

# **Provider update**

# Pharmacy Analgesic Opioid Prior Authorization Form

### **Instructions:**

- 1. Complete this form in its entirety. Any incomplete sections will result in a delay in processing. PA request can also be submitted electronically at www.covermymeds.com or www.availity.com.
- We review requests for prior authorization (PA) based on medical necessity only. If we
  approve the request, payment is still subject to all general conditions of Amerigroup District of
  Columbia, Inc., including current enrollee eligibility, other insurance, and program restrictions.
  We will notify the provider and the enrollee's pharmacy of our decision.
- 3. To help us expedite your authorization requests, please fax all the information required on this form to 844-487-9292 (retail) or 844-487-9294 (medical injectables).
- 4. Allow us at least 24 hours to review this request. If you have questions regarding a PA request, call us at **Phone Number**, Monday through Friday, 8 a.m. to 6 p.m. Eastern.
- 5. The pharmacy is authorized to dispense up to a 72-hour supply while awaiting the outcome of this request. Please contact the enrollee's pharmacy.
- 6. Access our website at www.provider.amerigroup.com/dc to view the Preferred Drug List.
- An ICD/diagnosis code is required for all requests. An HCPCS billing code is required for all medical injectable/oncology requests. If the billing facility is different from the requesting physician, the billing facility information will need to be completed.

### Enrollee information

Last name	First name	MI	Amerigroup ID #	Date of birth	Sex
					$\Box$ F $\Box$ M
Enrollee's place of residence:			Height	Weight	
□ Home □ Nursing facility					
□ Long-term care facility					
Administration site:					
□ Home □ Offic	e 🛛 Outpatient faci	lity			

#### Medication information (Please use a separate form for each medication request.)

Drug name and strength requested	SIG (dose, frequency, and duration)	Quantity
Diagnosis and/or indication	ICD code	HCPCS billing code

# PA type

Approval duration: <ul> <li>New prescription — approved for three months</li> <li>Continuation therapy (enrollee has been taking this medication) — approved for six months</li> </ul>					
Request type:					
□ Quantity limit □ Long-acting opioid	☐ High dose (≥90 cumulative MED/day) ☐ Nonpreferred				
$\Box$ Methadone for pain	Fentanyl     Other				

# Approval criteria (Please check all boxes that apply.)

**Note**: Any areas not filled out are considered not applicable to your enrollee and may affect the outcome of this request.

Y		Criteria			
Section A	<b>A.</b> <i>E</i>	nrollee meeting one of the following is not required to meet the PA criteria unless the requested			
agent is nonpreferred. For requests for nonpreferred long-acting agents, please					
proceed	to Se	ection F. For requests for nonpreferred short-acting agents, please proceed to			
Section	<i>G</i> .				
		Enrollee has a diagnosis of cancer-related pain and/or is actively undergoing cancer therapy.			
		If yes, please indicate specific diagnosis:			
		Enrollee has a diagnosis of terminal illness and is receiving palliative/end-of-life care.			
		If yes, please indicate specific diagnosis:			
		Enrollee has a diagnosis of sickle cell disease.			
		Enrollee is currently receiving care at a long-term care facility.			
Section 1	B. A	ll request types for enrollee not meeting one of the criteria under Section A			
		Prescriber has reviewed controlled dangerous substance (CDS) prescriptions in			
		Prescription Drug Monitoring Program (PDMP) (CRISP).			
		Enrollee has had/will have random urine drug screens before and during treatment.			
		Naloxone prescription was provided or offered to enrollee/enrollee's household.			
		Enrollee-prescriber pain management/opioid treatment agreement/contract signed			
		and in medical record.			
		Prescriber has certified the benefits of opioid treatment for the enrollee outweigh the			
		risks of treatment.			
Section	C. A	ll requests for long-acting agents			
		Enrollee has a diagnosis of pain severe enough to require daily, around -the-clock, long-term			
		opioid treatment.			
		If yes, please indicate specific diagnosis:			
		Enrollee has had an inadequate response to alternative treatment options such as			
		(but not limited to) non-opioid analgesics and immediate-release opioids.			
		Alternative treatment options would otherwise be inadequate to provide sufficient			
		management of pain.			
		Enrollee has contraindications to non-opioid analgesics (such as NSAID use in a			
		enrollee with active ulcer condition/gastrointestinal bleeding/renal failure).			
		Enrollee is 18 years of age or older.			

		Prescriber has consulted with the enrollee regarding risks of opioid therapy.			
		Clear treatment goals have been defined and outlined as part of overall plan.			
		Requested medication is being used as an as-needed analgesic.			
		Enrollee has one of the following conditions: significant respiratory depression, acute			
		or severe bronchial asthma or hypercarbia, or known or suspected paralytic ileus.			
Section	1 D. A	Additional for requests for initial therapy			
		Enrollee is currently taking a short-acting analgesic (for example, use of opioid analgesia as			
		an inpatient for post-surgical pain).			
	_	Enrollee is transitioning from one long-acting opioid analgesic to another long-acting			
		opioid analgesic.			
Section	. E. A	dditional for requests for continuation of therapy			
_		Long-acting opioid therapy has provided meaningful improvement in pain and/or			
		function compared to baseline.			
Section	F. A	dditional for requests for nonpreferred long-acting agents			
		g-acting agents are morphine sulfate ER tablets (generic MS Contin), methadone and			
		h (generic Duragesic).			
		Enrollee has had a trial (medication samples/coupons/discount cards are excluded from			
		consideration as a trial) and inadequate response or intolerance to two			
		preferred long-acting agents.			
		Enrollee has completed titration and is already maintained on a stable dose of the			
		requested drug.			
		Preferred long-acting opioids are unacceptable due to concomitant clinical situations			
		such as (but not limited to) known hypersensitivity to any ingredient not			
		also in the requested nonpreferred agent.			
		Please describe medical necessity for nonpreferred agents:			
		dditional for requests for nonpreferred short-acting agents			
-		d short-acting agents are all brand products, tapentadol (generic Nucynta) and			
oxymor	shone	e (generic Opana).			
_		Enrollee has had a trial (medication samples/coupons/discount cards are excluded from			
		consideration as a trial) and inadequate response or intolerance to one			
		preferred short-acting agent.			
		Please describe medical necessity for nonpreferred agents:			
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### Prescriber information

Last name	First name	MI	NPI (required)	DEA/license no.
Address where service was rendered			City	State
ZIP code Telephone number			Fax number	
Office contact name			Contact direct phone number	

### **Billing facility information**

Name		NPI /tax ID (required)	DEA/license no.
Address		City	State
ZIP code	Telephone number	Fax number	Office contact name

#### **Pharmacy information**

Name	Pharmacy NPI	Telephone number	Fax number

## Signature

I certify the information provided is accurate and complete to the best of my knowledge, and I understand that any falsification, omission, or concealment of material may be subject to civil or criminal liability.

Prescriber signature (or authorized representative) Date