

Prior Authorization Request Form for Opioid Dependence Treatment

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: _____	
Does the member have a history of contraindication to the prescribed medication?	<input type="checkbox"/> Yes <input type="checkbox"/> No <i>Submit documentation.</i>		
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 <i>Submit documentation.</i>		
<input type="checkbox"/> If requesting for daily quantity exceeding daily limit (Refer to https://www.dhs.pa.gov/providers/Pharmacy-Services/Pages/Quantity-Limits-and-Daily-Dose-Limits.aspx), please provide supporting information: _____			
Therapeutic Duplication: If concurrently prescribed a therapeutic duplicate (i.e. another Androgenic Agent or dose different from the agent being requested): <ul style="list-style-type: none"> <input type="checkbox"/> is being transitioned from one Androgenic Agent to another with the intent of discontinuing one of the medications <input type="checkbox"/> has a medical reason for concomitant use of the requested medications that is supported by peer-reviewed literature or national treatment guidelines 			
SUBMIT MEDICAL RECORD INFORMATION FOR EACH APPLICABLE ITEM.			
BUPRENORPHINE WITHOUT NALOXONE: <ul style="list-style-type: none"> <input type="checkbox"/> Member meets one of the following: <ul style="list-style-type: none"> <input type="checkbox"/> Prescribed for induction therapy <input type="checkbox"/> Pregnant <input type="checkbox"/> Breastfeeding <input type="checkbox"/> History of contraindication or intolerance to naloxone 			
NON-PREFERRED OPIOID DEPENDENCE TREATMENT: <ul style="list-style-type: none"> <input type="checkbox"/> Oral Buprenorphine, has a therapeutic failure, contraindication or intolerance to the preferred oral buprenorphine Opioid Dependence Treatment: _____ <input type="checkbox"/> Alpha-2 Adrenergic Agonist, has a therapeutic failure, contraindication or intolerance to the preferred alpha-2 adrenergic agonist Opioid Dependence Treatment: _____ <input type="checkbox"/> Non-Oral Buprenorphine, has a therapeutic failure, contraindication or intolerance to the preferred non-oral buprenorphine Opioid Dependence Treatment: _____ 			

REQUEST FOR ORAL BUPRENORPHINE ABOVE 24MG PER DAY:

- ☐ Prescribed daily dose is consistent with medically accepted prescribing practices and standard of care
- ☐ Documentation of an evaluation to determine the recommended level of care
- ☐ Documentation of member is in a substance abuse or behavioral health counseling or treatment program or an addiction recovery program
- ☐ Member has urine drug screen for drugs with potential for abuse
- ☐ For members already on buprenorphine, the member has a recent urine drug screen positive for buprenorphine and norbuprenorphine

RENEWAL REQUESTS:

- ☐ Member has experienced a positive clinical response as 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.)