

Zelapar (selegiline) ODT Effective 04/17/2019

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

Overview

Zelapar is FDA indicated as adjunct therapy in the management of patients with Parkinson disease being treated with levodopa/carbidopa who exhibit deterioration in the quality of their response to this therapy

Coverage Guidelines

Authorization may be granted when the following criteria are met, and documentation has been submitted:

- 1. Member has been started and stabilized on Zelapar within the past 180 days (Note: Physician samples are not considered adequate justification for started & stabilized) **AND**
- 2. Member is currently taking carbidopa/levodopa

OR

- 1. Member is unable to swallow pills or use a conventional dosage form AND
- 2. Member is not currently receiving other oral solid dosage forms

Limitations

1. Approvals are granted for 12 months.

References

1. Zelapar (selegiline) [prescribing information]. Bridgewater, NJ: Valeant Pharmaceuticals North America LLC; August 2016

Review History

04/23/07 - Reviewed

09/21/09 - Updated

09/27/10 - Reviewed

12/15/10 – Updated disclaimer

09/19/11 - Reviewed

09/24/12 - Reviewed

09/23/13 - Reviewed

09/22/14 - Reviewed

09/21/15 - Reviewed

09/19/16 - Reviewed

09/18/17 - Reviewed

04/17/19 - Reviewed

11/16/2022 – Reviewed for Nov P&T. Separated out MH vs Comm/Exch. No clinical changes.

