Mydayis® (amphetamine salts extended-release)
Effective 02/20/19

Plan	☒ MassHealth ☐Commercial/Exchange	**Program Type**	☒ Prior Authorization ☒ Quantity Limit ☐ Step Therapy
Benefit	☒ Pharmacy Benefit ☐ Medical Benefit (NLX)		
Specialty Limitations	N/A		
Contact Information	**Specialty Medications**		
	All Plans	Phone: 866-814-5506	Fax: 866-249-6155
	Non-Specialty Medications		
	MassHealth	Phone: 877-433-7643	Fax: 866-255-7569
	Commercial	Phone: 800-294-5979	Fax: 888-836-0730
	Exchange	Phone: 855-582-2022	Fax: 855-245-2134
	Medical Specialty Medications (NLX)		
	All Plans	Phone: 844-345-2803	Fax: 844-851-0882
Exceptions	N/A		

Overview
Mydayis is a central nervous system (CNS) stimulant prescription medicine used for the treatment of Attention Deficit Hyperactivity Disorder (ADHD) in people 13 years of age and older.

Coverage Guidelines
Authorization may be granted when the following criteria are met:

1. Member has a diagnosis of attention deficit hyperactivity disorder (ADHD) or autism spectrum disorder (ASD) **AND**
2. Member is at least 13 years of age or older **AND**
3. Prescriber has provided documentation of an inadequate response, adverse reaction, or contraindication to Adderall XR® (amphetamine salts extended-release) **AND**
4. Prescriber has provided documentation of an inadequate response, adverse reaction, or contraindication to Vyvanse® (lisdexamfetamine)

Please note: Additional criteria may apply for members under the age of 18. Please refer to the MassHealth Pediatric Behavioral Health Medication Initiative guideline for criteria.

Continuation of Therapy
Reauthorization requires physician documentation indicating a positive response to therapy.

Limitations
1. Approvals will be granted for 12 months.
2. A quantity limit of a maximum of 30 units per 30 days applies.

Appendix

Adverse reactions/contraindications to stimulants include but are not limited to weight loss or failure to gain weight, insomnia, agitation, increased anger and aggression, tic disorder, seizures w/ ADHD, traumatic brain injury w/ ADHD (trial of **both** a methylphenidate product and amphetamine product is required for approval), symptomatic heart disease in children (tachycardia or hypertension), symptomatic heart disease in adults (there should be at least 1 failed trial of another non-stimulant medication that may be used for ADHD such as bupropion.).

References

1. Mydayis Package Insert, Shire Laboratories, 2017.

Review History

03/01/18 – Effective (adopted MH RS)
02/20/19 – Reviewed in P&T Meeting

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