



Kaiser Permanente Health Plan of Mid-Atlantic States, Inc.  
Attention Deficit-Hyperactive (ADHD) & Narcolepsy  
Prior Authorization (PA)  
Pharmacy Benefits Prior Authorization Help Desk  
Length of Authorizations: Initial- 12 months; Continuation- 12 months

**Instructions:**

This form is used by Kaiser Permanente and/or participating providers for coverage of **Attention Deficit-Hyperactive (ADHD) & Narcolepsy**. Please complete all sections, incomplete forms will delay processing. 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 all sections are complete.

KP-MAS Formulary can be found at: [Pharmacy | Community Provider Portal | Kaiser Permanente](#)

**Medications:**

<ul style="list-style-type: none"><li>• COTEMPLA XR-ODT TBED (6 to 17 years)</li><li>• DAYTRANA PTCH (6 to 17 years)</li></ul>	<ul style="list-style-type: none"><li>• QUILLICHEW ER CHER (≥6 years)</li><li>• QUILLIVANT XR SRER (≥6 years)</li></ul>
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**1 – Patient Information**

Patient Name: \_\_\_\_\_ Kaiser Medical ID#: \_\_\_\_\_ Date of Birth: \_\_\_\_\_

**2 – Prescriber Information**

Prescriber Name: \_\_\_\_\_ Specialty: \_\_\_\_\_ NPI: \_\_\_\_\_  
Prescriber Address: \_\_\_\_\_  
Prescriber Phone #: \_\_\_\_\_ Prescriber Fax #: \_\_\_\_\_

**3 – Pharmacy Information**

Pharmacy Name: \_\_\_\_\_ Pharmacy NPI: \_\_\_\_\_  
Pharmacy Phone # \_\_\_\_\_ Pharmacy Fax #: \_\_\_\_\_

**4 – Drug Therapy Requested**

Drug 1: Name/Strength/Formulation: \_\_\_\_\_  
Sig: \_\_\_\_\_  
Drug 2: Name/Strength/Formulation: \_\_\_\_\_  
Sig: \_\_\_\_\_

**5– Diagnosis/Clinical Criteria**

- 1. Is this request for initial or continuing therapy?  
 Initial therapy                       Continuing therapy, State date: \_\_\_\_\_
  
- 2. Indicate the patient’s diagnosis for the requested medication: \_\_\_\_\_

**Clinical Criteria:**

- 1. Indicate the member’s age: \_\_\_\_\_.
  
- 2. **AND** member has a diagnosis of Attention Deficit Hyperactivity Disorder (ADHD) as confirmed by psychoeducational testing\*\*\*,  
 No  Yes
  
- 3. **AND** member has had an adequate trial\* (1 week) and/or intolerance\*\* or allergy to dextroamphetamine-amphetamine (generic Adderall XR), intermediate or long-acting methylphenidate (methylphenidate SR, methylphenidate CD, or methylphenidate ER), and dexmethylphenidate (generic Focalin XR)?  
 No  Yes

**For continuation of therapy, please respond to additional questions below:**

- 1. Has the member continued to meet the initial review criteria and has demonstrated positive clinical response to medication?  
 No  Yes

**NOTES:**

\*Adequate trial of a long-acting agent is further defined as wearing off that is not resolved by increasing the dose, AND adding a short-acting agent OR increasing frequency to twice daily OR clinically significant side effects related to the dosage form that cannot be resolved by adjusting the dose or timing  
\*\*Intolerance excludes adverse drug reactions that are expected, mild in nature, resolve with continued treatment and do not require medication discontinuation  
\*\*\*Criteria only applies for 18 years of age and older

**6 – Prescriber Sign-Off**

**Additional Information –**

- 1. **Please submit chart notes/medical records for the patient that are applicable to this request.**
- 2. **If member has not tried preferred agent(s) please provide rationale/explanation and any additional supporting information that should be taken into consideration for the requested medication:**

\_\_\_\_\_  
\_\_\_\_\_  
\_\_\_\_\_

**I certify that the information provided is accurate. Supporting documentation is available for State audits.**

<b>Prescriber Signature:</b>	<b>Date:</b>
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