

Neupro (rotigotine) transdermal system
Effective 08/01/2020

Plan	<input checked="" type="checkbox"/> MassHealth <input type="checkbox"/> Commercial/Exchange	Program Type	<input checked="" type="checkbox"/> Prior Authorization
Benefit	<input checked="" type="checkbox"/> Pharmacy Benefit <input type="checkbox"/> Medical Benefit (NLX)		<input type="checkbox"/> Quantity Limit <input type="checkbox"/> Step Therapy
Specialty Limitations	N/A		
Contact Information	Specialty Medications		
	All Plans	Phone: 866-814-5506	Fax: 866-249-6155
	Non-Specialty Medications		
	MassHealth	Phone: 877-433-7643	Fax: 866-255-7569
	Commercial	Phone: 800-294-5979	Fax: 888-836-0730
	Exchange	Phone: 855-582-2022	Fax: 855-245-2134
	Medical Specialty Medications (NLX)		
	All Plans	Phone: 844-345-2803	Fax: 844-851-0882
Exceptions	N/A		

Overview

Neupro Transdermal Patch is a dopamine agonist indicated for the treatment of Parkinson’s disease (PD) and moderate-to-severe primary Restless Legs Syndrome (RLS).

Coverage Guidelines

Authorization may be granted for members new to the plan who are currently receiving treatment with Neupro, excluding when the product is obtained as samples or via manufacturer’s patient assistance programs.

OR

Authorization may be granted when the following criteria has been met:

- Member has a diagnosis of PD or RLS **AND**
 - Member has a documented diagnosis of a swallowing disorder or difficulty swallowing tablets
- OR**
- Member has had a documented side effect, allergy, or treatment failure to a trial of an oral dopamine agonist (e.g., pramipexole, or ropinirole)

Limitations

1. Approvals will be granted for 36 months.

Dosing

Early stage PD	Max dose - 6mg per day	30 patches per 30 days
Advanced stage PD	Max dose - 8mg per day	30 patches per 30 days
RLS	Max dose - 3mg per day	30 patches per 30 days



References

- 1 Neupro (rotigotine) [prescribing information]. Smyrna, GA: UCB Inc; January 2019
- 2 Tarsey, Daniel. Pharmacologic treatment of Parkinson disease. In: Basow DS (Ed). UpToDate. Waltham (MA): UpToDate 2014. Available at: <http://www.utdol.com/utd/index.co>
- 3 Pahwa R, Factor SA, Lyons KE, Ondo WG, Gronseth G, Bronte-Stewart H, et al. Practice Parameter: Treatment of Parkinson disease with motor fluctuations and dyskinesia (an evidence-based review): Report of the Quality Standards Subcommittee of the American Academy of Neurology. *Neurology*. 2006;66(7):983-995.
- 4 Zesiewicz TA, Martinez-Martin P. Effects of rotigotine transdermal system on non-motor symptoms in Parkinson's disease: an overview. *Expert Review of Neurotherapeutics*. 2013;13(12):1329-42.
- 5 Esteve V, Carneiro J, Salazar G, et al. Effects of rotigotine on clinical symptoms, quality of life and sleep hygiene adequacy in haemodialysis-associated restless legs syndrome. *Nefrologia* 2018; 38:79
- 6 Iftikhar IH, Alghothani L, Trotti LM. Gabapentin enacarbil, pregabalin and rotigotine are equally effective in restless legs syndrome: a comparative meta-analysis. *Eur J Neurol* 2017; 24:1446
- 7 Mizuno Y, Nomoto M, Kondo T, Hasegawa K, Murata M, Takeuchi M, et al. Transdermal rotigotine in early stage Parkinson's disease: a randomized, double-blind, placebo-controlled trial. *Movement Disorders*. 2013;28(10):1447-50.

Review History

02/25/2008 - Reviewed

04/15/2008 - Implemented

08/13/2012 - Updated (Neupro reintroduced & new indication; 7/30/12 file)

11/25/2013 - Reviewed

11/24/2014 - Reviewed

11/27/2017 - Reviewed

04/17/2019 – Reviewed

05/20/2020 – reviewed May P&T Mtg; added started and stabilized statement; added dosing and QL to criteria. Effective 8/1/20.

09/21/2022 – Reviewed at Sept P&T; Separated Comm/Exch vs MH policy; no clinical updates

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