

Short-Acting Opioids

Patient Information	n:
Name:	
Member ID:	
Address:	
City, State, Zip:	
Date of Birth:	
Prescriber Informa	ition:
Name:	
NPI:	
Phone Number:	
Fax Number	
Address:	
City, State, Zip:	
Requested Medica	tion
Rx Name:	
Rx Strength	
Rx Quantity:	
Rx Frequency:	
Rx Route of	
Administration:	
Diagnosis and ICD (Code:
prescribed a medication quantities can be provided upon receipt of the construction of	tion benefit requires that we review certain requests for coverage with the prescriber. You have a for your patient that requires Prior Authorization before benefit coverage or coverage of additional ded. Please complete the following questions then fax this form to the toll-free number listed below. completed form, prescription benefit coverage will be determined based on the plan's rules. ase note that supporting clinical documentation is required for ALL PA acy prior authorization reviews can be subject to trial with additional are not listed within the criteria. The policies are subject to change based irements, MDH transmittals and updates to treatment guidelines.
1 Is the requ	est an INITIAL or CONTINUATION of therapy?
[] Initial (If o	checked, go to 2)
[] Continua	tion (If checked, go to 5)
opioid prod	est for one of the preferred single-agent or combination short-acting Yes No ucts? to question 6.]

If you have any questions, call: 1-888-258-8250

	[Note: Preferred agents include Morphine IR, Oxycodone IR, Hydromorphone IR, Tramadol IR, Hydrocodone/APAP, Oxycodone/APAP, Hydrocodone/IBU, APAP/Codeine, Tramadol/APAP]		
3	Has the patient tried at least THREE of the preferred formulary alternatives for at least 3 months in the last 365 days? [If yes, skip to question 6.]	Yes	No
4	Is the patient intolerant to or contraindicated to at least THREE preferred alternatives? ACTION REQUIRED: Submit supporting documentation. [NOTE: Must have clinical documentation of intolerance to or contraindication to at least THREE preferred alternatives.] [If yes, skip to question 6.] [If no, no further questions.]	Yes	No
5	Is the patient responding to treatment? [If no, no further questions.]	Yes	No
6	Is the patient currently an inpatient at an acute care hospital?	Yes	No
7	Is the patient being discharged from the hospital or emergency department?	Yes	No
8	Is the patient pregnant?	Yes	No
9	Is the patient undergoing active cancer treatment? If yes, please document the type of cancer [If yes, no further questions.]	Yes	No
10	Does the patient have sickle cell disease? [If yes, no further questions.]	Yes	No
11	Is the patient being treated as part of hospice care, long term care, skilled nursing facility care, or palliative care (diagnosis code Z51.5)? If yes, please document the diagnosis [If yes, no further questions.]	Yes	No
12	Are you an inpatient hospital, ambulatory surgery center, or emergency room prescriber, OR are you an outpatient prescriber providing ongoing care?		
	[] Inpatient hospital, ambulatory surgery, or emergency room prescriber (If checked, go to 13)		
	[] Outpatient prescriber providing ongoing care (If checked, go to 17)		
	[] None of the above. Please document prescriber's specialty and setting (If checked, go to 17)		

13	Have you discussed the risks/benefits associated with opioid use with patient or the patient's household?	Yes	No
14	Is the patient exempt from need for a Patient-Prescriber Pain Management/Opioid Treatment Agreement and random urine drug screen (UDS) because he/she is being discharged from the Hospital/Ambulatory surgery center (ASC)/Emergency room (ER) and opioid treatment prescribed by the discharging provider will be for less than 30 days or the need for further opioid use will be re-evaluated by an outpatient provider within 30 days?	Yes	No
15	Has a Naloxone prescription been provided or offered to the patient or the patient's household?	Yes	No
16	Have you reviewed the Controlled Substance Prescriptions in Prescription Drug Monitoring Program (PDMP) (Chesapeake Regional Information System for our Patients [CRISP])? [If yes, skip to question 22.] [If no, no further questions.]	Yes	No
17	Will the patient have random Urine Drug Screens?	Yes	No
18	Is the Patient/Prescriber Pain Management/Opioid Treatment Agreement signed and in medical record?	Yes	No
19	Have you discussed the risks/benefits associated with opioid use with patient or the patient's household?	Yes	No
20	Has a Naloxone prescription been provided or offered to the patient or the patient's household?	Yes	No
21	Have you reviewed the Controlled Substance Prescriptions in Prescription Drug Monitoring Program (PDMP) (Chesapeake Regional Information System for our Patients [CRISP])? [If no, no further questions.]	Yes	No
22	Has the patient's diagnosis been submitted? Please document the diagnosis	Yes	No
	[If no, no further questions.]		
23	Does the requested quantity exceed the quantity limits placed on this medication? Please document the requested quantity for the medication	Yes	No
	[If no, no further questions.]		
24	Has the provider evaluated the need for the requested quantity and assessed opportunities to wean opioid utilization when appropriate? ACTION REQUIRED: Submit supporting documentation.	Yes	No

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Please document the diagnoses, symptoms, and/or any other information important to this review:				
SECTION B: Physician Signature				
PHYSICIAN SIGNATI IRE	DATE			

FAX COMPLETED FORM TO: 1-833-896-0656

Disclaimer: An authorization is not a guarantee of payment. Member must be eligible at the time services are rendered. Services must be a covered Health Plan Benefit and medically necessary with prior authorization as per Plan policy and procedures.

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