

Apomorphine (Non-PDL)

Phone: 215-991-4300

Fax back to: 866-240-3712

Health Partners Plans manages the pharmacy drug benefit for your patient. Certain requests for coverage require review with the prescribing physician. Please answer the following questions and fax this form to the number listed above.

PLEASE NOTE: Any information (patient, prescriber, drug, labs) left blank, illegible, or not attached WILL DELAY the review process.

Patient Name:		Prescriber Name:	
HPP HPP Member Number:		Fax:	Phone:
Date of Birth:		Office Contact:	
Patient Primary Phone:		NPI:	PA PROMISe ID:
Address:		Address:	
City, State ZIP:		City, State ZIP:	
Line of Business: <input type="checkbox"/> Medicaid <input type="checkbox"/> CHIP		Specialty Pharmacy (if applicable):	
Drug Name:		Strength:	
Quantity:		Refills:	
Directions:			
Diagnosis Code:		Diagnosis:	
<i>HPP's maximum approval time is 12 months but may be less depending on the drug.</i>			

Please attach any pertinent medical history including labs and information for this member that may support approval.

Please answer the following questions and sign.

Q1. Is this a renewal request? If yes, go to 2. If not, go to 6.

Yes

No

Q2. Does the patient continue to need Apomorphine and meet the criteria identified for initial approval?

Yes

No

Q3. Does the patient tolerate the medication without significant or serious side effects (must attach documentation)?

Yes

No

Q4. Has the patient had an improvement in symptoms from baseline (must attach documentation)?

Yes

No

Q5. Is there documentation of a treatment plan including duration of treatment (must attach documentation)?

Yes

No

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Q6. Does the patient have a diagnosis of advance Parkinson's Disease (PD) with documented hypomobility "off" episodes ("end-of-dose wearing off" and unpredictable "on/off" episodes) (documentation must be attached)?

Yes

No

Q7. Is the medication being prescribed by or in consultation with a specialist (who specializes in the treatment of PD or a neurologist)?

Yes

No

Q8. Does the patient have a history of therapeutic failure, a contraindication to or intolerance of the preferred Antiparkinson's agents (such as carbidopa-levodopa, pramipexole, ropinirole, bromocriptine, amantadine, selegiline, trihexyphenidyl, bntropine,) (Must attach documentation)?

Yes

No

Q9. Will the initial "test" dose be given under medical supervision?

Yes

No

Q10. Will the medication ONLY be given via subcutaneous route of administration?

Yes

No

Q11. Will trimethobenzamide be started 3 days prior to the initial dose of Apomorphine, and continue as long as necessary to control nausea and vomiting (generally no longer than 2 months)?

Yes

No

Q12. Will this medicine be administered with 5HT3 antagonists (such as ondansetron) to control nausea?

Yes

No

Q13. Has renal function been evaluated and has medication been dose adjusted for renal impairment, if necessary?

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Yes

No

Q14. Has a cardiac evaluation been performed (including assessment of QTc interval)?

Yes

No

Q15. Has the patient been counseled on the risks of using alcohol, antihypertensive medications, and vasodilating medications while taking this medication

Yes

No

Q16. Will the patient abstain from alcohol while taking this medicine?

Yes

No

Q17. Is the treatment plan attached showing how the medication will be administered, duration of therapy, and other medications that will be continued?

Yes

No

Q18. Is each dose less than or equal to 0.6 mL with a dosing frequency of less than or equal to five times per day?

Yes

No

Q19. Additional Information:

Prescriber Signature

Date

Updated for 2024