



# New Hampshire Medicaid Fee-for-Service Program Prior Authorization Drug Approval Form

Morphine Milligram Equivalent (MME)

DATE OF MEDICATION REQUEST:     /     /

## SECTION I: PATIENT INFORMATION AND MEDICATION REQUESTED

LAST NAME:

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FIRST NAME:

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MEDICAID ID NUMBER:

--	--	--	--	--	--	--	--	--	--	--	--	--	--	--	--	--	--	--	--

DATE OF BIRTH:

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GENDER: ☐ Male ☐ Female

Drug Name:

Strength:

Dosing Directions:

Length of Therapy:

## SECTION II: PRESCRIBER INFORMATION

LAST NAME:

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FIRST NAME:

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SPECIALTY:

NPI NUMBER:

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PHONE NUMBER:

--	--	--	--	--	--	--	--	--	--	--	--	--	--	--	--	--	--	--	--

FAX NUMBER:

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## SECTION III: CLINICAL HISTORY

1. Is the prescriber a pain specialist, specialist within the same organ system as the primary pain diagnosis, or has one been consulted in this case? ☐ Yes ☐ No
2. For what condition is this medication being prescribed? Select all that apply.
  - ☐ Pain associated with acute sickle cell disease
  - ☐ Pain associated with cancer
  - ☐ Hospice or end-of-life care
  - ☐ Severe, persistent pain that requires continuous around-the-clock pain control for at least 10 days
  - ☐ Other: \_\_\_\_\_

(Form continued on next page.)



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Prior Authorization Drug Approval Form**

Morphine Milligram Equivalent (MME)

**DATE OF MEDICATION REQUEST:**     /     /

**PATIENT LAST NAME:**

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**PATIENT FIRST NAME:**

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**SECTION III: CLINICAL HISTORY (*Continued*)**

3. Has the patient tried and failed or is patient not a candidate for at least 3 of the following? ☐ Yes ☐ No  
Provide details below.

- ☐ Topical NSAIDS: \_\_\_\_\_
- ☐ Oral NSAIDS: \_\_\_\_\_
- ☐ Oral Acetaminophen: \_\_\_\_\_
- ☐ Transcutaneous electrical nerve stimulation: \_\_\_\_\_

4. Has the patient failed or had an adequate trial of a lower MME dose? ☐ Yes ☐ No

a. If yes, list treatment failures and provide dates:

\_\_\_\_\_

5. Do you attest that the NH Prescription Drug Monitoring Program has been reviewed in the last 60 days? ☐ Yes ☐ No

6. Do you attest that the risks associated with taking high-dose opioids have been reviewed with the patient? ☐ Yes ☐ No

7. Does the patient have a written pain agreement? ☐ Yes ☐ No

8. Do you attest that you had a discussion with the patient about attempting to taper the dose slowly at an individualized pace? ☐ Yes ☐ No

9. Do you attest that the patient is being monitored to mitigate overdose risk? ☐ Yes ☐ No

10. Will the patient be prescribed concurrent naloxone? ☐ Yes ☐ No

11. Does the patient have a history of severe asthma or other lung disease? ☐ Yes ☐ No

12. Will the patient require concurrent therapy with a benzodiazepine, sedative hypnotic or barbiturate? ☐ Yes ☐ No

Provide any additional information that would help in the decision-making process. **If additional space is needed, please use a separate sheet.**

**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.**

**PRESCRIBER'S SIGNATURE:** \_\_\_\_\_ **DATE:** \_\_\_\_\_

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MANAGEMENT<sup>SM</sup>