

Part B Prior	Authorization	Guidelines
--------------	---------------	------------

#### 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										
<ul> <li>New Start or Initial Request: (Clinical documentation required for all requests)</li> <li>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:     </li> </ul>										
<ul> <li>Continuation Requests: (Clinical documentation required for all requests)</li> <li>Provider has reviewed the attached "Criteria for Continuation" and attests the member meets ALL required PA Continuation criteria.</li> </ul>										
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#