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. F	or renewal requests, proceed to <u>Length of Authorization</u> .			
Drug Name/Form:				
Strength:				
Dosing Frequency:				
Length of Therapy:				
Quantity per Day:				
(Form continued on next page.)				

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Member's Last Name:	Member's First Name:
DIAGNOSIS AND MEDICAL INFORMATI	ON
If the physician does not have the necessa requesting additional information will be	ary information, the request will be denied and the fax form sent to the prescriber.
Coverage for these medications will be lin	nited to the following:
Absence of medical contraindications:	
No contraindications to use (i.e., un products); AND	ncontrolled hypertension, hyperthyroidism etc for stimulant based
No malabsorption syndromes, chole	estasis, pregnancy, and lactation (for orlistat); AND
☐ No history of an eating disorder (e.	g., anorexia, bulimia); AND
<u> </u>	avoir or ideation, personal or family history of medullary thyroid asia 2 syndrome (if requesting GLP-1 Receptor Agonists)
For all others except Imcivree®, additiona	l qualifying criteria are:
Participation in nutritional counseli	ng; AND
Participation in physical activity pro	ogram, unless medically contraindicated; AND
Commitment to continue the above	e weight-loss treatment plan.
The provider attests that the patient's obe	esity is disabling and life threatening (i.e., puts the patient at risk
Yes No	
The written documentation must include	the following:
specific reduced-calorie meal plan,	loss plan. An individualized weight-loss program should include a recommended routine physical activity, and behavioral dification as needed to improve adherence and outcomes; AND
Current accurate height and weight	measurements
Summarize details of previous weight-loss submitting a copy of the plan:	s treatment plans to include diet and exercise plans, in addition to
(Form continued on next page.)	

Member's Last Name:		Member's First Name:			
Assessment:					
Ot	Other diagnoses and risk factors:				
DI	RUG SPECIFIC CRITERIA				
N	OTE: Minimum ages are per FDA appro	ovals.			
1. For phentermine (mininum age 17), phendimetrazine tablet (mininum a capsule (mininum age 17), and orlistat (mininum age 12):					
	The member has a BMI of ≥ 30 kg	/m²; OR			
	The member has a BMI of ≥ 27 kg/m² with at least one weight-related comorbidity (e.g., coronary hea 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 ≥ 30 kg/m²				
3.	For Imcivree® (mininum age 6):				
	BMI ≥ 30 kg/m ² ; AND				
	Prescribed by or in consultation w	rith an endocrinologist or geneticist; AND			
	Member has Bardet-Biedl syndror	me (BBS); OR			
		n (POMC), proprotein convertase subtilisin/kexin type 1 (PCSK1), or as confirmed by a genetic test; AND			
	Member's genetic variants are int (VUS).	erpreted as pathogenic, likely pathogenic, or of uncertain significance			
4.	For GLP-1 receptor agonists indicated minimum age 18):	d for weight loss (Wegovy/Saxenda minimum age 12, Zepbound			
	\square BMI > 40 kg/m ² if no applicable ris	sk factors; OR			
	BMI > 37 kg/m ² with one or more diabetes; AND	of the following risk factors: dyslipidemia, hypertension, or type 2			
	Member has tried and failed one	of the non-GLP1 weight-loss medications*; OR			
	☐ Member is intolerant to all non-G	LP1 weight-loss medications*; AND			
	Member not concurrently on ano	ther GLP-1 receptor agonist; AND			
		ptor agonist, the member has tried and failed* the selected product www.virginiamedicaidpharmacyservices.com/provider/preferred-			

(Form continued on next page.)

Mer	nber's Last Name:	Member's First Name:
* De	efinitions of Accepted Drug Trial:	
•	Benzphetamine, diethylpropion, phendimetrazi 10 lbs	ne, phentermine: 3 month trial without a weight loss of
•	Orlistat: 6 month trial without a weight loss of 1	0 lbs
•	GLP-1 Receptor Agonist: 6 month trial without a	body weight reduction of 5%
LEN	GTH OF AUTHORIZATION	
□ I	nitial 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 be	elow (drug specific)
•	pound (lb.) weight loss during the initial 3 month	ne, phentermine – If the member achieves at least a 10 is of therapy, an additional 3-month SA may be granted. If months (waiting period of 6 months before next)
•		weight loss, an additional 6-month SA may be granted. 4 months (waiting period of 6 months before next
•	Imcivree® – If the member has experienced ≥ 5% those with continued growth potential), an additional states and the second seco	ional 1 year SA may be granted.
•	GLP-1 Receptor Agonists – If the member achieve compared to the most recent authorization, an a	res a weight loss of ≥ 5% reduction in body weight idditional 6-month SA may be granted.
All a	approvals are subject to the criteria on this form.	Existing authorizations will be honored until renewal.
	Attachments	

(Form continued on next page.)

Member's Last Name:	Member's First Name:	
Prescriber Signature (Required) By signature, the physician confirms the above inform and verifiable by member records.	Date ation is accurate	
Please include ALL requested information. Incomplete	e forms will delay the PA process.	

Submission of documentation does NOT guarantee coverage.