

ANALGESICS, OPIOID LONG-ACTING PRIOR AUTHORIZATION FORM (form effective 01/05/2021)

Prior authorization guidelines for Analgesics, Opioid Long-Acting and Quantity Limits/Daily Dose Limits are available on Geisinger Health Plan's website at <https://healthplan.geisinger.org/pharmacy/pharmacy.aspx?strip=true&style=OneGeisinger>

<input type="checkbox"/> New request <input type="checkbox"/> Renewal request		Total # pages: _____	Prescriber name:	
Name of office contact:			Specialty:	
Contact's phone number:		NPI:	State license #:	
Facility contact/phone:		Street address:		
Beneficiary name:		Suite #:	City/state/zip:	
Beneficiary ID#:	DOB:	Phone:	Fax:	

CLINICAL INFORMATION

Refer to <https://papdl.com/preferred-drug-list> for a list of preferred and non-preferred drugs in each class

Drug requested:	Strength:	Formulation (capsule, tablet, etc):
Directions:	Weight (if < 21 yrs): _____ lbs / kg	
Quantity per fill: _____ to last _____ days	Requested duration:	
Diagnosis <i>(submit documentation)</i> :	Dx Code <i>(required)</i> :	
Does the beneficiary have a history of trial and failure, contraindication, or intolerance other Short-acting Opioid(s)?	<input type="checkbox"/> Yes Please list: <input type="checkbox"/> No	
Is the beneficiary being treated for active cancer, sickle cell with crisis, or neonatal abstinence syndrome OR receiving hospice or palliative care services?	<input type="checkbox"/> Yes – <i>Submit documentation and send to GHP</i> <input type="checkbox"/> No – <i>Continue with form.</i>	
What is the anticipated duration of therapy with opioid analgesics?	Requested duration: _____ days / 1 mo / 2 mos / 3 mos	
Did the prescriber or prescriber's delegate search the PDMP to review the beneficiary's controlled substance prescription history before issuing this prescription for the requested agent?	<input type="checkbox"/> Yes – specify: _____ <input type="checkbox"/> No	
If taking a benzodiazepine, is there a plan for the tapering of the benzodiazepine(s)?	<input type="checkbox"/> Yes <input type="checkbox"/> No	
If taking a benzodiazepine, is there documentation that the member has been counseled on the increased risk of death associated with concurrent benzodiazepine and opioid use AND has been offered naloxone	<input type="checkbox"/> Yes <input type="checkbox"/> No	
If taking a buprenorphine agent or naltrexone for extended-release injection suspension (Vivitrol), are this medication and the long-acting opioid prescribed by the same prescriber or, if prescribed by different prescribers, are all prescribers are aware of the other prescriptions?	<input type="checkbox"/> Yes <input type="checkbox"/> No	
If taking a buprenorphine agent or naltrexone for extended-release injection suspension (Vivitrol), is there a need for therapy with a long-acting opioid and will the other therapy be suspended during the treatment for pain?	<input type="checkbox"/> Yes <input type="checkbox"/> No	

INITIAL requests

Check all of the following that apply to the beneficiary. **Submit detailed medical record documentation for EACH item.**

INITIAL requests:

- Member has a documentation of pain that is ALL of the following Caused by a medical condition Not neuropathic or migraine in type
- has documentation of a complete physical exam, including diagnostic testing/imaging results (cause, severity, location, etc), and documentation by a pain assessment tool measurement (e.g., a numeric or visual analog scale) **Result of numeric or visual analog scale:** _____
- has tried or cannot try non-pharmacologic techniques (i.e., behavioral, cognitive, physical, and/or supportive therapies)
- has tried or cannot try non-opioid drugs for the treatment of pain:
 - acetaminophen Lyrica (pregabalin)
 - Cymbalta (duloxetine) tricyclic antidepressants (e.g., amitriptyline, nortriptyline, ect.)
 - Gabapentin other (specify): _____
 - NSAIDs (e.g., ibuprofen, naproxen, meloxicam, etc.) _____
- Will use the requested opioid in combination with tolerated non-drug therapies and non-opioid drugs
- Has documentation of a trial of short-acting opioids
- Is opioid-tolerant (taking at least morphine 60 mg/day, transdermal fentanyl 25 mcg/hour, oxycodone 30 mg/day, oral hydromorphone 8 mg/day, or equal dose of another opioid for at least one week)
- Was assessed for the potential risk of misuse, abuse, and addiction based on family and social history obtained by prescriber
- Was counseled about the potential side effects of opioids, including the risk for misuse, abuse, and addiction
- If under 21 years of age**, a parent/guardian was counseled about the potential side effects of opioids, including the risk for misuse, abuse, and addiction
- Was evaluated for risk factors for opioid-related harm
 - Determined to be at high-risk for opioid-related harm
 - The prescriber considered prescribing naloxone for the beneficiary
- Was assessed for recent use of opioids (within the past 60 days)
- 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, and tramadol that is **consistent** with prescribed controlled substances **Date of last UDS Test:** _____

RENEWAL requests

Complete the sections below that are applicable to the beneficiary and this request and **SUBMIT DOCUMENTATION** for each item.

- Experienced an improvement in pain control and/or level of functioning while on the requested medication
- Will use the requested opioid in combination with tolerated non-drug therapies and non-opioid drugs
- Was recently evaluated by the prescriber for adverse events and warning signs of serious problems, such as overdose and opioid use disorder
- Was evaluated for risk factors for opioid-related harm
 - Determined to be at high-risk for opioid-related harm
 - The prescriber considered prescribing naloxone for the beneficiary
- 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, and tramadol that is **consistent** with prescribed controlled substances **Date of last UDS Test:** _____

Please submit to PromptPA <https://ghp.promptpa.com> OR fax to Geisinger Health Plan at 570-271-5610 the completed form with required clinical documentation.

Prescriber Signature:	Date:
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