

PATIENT INFORMATION
PATIENT NAME: _____ **PHONE:** _____

DATE OF BIRTH: _____ **SEX:** M F **HEIGHT:** _____ **WEIGHT:** _____ LBS KG

ALLERGIES: _____ **PREFERRED CLINIC:** _____

REFERRAL STATUS: NEW REFERRAL ORDER CHANGE ORDER RENEWAL

DIAGNOSIS & CLINICAL DOCUMENTATION

- | | |
|---|--|
| <input type="checkbox"/> G30.0 Alzheimer's disease with early onset | <input type="checkbox"/> G30.9 Alzheimer's disease, unspecified |
| <input type="checkbox"/> G30.1 Alzheimer's disease with late onset | <input type="checkbox"/> G31.84 Mild cognitive impairment, so stated |
| <input type="checkbox"/> G30.8 Other Alzheimer's disease | |

 ICD-10 CODE: _____ DESCRIPTION: _____

REQUIRED DOCUMENTATION

- | | | | | |
|--|--|---|---|---|
| <input type="checkbox"/> Insurance Information | <input type="checkbox"/> List of Medications | <input type="checkbox"/> Most Recent History & Physical | <input type="checkbox"/> Tried & Failed Therapies | <input type="checkbox"/> Cognitive Assessment & Score |
| <input type="checkbox"/> Functional Assessment & Score | <input type="checkbox"/> Confirmed Amyloid Pathology | <input type="checkbox"/> Recent MRI prior to initiating Kisunla | <input type="checkbox"/> Proof of CED registry submission | |

MEDICATION ORDER
***Referring provider responsible for obtaining MRI prior to infusion #2, #3, #4 and #7 for monitoring of ARIA**

- LOADING:** Kisunla® (donanemab) 350 mg IV in 50 ml NS over 30 minutes at week 0; 700 mg IV in 100 ml NS over 30 minutes at week 4; 1,050 mg IV in 100 ml NS over 30 minutes at week 8.
- MAINTENANCE:** Kisunla® (donanemab) 1,400 mg IV in 250 ml NS over 30 minutes every 4 weeks
 *Maintenance dosing scheduled 4 weeks from last loading dose (week 8 dose)

 REFILL X 12 MONTHS UNLESS OTHERWISE NOTED HERE: _____

Patient to be observed for 30 minutes following each administration. Administer per protocol.
 In the event of an adverse reaction occurring in the infusion clinic, utilize the Immersiv Health adverse reaction protocol.
PRE-MEDICATIONS
PO

- Acetaminophen: 650 mg
 Cetirizine: 10 mg
 Diphenhydramine: 25 mg

IV

- Methylprednisolone: 125 mg
 Diphenhydramine: 25 mg

 OTHER: _____ PO IV

PRESCRIBER INFORMATION
PROVIDER NAME: _____ **NPI #:** _____

EMAIL: _____ **PHONE:** _____ **FAX:** _____

ADDRESS (INCLUDE CITY, STATE, ZIP): _____

SUPERVISING PHYSICIAN: _____ **CONTACT NAME:** _____
(IF APPLICABLE)
SIGNATURE: _____ **DATE:** _____
(NO STAMPS)

SUBSTITUTION PERMITTED

DISPENSE AS WRITTEN