PRIOR AUTHORIZATION CRITERIA

DRUG CLASS INSOMNIA AGENTS

BRAND NAME (generic)

BELSOMRA (suvorexant)

DAYVIGO (lemborexant)

QUVIVIQ (daridorexant)

Status: CVS Caremark® Criteria Type: Initial Prior Authorization

POLICY

FDA-APPROVED INDICATIONS

Belsomra

Belsomra (suvorexant) is indicated for the treatment of insomnia characterized by difficulties with sleep onset and/or sleep maintenance.

Dayvigo

Dayvigo is indicated for the treatment of adult patients with insomnia, characterized by difficulties with sleep onset and/or sleep maintenance.

Quviviq

Quviviq is indicated for the treatment of adult patients with insomnia, characterized by difficulties with sleep onset and/or sleep maintenance.

COVERAGE CRITERIA

The requested drug will be covered with prior authorization when the following criteria are met:

 The requested drug is being prescribed for the treatment of insomnia characterized by difficulties with sleep onset and/or sleep maintenance

AND

The request is NOT for continuation of therapy

AND

 Potential factors contributing to sleep disturbances have been addressed or are currently being addressed (e.g., inappropriate sleep hygiene and sleep environment issues) as well as treatable medical/psychiatric disorders that are co-morbid with insomnia

AND

 The patient is 65 years of age or older OR

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- The patient is less than 65 years of age AND
 - The patient experienced an inadequate treatment response to any of the following: A) a generic non-benzodiazepine sedative-hypnotic (e.g., eszopiclone, zaleplon, zolpidem), B) a benzodiazepine (e.g., temazepam)

 OR
 - The patient experienced an intolerance to any of the following: A) a generic nonbenzodiazepine sedative-hypnotic (e.g., eszopiclone, zaleplon, zolpidem), B) a benzodiazepine (e.g., temazepam)
 OR
 - The patient has a contraindication that would prohibit a trial of ALL of the following A) a generic non-benzodiazepine sedative-hypnotic (e.g., eszopiclone, zaleplon, zolpidem), B) a benzodiazepine (e.g., temazepam)

OR

The request is for continuation of therapy

AND

- The patient has achieved or maintained a positive response to treatment from baseline AND
- The patient's need for continued therapy has been assessed AND
- Potential factors contributing to sleep disturbances continue to be addressed (e.g., inappropriate sleep hygiene, sleep environment issues, treatable medical/psychiatric comorbid disorders)

Quantity Limits apply.

30 tablets per 25 days* or 90 tablets per 75 days*.

*The duration of 25 days is used for a 30-day fill period and 75 days is used for a 90-day fill period to allow time for refill processing.

Duration of Approval (DOA):

• 1177-C: Initial therapy DOA: 6 months; Continuation of therapy DOA: 12 months

REFERENCES

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- 4. Lexicomp Online, AHFS DI (Adult and Pediatric) Online. Waltham, MA: UpToDate, Inc.; 2023. https://online.lexi.com. Accessed March 13, 2023.
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- 8. Edinger JD, Arnedt JT, Bertisch SM, et al. Behavioral and psychological treatment for chronic insomnia disorder in adults: an American Academy of Sleep Medicine clinical practice guideline. *J Clin Sleep Med.* 2021;17(2):255-262.
- 9. The 2019 American Geriatrics Society Beers Criteria Update Expert Panel. American Geriatrics Society 2019 Updated AGS Beers Criteria for Potentially Inappropriate Medication Use in Older Adults. *J Am Geriatr Soc* 00:1–21, 2019.

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