

CNS Stimulants Prior Authorization Request Form (Page 1 of 2)

Member Information <small>(required)</small>			Provider Information <small>(required)</small>		
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 <small>(required)</small>					
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 <small>(required)</small>					
Select the diagnosis below: <input type="checkbox"/> Attention deficit disorder (ADD) or attention deficit hyperactivity disorder (ADHD) <input type="checkbox"/> Obesity [Desoxyn (methamphetamine) & Evekeo only] <input type="checkbox"/> Binge eating disorder (BED) – moderate to severe [Vyvanse only] <input type="checkbox"/> Narcolepsy (confirmed by sleep study) <input type="checkbox"/> Other diagnosis: _____ ICD-10 Code(s): _____ <input type="checkbox"/> Depression (for augmentation of antidepressant therapy)					
ADD or ADHD: Have symptoms been present prior to 12 years of age? <input type="checkbox"/> Yes <input type="checkbox"/> No Does the patient have symptoms which interfere with or reduce the quality of academic or occupational functioning? <input type="checkbox"/> Yes <input type="checkbox"/> No Was the prescription for the requested medication written by or in consultation with a mental health specialist? <input type="checkbox"/> Yes <input type="checkbox"/> No					
Binge eating disorder (BED) – moderate to severe [Vyvanse only]: Has the patient had binge eating disorder for 3 months or longer? <input type="checkbox"/> Yes <input type="checkbox"/> No Does the patient have between 4 and 13 binge-eating episodes per week? <input type="checkbox"/> Yes <input type="checkbox"/> No Does the patient eat much more rapidly than normal? <input type="checkbox"/> Yes <input type="checkbox"/> No Does the patient eat until feeling uncomfortably full? <input type="checkbox"/> Yes <input type="checkbox"/> No Does the patient eat large amounts of food when not feeling physically hungry? <input type="checkbox"/> Yes <input type="checkbox"/> No Does the patient eat alone because of feeling embarrassed by how much he or she is eating? <input type="checkbox"/> Yes <input type="checkbox"/> No Does the patient feel disgusted with himself or herself, depressed, or very guilty after binge-eating? <input type="checkbox"/> Yes <input type="checkbox"/> No					
Reauthorization [Vyvanse only]: If this is a reauthorization request, answer the following: Is there documentation of positive clinical response (e.g., meaningful reduction in the number of binge eating episodes or binge days per week from baseline, improvement in the signs and symptoms of binge eating disorder) to Vyvanse therapy? <input type="checkbox"/> Yes <input type="checkbox"/> No					
Narcolepsy: Has a sleep study confirming the diagnosis of narcolepsy been performed and the results submitted with this form? <input type="checkbox"/> Yes <input type="checkbox"/> No If there is a reason that a sleep study would not be feasible, please explain in the comments section below.					
Obesity [Desoxyn (methamphetamine) & Evekeo only]: Is the requested medication used for short term (i.e., a few weeks) adjunct therapy for exogenous obesity? <input type="checkbox"/> Yes <input type="checkbox"/> No Is the patient refractory to alternative therapy (e.g., repeated diets, group programs, medications)? <input type="checkbox"/> Yes <input type="checkbox"/> No Does the patient have a body mass index (BMI) greater than or equal to 30kg/m ² ? <input type="checkbox"/> Yes <input type="checkbox"/> No Does the patient have a BMI greater than or equal to 27kg/m ² ? <input type="checkbox"/> Yes <input type="checkbox"/> No Does the patient have a weight-related comorbidity (e.g., hypercholesterolemia, hypertension, diabetes, sleep apnea)? <input type="checkbox"/> Yes <input type="checkbox"/> No					
Select the medications the patient has a failure, contraindication, or intolerance to: <input type="checkbox"/> Amphetamine-dextroamphetamine <input type="checkbox"/> Dexmethylphenidate <input type="checkbox"/> Dextroamphetamine ER <input type="checkbox"/> Methylphenidate <input type="checkbox"/> Amphetamine-dextroamphetamine extended-release (ER) <input type="checkbox"/> Dexmethylphenidate ER <input type="checkbox"/> Metadate ER <input type="checkbox"/> Methylphenidate ER <input type="checkbox"/> Adderall XR <input type="checkbox"/> Dextroamphetamine <input type="checkbox"/> Methamphetamine <input type="checkbox"/> Vyvanse <input type="checkbox"/> Other CNS stimulant(s). Please specify: _____					

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Quantity limit requests:

What is the quantity requested per DAY? _____

Is the prescription written by or in consultation with a mental health specialist? Yes No

Will adequate scientific literature, peer-reviewed medical literature, or national compendia supporting the use of higher doses be provided? Yes No

***Please note: Submission of information requested above is required for quantity limit requests for this drug.*

What is the reason for exceeding the plan limitations?

- Titration
- Loading dose purposes
- 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)
- Requested strength/dose is not commercially available
- Other: _____

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

Approved, through

Denied (documentation attached, if necessary)

Tracking:

1 st Attempt		2 nd Attempt		Letter Mailed:	
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