



## NC DHB Pharmacy Request for Prior Approval Long-Acting Opioid Analgesic

### Recipient Information

DMA-3571 (V01)

1. Recipient Last Name: \_\_\_\_\_ 2. First Name: \_\_\_\_\_  
3. Recipient ID #: \_\_\_\_\_ 4. Recipient Date of Birth: \_\_\_\_\_ 5. Recipient Gender: \_\_\_\_\_

### Payer Information

6. Is this a Medicaid or Health Choice Request? Medicaid: ☐ Health Choice: ☐

### Prescriber Information

7. Prescribing Provider #: \_\_\_\_\_ NPI: ☐ or Atypical: ☐  
8. Prescriber DEA #: \_\_\_\_\_  
Requester Contact Information: Name: \_\_\_\_\_ Phone #: \_\_\_\_\_ Ext: \_\_\_\_\_

### Drug Information

9a. Drug Name: \_\_\_\_\_ 9b. Is this request for a Non-Preferred Drug? ☐ Yes ☐ No  
10. Strength: \_\_\_\_\_ 11. Quantity Per 30 Days: \_\_\_\_\_  
12. Length of Therapy (in days): ☐ up to 30 ☐ 60 ☐ 90 ☐ 120 ☐ 180 ☐ 365 ☐ Other: \_\_\_\_\_

### Clinical Information

1. Does the patient have a diagnosis of malignant cancer or pain due to neoplasm? ☐ Yes ☐ No If yes, the patient is exempt from the prior authorization requirement
2. Does the beneficiary have a diagnosis of chronic pain syndrome of at least four (4) weeks duration? ☐ Yes ☐ No
3. **Is the requested daily dose in combination with other concurrent opioids less than or equal to 90mg of morphine or an equivalent dose?** ☐ Yes ☐ No
- Answer questions 3a and 3b when the response to question 3 is 'No'.
- 3a. Please supply the beneficiary's diagnosis and reason for exceeding dose per day limits.  
Please list: \_\_\_\_\_
- 3b. Please provide the duration (days supply) the beneficiary will exceed the limit of 90mg of morphine or an equivalent dose.  
Please list: \_\_\_\_\_
4. Is this an initial authorization request? Select 'Yes' for an initial authorization. Select 'No' for a reauthorization request.  
☐ Yes ☐ No
- 4a. If Yes, has the beneficiary tried a short-acting Opioid Analgesic in the past 45 days? ☐ Yes ☐ No
- 4b. If no, explain: \_\_\_\_\_
5. Has the prescriber reviewed and is adhering to the N.C. Medical Board statement on the use of controlled substances for the treatment of pain? ☐ Yes ☐ No
6. Is the prescribing clinician adhering, as medically appropriate, to the guidelines which include: (a) complete beneficiary evaluation, (b) establishment of a treatment plan (contract), (c) informed consent, (d) periodic review, and (e) consultation with specialists in various treatment modalities as appropriate? ☐ Yes ☐ No
7. Has the prescribing physician checked the beneficiary's utilization of controlled substances on the NC Controlled Substance Reporting System? ☐ Yes ☐ No
8. Has the prescribing clinician reviewed the current CDC Guideline for Prescribing Opioids for Chronic Pain? ☐ Yes ☐ No
- Non-Preferred Products:**
9. Does the patient have a documented history within the past year of two preferred long-acting Opioid Analgesics at a dose equal to or equivalent to the non-preferred long-acting Opioid Analgesic being prescribed? ☐ Yes ☐ No  
Please list: \_\_\_\_\_
10. Does the patient have a contraindication or allergy to ingredients in the preferred product? ☐ Yes ☐ No  
Please list: \_\_\_\_\_

Signature of Prescriber: \_\_\_\_\_ Date: \_\_\_\_\_

\*Prescriber signature mandatory

Fax this form to NCTracks: (855) 710-1969

Pharmacy PA Call Center: (866) 246-8505