

Leqembi (lecanemab-irmb)

Infusion Office Preference: _____

PATIENT INFORMATION

Date:	Patient Name:	DOB:
<input type="checkbox"/> NKDA Allergies:	Weight (lbs / kg):	Height:
Patient Status: <input type="checkbox"/> New to Therapy <input type="checkbox"/> Continuing Therapy - Last Treatment Date:	Next Due Date:	

DIAGNOSIS AND ICD 10 CODE

<input type="checkbox"/> Alzheimer's disease with early onset	ICD 10 Code: G30.0
<input type="checkbox"/> Mild Cognitive Impairment, So stated	ICD 10 Code: G31.84
<input type="checkbox"/> other ICD10 Code:	Description:

Please choose one G code from the items below, automatic secondary will be added

<input type="checkbox"/> G30.1 Alzheimer's disease late onset	Secondary • Z00.6 Encounter for examination for normal comparison and control in clinical research program
<input type="checkbox"/> G30.8 Other Alzheimer's disease	
<input type="checkbox"/> G30.9 Alzheimer's disease , unspecified	

This signed order form by the provider Patient demographics AND insurance info Clinical/Progress notes

Prescriber must indicate that the following requirements have been met (provide supporting documentation)

Beta Amyloid Pathology Confirmed via:

↳ Amyloid PET Scan **OR** CFS Analysis-Date: _____ Result: _____

Cognitive Assessment Used: _____ Date: _____ Result: _____

ApoE εε4 Genetic Test - Date: _____ Result: Homozygote Heterozygote Noncarrier

MEDICARE REGISTRATION # IF APPLICABLE _____

MEDICATION ORDERS(Note: Only one stage of treatment may be ordered at a time)

Stage 1 (Infusions #1 and #2) ✓ Leqembi 10mg/kg IV every two weeks x 2 doses. Each infusion to be given over one hour.
Required Documentation to Initiate this Phase:
MRI of brain within one year prior to first infusion. Date of MRI: _____

By checking this box, I confirm that Beta Amyloid Pathology has been confirmed via CSF or PET.

Stage 2 (Infusions #3 and #4) ✓ Leqembi 10mg/kg IV every two weeks x 2 doses. Each infusion to be given over one hour.
Required Documentation to Initiate this Phase:
 By checking this box, I confirm that patient has undergone MRI of brain before dose #3. I have reviewed the results and clear patient to proceed with infusions #3 and #4

Stage 3 (Infusions #5 and #6) ✓ Leqembi 10mg/kg IV every two weeks x 2 doses. Each infusion to be given over one hour.
Required Documentation to Initiate this Phase:
 By checking this box, I confirm that patient has undergone MRI of brain before dose #5. I have reviewed the results and clear patient to proceed with infusions #5 and #6.

Stage 4 (Infusions #7-13) ✓ Leqembi 10mg/kg IV every two weeks x 7 doses. Each infusion to be given over one hour.
Required Documentation to Initiate this Phase:
 By checking this box, I confirm that patient has undergone MRI of brain before dose #7. I have reviewed the results and clear patient to proceed with infusions #7 through #13.

Fax referral to 866-507-1164 or email to bionurses@metroinfusioncenter.com

All information contained in this order form is strictly confidential and will become 01/29/25part of the patient's medical record.

*** Stage 5 (Infusions #14 & beyond) MUST SELECT ONE DOSING BELOW**

Leqembi 10mg/kg IV every **two weeks** over one hour **OR** Leqembi 10mg/kg IV every **four weeks** over one hour.

Required Documentation to Initiate this Phase:

By checking this box, I confirm that patient has undergone MRI of brain before dose #14. I have reviewed the results and clear patient to proceed with infusions #14 and beyond as ordered above

PRE-INFUSION :

- Confirm baseline MRI results prior to initiation of treatment
- Confirm MRI completed and reviewed by prescriber prior to the 5th, 7th, and 14th treatment
- Measure and record weight prior to each treatment to determine dose
- Hold infusion and notify provider if patient reports:
 - Headache
 - Dizziness
 - Nausea
 - Vision Changes
 - New or worsening confusion

Post -INFUSION :

- Educate patient/care partner to report headache, dizziness, nausea, vision changes, or new/worsening confusion.
- Fax infusion record to provider below :

PRESCRIBER INFORMATION

Provider Name (print)

Provider Signature**:

Date:

Office Phone

Office Fax

****Amyloid Related Imaging Abnormalities (ARIA): Enhanced clinical vigilance for ARIA is recommended during the first 14 weeks of treatment with LEQEMBI. Referring physician to monitor patient**

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Revised 03/23/26