AETNA BETTER HEALTH[®] OF VIRGINIA REQUEST FORM WEIGHT-LOSS MANAGEMENT Fax back to: 1-855-799-2553

If the following information is not complete, correct, or legible, the PA process can be delayed. Please use one form per member.

| MEMBER INFORMATION | |
|------------------------------------------|---------------------------------------------------------|
| Last Name: | First Name: |
| Medicaid ID Number: | Date of Birth: |
| | Weight in Kilograms: |
| PRESCRIBER INFORMATION | |
| Last Name: | First Name: |
| NPI Number: | |
| Phone Number: | Fax Number: |
| DRUG INFORMATION | |
| For initial requests, continue below. Fo | r renewal requests, proceed to Length of Authorization. |
| Drug Name/Form: | |
| Strength: | |
| Dosing Frequency: | |
| Length of Therapy: | |
| Quantity per Day: | |
| (Form continued on next name) | |

Member's Last Name:

Member's First Name:

DIAGNOSIS AND MEDICAL INFORMATION

| If the physician does not have the necessary information, the request will be denied and the fax form |
|-------------------------------------------------------------------------------------------------------|
| requesting additional information will be sent to the prescriber. |

Coverage for these medications will be limited to the following:

Absence of medical contraindications:

| No contraindications to use (i.e., uncontrolled hypertension, hyperthy | nyroidism | etc for stimulant based |
|------------------------------------------------------------------------|-----------|-------------------------|
| products); AND | | |
| | | _ |

No malabsorption syndromes, cholestasis, pregnancy, and lactation (for orlistat); AND

No history of an eating disorder (e.g., anorexia, bulimia); AND

No acute pancreatitis, suicidal behavoir or ideation, personal or family history of medullary thyroid cancer or multiple endocrine neoplasia 2 syndrome (if requesting GLP-1 Receptor Agonists)

For all others except Imcivree[®], additional qualifying criteria are:

Participation in nutritional counseling; AND

Participation in physical activity program, unless medically contraindicated; AND

Commitment to continue the above weight-loss treatment plan.

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

The 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; **AND**

Current accurate height and weight measurements

Summarize details of previous weight-loss treatment plans to include diet and exercise plans, in addition to submitting a copy of the plan:

AETNA BETTER HEALTH® OF VIRGINIA REQUEST FORM: Weight-Loss Management

Member's Last Name:

Member's First Name:

Assessment:

Other diagnoses and risk factors:

DRUG SPECIFIC CRITERIA

| NC | NOTE: Minimum ages are per FDA approvals. | | | | | | |
|----|----------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------|--|--|--|--|--|--|
| 1. | For phentermine (mininum age 17), phendimetrazine tablet (mininum age 18), phendimetrazine ER capsule (mininum age 17), and orlistat (mininum age 12): | | | | | | |
| | The member has a BMI of \geq 30 kg/m ² ; OR | | | | | | |
| | The member has a BMI of ≥ 27 kg/m ² with at least one weight-related comorbidity (e.g., coronary heart disease, dyslipidemia, hypertension, sleep apnea, type 2 diabetes) | | | | | | |
| 2. | For benzphetamine (mininum age 17) and diethylpropion (mininum age 16): | | | | | | |
| | The member has a BMI of \geq 30 kg/m ² | | | | | | |
| 3. | For Imcivree [®] (mininum age 6): | | | | | | |
| | BMI ≥ 30 kg/m²; 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 minimum age 12, Zepbound minimum 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, or type 2 diabetes; AND | | | | | | |
| | 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 agonist; AND | | | | | | |
| | If for an FDA-indicated GLP-1 receptor agonist, the member has tried and failed* the selected product as indicated on the PDL: <u>https://www.virginiamedicaidpharmacyservices.com/provider/preferred-drug-list/</u> | | | | | | |

Member's Last Name:

Member's First Name:

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

LENGTH OF AUTHORIZATION

| | Initial | Request: | Varies | (drug | specific) |
|--|---------|-----------------|--------|-------|-----------|
|--|---------|-----------------|--------|-------|-----------|

- Benzphetamine, diethylpropion, phendimetrazine, phentermine 3 months
- GLP-1 receptor agonists 6 months
- Orlistat 6 months
- Imcivree[®] 4 months

Renewal Request: See additional requirements below (drug specific)

- Benzphetamine, diethylpropion, phendimetrazine, phentermine If the member achieves at least a 10 pound (lb.) weight loss during the initial 3 months of therapy, an additional 3-month SA may be granted. Maximum length of continuous drug therapy is 6 months (waiting period of 6 months before next request).
- **Orlistat** If the member achieves at least a 10 lb weight loss, an additional 6-month SA may be granted. Maximum length of continuous drug therapy is 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 SA 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.

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

Attachments

Member's Last Name:

Member's First Name:

Prescriber Signature (Required)

Date

By signature, the physician confirms the above information is accurate and verifiable by member records.

Please include ALL requested information. Incomplete forms will delay the PA process.

Submission of documentation does NOT guarantee coverage.