

Prior Authorization Request Form for Opioid Use Disorder Treatments

FAX this completed form to (844) 205-3386

OR Mail requests to: Pharmacy Department | 5 River Park Place East, Suite 210 | Fresno, CA 93720 OR Prior authorization may be completed at https://www.covermymeds.com/main/prior-authorization-forms/

OR Prior authorization may be co	ompieted at <u>nttps://www.co</u>	overmymeus.com/main/j	orior-authorization-torms/	
□New request □Renewal request total	# pages: Presc	Prescriber name:		
Name of office contact:		Specialty:		
Contact's phone number:		PI: State license #:		
Facility contact name/phone:		Street address:		
Member name: Cit		City/state/zip:		
Member ID#:	OB: Phone	9:	Fax:	
CLINICAL INFORMATION				
Drug requested:		Strength:	Dosage form:	
Directions:		Quantity:	Requested duration:	
Diagnosis (submit documentation):			Dx code (required):	
 Pennsylvania law requires prescribers to query the PA PDMP each time a patient is prescribed an opioid drug product or benzodiazepine. Naloxone is available at Pennsylvania pharmacies via standing order from the Secretary of the Department of Health. Pennsylvania Medical Assistance beneficiaries may obtain naloxone <u>free-of-charge</u> through their prescription drug benefit. Complete all sections that apply to the member and this request. Check all that apply and <u>submit documentation</u> for each item. 				
1. For a NON-PREFERRED SUBLINGUAL buprenorphine product (eg, film, tablet): Tried and failed or has a contraindication or an intolerance to the preferred SUBLINGUAL buprenorphine Opioid Use Disorder Treatments (Refer to https://papdl.com/preferred-drug-list for a list of preferred and non-preferred drugs in this class.) (medication, start and end date):				
2. For a non-preferred NON-SUBLINGUAL buprenorphine product (eg, injection): Tried and failed or has a contraindication or an intolerance to the preferred NON-SUBLINGUAL buprenorphine Opioid Use Disorder Treatments (Refer to https://papdl.com/preferred-drug-list for a list of preferred and non-preferred drugs in this class.) (medication, start and end date):				
3. For Lucemyra (lofexidine): Tried and failed or has a contraindication or an intolerance to clonidine tablet:				
4. For a SUBLINGUAL buprenorphine product ABOVE THE DAILY DOSE LIMIT OF 24 MG of buprenorphine per day: Is prescribed a daily dose consistent with medically accepted prescribing practices and standards of care Had an unsatisfactory clinical response (eg, uncontrolled withdrawal or cravings) at the current quantity limit of 24 mg per day If already established on buprenorphine, has results of a recent UDS demonstrating compliance with sublingual buprenorphine therapy				

ADDITIONAL RATIONALE FOR REQUEST / PERTINENT CLINICAL I	NFORMATION		
PLEASE <u>FAX</u> COMPLETED FORM WITH <u>REQUIRED CLINICAL DOCUMENTATION</u> TO (844) 205-3386			
Prescriber Signature:	Date:		

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