

## Prior Authorization Guideline

<b>Guideline Name</b>	Sublocade Extended Release Monthly Injection
<b>Formulary</b>	<ul style="list-style-type: none"> <li>• Arizona Health Care Cost Containment (AZM)</li> </ul>

### Guideline Note:

Effective Date:	10/1/2019
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### 1. Criteria

Product Name: Sublocade	
Approval Length	6 Month(s)
Therapy Stage	Initial Authorization
Guideline Type	Prior Authorization
<p>Buprenorphine extended-release injection (Sublocade) is proven and/or medically necessary for the treatment of moderate to severe opioid use disorder in patients who meet ALL of the following:</p> <p><b>Approval Criteria</b></p> <p><b>1</b> - Patient has severe Opioid Use Disorder (OUD) as defined by the DSM-5 OUD Diagnostic Tool and has a demonstrated history of non-adherence to oral medications</p> <p style="text-align: center;"><b>AND</b></p> <p><b>2</b> - Patient is currently maintained on 8mg to 24mg per day dose of oral, sublingual, or transmucosal buprenorphine product equivalent for at least 7 days prior to initiation of extended-release buprenorphine injection</p> <p style="text-align: center;"><b>AND</b></p> <p><b>3</b> - Patient has not, nor will receive supplemental, oral, sublingual, or transmucosal buprenorphine</p> <p style="text-align: center;"><b>AND</b></p> <p><b>4</b> - Patient is receiving psychosocial interventions as part of a comprehensive medication assisted treatment (MAT) program</p> <p style="text-align: center;"><b>AND</b></p>	

**5** - Prescriber meets DATA 2000 requirements and has been assigned a unique identification number specific to the prescription of medication assisted therapy (DEA-X)

**AND**

**6** - Prescriber checks the Arizona State Board of Pharmacy Controlled Substance Prescription Monitoring Program (CSPMP) database prior to each monthly injection

**AND**

**7** - Sublocade dosing is in accordance with the U. S. Food and Drug Administration approved labeling: 300mg subcutaneously monthly for the first 2 months, followed by a maintenance dose of 100 mg or 300mg monthly

Product Name: Sublocade

Approval Length	12 Month(s)
Therapy Stage	Reauthorization
Guideline Type	Prior Authorization

**Approval Criteria**

**1** - Physician documentation that the patient has experienced a positive clinical response to buprenorphine extended-release therapy, as defined by the provider

**AND**

**2** - Patient has not, nor will receive supplemental, oral, sublingual, or transmucosal buprenorphine

**AND**

**3** - Patient is receiving psychosocial interventions as part of a comprehensive medication assisted treatment (MAT) program

**AND**

**4** - Prescriber meets DATA 2000 requirements and has been assigned a unique identification number specific to the prescription of medication assisted therapy (DEA-X)

**AND**

**5** - Prescriber checks the Arizona State Board of Pharmacy Controlled Substance Prescription Monitoring Program (CSPMP) database prior to each monthly injection

**AND**

**6** - Sublocade dosing is in accordance with the U. S. Food and Drug Administration approved labeling: maintenance dose of 100mg or 300mg monthly

## **2. Revision History**

Date	Notes
10/1/2019	New custom program.