



Department of Vermont Health Access  
 NOB 1 South, 280 State Drive  
 Waterbury, Vermont 05671-1010

**~ HUB (OTP) BUPRENORPHINE Prior Authorization Form ~**

All requests for Suboxone® Film > 24mg, Buprenorphine/Naloxone tablets, and Buprenorphine monotherapy must be reviewed by the Change Healthcare Clinical Call Center. Documentation must accompany this form. For questions, please contact the Change Healthcare help desk at 1-844-679-5363.

**Submit request via Fax: 844-679-5366**

Prescribing physician:	Member:
Name: _____	Name: _____
NPI: _____	Medicaid ID#: _____
Specialty: _____	Date of Birth: _____ Sex: _____
Phone#: _____	Diagnosis: _____
Fax#: _____	Date of Admission to HUB: _____
Address: _____	
Contact Person at HUB (OTP): _____	

**CHECK HERE IF PATIENT IS ADAP UNINSURED**

**Request is from the following HUB location:** \_\_\_\_\_ / \_\_\_\_\_

**Name** **NPI**

<input type="checkbox"/> <b>Suboxone® Film &gt; 24 mg</b> Dose per day requested: _____mg
* Clinical note/letter from prescriber that documents the prescriber’s clinical rationale for requesting Suboxone® Film >24mg must be attached (REQUIRED). Requests for doses >24mg will require review by DVHA Medical Director.
<input type="checkbox"/> <b>Buprenorphine/Naloxone tablets</b> Dose per day requested: _____mg
* FDA Medwatch form documenting a provider observed reaction to Suboxone® Film severe enough to require discontinuation including measures taken to mitigate/manage symptoms must be attached (REQUIRED)
<input type="checkbox"/> <b>Buprenorphine (mono formulation)</b> Dose per day requested: _____mg
<input type="checkbox"/> Pregnancy DUE DATE: _____ <input type="checkbox"/> Pregnancy test/ultrasound result/lab attached (REQUIRED)
<input type="checkbox"/> Breastfeeding an opiate dependent baby (baby is being administered morphine or methadone for opiate withdrawal symptoms) *Clinical note/letter from a pediatrician/neonatologist must be attached (REQUIRED)
<input type="checkbox"/> Using buprenorphine mono to switch from methadone to Suboxone® Dates buprenorphine mono will be administered: _____
<input type="checkbox"/> Using buprenorphine mono due to provider observed reaction to both Suboxone® Film and Buprenorphine/Naloxone tablets severe enough to require discontinuation * FDA Medwatch form documenting reaction and measures taken to mitigate/manage symptoms must be attached (REQUIRED)

**Prescriber Signature:** \_\_\_\_\_ **(stamps not acceptable)** **Date of request:** \_\_\_\_\_

