



AETNA BETTER HEALTH®  
Coverage Policy/Guideline

Name: Nuvigil (armodafinil)

Page: 1 of 3

Effective Date: 10/24/2023

Last Review Date: 10/24/2023

Applies to:	<input type="checkbox"/> Illinois	<input type="checkbox"/> Florida	<input checked="" type="checkbox"/> Florida Kids
	<input checked="" type="checkbox"/> New Jersey	<input type="checkbox"/> Maryland	<input type="checkbox"/> Michigan
	<input checked="" type="checkbox"/> Pennsylvania Kids	<input type="checkbox"/> Virginia	<input type="checkbox"/> Texas

### Intent:

The intent of this policy/guideline is to provide information to the prescribing practitioner outlining the coverage criteria for Nuvigil (armodafinil) under the patient's prescription drug benefit.

### Description:

Nuvigil (armodafinil) is indicated to improve wakefulness in adult patients with excessive sleepiness associated with obstructive sleep apnea (OSA), narcolepsy, or shift work disorder (SWD).

### Limitations of Use

In OSA, Nuvigil (armodafinil) is indicated to treat excessive sleepiness and not as treatment for the underlying obstruction. If continuous positive airway pressure (CPAP) is the treatment of choice for a patient, a maximal effort to treat with CPAP for an adequate period of time should be made prior to initiating Nuvigil (armodafinil) for excessive sleepiness.

### Applicable Drug List:

Armodafinil

### Policy/Guideline:

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

- The patient has a diagnosis of narcolepsy
  - AND**
  - The request is for continuation of therapy
    - AND**
    - The patient had a positive response to treatment
  - OR**
  - The requested drug is being prescribed by, or in consultation with, a sleep specialist
    - AND**
    - The diagnosis is confirmed by sleep lab evaluation
- OR**
- The patient has a diagnosis of shift work disorder (SWD)
  - AND**
  - The request is for continuation of therapy
    - AND**
    - The patient had a positive response to treatment
  - AND**



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- The patient is still a shift-worker

**OR**

- The requested drug is being prescribed by, or in consultation with, a sleep specialist

**AND**

- A sleep log and actigraphy monitoring have been completed for at least 14 days and shows a disrupted sleep and wake pattern

**AND**

- Symptoms have been present for 3 or more months

**OR**

- The patient has a diagnosis of obstructive sleep apnea (OSA)

**AND**

- The request is for continuation of therapy

**AND**

- The patient had a positive response to treatment

**AND**

- The patient is compliant with using continuous positive airway pressure (CPAP) or bilevel positive airway pressure (BIPAP)

**OR**

- The requested drug is being prescribed by, or in consultation with, a sleep specialist

**AND**

- The diagnosis has been confirmed by polysomnography

**AND**

- The patient has been receiving treatment for the underlying airway obstruction (continuous positive airway pressure [CPAP] or bilevel positive airway pressure [BIPAP]) for at least one month

**AND**

- Treatment with continuous positive airway pressure (CPAP) or bilevel positive airway pressure (BIPAP) will continue

#### Approval Duration and Quantity Restrictions:

**Approval:** 12 months

#### Quantity Level Limit:

- Nuvigil (armodafinil) 50 mg: 60 tablets / 25 days\* or 180 tablets / 75 days\*
- Nuvigil (armodafinil) 150 mg, 200 mg, 250 mg: 30 tablets / 25 days\* or 90 tablets / 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.*



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