

LEQEMBI ORDER FORM

PATIENT INFORMATION

Last Name: _____ First Name: _____ MI: _____ DOB: _____

HT: _____ in WT: _____ kg Sex: Male Female Allergies: NKDA, _____

Physician Name _____ Contact Name _____ Contact Phone # _____

NPI #: _____ Tax ID#: _____ Fax #: _____

Does patient have venous access? YES NO If yes, what type MEDIPORT PIV PICC LINE OTHER: _____

a) ALL MEDIPORTS / IV ACCESSES WILL BE FLUSHED WITH HEPARIN OR SALINE PER HOSPITAL PROTOCOL PRN

PLEASE SELECT FROM BELOW:

- _____ Perform port flush every _____ weeks per hospital policy.
- _____ Perform IV site care per hospital protocol.
- _____ Activase 2mg IVP per hospital protocol.

DUAL DIAGNOSIS IS REQUIRED – SELECT ONE OPTION OF BOTH CONDITIONS THAT APPLY FROM BELOW:

- | | | |
|--|--|---|
| <input type="checkbox"/> G30.0 Alzheimer's Disease, Early Onset
<input type="checkbox"/> G30.1 Alzheimer's Disease, Late Onset
<input type="checkbox"/> G30.8 Other Alzheimer's disease
<input type="checkbox"/> G30.9 Alzheimer's disease, unspecified
<input type="checkbox"/> G31.84 Mild Cognitive Impairment, So Stated
<input type="checkbox"/> Other: _____ (ICD 10 + Description) | ← G30.X codes require
secondary F02.8X code → | <input type="checkbox"/> F02.80 Dementia without behavioral disturbance
<input type="checkbox"/> F02.81 Dementia with behavioral disturbance |
|--|--|---|

Prescriber must indicate the following requirements have been met (please provide documentation):

- | | |
|---|---|
| <input type="checkbox"/> Beta Amyloid Pathology Confirmed Via
<input type="checkbox"/> Amyloid PET Scan Date: _____ OR
<input type="checkbox"/> Cognitive Assessment Used: _____ Date: _____ Result: _____
<input type="checkbox"/> ApoE ε4 Genetic Test Date: _____ Result: <input type="checkbox"/> Homozygote <input type="checkbox"/> Heterozygote <input type="checkbox"/> Noncarrier | <input type="checkbox"/> CSF Analysis Date: _____ Result: _____
<input type="checkbox"/> F02.81 Dementia with behavioral disturbance |
|---|---|

PRESCRIPTION ORDERS

Leqembi	10mg X _____ kg = _____ mg	IV Over At Least 60 Minutes	Every 2 Weeks (at least 14 days apart)	12 Months
<small>DRUG</small>	<small>DOSE</small>	<small>ROUTE</small>	<small>FREQUENCY</small>	<small>DURATION</small>

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/caregiver to report headache, dizziness, nausea, vision changes, or new/worsening confusion.

Physician's Signature _____ Time _____ Date _____
*Signature Must Be Clear and Legible

Cosignature (If Required) _____ Time _____ Date _____
*Signature Must Be Clear and Legible

Fax completed form, supporting documentation, facesheet, and insurance cards to the Outpatient Infusion Center at 1 (877) 249-1191.