve adherence and outcomes.
- ❑ Current dated accurate height and weight measurements

Document details of <u>previous weight loss treatment plans</u> to include diet and exercise plans, in addition to submitting a copy of the plan:

_____

_____

_____

_____

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Drug Specific Requirements (Minimum ages are per FDA approvals):

1. For phentermine (min age 17), phendimetrazine tablet (min age 18), phendimetrazine ER capsule (min age 17) and orlistat (min age 12):
   - ❑ Body mass index (BMI) ≥ 30kg/m^2, if no applicable risk factors; **OR**
   - ❑ Body mass index (BMI) ≥ 27 with **at least one** weight-related comorbidity:

❑ coronary heart disease	❑ dyslipidemia	❑ hypertension	❑ sleep apnea	❑ Type II Diabetes

2. For benzphetamine (min age 17), diethylpropion (min age 16):
   - ❑ Body mass index (BMI) ≥ 30kg/m^2
3. For Imcivree® (min age 6)
   - ❑ BMI ≥ 30 kg/m^2; **AND**
   - ❑ Prescribed by or in consultation with an endocrinologist or geneticist; **AND**
   - ❑ Member has Bardet-Biedl syndrome (BBS); **OR**
   - ❑ Member has proopiomelanocortin (POMC), proprotein convertase subtilisin/kexin type 1 (PCSK1), or leptin receptor (LEPR) deficiency, as confirmed by a genetic test; **AND**
   - ❑ Member's genetic variants are interpreted as pathogenic, likely pathogenic, or of uncertain significance (VUS)
4. **For GLP-1 receptor agonists indicated for weight loss (Wegovy/Saxenda min age 12, Zepbound Min age 18):**
   - ❑ BMI > 40 kg/m^2, if no applicable risk factors; **OR**
   - ❑ BMI > 37 kg/m^2 with one or more of the following risk factors:

❑ dyslipidemia	❑ hypertension	❑ Type II Diabetes

   - ❑ Member has tried and failed one of the non-GLP1 weight-loss medications*; **OR**
   - ❑ Member is intolerant to all non-GLP1 weight-loss medications*; **AND**
   - ❑ Member not concurrently on another GLP-1 receptor agonists; **AND**
   - ❑ The member has tried and failed* the selected product, **Saxenda®**, as indicated on the PDL: https://www.virginiamedicaidpharmacyservices.com/provider/preferred-drug-list/

*Definitions of Accepted Drug Trial	
Drug	**Trial**
Benzphetamine, diethylpropion, phendimetrazine, phentermine	3-month trial without a weight loss of 10lbs
Orlistat	6-month trial without a weight loss of 10lbs
GLP-1 Receptor Agonist	6-month trial without a body weight reduction of 5%

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Length of Authorization:

- **Initial request:** Varies **(drug specific)**
  - Benzphetamine, diethylpropion, phendimetrazine, phentermine **-- 3 months**
  - GLP-1 receptor agonists – **6 month**
  - Xenical®(orlistat) **- 6 months**
  - Imcivree™ **-- 4 months**
- **Renewal requests**: **Renewals will no longer be granted once a member reaches a BMI < 25. See additional requirements below (drug specific):**
  - **Benzphetamine, diethylpropion, phendimetrazine, phentermine** – If patient achieves at least a 10-lb weight loss during initial 3 months of therapy, an additional 3-month prior authorization may be granted. Maximum length of continuous drug therapy = 6 months **(waiting period of 6 months before next request)**
  - **Alli®/Xenical®** (orlistat)- If patient achieves at least a 10-lb weight loss, an additional 6-month prior authorization may be granted. Maximum length of continuous drug therapy = 24 months (waiting period of 6 months before next request)
  - **Imcivree™** – If the member has experienced ≥ 5% reduction in body weight (or ≥ 5% of baseline BMI in those with continued growth potential), an additional 1-year prior authorization may be granted
  - **GLP-1 Receptor Agonists** – If the member achieves a weight loss of ≥ 5% reduction in body weight compared to the most recent authorization, an additional 6-month SA may be granted

Current measurements: BMI (Adult) or % of 95th percentile weight (12-17 y/o):

Height: _____ Weight: _____ BMI: _____ DATE: _____ (attach chart notes)

Previous authorization measurements: BMI (Adult) or % of 95th percentile weight (12-17 y/o):

Height: _____ Weight: _____ BMI: _____ DATE: _____ (attach chart notes)

All approvals are subject to the criteria on this form. Existing authorizations will be honored until renewal.

*****Use of samples to initiate therapy does not meet step edit/ preauthorization criteria.*****

****Previous therapies will be verified through pharmacy paid claims or submitted chart notes.****