

Part B Prior	Authorization	Guidelines
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Dopamine Agonists: Apokyn / Kynmobi (apomorphine) J0364 Prior Authorization Request Medicare Part B Form

Instructions: * Indicates required information – Form may be returned if required information is not provided. Please fax this request to the appropriate fax number listed at the bottom of the page.

NEW START - Start Date:			Continuation (within 365 days): Date of last treatment							
		r Clinic name: _					/ Fax			
MEMBER INFORMATION										
*Name:*ID#:*DOB:										
PRESCRIBER INFORMATION										
*Nai	me:	DM	ID 🗆 F		OO □NP □PA	*Phone	e:			
*Ado	dress:					*Fax:_				
DISPENSING PROVIDER / ADMINISTRATION INFORMATION										
*Name: Phone:										
*Address: Fax:										
		PROCEDURE / I	PROD	UCT IN	FORMATION			Find Data if		
HC	PC Code	Name of Drug	Dos	e (Wt: _	kg Ht:)	Frequency	End Date if known		
Chart notes attached. Other important information:										
Diagnosis: ICD10: Description:										
□ Provider attests the diagnosis provided is an FDA-Approved indication for this drug										
CLINICAL INFORMATION										
 New Start or Initial Request: (Clinical documentation required for all requests) Provider has reviewed the attached "Criteria for Approval" and attests the member meets ALL required PA criteria. If not, please provide clinical rationale for formulary exception: 										
 Continuation Requests: (Clinical documentation required for all requests) Provider has reviewed the attached "Criteria for Continuation" and attests the member meets ALL required PA Continuation criteria. 										
Patient had an <u>adequate response</u> or <u>significant improvement</u> while on this medication. If not, please provide clinical rationale for continuing this medication:										
ACKNOWLEDGEMENT										
Request By (Signature Required):										



Prior Authorization Group – Dopamine Agonist PA

Drug Name(s): APOKYN KYNMOBI APOMORPHINE

Criteria for approval of Prior Authorization Drug:

- 1. Prescribed for an approved FDA diagnosis (as listed below):
- 2. Member does not have any clinically relevant contraindications, or CMS/Plan exclusions, to the requested drug.
- If the member meets all these criteria, they may be approved by the Plan for the requested drug.
- Quantity limits and Tiering will be determined by the Plan.

Exclusion Criteria: N/A

Prescriber Restrictions: N/A

Coverage Duration: Approvals will be for 12 months

FDA Indications:

Apokyn, Kynmobi

• Parkinson's disease, Acute, intermittent treatment of hypomobility "off" episodes

Off-Label Uses:

- Erectile dysfunction
- Induction of emesis, For treatment of acute poisoning
- Parkinsonism; Diagnosis

Age Restrictions:

Safety and effectiveness have not been established in pediatric patients

Other Clinical Considerations:

CI: Concomitant use with serotonin 5-hydroxytryptamine-3 (5-HT(3)) receptor antagonists including antiemetics

Resources:

https://www.micromedexsolutions.com/micromedex2/librarian/CS/908395/ND_PR/evidencexpert/ND_P/evidencexpert /DUPLICATIONSHIELDSYNC/DECC9E/ND_PG/evidencexpert/ND_B/evidencexpert/ND_AppProduct/evidencexpert/ND_T/ evidencexpert/PFActionId/evidencexpert.DoIntegratedSearch?SearchTerm=apomorphine#