

Zurzuvaе (zuranolone)
Effective 03/01/2024

Plan	<input type="checkbox"/> MassHealth UPPL <input checked="" type="checkbox"/> Commercial/Exchange	Program Type	<input checked="" type="checkbox"/> Prior Authorization <input checked="" type="checkbox"/> Quantity Limit <input type="checkbox"/> Step Therapy
Benefit	<input checked="" type="checkbox"/> Pharmacy Benefit <input type="checkbox"/> Medical Benefit		
Specialty Limitations	N/A		
Contact Information	Specialty Medications		
	All Plans	Phone: 877-519-1908	Fax: 855-540-3693
Contact Information	Non-Specialty Medications		
	All Plans	Phone: 800-711-4555	Fax: 844-403-1029
Exceptions	N/A		

Overview

Zurzuvaе is indicated for postpartum depression in adults.

Coverage Guidelines

Authorization may be granted for members new to the plan who are currently receiving treatment with the requested medication, excluding when the product is obtained as samples or via manufacturer’s patient assistance programs.

OR

Authorization may be granted when the following criteria is met:

1. The member has a diagnosis of postpartum depression.
2. Prescriber is a specialist (e.g., obstetrician-gynecologist/family medicine or psychiatrist) or consult is provided from a specialist.
3. Member is 18 years of age or older.
4. Member is not currently pregnant.
5. Member is ≤12 months postpartum (date of delivery is required)
6. Medical charts documenting ONE of the following:
 - a. Member has had treatment failure, intolerance, or contraindication to an oral antidepressant.
 - b. Clinical rationale that a trial of oral antidepressant is not appropriate due to severity of depression.

Continuation of Therapy

Authorization may be granted for continued treatment in members when the following criteria are met:

1. The member has a diagnosis of postpartum depression.
2. Member is not currently pregnant.
3. Member is ≤12 months postpartum (date of delivery is required)
4. Last day of treatment with requested agent is ≥ 45 days prior to current request.

Limitations

1. Initial approvals and reauthorizations will be granted for 14 days

2. The following quantity limits apply:

Zurzuvae 20mg and 25mg	28 tablets per 14 days
Zurzuvae 30mg	14 tablets per 14 days

References

1. Zurzuvae® [package insert]. Cambridge (MA): Biogen, Inc.; 2023 Aug.

Review History

2/14/2023: Created and Reviewed at Feb P&T, Effective 3/1/2024

