

# **PRIOR AUTHORIZATION POLICY**

**POLICY:** Parkinson's Disease – Kynmobi Prior Authorization Policy

Kynmobi<sup>™</sup> (apomorphine sublingual film – Sunovion)

**REVIEW DATE:** 03/13/2024

#### INSTRUCTIONS FOR USE

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# CIGNA NATIONAL FORMULARY COVERAGE:

#### **OVERVIEW**

Kynmobi, a non-ergoline dopamine agonist, is indicated for the acute, intermittent treatment of "off" episodes in patients with **Parkinson's disease**.<sup>1</sup>

# **Guidelines**

The International Parkinson and Movement Disorder Society published an evidence-based review for treatment for motor symptoms of Parkinson's disease (2018).<sup>2</sup> Kynmobi is not addressed. The review categorically divides treatment recommendations by Parkinson's disease characteristics. Apomorphine subcutaneous is noted to be efficacious and clinically useful in treatment for motor fluctuations.

### **POLICY STATEMENT**

Prior Authorization is recommended for prescription benefit coverage of Kynmobi. All approvals are provided for the duration noted below. Because of the specialized skills required for evaluation and diagnosis of patients treated with Kynmobi as well as the monitoring required for adverse events and long-term efficacy, approval requires Kynmobi to be prescribed by or in consultation with a physician who specializes in the condition being treated.

• Kynmobi™ (apomorphine sublingual film – Sunovion) is(are) covered as medically necessary when the following criteria is(are) met for FDA-approved indication(s) or other uses with supportive evidence (if applicable):

# **FDA-Approved Indication**

- **1. Parkinson's Disease.** Approve for 1 year if the patient meets all of the following (A, B, C, <u>and</u> D):
  - **A)** Patient is experiencing "off" episodes such as muscle stiffness, slow movements, or difficulty starting movements; AND
  - **B)** Patient is currently receiving carbidopa/levodopa therapy; AND
  - **C)** Patient has previously tried one other treatment for "off" episodes and meets ONE of the following (i <u>or</u> ii):
    - i. Patient had significant intolerance, according to the prescriber; OR
    - **ii.** Patient had inadequate efficacy, according to the prescriber; AND Note: Examples of treatments for "off" episodes include entacapone, rasagiline, pramipexole, ropinirole, tolcapone, cabergoline, selegiline, Apokyn (apomorphine subcutaneous injection), Ongentys (opicapone capsules), or Xadago (safinamide tablets).
  - **D)** The medication is prescribed by or in consultation with a neurologist.

# **CONDITIONS NOT COVERED**

- Kynmobi™ (apomorphine sublingual film Sunovion) is(are) considered experimental, investigational or unproven for ANY other use(s) including the following (this list may not be all inclusive; criteria will be updated as new published data are available):
- **1. Concurrent Use with a Serotonin 5-HT**<sub>3</sub> **Antagonist.** Administration of Kynmobi in conjunction with a serotonin 5-HT<sub>3</sub> antagonist (e.g., ondansetron, granisetron, dolasetron, palonosetron, alosetron) can result in extreme lowering of blood pressure and loss of consciousness.<sup>1</sup>

### REFERENCES

- 1. Kynmobi<sup>™</sup> sublingual film [prescribing information]. Marlborough, MA: Sunovion; September 2022.
- 2. Fox SH, Katzenschlager R, Lim SY, et al. International Parkinson and movement disorder society evidence-based medicine review: Update on treatments for the motor symptoms of Parkinson's disease. *Mov Disord*. 2018;33(8):1248-1266.

#### HISTORY

Type of Revision	Summary of Changes	Review Date
Annual	No criteria changes.	07/26/2023
Revision		

Annual	No criteria changes.	03/14/2024
Revision		

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