



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

## Short-Acting Fentanyl Analgesic Medications

**DATE OF MEDICATION REQUEST:** / /

SECTION I: PATIENT INFORMATION AND MEDICATION REQUESTED									
LAST NAME:					FIRST NAME:				
MEDICAID ID NUMBER:					DATE OF BIRTH:				
GENDER: <input type="checkbox"/> Male <input type="checkbox"/> Female									
Drug Name					Strength				
Dosing Directions					Length of Therapy				
SECTION II: PRESCRIBER INFORMATION									
LAST NAME:					FIRST NAME:				
SPECIALTY:					NPI NUMBER:				
PHONE NUMBER:					FAX NUMBER:				
SECTION III: CLINICAL HISTORY									
1. Is the medication being prescribed for the treatment of breakthrough cancer pain? <input type="checkbox"/> Yes <input type="checkbox"/> No									
2. For what condition is this medication being prescribed? _____									
3. What is the patient's age? _____									
4. Is the patient already receiving and tolerant to opioid therapy? <input type="checkbox"/> Yes <input type="checkbox"/> No									
5. Has the patient tried and failed immediate-release narcotics for breakthrough pain? <input type="checkbox"/> Yes <input type="checkbox"/> No									
Please list treatment failures and dates: _____									
6. Has an oncologist, pain specialist, palliative care specialist, or hospice specialist been consulted on <input type="checkbox"/> Yes <input type="checkbox"/> No									

*(Form continued on next page.)*



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

Short-Acting Fentanyl Analgesic Medications

DATE OF MEDICATION REQUEST:    /    /

PATIENT LAST NAME:

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

PATIENT FIRST NAME:

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

## SECTION III: CLINICAL HISTORY (CONTINUED)

7. Are you enrolled in the transmucosal immediate-release fentanyl Risk Evaluation and Mitigation Strategies (TIRF REMS) Access program?  Yes  No  
***Prescribers, pharmacies, and patients must be enrolled in the TIRF REMS Access program.***

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

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

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

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

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

13. Will the patient be prescribed concurrent naloxone?  Yes  No

Provide current opioid (pain management) treatment (drug, dose, frequency, duration):

---

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: \_\_\_\_\_

Phone: 1-866-675-7755

Fax: 1-888-603-7696