



Request for Prior Authorization
CNS STIMULANTS AND ATOMOXETINE

(PLEASE PRINT - ACCURACY IS IMPORTANT)

Form with fields for IA Medicaid Member ID #, Patient name, DOB, Patient address, Provider NPI, Prescriber name, Phone, Prescriber address, Fax, Pharmacy name, Address, Phone, Pharmacy NPI, Pharmacy fax, NDC.

Prior Authorization (PA) is required for CNS stimulants and atomoxetine for patients 21 years of age or older. Prior to requesting PA for any covered diagnosis, the prescriber must review the patient's use of controlled substances on the Iowa Prescription Monitoring Program (PMP) website at https://pmp.iowa.gov/IAPMPWebCenter/. Payment for CNS stimulants and atomoxetine will be considered under the following conditions: 1) Attention Deficit Disorder (ADD) or Attention Deficit Hyperactivity Disorder (ADHD) meeting the DSM-5 criteria and confirmed by a standardized rating scale (such as Conners, Vanderbilt, Brown, Snap-IV). Symptoms must have been present before twelve (12) years of age and there must be clear evidence of clinically significant impairment in two or more current environments (social, academic, or occupational). Documentation of a recent clinical visit that confirms the patient continues to require medication to treat the symptoms of ADD/ADHD will be required for renewals or patients newly eligible that are established on medication to treat ADD/ADHD. 2) Narcolepsy with diagnosis confirmed with a recent sleep study (ESS, MSLT, PSG). 3) Excessive sleepiness from obstructive sleep apnea/hypopnea syndrome (OSAHS) with documentation of non-pharmacological therapies tried (weight loss, position therapy, CPAP at maximum titration, BiPAP at maximum titration or surgery) and results from a recent sleep study (ESS, MSLT, PSG) with the diagnosis confirmed by a sleep specialist.

Payment for a non-preferred agent will be authorized only for cases in which there is documentation of previous trial and therapy failure with a preferred agent. \* If a non-preferred long-acting medication is requested, a trial with the preferred immediate release and extended release product of the same chemical entity (methylphenidate class) or chemically related agent (amphetamine class) is required. The required trials may be overridden when documented evidence is provided that the use of these agents would be medically contraindicated.

Requests for Vyvanse for Binge Eating Disorder must be submitted on the Binge Eating Disorder Agents PA form.

Preferred

- Adzenys XR ODT
Amphetamine Salt Combo
Daytrana
Dexmethylphenidate Tabs
Focalin XR
Metadate CD
Methylin Solution
Methylphenidate IR Tabs
Methylphenidate ER Tabs (18, 27, 36, 54mg)

- Modafinil
Quillichew ER
Quillivant XR
Ritalin LA
Strattera
Vyvanse

Non-Preferred

- Adderall
Adderall XR\*
Amphetamine ER\*
Aptensio XR\*
Concerta
Desoxyn
Dexedrine\*
Dextroamphetamine ER Cap\*
Dexmethylphenidate
Dexmethylphenidate ER\*
Dyanavel XR
Evekeo

- Focalin
Methylin Chew
Methylphenidate ER Tabs (10mg & 20mg)\*
Methylphenidate ER Capsules\*
Nuvigil
Procentra
Provigil
Ritalin
Ritalin SR\*

Strength Dosage Instructions Quantity Days Supply

**Request for Prior Authorization  
CNS STIMULANTS AND ATOMOXETINE**

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**Diagnosis:**

- Attention Deficit Disorder (ADD)**
- Attention Deficit Hyperactivity Disorder (ADHD)**

Age of patient at onset of symptoms: \_\_\_\_\_

Date of most recent clinical visit confirming initial or continued diagnosis: \_\_\_\_\_

Rating scale used to determine diagnosis: \_\_\_\_\_

Documentation of clinically significant impairment in two or more **current** environments (social, academic, or occupational).

Current Environment 1 & description: \_\_\_\_\_

Current Environment 2 & description: \_\_\_\_\_

- Narcolepsy (Please provide results from a recent ESS, MSLT, and PSG)**

- Excessive sleepiness from obstructive sleep apnea/hypopnea syndrome (OSAHS)**

Have non-pharmacological treatments been tried?  No  Yes *If Yes, please indicate below:*

Weight Loss

Position therapy

CPAP Date: \_\_\_\_\_

Maximum titration?  Yes  No

BiPAP Date: \_\_\_\_\_

Maximum titration?  Yes  No

Surgery Date: \_\_\_\_\_

Specifics: \_\_\_\_\_

Diagnosis confirmed by a sleep specialist?  Yes  No

- Other (specify)** \_\_\_\_\_

**Prescriber review of patient's controlled substances use on the Iowa PMP website:**

No  Yes Date Reviewed: \_\_\_\_\_

Please document prior psychostimulant trial(s) and failures(s) including drug name(s) strength, dose, exact date ranges and failure reasons: \_\_\_\_\_

**Other** - Please provide all pertinent medication trial(s) relating to the diagnosis including drug name(s) strength, dose and exact date ranges:

Reason for use of Non-Preferred drug requiring approval: \_\_\_\_\_

Prescriber signature (Must match prescriber listed above.)	Date of submission
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**IMPORTANT NOTE:** *In evaluating requests for prior authorization the consultant will consider the treatment from the standpoint of medical necessity only. If approval of this request is granted, this does not indicate that the member continues to be eligible for Medicaid. It is the responsibility of the provider who initiates the request for prior authorization to establish by inspection of the member's Medicaid eligibility card and, if necessary by contact with the county Department of Human Services, that the member continues to be eligible for Medicaid.*