

Xermelo
Effective 07/01/2023

Plan	<input type="checkbox"/> MassHealth UPPL <input checked="" type="checkbox"/> Commercial/Exchange	Program Type	<input checked="" type="checkbox"/> Prior Authorization <input type="checkbox"/> Quantity Limit <input type="checkbox"/> Step Therapy
Benefit	<input checked="" type="checkbox"/> Pharmacy Benefit <input type="checkbox"/> Medical Benefit		
Specialty Limitations	This medication has been designated specialty and must be filled at a contracted specialty pharmacy.		
Contact Information	Medical and Specialty Medications		
	All Plans	Phone: 877-519-1908	Fax: 855-540-3693
Exceptions	Non-Specialty Medications		
	All Plans	Phone: 800-711-4555	Fax: 844-403-1029
Exceptions	N/A		

Overview

Telotristat etiprate is a prodrug of telotristat, which is a tryptophan hydroxylase inhibitor. Used in combination with a somatostatin analog (SSA) for treatment of diarrhea associated with carcinoid syndrome in adults that SSA therapy alone has shown inadequate

Coverage Guidelines

Authorization may be granted for members currently being treated with Xermelo, except when the medication has been obtained through physician samples or manufacturer's assistance program.

OR

Authorization may be granted if the member meets all following criteria and documentation has been submitted:

1. The member is at least 18 years of age
2. The member has diagnosis of carcinoid syndrome diarrhea
3. The member has had an inadequate response to somatostatin analog (SSA) therapy
4. Xermelo will be used in combination with SSA therapy

Continuation of Therapy

Reauthorization will be granted when all initial criteria is met and provider attest to improvement in member's condition

Limitations

1. Approvals will be granted for 12 months
2. The following quantity limits apply:

Xermelo 250mg	90 tablets per 30 days
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References

1. Xermelo (telotristat ethyl) [prescribing information]. Deerfield, IL: TerSera Therapeutics LLC; October 2020

2. Kulke MH, Horsch D, Caplin ME, et al. Telotristat ethyl, a tryptophan hydroxylase inhibitor for the treatment of carcinoid syndrome. *J Clin Oncol*. 2017;35(1):14-23.[PubMed 27918724]

Review History

02/20/2019 – Approved by P&T

01/20/2021 – Reviewed by P&T. Effective 09/01/2021.

05/10/2023 – Reviewed and Updated for May P&T; added continuation of therapy criteria. Effective 7/1/23

