

## Intent:

The intent of this policy/guideline is to provide information to the prescribing practitioner outlining the coverage criteria for amphetamine products under the member's prescription drug benefit.

## **Description:**

## Adderall

Adderall is indicated for the treatment of Attention-Deficit Hyperactivity Disorder (ADHD) and Narcolepsy.

## Adderall XR

Adderall XR is indicated for the treatment of Attention-Deficit Hyperactivity Disorder (ADHD).

# Adzenys ER, Adzenys XR-ODT, Desoxyn, Dyanavel XR

These products are indicated for the treatment of Attention-Deficit Hyperactivity Disorder (ADHD) in patients 6 years and older.

# Dexedrine

## Narcolepsy

<u>Attention Deficit Disorder with Hyperactivity</u> as an integral part of a total treatment program that typically includes other measures (psychological, educational, social) for patients (ages 6 years to 16 years) with this syndrome.

# Dextroamphetamine, ProCentra, Zenzedi

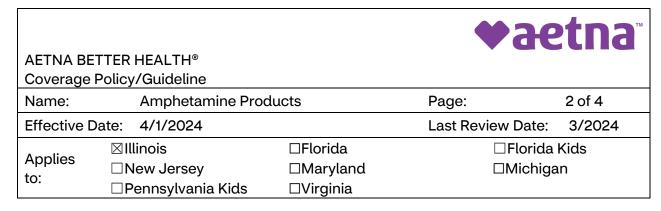
## <u>Narcolepsy</u>

<u>Attention Deficit Disorder with Hyperactivity</u> as an integral part of a total treatment program which typically includes other remedial measures (psychological, educational, social) for a stabilizing effect in pediatric patients (ages 3 to 16 years) with a behavioral syndrome characterized by the following group of developmentally inappropriate symptoms: moderate to severe distractibility, short attention span, hyperactivity, emotional lability, and impulsivity.

# Evekeo

## **Narcolepsy**

<u>Attention Deficit Disorder with Hyperactivity</u> as an integral part of a total treatment program which typically includes other remedial measures (psychological, educational, social) for a stabilizing effect in children with behavioral syndrome characterized by the following group



of developmentally inappropriate symptoms: moderate to severe distractibility, short attention span, hyperactivity, emotional lability, and impulsivity.

<u>Exogenous Obesity</u> as a short term (a few weeks) adjunct in a regimen of weight reduction based on caloric restriction for patients refractory to alternative therapy, e.g., repeated diets, group programs, and other drugs. The limited usefulness of amphetamines should be weighed against possible risks inherent in use of the drug.

## **Evekeo ODT**

Evekeo ODT is indicated for the treatment of Attention-Deficit Hyperactivity Disorder (ADHD) in pediatric patients 3 to 17 years of age.

## **Mydayis**

Mydayis is indicated for the treatment of Attention-Deficit Hyperactivity Disorder (ADHD) in patients 13 years and older.

## <u>Xelstrym</u>

Xelstrym is indicated for the treatment of Attention-Deficit Hyperactivity Disorder (ADHD) in adults and pediatric patients 6 years and older.

Compendial Uses Narcolepsy

# Applicable Drug List:

Reference Non-Preferred drugs and Preferred with Prior Authorization drugs on the Illinois Medicaid Preferred Drug List

## **Policy/Guideline:**

## Documentation for Initial Requests for all indications:

For non-preferred medication requests, the patient is unable to take three (3) formulary alternatives for the given diagnosis due to a trial and inadequate treatment response or intolerance, or a contraindication. Documentation is required for approval.

## **Criteria**:

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

• The patient has a diagnosis of Attention-Deficit/Hyperactivity Disorder (ADHD) or Attention Deficit Disorder (ADD) **AND** 



## **AETNA BETTER HEALTH®**

Coverage Policy/Guideline

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Effective Date: 4/1/2024			Last Review Date:	3/2024			
Applies to:	⊠Illinois	□Florida	🗆 Florida Kids				
	□New Jersey	□Maryland	□Michigan				
	🗆 Pennsylvania Kids	□Virginia					

 The diagnosis has been appropriately documented (e.g., evaluated by a complete clinical assessment, using DSM-5, standardized rating scales, interviews/questionnaires) AND

- The patient is 6 years of age or older OR
- The patient is 5 years of age or younger AND
  - The patient continues to have ADHD/ADD (Attention-Deficit/Hyperactivity Disorder or Attention Deficit Disorder) symptoms despite participating in evidence-based behavioral therapy (e.g., parent training in behavior management (PTBM), behavioral classroom interventions)

## OR

- The request is for continuation of therapy **AND**
- The patient achieved or maintained improvement in their signs and symptoms of ADHD/ADD (Attention-Deficit/Hyperactivity Disorder or Attention Deficit Disorder) from baseline

## AND

 The patient's need for continued therapy has been assessed within the previous year

# OR

• The patient has a diagnosis of narcolepsy

# AND

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

# AND

 $\circ$   $\;$  The diagnosis has been confirmed by a sleep study  $\;$ 

OR

- The request is for continuation of therapy **AND**
- The patient achieved or maintained improvement in daytime sleepiness with narcolepsy from baseline

# **Approval Duration and Quantity Restrictions:**

**Approval:** Initial and Renewal - Approve 12 months **Quantity Level Limit:** Reference Formulary for drug specific quantity level limits

	ETTER HEALTH® Policy/Guideline		<b>*a</b>	etna™
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Applies to:	⊠Illinois □New Jersey □Pennsylvania Kids	□Florida □Maryland □Virginia	□Florida Kids □Michigan	

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