

Tarpeyo (budesonide controlled-release) Effective 07/01/2022

Plan	 □ MassHealth UPPL ⊠Commercial/Exchange 		Prior Authorization
Benefit	 Pharmacy Benefit Medical Benefit 	Program Type	 Quantity Limit Step Therapy
Specialty Limitations	This medication has been designated specialty and must be filled at a contracted specialty pharmacy.		
	Medical and Specialty Medications		
Contact	All Plans	Phone: 877-519-1908	Fax: 855-540-3693
Information	Non-Specialty Medications		
	All Plans	Phone: 800-711-4555	Fax: 844-403-1029
Exceptions	N/A		

Overview

Tarpeyo (budesonide controlled-release) is FDA approved to reduce proteinuria in adults with primary immunoglobulin A nephropathy (IgAN) at risk of rapid progression of disease, generally a urine protein-to-creatinine ratio \geq 1.5 g/g.

Coverage Guidelines

Authorization may be granted for members new to AllWays Health Partners who are currently receiving treatment and is stable with Tarpeyo, excluding when the product is obtained as samples or via manufacturer's patient assistance programs.

OR

Authorization may be granted when ALL of the following criteria is met:

- 1. The member is at least 18 years of age
- 2. Member has a diagnosis of primary immunoglobulin A nephropathy (IgAN)
- 3. Member is using Tarpeyo to reduce proteinuria
- 4. MD documented urine protein-to-creatinine ratio ≥ 1.5 g/g
- 5. The member has had intolerance, adverse events, or contraindication to generic budesonide capsules/tablets

Continuation of Therapy

Reauthorization of may be granted for all members who have a positive response to therapy as evidence by low disease activity or improvement in signs and symptoms of the condition.

Limitations

- 1. Initial approvals and reauthorizations will be granted for 24 months
- 2. The following quantity limits apply:

Tarpeyo capsule120 capsules per 30 days

Mass General Brigham Health Plan includes Mass General Brigham Health Plan, Inc. and Mass General Brigham Health Insurance Company.

References

1. Tarpeyo (budesonide) [prescribing information]. Stockholm Sweden: Colliditas Therapeutics AB; December 2021.

Review History

05/18/2022 – Created and reviewed for May P&T. Effective 07/01/2022.