



Opioid Dependence Therapy Agents Prior Authorization Form

Member information		
1. Member last name:	2. Mem	ber first name:
3. Member ID #: 4	I. Member date of birth:	5. Member gender:
Prescriber information		
6. Prescribing provider NPI #:		
7. Prescriber contact information		
Name:		
Phone:	Ext:	
Drug information		
8. Drug name:	9. Stren	ıgth:
10. Quantity per 30 days:		
11. Length of therapy (in days): □ up to 30 days □ 60 days □ 90 days □ 120 days □ 180 days □ 365 days □ Other:		
Clinical information		
For coverage of Buprenorphine/Naloxone SL Films, and Zubsolv:		
1. Has the member failed one preferred drug? \Box Yes \Box No		
Please list:		
1a. □ Allergic reaction 1b. □ Drug-to-drug interaction. Please 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? \Box Yes \Box 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 beneficiary have a diagnosis of opioid withdrawal symptoms? Yes No (trial and failure of		
preferreds are not required)		

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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 **844-376-2318**. Healthy Blue Pharmacy PA Call Center: **844-594-5072**