

Nasal and Oral Fentanyl Products Prior Authorization Request Form (Page 1 of 2)

Member Information (required)			Provider Information (required)		
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 (required)					
Medication Name:			Strength:		Dosage Form:
<input type="checkbox"/> Check if requesting brand			Directions for Use:		
<input type="checkbox"/> Check if request is for continuation of therapy					
Clinical Information (required)					
For states, such as AR, that have a terminal illness mandate, and for patients who have a terminal illness, please answer the following:					
Will the requested medication be used for the treatment of a terminal condition or associated symptoms? <input type="checkbox"/> Yes <input type="checkbox"/> No					
If "YES", please indicate the patient's estimated life expectancy:					
<input type="checkbox"/> Less than 6 months <input type="checkbox"/> Less than _____ months (please specify)					
Select the diagnosis below:					
Does the patient have a diagnosis of cancer? <input type="checkbox"/> Yes <input type="checkbox"/> No					
If yes to the above, is the requested medication used to manage breakthrough pain due to cancer? <input type="checkbox"/> Yes <input type="checkbox"/> No					
<input type="checkbox"/> Other diagnosis: _____ ICD-10 Code(s): _____					
Select any of the following which the patient has at least a one week history of:					
<input type="checkbox"/> An alternative opioid at an equianalgesic dose (e.g., oral methadone greater than or equal to 20 mg/day)					
<input type="checkbox"/> Duragesic (fentanyl transdermal patch) at doses greater than or equal to 25 µg/hour					
<input type="checkbox"/> Morphine sulfate at doses of greater than or equal to 60 mg/day					
<input type="checkbox"/> Oral oxymorphone at a dose of greater than or equal to 25 mg/day					
<input type="checkbox"/> Oral hydromorphone at a dose of greater than or equal to 8 mg/day					
<input type="checkbox"/> Oxycodone at a dose of greater than or equal to 30 mg/day					
Select the medications the patient has a failure, contraindication, or intolerance to:					
<input type="checkbox"/> Fentanyl lozenge (generic Actiq)					
<input type="checkbox"/> Hydromorphone immediate-release (IR)					
<input type="checkbox"/> Morphine sulfate IR					
<input type="checkbox"/> Oxymorphone IR					
<input type="checkbox"/> Oxycodone IR					
Current treatment:					
Is the patient currently taking a long-acting opioid around the clock for cancer pain? <input type="checkbox"/> Yes <input type="checkbox"/> No					

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Prescriber specialty:
Select which of the following specialists the medication was prescribed by:
 Hematologist Hospice care specialist Oncologist Pain specialist Palliative care specialist
 Other _____

Select which of the following specialists the medication was prescribed in consultation with:
 Hematologist Hospice care specialist Oncologist Pain specialist Palliative care specialist
 Other _____

Quantity limit requests:
 What is the quantity being requested per DAY: _____

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
 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 opioid therapy
 Patient physical examination
 Verification that the risks and benefits of the use of the controlled substance have been discussed with the patient, significant other(s), and/or guardian

Chart documentation:
 Will chart documentation be submitted to ProAct® 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.

Are there any other comments, diagnoses, symptoms, medications tried or failed, and/or any other information the physician feels is important to this review?

Please note: This request may be denied unless all required information is received.

Please fax this form to 1-844-712-8129 to initiate a prior authorization review for the member and medication above.

Please note: plan benefits may limit or exclude coverage of specific medications including those requested on this form.

I certify, to the best of my knowledge, the statements and information provided on this form are factual and correct.

Provider/Representative (and Title): _____ Date: _____

PROACT INTERNAL USE ONLY:				
Clinical Review Decision				
Approved, through				
Denied (documentation attached, if necessary)				
Tracking:				
1 st Attempt		2 nd Attempt	Letter Mailed:	