

Opioids & Medication Limits Prior Authorization Request Form (Page 1 of 3)

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Member Information <small>(required)</small>			Provider Information <small>(required)</small>		
Member Name:			Provider Name:		
Insurance ID#:			NPI#:		Specialty:
Date of Birth:			Office Phone:		
Street Address:			Office Fax:		
City:	State:	Zip:	Office Street Address:		
Phone:			City:	State:	Zip:
Medication Information <small>(required)</small>					
Medication Name:			Strength:		Dosage Form:
<input type="checkbox"/> Check if requesting brand			Directions for Use:		
Clinical Information <small>(required)</small>					
<u>Answer ALL questions on this page for ALL requests</u>					
What is the patient's diagnosis for the medication being requested? _____					
ICD-10 Code(s): _____					
What medication(s) has the patient tried and had an inadequate response to? (Please specify <u>ALL</u> medication(s)/strengths tried, length of trial, and reason for discontinuation of each medication)					
What medication(s) does the patient have a contraindication or intolerance to? (Please specify <u>ALL</u> medication(s) with the associated contraindication to or specific issues resulting in intolerance to each medication)					
Cancer Pain/Sickle Cell Disease/End of life care (palliative care): Is the requested drug being prescribed for pain associated with <u>active</u> cancer, sickle cell disease or end of life/palliative care? <input type="checkbox"/> Yes <input type="checkbox"/> No Is the patient currently enrolled in hospice? <input type="checkbox"/> Yes <input type="checkbox"/> No Does the provider confirm that the opioid is NOT used to manage symptoms associated with the patient's terminal condition or condition(s) related to the terminal illness? <input type="checkbox"/> Yes <input type="checkbox"/> No Is the prescriber affiliated with the hospice provider? <input type="checkbox"/> Yes <input type="checkbox"/> No If the prescriber is NOT affiliated with the hospice provider, does the prescriber attest coordination with the hospice provider confirming that the medication is unrelated to the terminal illness or related conditions? <input type="checkbox"/> Yes <input type="checkbox"/> No					
Long Term Care facility: Is the patient in a long-term care facility (e.g., hospital or skilled nursing facility where patient is receiving skilled nursing care)? <input type="checkbox"/> Yes <input type="checkbox"/> No					

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Opoid Care Management & Morphine Milligram Equivalent Requests

At the time of dispensing, high cumulative opoid dosing has been identified prompting a safety review. This review is to ensure the cumulative opoid utilization is safe and appropriate for your patient. The cumulative morphine milligram equivalent (MME) value is calculated based on the number of opoid drugs prescribed by one or more prescribers over a period of time. The MME value includes incoming claims and claim history. **Your patient has exceeded the established daily cumulative MME dosage and is receiving opoid drugs from 2 or more prescribers.**

Opoid Care Coordination: For cumulative opoid doses between 90-200 MME

Does the provider attest that in his/her clinical judgment, the requested current cumulative dosage exceeding 90 Morphine Milligram Equivalent (MME) is medically necessary? Yes No

Cumulative MME: For cumulative opoid dosage greater than 200 MME

Does the provider attest that in his/her clinical judgment, the current daily Morphine Milligram Equivalent (MME) dosage exceeding the current daily MME threshold is medically necessary? Yes No

Will there be a dose escalation in the patient's opoid utilization in the next 90 days? Yes No

7 Day Supply Limit Requests

At the time of dispensing, your patient was identified as new to opoid therapy. They are allowed a 7 day supply for a first fill based on a lack of previous history over a 120 day period, prompting a safety review. This review is to ensure the opoid utilization is safe and appropriate for your patient. **Your patient has exceeded the initial 7 day supply limit.**

Does the provider attest that in his/her clinical judgment, the requested day supply exceeding the current 7 day supply limit is medically necessary? Yes No

Does the provider attest that in his/her clinical judgment, the requested day supply exceeding the current 7 day supply limit is medically necessary because the patient will be on an intermittent schedule (e.g., the patient will have multiple surgeries during the year where a 10-day course of opoid therapy is needed with each surgery)? Yes No

Concurrent use of opoids plus benzodiazepines

At the time of dispensing, your patient was identified as utilizing an opoid AND a benzodiazepine, prompting a safety review. This review is to ensure the opoid utilization is safe and appropriate for your patient. **Your patient is taking both an opoid and a benzodiazepine.**

Does the provider attest that in his/her clinical judgment, the requested concurrent use of opoid plus benzodiazepine, is safe and medically necessary? Yes No

Does the provider attest that either the benzodiazepine or opoid drug interacting with each other will be discontinued? Yes No

Duplicate long-acting opoid use

At the time of dispensing, your patient was identified as utilizing 2 or more long-acting opoids concurrently, prompting a safety review. This review is to ensure the opoid utilization is safe and appropriate for your patient. **Your patient is currently on 2 or more long-acting opoids.**

Does the provider attest that in his/her clinical judgment, the overlap of two or more long-acting opoid therapy, is medically necessary? Yes No

Does the provider attest that therapy will change to include only one long-acting opoid drug? Yes No

Opoid-Medication Assisted Treatment (MAT) Combination Therapy

At the time of dispensing, your patient was identified as utilizing an opoid after medication assisted treatment (MAT) [buprenorphine], prompting a safety review. This review is to ensure the opoid utilization is safe and appropriate for your patient. **Your patient was currently prescribed an opoid while on MAT (buprenorphine) therapy.**

Does the provider attest that in his/her clinical judgment, the requested concurrent use of opoid plus buprenorphine containing products (used for MAT of opoid dependence), is safe and medically necessary for this patient? Yes No

Does the provider attest that either the buprenorphine or opoid drug interacting with each other will be discontinued? Yes No

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Opoid-APAP Combination Therapy

At the time of dispensing, your patient was identified as utilizing a combination of opoid plus Acetaminophen (APAP), where the APAP exceeds 4000mg prompting a safety review. This review is to ensure the opoid utilization is safe and appropriate for your patient. **Your patient is currently on an opoid-APAP combination therapy, where the APAP exceeds 4000mg.**

Does the provider attest that in his/her clinical judgment, the requested combination of opoid plus APAP therapy, where the APAP exceeds 4000mg is medically necessary? Yes No

Opoid-Prenatal Vitamin Combination Therapy

At the time of dispensing, your patient was identified as utilizing an opoid plus prenatal vitamin therapy, prompting a safety review. This review is to ensure the opoid utilization is safe and appropriate for your patient. **Your patient is currently on an opoid plus prenatal vitamin therapy.**

Does the provider attest that in his/her clinical judgment, the requested combination of opoid plus prenatal vitamin therapy is medically necessary? Yes No

Does the provider attest that the patient has discontinued the opoid or that the patient is not pregnant? Yes No

Quantity Limits

What is the quantity requested per DAY? _____

What is the reason for exceeding the plan limitations?

- Titration or loading-dose purposes
- Patient is on a dose-alternating schedule (e.g., one tablet in the morning and two tablets at night, one to two tablets at bedtime)
- Requested strength/dose is not commercially available
- There is a medically necessary justification why the patient cannot use a higher commercially available strength to achieve the same dosage and remain within the same dosing frequency. **Please specify:** _____

Does the patient's diagnosis include malignant (cancer) pain? Yes No

Was the requested medication prescribed by a pain specialist or by pain management consultation? Yes No

Is the requested medication being used to treat postoperative pain? Yes No

If Yes, answer the following:

Is the requested medication being prescribed for pain related to a dental procedure? Yes No

Is the requested dose being prescribed the same dose that the patient was stable on prior to discharge? Yes No

Select all of the following that have been maintained and documented in chart notes*:

- A description of the nature and intensity of the pain
- An appropriate patient medical history and physical examination
- An updated, comprehensive treatment plan (the treatment plan should state objectives that will be used to determine treatment success, such as pain relief or improved physical and/or psychosocial function)
- Appropriate dose escalation
- Ongoing, periodic review of the course of opoid therapy
- Verification that the risks and benefits of the use of the requested drug have been discussed with the patient, significant other(s), and/or guardian

Chart documentation:

Will chart documentation be submitted to *Optum Rx*[®] with this form, confirming the above information? Yes No

**Please note: Chart documentation of the above is required to be submitted for quantity limit requests for this drug.*

Please note:

This request may be denied unless all required information is received within established Medicare timelines.

If the patient is not able to meet the above standard prior authorization requirements, please call 1-800-711-4555.

For urgent or expedited requests please call 1-800-711-4555.

This form may be used for non-urgent requests and faxed to 1-844-403-1028.