

# Prior Authorization Request Form for Stimulant and Related Agents

**FAX this completed form to (877) 386-4695**

**OR Mail requests to: Envolve Pharmacy Solutions PA Department | 5 River Park Place East, Suite 210 | Fresno, CA 93720**

I. PROVIDER INFORMATION		II. MEMBER INFORMATION	
Prescriber Name:		Member Name:	
Prescriber Specialty:		Identification #:	
Office Contact Name:		Group #:	
Group Name:		Date of Birth:	
Fax #:		Medication Allergies:	
Phone #:			
III. DRUG INFORMATION (One drug request per form)			
Drug name and strength:	Dosage Interval (sig):	Qty. per Day:	
IV. REQUIRED DOCUMENTATION (Detailed medical record documentation demonstrating evidence for each item must be submitted with prior authorization request)			
Specify diagnosis & diagnosis code relevant to this request:		Dx/Dx Code: _____	
Did the prescriber or prescriber's delegate search the PDMP to review the member's controlled substance prescription history before issuing this prescription for the requested agent?		<input type="checkbox"/> Yes <input type="checkbox"/> No	
<b>Requests for all non-preferred medications:</b> Does the member have a history of trial and failure of or contraindication or intolerance to the preferred Stimulant and Related agents? Refer to <a href="https://papdl.com/preferred-drug-list">https://papdl.com/preferred-drug-list</a> for a list of preferred and non-preferred medications in this class.		<input type="checkbox"/> Yes <input type="checkbox"/> No	
<input type="checkbox"/> Member has a current history (within past 90 days) of using the prescribed the requested non-preferred Stimulant and Related agent: _____ <input type="checkbox"/> Member is not receiving concurrent treatment with a sedative hypnotic <input type="checkbox"/> Member was assessed for potential risk of misuse, abuse, or addiction based on family and social history obtained by prescribing provider <input type="checkbox"/> Member has been educated on the potential adverse effects of stimulants, including the risk for misuse, abuse, and addiction <input type="checkbox"/> For member's with a history of comorbid substance dependency, abuse, or diversion has results of a recent urine drug screen testing for licit and illicit drugs with the potential for abuse (including oxycodone, fentanyl and tramadol) <input type="checkbox"/> If requesting for daily quantity exceeding daily limit (Refer to <a href="https://www.dhs.pa.gov/providers/Pharmacy-Services/Pages/Quantity-Limits-and-Daily-Dose-Limits.aspx">https://www.dhs.pa.gov/providers/Pharmacy-Services/Pages/Quantity-Limits-and-Daily-Dose-Limits.aspx</a> ), please provide supporting information: _____			
<b>Therapeutic Duplication:</b> If concurrently prescribed a therapeutic duplicate (i.e. stimulant different from the agent being requested): <input type="checkbox"/> Member is transitioned from one stimulant and related agent to another with the intent of discontinuing one of the medications <input type="checkbox"/> Member has a medical reason for concomitant use of the requested medications that is supported by peer-reviewed literature or national treatment guidelines			
<b>SUBMIT MEDICAL RECORD INFORMATION FOR EACH APPLICABLE ITEM.</b> <b>ATTENTION DEFICIT HYPERACTIVITY DISORDER (ADHD):</b> <input type="checkbox"/> Member has a diagnosis of ADHD confirmed according to the current Diagnostic and Statistical Manual of Mental Disorders (DSM) criteria			

**MODERATE TO SEVERE BINGE EATING DISORDER:**

- ☐ Member has a diagnosis of moderate to severe binge eating disorder confirmed according to the current Diagnostic and Statistical Manual of Mental Disorders (DSM) criteria
- ☐ Member does not have ADHD and has documented history of therapeutic failure, contraindication or intolerance to at least 1 of the following: (medication, start date and end date)
  - ☐ Topiramate: \_\_\_\_\_
  - ☐ Selective Serotonin Reuptake Inhibitor (SSRI): \_\_\_\_\_
- ☐ Member has a documentation of a referral for cognitive behavioral therapy or other psychotherapy

**NARCOLEPSY:**

- ☐ Member has a diagnosis of narcolepsy confirmed according to the most recent consensus treatment guidelines (e.g. American Academy of Sleep Medicine International Classification of Sleep Disorders)

**RENEWAL REQUESTS:**

- ☐ Documentation of tolerability and experienced a positive clinical response to requested medication evidenced by: \_\_\_\_\_

**IV. ADDITIONAL RATIONALE FOR REQUEST / PERTINENT CLINICAL INFORMATION :**

Appropriate clinical information to support the request on the basis of medical necessity must be submitted.	Provider Signature:	Date:
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Envolve Pharmacy Solutions will respond via fax or phone within 24 hours.

Requests for prior authorization (PA) requests must include member name, ID#, and drug name. Please include lab reports with requests when appropriate (e.g., Culture and Sensitivity; Hemoglobin A1C; Serum Creatinine; CD4; Hematocrit; WBC, etc.)