

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

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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 requested drug below:		
<input type="checkbox"/> Amphetamine-dextroamphetamine (Adderall) <input type="checkbox"/> Amphetamine-dextroamphetamine ER (Adderall XR) <input type="checkbox"/> Cotempla XR-ODT <input type="checkbox"/> Daytrana <input type="checkbox"/> Dexmethylphenidate (Focalin) <input type="checkbox"/> Dexmethylphenidate ER (Focalin XR)	<input type="checkbox"/> Dextroamphetamine ER (Dexedrine) <input type="checkbox"/> Dextroamphetamine solution (Procentra) <input type="checkbox"/> Dextroamphetamine tablet (Zenzedi) <input type="checkbox"/> Metadate ER <input type="checkbox"/> Methamphetamine (Desoxyn) <input type="checkbox"/> Methylphenidate (Ritalin) <input type="checkbox"/> Methylphenidate chewable tablet	<input type="checkbox"/> Methylphenidate ER (Concerta) <input type="checkbox"/> Methylphenidate ER capsule (Ritalin LA) <input type="checkbox"/> Methylphenidate ER tablet <input type="checkbox"/> Methylphenidate solution (Methylin) <input type="checkbox"/> Mydayis <input type="checkbox"/> Vyvanse capsule <input type="checkbox"/> Vyvanse chewable tablet

Select the diagnosis below:

Attention deficit hyperactivity disorder (ADHD)/Attention deficit disorder (ADD)
Please complete corresponding section below and submit any documentation as requested

Narcolepsy
Please submit documentation (i.e., polysomnography and MSLT) to support diagnosis

Other diagnosis: _____ ICD-10 Code(s): _____
Please submit documentation (i.e., chart notes, previous therapies tried) to support diagnosis

Please note: Non-FDA approved indications - Submit two (2) peer reviewed clinical studies documenting the safety and efficacy of the specified drug for that particular indication.

Attention deficit hyperactivity disorder (ADHD)/Attention deficit disorder (ADD):

Select if there is documentation of one of the following:

Existence of 5 or more Inattentive Symptoms for a minimum of 6 months

Existence of 5 or more Hyperactive-Impulsive Symptoms for a minimum of 6 months

Existence of 10 or more Combined Symptoms for a minimum of 6 months (including 5 or more inattentive symptoms AND 5 or more hyperactive-impulsive symptoms)

Is there documentation that symptoms impair or compromise normal functioning? Yes No

Is there documentation that symptoms are present in two (2) or more settings/environments? Yes No

If **yes**, indicate the settings: 1. _____ 2. _____

Are the patient's symptoms better explained by another disorder (e.g., Schizophrenia, Mood Disorder, Anxiety Disorder, Substance Abuse, Dissociative Disorder, or Personality Disorder)? Yes No

Has the provider submitted the patient-specific DSM symptoms, criteria, psychological evaluation, and/or standardized rating scale used to make or verify the diagnosis? Yes No

Please note: The patient-specific DSM symptoms, criteria, psychological evaluation, and/or standardized rating scale used to make or verify the diagnosis must be submitted with this form for approval

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Quantity limit requests:

What is the quantity requested per DAY? _____

Previous therapies failed and/or therapies currently used in combination with the requested medication (*List ALL medications tried or authorization process will be delayed*): _____

Is the prescribed dose higher than the maximum dose recommendation in FDA-approved labeling (i.e., the package insert)? Yes No

If **yes**, please provide documentation to support the safety and efficacy of the higher dose (such as evidence from practice guidelines or clinical trials from peer-reviewed medical literature).

**** Please note: Chart documentation of the above is required to be submitted along with this fax**

****Use of samples to initiate therapy does not meet step edit/ preauthorization criteria.****

Previous therapies will be verified through pharmacy paid claims or submitted chart notes.

Prescriber Signature: _____ Date: _____

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.
For urgent or expedited requests please call 1-800-711-4555.
This form may be used for non-urgent requests and faxed to 1-800-527-0531.