

Chronic Opioid Prior Authorization Form

The info requested in this form, although extensive, is based on best practice standards and the CDC Chronic Pain Opioid Guidelines. It is intended to facilitate the safe and effective treatment, improve outcomes, and reduce adverse events including opioid use disorder and/or overdose.

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Date	Diagnosis:									
		Reques	sted Medica	ation(s)						
Medication	Strength	Directi	Direction for use		Quantity requested per 30 days					
					□ 30 □ 60 □	□ 90	□ 120 □ O	ther:		
					□ 30 □ 60 □	□ 90	□ 120 □ O	ther:		
					□30 □60 □	□ 90 □	□ 120 □ O	ther:		
		Patie	ent Inform	ation						
Patient's Last Name:	First:	Middle:		Membe	er ID Number: Date of Birth:					
Street Address:		·	City				1			
State:	Zip Code:	Sex	□ M	□ F	Race/Ethnicity:					
		Prescri	iber Infor	mation	1					
Prescriber's Last Name	First:	Middle	e	Prescribe	er's NPI #:	Pres	scriber's DI	EA #:		
Prescriber's Specialty:					d cash for this off management	fice visit	t or for the		No	
Street Address:			City:							
State:	Zip:		Phone Num	umber:		Fax Nur	mber:			
		Pharm	acy Infor	mation						
Name:			Phone Nun	nber:						
		Medic	al inform	ation						
Please attach or list pat (Non-Pharmacological,						included	for each.			
Current treatments				Previously failed pain treatments of any/all types						
Is the patient pregnant?						☐ Yes	□ No			
Is the patient allergic to	e list and des	cribe reac	tions in 2 to 3 wor	ds)		☐ Yes	□ No			
Does the patient have n (If No, please provide G			tively)					☐ Yes	□ No	

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Physical exam findings relevant to pain diagnosis (Please briefly describe after Height, Weight, & Vital Signs)											
Height:	ght: Weight: Blood Pressure:		Heart Rate: Respire		Respiratory	tory Rate:					
Laboratory findings relevant to pain diagnosis (Please attach and/or briefly describe)											
Radiological findings (MRI, X-Ray, or Ultras	ound) relevant to the j	pain diagnosis (l	Please briefly describe)							
Has the patient experienced a decrease in his/her daily function (i.e. ability to climb stairs, complete house work, perform tasks, etc.) beyond a subjective increase in daily pain?							□ No				
Has the patient been screened for risk of substance-use disorder? (Please indicate risk screening tool, result, and attach)											
☐ Opioid Risk Tool (O☐ Drug Abuse Screeni			-	oid Misuse Measure (COM) Drug Use Questionnaire (I							
☐ Diagnosis, Intractab	oility, Risk, & Efficacy	Score (DIRE)	☐ Pain Medica	ation Questionnaire (PMQ)							
What was the patient's	risk of substance abus	se based on the above	screening tool?	□Low	✓ □ Moder	ate 🗆	High				
Does the patient currently have an up-to-date & signed Patient & Provider Agreement (Please Attach) including:											
Treatment time fr	of reducing pain and i ame with a planned er ociated risks of opioid	nd point as appropriate				□ Yes	□ No				
Has the patient been educated on the proper storage/disposal of controlled substances?							□ No				
Patient's opioid daily dose is >50MME/day. The CDC Opioid guidelines recommend education & utilization of naloxone.											
Has the patient been e	ducated on being a car	ndidate for carrying na	aloxone?			□ Yes	□ No				
Has the patient been p	rescribed naloxone?					☐ Yes	□ No				
	quires initial and at lea wed immediately pric	or to the prescribing of	of the requested	-	DMP). Has	□ Yes	□ No				
This confirmatory inform	nation will be shared wit	h the WV PDMP adminis	stration.								
Has a Urine Drug Screening been completed prior to the prescribing of the requested opioid medication?							□ No				
Were the results consis	stent with current trea	tment and devoid of il	licit substance?	(Please submit results with	each request.)	□ Yes	□ No				
Practitioner Signat	ure:			Date							
(If a signature stamp is used,	then the prescribing practition	oner must initial the signatur	re, signatures by age	nts of the practitioner are not accep	table)						

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