

Kaiser Permanente Health Plan of Mid-Atlantic States, Inc.
Weight Loss Agents Prior Authorization (PA)
Pharmacy Benefits Prior Authorization Help Desk
Length of Authorizations: Initial- 6 months; Continuation- 6 months

Instructions:

This form is used by Kaiser Permanente and/or participating providers for coverage of **Weight Loss Agents.** Please complete and fax this form back to Kaiser Permanente within 24 hours [fax: 1-866-331-2104]. If you have any questions or concerns, please call 1-866-331-2103. Requests will not be considered unless this form is complete. The KP-MAS Formulary can be found at: Pharmacy | Community Provider Portal | Kaiser Permanente

	1 – Patient Information	
Patient Name:	Kaiser Medical ID#:	Date of Birth:
	2 – Provider Information	
Provider Name:	Provider NPI:	
Provider Address:		
Provider Phone #:	Provider Fax #:	
Please check the boxes that apply: Initial Request- Complete question Continuation of Therapy Request-		
	3 – Pharmacy Information	
Pharmacy Name:	Pharmacy NPI:	
Pharmacy Phone #	Pharmacy Fax #:	
	4 – Drug Therapy Requested	
	:	
Drug 2: Name/Strength/Formulation	:	
	5 – Diagnosis	
☐ Initial Request		
Body Mass Index (BMI) Requirement	s: Adult Patients:	
☐ BMI ≥ 30, without risk factors		
□ BMI > 27, with > 2 of the following risk factors (select below)		

	ry Heart Disease		
□ Dyslipio			
	□ Hypertension		
□ Sleep A			
□ Type 2	Diabetes		
Pediatric Pa Imcivree	tients:		
□ BMI ≥ 30 or ≥ 95th percentile on pediatric growth chart			
Wegovy®, S	axenda®, and Zepbound		
□ Patient 12	2–18 years of age, with a BMI of ≥ 140% of the 95th percentile by age and sex		
□ Patient 12	2–18 years of age, with an initial BMI that is \ge 120% of the 95th percentile by age and sex with two or more of the		
following ris	sk factors: coronary heart disease, dyslipidemia, hypertension, sleep apnea, type 2 diabetes		
	6 – Clinical Criteria		
1 Λαο	requirements:		
_	Is the member ≥ 16 years old? (See specific age-related indications for Saxenda, Wegovy, Imcivree, Zepbound)		
	□ No □ Yes		
	Saxenda, Wegovy is the patient ≥ 12 years old?		
	□ No □ Yes		
	Imcivree, is the patient ≥ 6 years old?		
	□ No □ Yes		
	Zepbound, Contrave is the patient ≥ 18 years old?		
	□ No □ Yes		
	es the patient have any of the following contraindications? Alabsorption Syndromes Cholestasis Pregnant and/or Breastfeeding History of Eating Disorders		
	ronic Opioid Use (contraindicated with Contrave)		
	(enical)		
•	es provider attest that the patient's obesity is disabling and life threatening (i.e., puts the patient at risk for high-		
	bidity conditions): □ No □ Yes		
Imcivree ad	ditional Criteria:		
	Is therapy prescribed by or in consultation with an endocrinologist or geneticist? And,		
	□ No □ Yes		
	Does the member have proopiomelanocortin (POMC), proprotein convertase subtilisin/kexin type 1 (PCSK1), or		
	leptin receptor (LEPR) deficiency, as confirmed by a genetic test; AND		
	□ No □ Yes		
	Member's genetic variants are interpreted as pathogenic, likely pathogenic, or of uncertain significance (VUS) \Box No \Box Yes		
	Does the member have Bardet-Biedl syndrome (BBS)		
	□ No □ Yes		
	egovy, Zepbound additional Criteria:		
	Has member tried and failed a 30-day therapy of a non-GLP-1 weight loss drug in the last 6 months? □ No □ Yes; If yes, list therapy and outcome:		
	Does the member have a documented intolerance or allergy to a non-GLP-1 weight loss drug:		
•	□ No □ Yes; If yes, list therapy and outcome:Attest that		
	member is not concurrently on Victoza® or Ozempic® or other GLP-1 inhibitors □		

Contra	ve additional Criteria:		
	Member is not concurrently using opioids		
4.	4. Please provide documentation of the patient's current medical status, including:		
	Nutritional or Dietetic Assessment □ No □ Yes		
	Describe Current Weight Loss Plan with Diet and Exercise Components:		
5.	Initial Request Length of Approval Varies (Drug Specific):		
	Benzphetamine, diethylpropion, phendimetrazine, phentermine, Qsymia, Contrave®, – 3 months		
	 Wegovy/Zepbound™, Alli®/Xenical - 6 months 		
	Saxenda® and Imcivree™ – 4 months		
□ <u>Cont</u>	inuation of Therapy:		
Note:	Applicable to all WL therapies:		
	 Members lacking a weight loss response may still be considered for renewal with two or more of the following weight related risk factors: coronary heart disease, dyslipidemia, hypertension, sleep apnea, type 2 diabetes. At this time, authorization requests over one year are subject to initial criteria including all documentation. 		
	• In the event of an FDA recognized shortage, approved members will be eligible for the full allotment of approved drug once the shortage is resolved.		
1.	Check all that apply		
	Continue to meet Initial Therapy criteria □ Yes □ No BMI ≥ 24 □ Yes □ No		
_	Documentation of continued weight loss while on requested therapy		
	Initial BMI: Initial Weight:Initial Height: Current BMI or % of 95th percentile weight (12–18 y.o.):		
	urrent Weight: Current Height:		
	Other Diagnosis/Risk Factors:		
_	Current Medications:		
	Is there any contraindication for this use, malabsorption syndromes, cholestasis, pregnancy, and/or lactation?		
	□ No □ Yes		
3.	Summarize details of previous weight-loss treatment plans to include diet and exercise plans, in addition to		
	submitting a copy of the plan consistent with Question 4:		
4	Continuation of Therapy additional requirements (Drug Specific):		
du	nzphetamine, diethylpropion, phendimetrazine, phentermine – If the member achieves at least a 10 lb. weight loss ring the initial 3 months of therapy, an additional 3-month SA may be granted. Maximum length of continuous drug erapy is 6 months (waiting period of 6 months before next request).		

next request).

Qsymia® – If the member achieves a weight loss of at least 3% of baseline weight, an additional 3-month SA may be granted. For a subsequent renewal, member must meet a weight loss of at least 5% of baseline weight to qualify for an additional 6-month SA. Maximum length of continuous drug therapy is 12 months (waiting period of 6 months before

- **Xenical**®/Alli 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).
- Contrave® Approve for 6 months with each renewal if weight reduction continues.
- Saxenda® If the member achieves a weight loss of at least 4% of baseline weight, an additional 6-month SA may be granted as long as weight reduction continues.
- ImcivreeTM If the member has experienced $\geq 5\%$ reduction in body weight (or $\geq 5\%$ of baseline BMI in those with continued growth potential), an additional 1 year SA may be granted.

Wegovy/Zepbound™ - If the member achieves a weight loss of at least 5% of baseline weight, an additional 6 month SA may be granted.

7 – Provider Sign-Off		
Additional Information –		
 Please submit chart notes/medical records for the patient that are applicable to the patient that are applicable to the provide rational of the provide rational of the requested medication information that should be taken into consideration for the requested medication in the requested medication in the patient that are applicable to the patient that are applicable that are applicable to the patient that are applicable to the patient that are applicable to the patient that are applicable	and any additional supporting	
I certify that the information provided is accurate. Supporting documentation is	available for State audits.	
Provider Signature:	Date:	
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