

LONG-ACTING OPIOID ANALGESICS PRIOR AUTHORIZATION FORM



Keystone First

PERFORMRxSM
Next Generation Pharmacy Benefits

(form effective 1/5/21)

Fax to PerformRxSM at **1-215-937-5018**, or to speak to a representative call **1-800-588-6767**.

PRIOR AUTHORIZATION REQUEST INFORMATION

<input type="checkbox"/> New request	<input type="checkbox"/> Renewal request	Total # pages:	Name/phone of office or LTC facility contact:
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PATIENT INFORMATION

Patient name:	Patient ID#:	DOB:
Street address:	Apt. #:	City/state/zip:

PRESCRIBER INFORMATION

Prescriber name:	Specialty:	NPI:
Street address:	Suite #:	City/state/zip:
Phone:	Fax:	

MEDICATION REQUESTED (Names in parentheses are the brand name equivalents for reference purposes.)

Preferred Agents

Butrans patch
 Fentanyl patch 12 mcg, 25 mcg, 50 mcg, 75 mcg, 100 mcg (Duragesic)
 Morphine ER capsule (Kadian)

Morphine ER tablet (MS Contin)
 Xtampza ER

Non-Preferred Agents

<input type="checkbox"/> Arymo ER tablet	<input type="checkbox"/> fentanyl patch (37.5, 62.5, 87.5 mcg)	<input type="checkbox"/> Morphabond ER tablet	<input type="checkbox"/> oxymorphone ER tablet (Opana)
<input type="checkbox"/> Belbuca film	<input type="checkbox"/> hydromorphone ER tablet (Exalgo)	<input type="checkbox"/> morphine ER capsule (Avinza)	<input type="checkbox"/> tramadol ER capsule (ConZip)
<input type="checkbox"/> buprenorphine patch (Butrans)	<input type="checkbox"/> Hysingla ER tablet	<input type="checkbox"/> MS Contin tablet	<input type="checkbox"/> tramadol ER tablet (Ultram ER)
<input type="checkbox"/> Dolophine tablet	<input type="checkbox"/> Kadian ER capsule	<input type="checkbox"/> Nucynta ER tablet	<input type="checkbox"/> tramadol ER biphasic tablet (Ryzolt)
<input type="checkbox"/> Duragesic patch	<input type="checkbox"/> methadone tablet	<input type="checkbox"/> oxycodone ER tablet (OxyContin)	<input type="checkbox"/> Zohydro ER capsule
<input type="checkbox"/> Exalgo tablet	<input type="checkbox"/> methadone solution	<input type="checkbox"/> OxyContin tablet	
<input type="checkbox"/> fentanyl patch (37.5, 62.5, 87.5 mcg)			

Strength:	Qty per fill:	to last	days	Duration:	days / 1 mo / 2 mos / 3 mos
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Directions:

Weight (if <21 yrs):	lbs / kg	Diagnosis (submit documentation):	Dx code (required):
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1. Did the prescriber or prescriber's delegate search the PDMP to review the patient's controlled substance prescription history before issuing this prescription for the requested agent? <input type="checkbox"/> Yes – <i>Submit documentation</i> <input type="checkbox"/> No	2. Is the patient taking a benzodiazepine? <u>Submit patient's current medication list.</u> <input type="checkbox"/> Yes – List and provide medical justification: <input type="checkbox"/> No
3. For initial requests for a NON-PREFERRED agent, does the patient have a history of trial and failure, contraindication, or intolerance to the preferred Long-Acting Opioids listed above? <input type="checkbox"/> Yes <input type="checkbox"/> No Check drugs tried: <input type="checkbox"/> fentanyl patch <input type="checkbox"/> morphine ER tablet <input type="checkbox"/> morphine ER capsule <input type="checkbox"/> Butrans patch <input type="checkbox"/> Xtampza ER	
4. What is the anticipated duration of therapy with opioid analgesics? Specify duration:	<i>Submit documentation.</i>
5. Is the patient 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.</i> <input type="checkbox"/> No – Continue to the next question.	

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

INITIAL requests:

has documentation of a complete physical exam, including diagnostic testing/imaging results, and pain assessment (cause, severity, location, etc)
 has tried or cannot try non-drug pain management modalities (eg, behavioral, cognitive, physical, and/or supportive therapies)
 has tried or cannot try non-opioid drugs for the treatment of pain – check drugs tried: acetaminophen NSAIDs other: _____
 the requested opioid medication will be used in combination with tolerated non-drug therapies and non-opioid medications
 was assessed for recent (within the past 60 days) opioid use
 has documentation of a trial of short-acting opioids
 is opioid-tolerant
 was assessed for the potential risk of misuse, abuse, and addiction based on family and social history obtained by prescriber
 was counseled regarding potential side effects of opioids including risk of misuse, abuse, addiction (if <21 yo, parent/guardian may be counseled)
 was evaluated for risk factors for opioid-related harm if identified to be at high risk, the prescriber considered prescribing naloxone
 has a recent UDS testing for illicit and licit substances of abuse (with specific testing for oxycodone, fentanyl, tramadol, and carisoprodol). Date of last UDS: _____

For therapeutic duplication only:

is being transitioned to or from another long-acting opioid with the intent of discontinuing one of the medications; please provide explanation: _____
 has a medical reason for concomitant use of the requested medications; please provide explanation including peer-reviewed literature or national guidelines that support the duplication. _____

For requests exceeding the quantity limit (check all that apply):

has severe pain as documented by a pain assessment tool measurement
 medication is being prescribed by or in consultation with an appropriate specialist; list specialty: _____
 pain is inadequately controlled at the current quantity limit
 pain is inadequately controlled by, or there is intolerance or contraindication to, other long-acting opioid analgesics
 titrating dose or converting from other opioid-containing medications; provide explanation: _____

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RENEWAL requests:

- experienced an improvement in pain control and level of functioning while on the requested agent
- the requested opioid medication will be used in combination with tolerated non-drug therapies and non-opioid medications
- is being monitored 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 if identified to be at high risk, the prescriber considered prescribing naloxone
- has a recent UDS testing for illicit and licit substances of abuse (with specific testing for oxycodone, fentanyl, tramadol, and carisoprodol). Date of last UDS: _____

For requests exceeding the quantity limit (check all that apply):

- has severe pain as documented by a pain assessment tool measurement
- medication is being prescribed by or in consultation with an appropriate specialist, list specialty: _____
- pain is inadequately controlled at the current quantity limit
- pain is inadequately controlled by, or there is intolerance or contraindication to, other long-acting opioid analgesics
- titrating dose or converting from other opioid-containing medications, provide explanation: _____

PLEASE FAX COMPLETED FORM WITH REQUIRED CLINICAL DOCUMENTATION

Prescriber signature:

Date:

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