

PRIOR AUTHORIZATION REQUEST FORM

Individual and Family Plans

Apomorphine

Fax back to: (833) 605-4407 Phone: (215) 991-4300

Jefferson Health 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 NO	TE: Any information (patient, prescriber, drug, la	ibs) left blank, illegible, or not attached v	VILL delay the review process.	
Patient Name:		Prescriber Name:		
Member Number:		Fax: Phone:		
Date of Birth:		Office Contact:		
Line of Business:	□ Exchange - PA	NPI:	State Lic ID:	
Address:		Address:		
City, State ZIP:		City, State ZIP:		
Primary Phone:		Specialty/facility name (if applicab	le):	
	<u>DITED REVIEW</u> : By checking this box and signing below, I lee's ability to regain maximum function.	certify that the standard review timeframe ma	y seriously jeopardize the life or health of	
Drug Name:				
Strength:				
Directions / SIG:				
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 15. If NO, go to 2.				
☐ Yes		□ No		
Q2. Does the patient have a diagnosis of advanced 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		
	edication being prescribed by or in o	consultation with a specialist	(who specializes in	
☐ Yes		□ No		
Q4. Is there documentation of an inadequate response, intolerance, or contraindication to conventional oral therapies(such as carbidopa-levodopa, pramipexole, ropinirole, bromocriptine, amantadine, selegiline, trihexyphenidyl, benztropine, entacapone, tolcapone) (Must attach documentation)?				
☐ Yes		□ No		

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Patient Name:	Prescriber Name:		
Q5. Will the initial "test" dose be given under medical supervision?			
☐ Yes	□ No		
Q6. Will the medication ONLY be given via subcutaneous route of administration?			
□Yes	□ No		
Q7. Will trimethobenzamide be started 3 days prior to the initial dose of Apokyn, and continue as long as necessary to control nausea and vomiting (generally no longer than 2 months)?			
□ Yes	□ No		
Q8. Will this medicine be administered with 5HT3 antagonists (such as ondansetron) to control nausea?			
□ Yes	□ No		
Q9. Has renal function been evaluated and has medication been dose adjusted for renal impairment, if necessary?			
□ Yes	□ No		
Q10. Has a cardiac evaluation been performed (including assessment of QTc interval)?			
☐ Yes	□ No		
Q11. Has the patient been counseled on the risks of using alcohol, antihypertensive medications, and vasodilating medications while taking this medication?			
☐ Yes	□ No		
Q12. Will the patient abstain from alcohol while taking this medicine?			
□ Yes	□ No		
Q13. Is the treatment plan attached showing how the medication will be administered, duration of therapy, and other medications that will be continued?			

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Patient Name:	Prescriber Name:		
☐ Yes	□ No		
Q14. Is each dose less than or equal to 0.6 mL v five times per day?	vith a dosing frequency of less than or equal to		
☐ Yes	□ No		
Q15. Does the patient continue to need Apokyn@approval?	and meet the criteria identified for initial		
☐ Yes	□ No		
Q16. Does the patient tolerate the medication without significant or serious side effects (must attach documentation)?			
☐ Yes	□ No		
Q17. Has the patient had an improvement in syndocumentation)?	nptoms from baseline (must attach		
☐ Yes	□ No		
Q18. Is there documentation of a treatment plan including duration of treatment (must attach documentation)?			
☐ Yes	□ No		
Q19. Additional Information:			
Prescriber Signature	Date		

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