

PROVIDER ORDER FORM

Donanemab-azbt (Kisunla) Infusions

- Please complete form and fax with latest clinical documentation to 360.299.4237 ٠
- 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. _____ Height: _____ cm 🗌 in Weight:_____ 🗌 kg 🗌 lbs Allergies: ______ _____ ICD-10 Code: _____

Pre-auth Done? 🗌 Not required 🗌 Yes Authorization #: ______ Authorization Dates: ______

Medication Orders:

Diagnosis: _____

Donanemab-azbt (Kisunla): refill as directed for one year

Initial Dose: Infuse 700 mg IV over 30 minutes every 4 weeks for 3 infusions

Maintenance Dose: Infuse 1400 mg IV over 30 minutes every 4 weeks

Prior to the 2nd, 3rd, 4th and 7th infusions, 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 ɛ4 status. Homozygotes are at higher risk of severe ARIA.
- Confirm MRI results and prescriber approval to continue at 2nd, 3rd, 4th and 7th treatment
- Monitor for infusion and hypersensitivity reactions during infusion and for at least 30 minutes post infusion
- Hold infusion/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 Kisunla infusions.
- 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 #