

PROVIDER ORDER FORM

lecanemab-irmb (Leqembi) Infusions

- Please complete form and fax with latest clinical documentation to 360.299.4237
- Documentation must include a recent MRI of the brain and confirmation of amyloid beta pathology.
- Testing for ApoE ε4 status should be included for ARIA risk assessment.

Patient Name (First, Middle, Last): ______ D.O.B. _____ D.O.B.

| Height: cm in Weight: kg lbs | Allergies: |
|---|---------------------------|
| Diagnosis: ICD-10 Code: | |
| Pre-auth Done? 🗌 Not required 🗌 Yes Authorization | n #: Authorization Dates: |

Medication Orders:

Lecanemab-irmb (Leqembi): Infuse 10 mg/kg (actual body weight) (_____mg) IV over 1 hour every 2 weeks for one year

- Adjust dose for weight changes.
- Dilute medication in 250 mL 0.9% NS, use a 0.2 micron in-line filter.
- Before the 5th, 7th and 14th dose, an MRI report and prescriber's written approval to proceed; Must be provided and entered in the patient chart at Island Infusion Center.

Other

In Case of Infusion Reaction, Orders will be carried out per Island Infusion Center Policy unless crossed out.

- Stop infusion, obtain VS and assessment, call Provider*
- NS 0.9% 1000 mL/hour IV for blood pressure < 100/60.
- O2 per nasal cannula to maintain saturation >92%.
- Diphenhydramine 25 mg IV x1 if patient was premedicated with 25 mg within 2 hours of reaction, otherwise give 50 mg IV x1.
- Methylprednisolone 125 mg IV x1.
- Epinephrine 0.3 mg IM for severe reaction x1. (May repeat x1 in 3-5 minutes for continued severe symptoms.)

Nursing Protocol

- Confirm baseline MRI prior to first treatment
- Verify ApoE ε4 status. Homozygotes are at higher risk of severe ARIA.
- Measure and record weight prior to each treatment to determine dose
- Confirm MRI results and prescriber approval to continue at 5th, 7th and 14th treatment
- Monitor for infusion and hypersensitivity reactions during infusion
- Hold infusion and notify provider for headache, dizziness, nausea, vision changes, confusion, gait difficulty, stroke symptoms, seizures
- Educate patient and caregiver to report the above symptoms, that these symptoms mimic those of ischemic stroke, and to inform all medical providers that she is receiving Leqembi infusions.
- Required FDA medication guide will be provided to patient and caregiver.
- Patient will be instructed to consult with her prescriber before starting any new medications with bleeding risk.

| PROVIDER SIGNATURE | DATE |
|---------------------|-----------------------------|
| PRINT PROVIDER NAME | *PROVIDER EMERGENCY PHONE # |