

**Beneficiary Information**

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

**Prescriber Information**

6. Prescriber Name: \_\_\_\_\_ NPI #: \_\_\_\_\_  
Mailing address: \_\_\_\_\_ City: \_\_\_\_\_ State: \_\_\_\_\_ ZIP: \_\_\_\_\_  
7. Requester Contact Information: \_\_\_\_\_  
Name: \_\_\_\_\_ Phone #: \_\_\_\_\_ Fax #: \_\_\_\_\_

**Drug Information**

8. Drug Name: \_\_\_\_\_ 9. Strength: \_\_\_\_\_ 10. Quantity Per 30 Days: \_\_\_\_\_  
11. Length of Therapy:  up to 30 days  60 days  90 days  120 days  180 days  Other: \_\_\_\_\_

**Clinical Information**

1. Does the beneficiary have a diagnosis of malignant cancer or pain due to neoplasm? Yes\_\_\_ No\_\_\_  
\*If yes, the beneficiary is exempt from the prior authorization requirement.  
2. Does the patient have Sickle Cell Disease? Yes\_\_\_ No\_\_\_  
3. Is this an initial authorization request? ('Yes' for an initial authorization; 'No' for a reauthorization request.) Yes\_\_\_ No\_\_\_  
3a. If No, please attach documentation as to why the beneficiary needs continued opioid treatment and current plan of care.  
4. **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\_\_\_ Please answer questions 4a and 4b when the response to question 4 is 'No'.  
4a. Please supply the beneficiary's diagnosis and reason for exceeding dose per day limits. Please list: \_\_\_\_\_  
\_\_\_\_\_  
4b. Please provide the duration (day supply) the beneficiary will exceed the limit of 90mg of morphine or an equivalent dose.  
Please list: \_\_\_\_\_  
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 short-acting Opioid Analgesics at a dose equal to or equivalent to the non-preferred short-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**

I certify that the information provided is accurate and complete to the best of my knowledge, and I understand that any falsification, omission, or concealment of material fact may subject me to civil or criminal liability.

Fax this form to: 1-877-234-4274, or call Pharmacy Prior Authorization: 1-866-885-1406