PRIOR AUTHORIZATION CRITERIA

BRAND NAME (generic)

DRUG CLASS

INSOMNIA AGENTS

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

Insomnia (Belsomra, Dayvigo, Quviviq) PA with Limit Policy 1177-C UDR 04-2023

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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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- 3. Quviviq [package insert]. Radnor, Pennsylvania: Idorsia Pharmaceuticals US Inc.; October 2022.
- 4. Lexicomp Online, AHFS DI (Adult and Pediatric) Online. Waltham, MA: UpToDate, Inc.; 2023. https://online.lexi.com. Accessed March 13, 2023.
- 5. Micromedex (electronic version). Merative, Ann Arbor, Michigan, USA. Available at: https://www.micromedexsolutions.com/ (cited: 03/13/2023).
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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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