



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.