

Opioid Dependence Therapy Agents

Member Information

1. Last Name: _____ 2. First Name: _____
3. Trillium ID #: _____ 4. Date of Birth: _____ 5. Gender: _____

Prescriber Information

1. Prescriber Name: _____ 2. NPI #: _____
3. Requestor Name (Nurse/Office Staff): _____
4. Mailing Address: _____ City: _____ State: _____ Zip: _____
3. Phone #: _____ Ext. _____ Fax #: _____

Drug Information

1. Drug Name: _____ 2. Strength: _____ 3. Quantity Per 30 Days: _____
4. Length of Therapy: up to 30 Days 60 Days 90 Days 120 Days 180 Days 365 Days

Clinical Information

For Coverage of Buprenorphine/Naloxone SL Films, and Zubsolv:

1. Has the member failed one preferred drug? Yes No Please List: _____
 - a. Was the failure due to an allergic reaction? Yes No
 - b. Was the failure due to a drug-to-drug interaction? Yes NoPlease describe reaction: _____
2. Previous episode of an unacceptable side effect or therapeutic failure.
Please provide clinical information: _____
3. Clinical contraindication, co-morbidity, or unique member circumstance as a contraindication to preferred drug(s).
Please provide clinical information: _____
4. Age specific indications. Please give member age and explain:

5. Unique clinical indication supported by FDA approval or peer reviewed literature.
Please explain and provide a general reference: _____
6. Unacceptable clinical risk associated with therapeutic change.
Please explain: _____

For Coverage of Buprenorphine Sublingual Tablets:

7. Does the member have a diagnosis of Opioid Dependence? Yes No
8. Is the member unable to use Suboxone Film? Yes No (If Yes, please specify one or more of the following conditions)
 Member is pregnant: Please Provide Estimated Due Date: _____ Max Length of Therapy is 270 Days
 Member is breast feeding Max Length of Therapy is 60 Days (can be renewed)
 Member has an allergy to naloxone (rashes, hives, pruritis, bronchospasm, angioneurotic edema and anaphylactic shock) Max Length of Therapy is 365 Days
 Other condition Please List: _____
9. Has the prescriber reviewed the controlled substances reporting system database prior to writing the prescription to ensure that concomitant opioid use is not occurring? Yes No
10. Is the maximum daily dose less than or equal to 32 mg/day? Yes No

For Coverage of Lucemyra Tablets:

11. Does the member have a diagnosis of opioid withdrawal symptoms? Yes No
(trial and failure of preferred are not required)

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.

Fax this form to PerformRx at (833) 726-7628 or call Pharmacy PA Call Center: (855) 662-0277