ANALGESICS, OPIOID SHORT-ACTING PRIOR AUTHORIZATION FORM



Prescriber name:



(form effective 1/6/2025)

☐ Renewal request

 \square New request

Fax to PerformRx $^{\text{SM}}$ at **1-866-497-1387**, or to speak to a representative, call **1-800-588-6767**.

of pages:

Name of office contact:		Specialty:			
Contact's phone number:		NPI:	NPI: State license #:		
LTC facility contact/phone:		Street	Street address:		
Beneficiary name:		City/state/zip:			
Beneficiary ID#:	DOB:	Phone	:	Fax:	
CLINICAL INFORMATION					
Drug requested:		Strength:		Formulation (capsule, tablet, etc.):	
Directions:				Weight (if <21 years of age):	
Quantity per fill: to last day		S	Requested duration:		
Diagnosis (submit documentation):			Dx code (required):		
Pennsylvania law requires prescribers to query the <u>PA PDMP</u> 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 beneficiary and this request. Check all that apply and <u>submit documentation</u> for each item.					
	INITIAL	reau	ests		
1. For a transmucosal fentanyl product: Has a diagnosis of cancer					
3. For a non-preferred Analgesic, Opioid Short-Acting (See the Preferred Drug List for the list of preferred and non-preferred Analgesics, Opioid Short-Acting at: https://papdl.com/preferred-drug-list): Tried and failed or has a contraindication or an intolerance to the preferred Analgesics, Opioid Short-Acting List preferred medications tried:					
 4. For a beneficiary with a concurrent prescription for a buprenorphine agent indicated for the treatment of opioid use disorder (OUD) OR Vivitrol (naltrexone extended-release suspension for injection): □ Both prescriptions are prescribed by the same prescriber □ Prescriptions are prescribed by different prescribers and all prescribers are aware of the other prescription(s) □ Not applicable — beneficiary is not taking a buprenorphine agent indicated for the treatment of OUD or Vivitrol 					

INITIAL requests (continued)					
5. For <u>all</u> Analgesics, Opioid Short-Acting:					
☐ Has a diagnosis of active cancer, sickle cell with crisis, or neonatal abstinence syndrome					
☐ Is receiving palliative care or hospice services					
☐ Is receiving treatment post-operatively or following a traumatic injury					
☐ Has documentation of pain that is all of the following:					
☐ Caused by a medical condition					
☐ Moderate to severe					
□ Not migraine in type (does NOT apply to nasal butorphanol)					
☐ Tried and failed or has a contraindication or an intolerance to non-opioid analgesics appropriate for the beneficiary's condition:					
□ acetaminophen □ duloxetine (e.g., Cymbalta, Drizalma)					
☐ gabapentinoids (e.g., gabapentin, pregabalin [Lyrica])					
□ NSAIDs (e.g., ibuprofen, naproxen, meloxicam, etc.)					
□ tricyclic antidepressants (e.g., amitriptyline, nortriptyline, etc.)					
other (specify):					
□ Was assessed for the potential risk of opioid misuse or opioid use disorder by the prescriber					
6. For a beneficiary with a concurrent prescription for a benzodiazepine:					
☐ The benzodiazepine is being tapered					
☐ The opioid is being tapered ☐ Concomitant use of the benzodiazepine and opioid is medically necessary					
□ Not applicable — beneficiary is not taking a benzodiazepine					
For a beneficiary who has received opioid treatment for the past 3 months:					
Has results of a recent urine drug screen (UDS) testing for licit and illicit drugs with the potential for abuse, including specific testing for oxycodone, fentanyl,					
buprenorphine, and tramadol, that is consistent with prescribed controlled substances					
RENEWAL requests					
1. For <u>all</u> Analgesics, Opioid-Short Acting:					
☐ Has a diagnosis of active cancer, sickle cell with crisis, or neonatal abstinence syndrome					
☐ Is receiving palliative care or hospice services					
Experienced an improvement in pain control and/or level of functioning while on the requested medication Has results of a recent urine drug screen (UDS) testing for licit and illicit drugs with the potential for abuse, including specific testing for over the control of the c	woodone fontanyl				
buprenorphine, and tramadol, at least every 12 months that is consistent with prescribed controlled substances	kycodone, remanyi,				
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2. For a beneficiary with a concurrent prescription for a benzodiazepine:					
☐ The benzodiazepine is being tapered					
☐ The opioid is being tapered					
☐ Concomitant use of the benzodiazepine and opioid is medically necessary					
□ Not applicable — beneficiary is not taking a benzodiazepine					
DI FACE FAY COMPLETED FORM WITH DEGUIDED CLINICAL DOCUMENTATION					
PLEASE <u>FAX</u> COMPLETED FORM WITH <u>REQUIRED CLINICAL DOCUMENTA</u>					
Prescriber signature:	Date:				

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