

Cynthia A. Persily, Ph.D. Cabinet Secretary Cynthia Beane Commissioner

Office of Pharmacy Services Prior Authorization Criteria Veozah[®] (fezolinetant) *Effective 9/25/2024* Prior Authorization Request Form

VEOZAH is a neurokinin 3 (NK3) receptor antagonist indicated for the treatment of moderate to severe vasomotor symptoms due to menopause.

CRITERIA FOR APPROVAL:

- 1. The patient has a diagnosis of menopause with moderate to severe vasomotor symptoms; **AND**
- 2. Patient is within the age range as recommended by the FDA label; AND
- 3. Documentation detailing the frequency and severity of the vasomotor symptoms must be provided; **AND**
- 4. The patient has had a 30-day trial of <u>ONE</u> hormone replacement therapy (HRT) agent (unless contraindicated) **AND** has had a 30-day trial of <u>ONE</u> non-hormonal therapy (such as an SSRI, SNRI, gabapentin, pregabalin or clonidine) which failed to provide sufficient relief OR <u>TWO</u> 30-day trials of non-hormonal therapies (ONLY if a hormonal therapy cannot be tolerated) which failed to provide sufficient relief; **AND**
- 5. Patient must have baseline liver function tests prior to initiating therapy* and every month for the first three months after patients start treatment, and then at months 6 and 9 of treatment; **AND**
- 6. Patient does not have severe renal impairment or end-stage renal disease.

Initial approvals may be authorized for 90 days. Further approvals may be granted for 1 year after all the continuation of therapy criteria has been met.

*Do not start VEOZAH if concentration of ALT or AST is equal to or exceeds two times the ULN

9/25/2024 DUR Board Meeting



CONTINUATION OF THERAPY CRITERIA:

- 1) Patient continues to meet all initial approval criteria; **AND**
- 2) Demonstrate continued documented compliance; AND
- Documentation of positive clinical response to therapy must be provided (such as decrease in frequency of symptoms and/or improvement in severity of vasomotor symptoms); AND
- 4) Patient has not experienced any treatment-restricting adverse effects (e.g., ALT or AST > 3 times the ULN).

9/25/2024 DUR Board Meeting

