# Belsomra (suvorexant) Dayvigo (lemborexant) Quviviq (daridorexant)

## I. CRITERIA FOR APPROVAL

An authorization may be granted when all the following criteria are met:

- A. The member has a diagnosis of insomnia disorder as characterized by the DSM-V (difficulty with sleep onset or sleep maintenance), **AND**
- B. Member is at least 18 years of age, AND
- C. 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**
- D. Member meets one of the following:
  - 1. Member has experienced an inadequate treatment response, intolerance, or contraindication to at least three formulary alternatives (i.e., zolpidem, trazodone, temazepam, etc.),
  - 2. Member has a documented history of known substance use disorder and/or is receiving current treatment for addiction, in which case most formulary alternatives may be deemed inappropriate,
  - 3. Provider is a sleep specialist with appropriate reasoning and supporting documentation for not prescribing a formulary alternative
- E. If requesting Quviviq, member has also experienced an inadequate treatment response, intolerance, or contraindication to Dayvigo or Belsomra

### **II. CONTINUATION OF THERAPY**

Authorization of 12 months may be granted for all members who meet all initial criteria and who achieve and maintain positive clinical response after at least 6 months of therapy with Belsomra, Dayvigo, or Quviviq as evidenced by an improvement in sleep onset and/or sleep maintenance.

### III. QUANTITY LIMIT

• Belsomra, Dayvigo, Quviviq: 1 tablet per day

### **IV. COVERAGE DURATION**

• 12 months

### **V. REFERENCES**

- 1. Belsomra [package insert]. Whitehouse Station, New Jersey: Merck Sharp & Dohme Corp., a subsidiary of Merck & Co., Inc.; March 2025.
- 2. Dayvigo [package insert]. Woodcliff Lake, New Jersey: Eisai; February 2025.
- 3. Quviviq [package insert]. Randor, Pennsylvania: Idorsia Pharmaceuticals US Inc.; October 2024.

