

Alzheimer's Agents
Kisunla (donanemab-azbt)
Effective 05/12/2025

Plan	<input checked="" type="checkbox"/> MassHealth UPPL <input type="checkbox"/> Commercial/Exchange	Program Type	<input checked="" type="checkbox"/> Prior Authorization
Benefit	<input type="checkbox"/> Pharmacy Benefit <input checked="" type="checkbox"/> Medical Benefit		<input type="checkbox"/> Quantity Limit <input type="checkbox"/> Step Therapy
Specialty Limitations	N/A		
Contact Information	Medical and Specialty Medications		
	All Plans	Phone: 877-519-1908	Fax: 855-540-3693
Contact Information	Non-Specialty Medications		
	All Plans	Phone: 800-711-4555	Fax: 844-403-1029
Notes	Kisunla is also available on the pharmacy benefit. Please see the MassHealth Drug List for coverage and criteria. Additional agents from this class are available through the pharmacy benefit. Please see the MassHealth Drug List for coverage and criteria.		

Overview

Kisunla (donanemab-azbt) is an amyloid beta-directed antibodies indicated for the treatment of Alzheimer's disease. Treatment with Kisunla should be initiated in patients with mild cognitive impairment or mild dementia stage of disease.

Coverage Guidelines

Authorization may be reviewed on a case by case basis for members who are new to the plan currently receiving treatment with requested medication excluding when the product is obtained as samples or via manufacturer's patient assistance programs.

OR

Authorization may be granted for members when all the following criteria are met:

- Diagnosis of **ONE** of the following:
 - Mild cognitive impairment (MCI)
 - Mild dementia associated with Alzheimer's Disease (AD)
- Prescriber is a specialist in the treatment of dementia or Alzheimer's disease (e.g., neurologist, geriatric psychiatrist, geriatrician who specializes in treating dementia)
- Test results indicating clinically significant AD neuropathology based on **ONE** of the following:
 - Amyloid PET
 - Cerebral Spinal Fluid (CSF) biomarkers
- Member has had a brain magnetic resonance imaging (MRI) within the last 12 months
- Appropriate dosing

6. Baseline cognitive function test (dated within the last three months) based on **ONE** of the following objective assessments:
 - a. Mini Mental State Exam (MMSE) score ≥ 20
 - b. Montreal Cognitive Assessment (MoCA) score ≥ 15
 - c. Saint Louis University Mental Status Examination (SLUMS) score ≥ 16.1

Continuation of Therapy

1. Appropriate dosing
2. Attestation that all MRI monitoring has been completed in accordance with the FDA approved label
3. Current cognitive function test (within the past three months) based on ONE of the following objective assessments:
 - a. Mini Mental State Exam (MMSE)
 - b. Montreal Cognitive Assessment (MoCA)
 - c. Saint Louis University Mental Status Examination (SLUMS)

Limitations

1. Initial approvals will be granted for 6 months
2. Reauthorizations will be granted for 12 months

References

1. Kisunla [package insert]. Indianapolis (IN): Eli Lilly and Company; 2024 July.

Review History

12/11/24 – Created criteria for P&T. Kisunla will be managed as dual benefit, therefore medical criteria was created to be posted on MGBHP website (note: no preferred product requirement). Rx will be available on MHDL. Effective 2/28/2025

04/09/25 – Reviewed and updated for P&T. Removed medical record requirement. Effective 05/12/25

