

Anorexiant Prior Authorization Request Form (Page 1 of 2)

Member Information (required)			Provider Information (required)		
Member Name:			Provider Name:		
Insurance ID#:			NPI#:		Specialty:
Date of Birth:			Office Phone:		
Street Address:			Office Fax:		
City:	State:	Zip:	Office Street Address:		
Phone:			City:	State:	Zip:
Medication Information (required)					
Medication Name:			Strength:		Dosage Form:
<input type="checkbox"/> Check if requesting brand			Directions for Use:		
<input type="checkbox"/> Check if request is for continuation of therapy					
Clinical Information (required)					
Select the diagnosis below: <input type="checkbox"/> Appetite suppression <input type="checkbox"/> Weight loss <input type="checkbox"/> Other diagnosis: _____ ICD-10 Code(s): _____					
Lifestyle modification: Is the requested medication being used as an adjunct to lifestyle modification (e.g., dietary or caloric restriction, exercise, behavioral support, community based program)? <input type="checkbox"/> Yes <input type="checkbox"/> No Body Mass Index (BMI): What is the patient's current BMI? _____ kg/m ² Comorbidities: Does the patient have a weight-related comorbidity (e.g., hypercholesterolemia, hypertension, diabetes, sleep apnea)? <input type="checkbox"/> Yes <input type="checkbox"/> No					
Medication history: Is the requested medication being used in combination with another anti-obesity agent? <input type="checkbox"/> Yes <input type="checkbox"/> No For Belviq, Belviq XR, Qsymia and Saxenda requests, also answer the following: Has the patient failed to lose greater than or equal to 5% of baseline body weight after at least 16 weeks (one full course) of Contrave therapy? <input type="checkbox"/> Yes <input type="checkbox"/> No Does the patient have an intolerance or contraindication to Contrave therapy? <input type="checkbox"/> Yes <input type="checkbox"/> No					
Reauthorization: Has the patient had weight loss of greater than or equal to 5% of baseline body weight? <input type="checkbox"/> Yes <input type="checkbox"/> No Is the patient continuing to practice lifestyle modification? <input type="checkbox"/> Yes <input type="checkbox"/> No Is the requested medication being used in combination with another anti-obesity agent? <input type="checkbox"/> Yes <input type="checkbox"/> No					
Quantity limit requests: What is the quantity requested per DAY? _____ What is the reason for exceeding the plan limitations? <input type="checkbox"/> Titration or loading dose purposes <input type="checkbox"/> Patient is on a dose-alternating schedule (e.g., one tablet in the morning and two tablets at night, one to two tablets at bedtime) <input type="checkbox"/> Requested strength/dose is not commercially available <input type="checkbox"/> Other: _____					

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Are there any other comments, diagnoses, symptoms, medications tried or failed, and/or any other information the physician feels is important to this review?

Please note: This request may be denied unless all required information is received.

Please fax this form to 1-844-712-8129 to initiate a prior authorization review for the member and medication above.

Please note: plan benefits may limit or exclude coverage of specific medications including those requested on this form.

I certify, to the best of my knowledge, the statements and information provided on this form are factual and correct.

Provider/Representative (and Title): _____ Date: _____

PROACT INTERNAL USE ONLY:

Clinical Review Decision	
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	Approved, through
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	Denied (documentation attached, if necessary)
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Tracking:				
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1 st Attempt		2 nd Attempt		Letter Mailed:	
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