

ARYMO ER (extended-release morphine sulfate, crush resistant), buprenorphine buccal film (BELBUCA), BELBUCA (buprenorphine buccal film), buprenorphine patch (BUTRANS), BUTRANS (buprenorphine transdermal), CONZIP (tramadol extended-release), EMBEDA (morphine sulfate/naltrexone, abuse deterrent), EXALGO (extended-release hydromorphone, abuse deterrent), fentanyl transdermal (DURAGESIC), Fentanyl transdermal 37.5mcg/hr, 62.5mcg/hr, 87.5mcg/hr, hydrocodone extended-release, abuse deterrent, non-abuse deterrent (Zohydro ER), hydromorphone extended release, abuse deterrent (EXALGO), HYSINGLA ER (extended-release hydrocodone, abuse deterrent), levorphanol (LEVO-DROMORAN), methadone (DOLOPHINE), methadone oral solution, MORPHABOND ER (morphine extended-release), morphine extended-release (AVINZA, KADIAN), NUCYNTA ER (extended-release tapentadol, abuse deterrent), oxycodone, extended release (OXYCONTIN), OXYCONTIN (oxycodone extended release, abuse deterrent), oxymorphone, extended-release (OPANA ER) RYZOLT (tramadol extended-release), TRAMADOL 150mg extended-release capsule, tramadol extended-release (CONZIP, RYZOLT, ULTRAM ER), ULTRAM ER (tramadol extended-release), XTAMPZA ER (extended-release oxycodone, abuse deterrent), **ZOHYDRO ER (extended-release hydrocodone, abuse deterrent)**

Diagnosis Considered for Coverage:

 Treatment of severe pain requiring daily, around-the-clock, long-term opioid therapy

Coverage Criteria:

For pain related to a current diagnosis with cancer:

- Prescribed by an oncologist, and
- Total quantity of dosage form has been consolidated to the least number of higher strength.

<u>For those with pain not due to cancer, and currently NOT on an extended-release</u> opioid:

- The cause of the pain cannot be removed, or treated by other mode of therapy, and
- Pain occurs every day, and lasted for at least 3 months, and
- Patient requires around-the-clock, long-term therapy with opioids, and
- Inadequate response or intolerable side effect to 2 preferred non-opioid analgesics used around-the-clock, and
- Inadequate response or intolerable side effect to 2 preferred immediaterelease opioids used around-the-clock, **and**
- Doctor has a specific plan for evaluating patient's response to therapy, monitoring for potential misuse, monitoring for side effects, and tapering down opioid use, **and**
- Will not be used together with another long-acting opioid, and
- Total dose of all opioid being used does not exceed 120 mg morphine equivalent dose per day, and
- Total quantity of dosage form has been consolidated to the least number of higher strength, and
- For requests except Butrans, Fentanyl transdermal patches, Xtampza ER,
 Conzip, Ryzolt, and Ultram ER: Inadequate response or intolerable side
 effect to generically available extended-release morphine (MS Contin), and
- For requests for Conzip, Ryzolt, and Ultram ER: Adequate trial with immediate-release tramadol therapy, and
- For Fentanyl (12 mcg/hr, 25 mcg/hr, 75 mcg/hr, 100 mcg/hr) transdermal patch: Inadequate response or intolerable side effect to generically available extended-release morphine (MS Contin), and
- For Fentanyl (37.5 mcg/hr, 62.5 mcg/hr, 87.5 mcg/hr) transdermal patch: Medical rationale why preferred generically available fentanyl (12 mcg/hr, 25 mcg/hr, 50 mcg/hr, 75 mcg/hr, or 100 mcg/hr) transdermal patch formulations cannot be used, and
- For Xtampza ER: Inadequate response or intolerable side effect to generically available extended-release morphine (MS Contin), AND medical rationale why patient is unable to use generically available extended-release oxycodone (Oxycontin).

For those with pain not due to cancer, and currently on an extended-release opioid:

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- Doctor has a specific plan for evaluating patient's response to therapy, monitoring for potential misuse, monitoring for side effects, and tapering down opioid use, and
- Will not be used together with another long-acting opioid, and
- Total dose of all opioid being used does not exceed 120 mg morphine equivalent dose per day, and
- Total quantity of dosage form has been consolidated to the least number of higher strength, **and**
- For requests except Butrans, Fentanyl transdermal patches, Xtampza ER, Conzip, Ryzolt, and Ultram ER: Inadequate response or intolerable side effect to generically available extended-release morphine (MS Contin), and
- For requests for Conzip, Ryzolt, and Ultram ER: Adequate trial with immediate-release tramadol therapy, and
- For Fentanyl (12 mcg/hr, 25 mcg/hr, 75 mcg/hr, 100 mcg/hr) transdermal patch: Inadequate response or intolerable side effect to generically available extended-release morphine (MS Contin), and
- For Fentanyl (37.5 mcg/hr, 62.5 mcg/hr, 87.5 mcg/hr) transdermal patch: Medical rationale why preferred generically available fentanyl (12 mcg/hr, 25 mcg/hr, 50 mcg/hr, 75 mcg/hr, or 100 mcg/hr) transdermal patch formulations cannot be used, and
- For Xtampza ER: Inadequate response or intolerable side effect to generically available extended-release morphine (MS Contin), AND medical rationale why patient is unable to use generically available extended-release oxycodone (Oxycontin).

For brand-name Avinza, Belbuca, Conzip, Duragesic, Exalgo, Kadian, Methadose, Opana ER, Oxycontin, Ryzolt, Ultram ER, Zohydro ER:

- Meets above coverage criteria for generic, and
- Allergic or intolerable side effect to the generic formulation not expected with the brand.

Coverage Duration: up to 30 days

Effective Date: 6/01/2022