

Commercial & HealthCare Exchange PA Criteria

Effective: June 3, 2020

Prior Authorization: Dayvigo

<u>Products Affected:</u> Dayvigo (lemborexant tablets)

<u>Medication Description</u>: Dayvigo is an orexin receptor antagonist indicated for the treatment of insomnia, characterized by difficulties with sleep onset and/or sleep maintenance. The orexin neuropeptide signaling system is a central promoter of wakefulness. Blocking the binding of wake-promoting neuropeptides orexin A and orexin B to orexin receptors (OXR) OX1R and OX2R is thought to suppress wake drive.

Covered Uses: Insomnia

Exclusion Criteria: Narcolepsy

Required Medical Information:

1. Diagnosis

2. Previous therapies tried/failed

Age Restrictions: 18 years of age and older

Prescriber Restrictions: N/A

Coverage Duration: 12 months

Other Criteria:

- 1. Clinical diagnosis of insomnia; AND
- 2. Previous trial and failure, contraindication, or intolerance to **TWO** of the following:
 - a. eszopiclone
 - b. zolpidem/zolpidem ER
 - c. zaleplon
 - d. ramelteon

References:

1. Dayvigo tablets [package insert]. Eisai, Inc, Woodcliff Lake, NJ





Policy Revision history

Rev #	Type of Change	Summary of Change	Sections Affected	Date
1	New Policy	New Policy	All	6/3/2020