

Alzheimer's Agents
Leqembi (lecanemab-irmb)
Effective 06/01/2025

Plan	<input checked="" type="checkbox"/> MassHealth <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 Benefit	Phone: 833-895-2611	Fax: 888-656-6671	
	Pharmacy Benefit	Phone: 800-711-4555	Fax: 844-403-1029	
Exceptions	Leqembi 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

Leqembi (lecanemab-irmb) is indicated for the treatment of Alzheimer's disease. Treatment with Leqembi should be initiated in patients with mild cognitive impairment or mild dementia stage of disease, the population in which treatment was initiated in clinical trials. It initially received accelerated approval similar to Aduhelm, but subsequently received a traditional approval after a clinical benefit was established. Of note, while Leqembi appears to be a safer alternative than Aduhelm, there is a black box warning for amyloid-related imaging abnormalities (ARIA), particularly those patients who are apolipoprotein E (ApoE) ε4 homozygotes.

Coverage Guideline

Authorization may be granted for new members to the plan who are currently receiving treatment with requested medication excluding when the product is obtained as samples or via manufacturer's patient assistance programs, when continuation of therapy criteria is met.

OR

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

1. Diagnosis of ONE of the following:
 - a. Mild cognitive impairment (MCI)
 - b. Mild dementia associated with Alzheimer's Disease (AD)
2. Prescriber is a specialist in the treatment of dementia or Alzheimer's disease (e.g., neurologist, geriatric psychiatrist, geriatrician who specializes in treating dementia)
3. Test results indicating clinically significant AD neuropathology based on ONE of the following:
 - a. Amyloid PET
 - b. Cerebral Spinal Fluid (CSF) biomarkers
4. Member has had a brain magnetic resonance imaging (MRI) within the last 12 months
5. Appropriate dosing
6. Baseline cognitive function test (within the last three months) based on ONE of the following objective assessments:
 - a. Mini Mental State Exam (MMSE) score ≥ 22

- b. Montreal Cognitive Assessment (MoCA) score ≥ 15
- c. Saint Louis University Mental Status Examination (SLUMS) score ≥ 16.1

Continuation of Therapy

First reauthorization:

- 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)

Subsequent reauthorization (after completion of 18 months of treatment):

- 1. Attestation that all MRI monitoring has been completed in accordance with the FDA approved label
- 2. 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)
- 3. ONE of the following:
 - a. Dosing frequency reduced to every four weeks
 - b. Clinical rationale for continuing biweekly dosing

Limitations

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

References

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Review History

03/16/2022 – Reviewed and Created for March P&T; Match MH criteria Effective 05/01/2022.

01/11/2023 - Reviewed and updated for Jan P&T. Matched MH UPPL criteria. Adlarity was added to pharmacy benefit with PA and QL. Updated approval durations. Effective 3/1/23.

09/13/2023 – Reviewed and updated for P&T. Added Leqembi to criteria. Aduhelm initial criteria: Clarified provider specialty, SLUMS added as another assessment options, changed timeline of MRI scan from 3 to 12 months, and preferred trial of Leqembi. Aduhelm reauth criteria further simplified to require current objective assessments and attestation that all MRI monitoring has been completed. Effective 10/2/23

05/15/25 – Reviewed and updated for P&T. Updated formatting and references. Removed Aduhelm due to removal from the market. Adlarity is removed as it is available on the pharmacy benefit and criteria is available on MHD. Leqembi remains due to dual benefit. Criteria was updated to remove requirement of medical records and info on PA is acceptable. Reauthorization criteria has been split based on first and subsequent requests. Effective 6/1/25

