



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

~ HUB (OTP) BUPRENORPHINE Prior Authorization Form ~

All requests for buprenorphine containing products > 24mg 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"/> Suboxone® Film > 24 mg 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"/> Buprenorphine/Naloxone tablets > 24 mg Dose per day requested: _____ mg	
* Clinical note/letter from prescriber that documents the prescriber’s clinical rationale for requesting buprenorphine/naloxone tablets >24mg must be attached (REQUIRED). Requests for doses >24mg will require review by DVHA Medical Director.	
<input type="checkbox"/> Buprenorphine tablets (monotherapy) Dose per day requested: _____ mg	
<input type="checkbox"/> Using buprenorphine monotherapy for up to two weeks to switch from a high potency opioid (methadone/fentanyl) to Suboxone®	
<input type="checkbox"/> Using buprenorphine mono due to a current or past intolerance to preferred/combination products that cannot be resolved or mitigated through alternative efforts	
<input type="checkbox"/> Other _____	
* Please provide clinical justification explaining why the member cannot use the preferred buprenorphine formulations	
* > 24 mg Clinical note/letter from prescriber that documents the prescriber’s clinical rationale for requesting buprenorphine tablets (mono formulation) >24mg must be attached (REQUIRED). Requests for doses >24mg will require review by DVHA Medical Director.	

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

