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## Prior Authorization Criteria for Zulresso Begins July 1

### Background:

Zulresso is indicated for adult females (18 years and older) for the treatment of postpartum depression and must be prescribed by, or in consultation with, a psychiatrist or obstetrics/gynecologist. It is only available through a restricted program under a Risk Evaluation and Mitigation Strategy (REMS) due to the risk of excessive sedation or sudden loss of consciousness. For these reasons, Zulresso will be a clinician-administered drug covered through the medical benefit.

- Health-care facilities must enroll in the Zulresso REMS program.
- Patients must enroll in the Zulresso REMS program prior to Zulresso treatment. Certified facilities must ensure that Zulresso is only administered to clients who are enrolled in the Zulresso REMS program.
- Specialty pharmacies must be certified in the Zulresso REMS program and must only dispense to health-care facilities certified to administer Zulresso.

### Key Details:

Beginning July 1, prior authorization is required for Zulresso for Medicaid and CHIP. For approval of therapy, an individual must meet the following criteria:

- Diagnosis of postpartum depression (diagnosis code F530) with a HAM-D total score of at least 20, or as scored by an alternative comparable rating scale that measures depressive symptoms.
- Onset of the major depressive episode is within the third trimester and no later than the first four weeks postpartum.
- Six months or less postpartum at screening.
- No active psychosis or history of bipolar disorder or schizophrenia.
- Have not received treatment with Zulresso for the current postpartum depressive episode.
- Have continuous pulse oximetry monitoring during the infusion period due to risk of serious harm and be accompanied when interacting with their child(ren) as the drug can cause loss of consciousness.
- A health-care provider must be available on site for continuous monitoring of the client for the duration of the infusion.

### Contact:

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