



STANDARD COMMERCIAL DRUG FORMULARY
PRIOR AUTHORIZATION GUIDELINES

FEZOLINETANT

Generic	Brand	HICL	GCN	Medi-Span	Exception/Other
FEZOLINETANT	VEOZAH	48921		GPI-10 (3060603000)	

GUIDELINES FOR USE

INITIAL CRITERIA (FOR RENEWAL CRITERIA SEE BELOW)

1. Does the patient have a diagnosis of moderate to severe menopausal vasomotor symptoms (VMS) and meet **ALL** of the following criteria?
 - The patient experiences 7 or more hot flashes per day
 - The patient had a trial of or contraindication to hormonal therapy (e.g., estradiol transdermal patch [Minivelle, Climara], oral conjugated estrogens [Premarin], micronized progesterone [Prometrium])

If yes, **approve for 12 months by HICL or GPI-10 with a quantity limit of #1 per day.**

If no, do not approve.

INITIAL DENIAL TEXT: *Some terms are already pre-defined in parenthesis. Please use these definitions if the particular text you need to use does not already have definition(s) in it.

Our guideline named **FEZOLINETANT (Veozah)** requires the following rule(s) be met for approval:

- A. You have moderate to severe menopausal vasomotor symptoms (VMS: a type of symptom related to menopause)
- B. You experience 7 or more hot flashes per day
- C. You had a trial of or contraindication (harmful for) to hormonal therapy (such as estradiol transdermal patch [Minivelle, Climara], oral conjugated estrogens [Premarin], micronized progesterone [Prometrium])

Your doctor told us **[INSERT PT SPECIFIC INFO PROVIDED]**. We do not have information showing you **[INSERT UNMET CRITERIA]**. This is why your request is denied. Please work with your doctor to use a different medication or get us more information if it will allow us to approve this request.

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GUIDELINES FOR USE (CONTINUED)

RENEWAL CRITERIA

1. Does the patient have a diagnosis of moderate to severe menopausal vasomotor symptoms (VMS) and meet **ALL** of the following criteria?
 - The patient has a continued need for VMS treatment (i.e., persistently symptomatic with hot flashes)
 - The patient had a reduction in VMS frequency OR severity due to Veozah (fezolinetant) treatment

If yes, **approve for 12 months by HICL or GPI-10 with a quantity of #1 per day.**

If no, do not approve.

RENEWAL DENIAL TEXT: *Some terms are already pre-defined in parenthesis. Please use these definitions if the particular text you need to use does not already have definition(s) in it.

Our guideline named **FEZOLINETANT (Veozah)** requires the following rule(s) be met for renewal:

- A. You have moderate to severe menopausal vasomotor symptoms (VMS: a type of symptom related to menopause)
- B. You have a continued need for VMS treatment (you still experience persistent hot flashes)
- C. You had a reduction in VMS frequency OR severity due to Veozah (fezolinetant) treatment

Your doctor told us **[INSERT PT SPECIFIC INFO PROVIDED]**. We do not have information showing you **[INSERT UNMET CRITERIA]**. This is why your request is denied. Please work with your doctor to use a different medication or get us more information if it will allow us to approve this request.

RATIONALE

For further information, please refer to the Prescribing Information and/or Drug Monograph for Veozah.

REFERENCES

- Veozah [Prescribing Information]. Northbrook, IL: Astellas Pharma US, Inc.; May 2023.

Library	Commercial	NSA
Yes	Yes	No

Part D Effective: N/A

Commercial Effective: 06/12/23

Created: 05/23

Client Approval: 05/23

P&T Approval: 01/23