

Zelapar (selegiline) ODT
Effective 04/17/2019

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	N/A		
Contact Information	Medical Benefit Pharmacy Benefit	Phone: 833-895-2611 Phone: 800-711-4555	Fax: 888-656-6671 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.

