

| Market Applicability | | | | | | | | | | | | | | |
|----------------------|----|----------|--------|--------|----|----|----|----|----|----|----|----|----|----|
| Market | DC | FL & FHK | FL MMA | FL LTC | GA | KS | KY | MD | NJ | NV | NY | TN | TX | WA |
| Applicable | NA | NA | NA | NA | X | NA | NA | NA | NA | NA | NA | NA | NA | NA |

*FHK- Florida Healthy Kids

Buprenorphine/Naloxone – Georgia Medicaid

Based on Georgia Medicaid Fee For Service Prior Authorization Criteria:

https://dch.georgia.gov/sites/dch.georgia.gov/files/Opioid_Dependency_Agents.pdf

| Override(s) | Approval Duration |
|---------------------------------------|-------------------|
| Prior Authorization Quantity Limit | 6 months |

| Medications | Status | Strength | Quantity Limit |
|--|---------------|----------------|--------------------|
| Buprenorphine with naloxone Sublingual Tablet *Indicates FDA maximum recommended dose for specific drug and dosage strength – 24 mg/6 mg | Preferred | 8mg – 2mg | 3 tablets per day |
| | | 2mg – 0.5mg | 12 tablets per day |
| Suboxone (buprenorphine with naloxone) Sublingual Film *Indicates FDA maximum recommended dose for specific drug and dosage strength – 24 mg/6 mg | Preferred | 2mg – 0.5mg | 12 films per day |
| | | 4mg – 1mg | 64 films per day |
| | | 8mg – 2mg | 3 films per day |
| | | 12mg – 3 mg | 2 films per day |
| Zubsolv (buprenorphine with naloxone) *Indicates FDA maximum recommended dose for specific drug and dosage strength – 17.2 mg/4.2 mg | Non-Preferred | 0.7mg-0.18mg | 23 tablets per day |
| | | 2.9mg – 0.7mg | 4 tablets per day |
| | | 11.4mg – 2.9mg | 1 tablet per day |
| | | 1.4mg – 0.36mg | 12 tablets per day |
| | | 5.7mg – 1.4mg | 3 tablets per day |
| | | 8.6mg – 2.1 mg | 2 tablets per day |

This policy does not apply to health plans or member categories that do not have pharmacy benefits, nor does it apply to Medicare. Note that market specific restrictions or transition-of-care benefit limitations may apply.

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| Market | DC | FL & FHK | FL MMA | FL LTC | GA | KS | KY | MD | NJ | NV | NY | TN | TX | WA |
| Applicable | NA | NA | NA | NA | X | NA | NA | NA | NA | NA | NA | NA | NA | NA |

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| | | | |
|---|---------------|---------------|------------------------|
| Bunavail (buprenorphine with naloxone) *Indicates FDA maximum recommended dose for specific drug and dosage strength – 12.6 mg/2.1 mg | Non-Preferred | 2.1mg – 0.3mg | 6 buccal films per day |
| | | 4.2mg – 0.7mg | 3 buccal films per day |
| | | 6.3mg – 1mg | 2 buccal films per day |

APPROVAL CRITERIA

Requests for all buprenorphine with naloxone products may be approved for individuals who meet the following criteria:

- I. Individual is 16 years of age or older; **AND**
- II. Individual has been diagnosed with opioid dependence; **AND**
- III. Prescriber must have an “X” DEA number (unique DATA [Drug Addiction Treatment Act 2000] waiver identification number); **AND**
- IV. Individual will not use concurrently with opioids. (Note: If concurrent use is necessary, written documentation from the prescriber regarding the medical necessity of this therapy to be used along with the requested buprenorphine/naloxone is required.)

Requests for all non-preferred buprenorphine with naloxone products (Bunavail, and Zubsolv) may be approved for individuals who meet the following criteria (in addition to I-IV above):

- I. Individual meets one of the following:
 - A. Individual has been on the requested product in the previous 180 days **OR**
 - B. Individual has had a trial of one preferred buprenorphine with naloxone agent (current preferred agents: buprenorphine/naloxone sublingual tablets, Suboxone film) in the previous 180 days.

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