

Kisunla (donanemab-azbt)

REFERRAL STATUS: New Referral Dose or Frequency Change Order Renewal
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 – please choose one G code from the items below, automatic secondary will be added		
<input type="checkbox"/> G30.0 Alzheimer's disease with early onset <input type="checkbox"/> G30.1 Alzheimer's disease with late onset <input type="checkbox"/> G30.8 Other Alzheimer's disease <input type="checkbox"/> G31.84 Mild cognitive impairment, so stated <input type="checkbox"/> G30.9 Alzheimer's disease, unspecified	Secondary Diagnosis: <ul style="list-style-type: none">• Z00.6 Encounter for examination for normal comparison and control in clinical research program	
<input type="checkbox"/> This signed order form by the provider <input type="checkbox"/> Patient demographics AND insurance info <input type="checkbox"/> Clinical/Progress notes		
Prescriber must indicate the following requirements have been met to confirm diagnosis and that Patient has evidence of AD neuropathology and has been assessed for baseline ARIA risk via MRI (provide supporting documentation)		
<input type="checkbox"/> Amyloid pathology confirmed via: _____ (Kisunla is not a treatment option for this Pt, if checked) ➔ <input type="checkbox"/> Amyloid PET Scan OR <input type="checkbox"/> CSF analysis OR <input type="checkbox"/> Blood plasma Date: _____ Result: <input type="checkbox"/> Amyloid Positive <input type="checkbox"/> Amyloid Negative ↵		
<input type="checkbox"/> Recent MRI obtained prior to initiating Kisunla (including FLAIR and T2/GRE or SWI) to assess ARIA risk ➔ <input type="checkbox"/> Prescriber has verified that this Patient does not have evidence of prior ARIA-H Date: _____		
<input type="checkbox"/> Completion of cognitive assessment type: ➔ <input type="checkbox"/> MMSE <input type="checkbox"/> MoCA <input type="checkbox"/> CDR <input type="checkbox"/> Other Date: _____		
<input type="checkbox"/> Completion of functional assessment type: ➔ <input type="checkbox"/> FAQ <input type="checkbox"/> FAST <input type="checkbox"/> Other Date: _____		
<input type="checkbox"/> Completion of CMS approved CED registry (only required for Patients with Medicare) ClinicalTrials.gov Registry Number: NCT _____ CED Submission Date: _____ Submission Number (if applicable): _____		
MEDICATION ORDERS		
Initial dosing	<input type="checkbox"/> Kisunla 350 mg IV at 0, 700mg IV at week 4, 1050mg IV at week 8	
Maintenance dosing	<input type="checkbox"/> Kisunla 1400 mg IV every 4 weeks thereafter	
Refills*: <input type="checkbox"/> None <input type="checkbox"/> X6 months <input type="checkbox"/> X1 year <input type="checkbox"/> Other: _____ <i>*(if not indicated order will expire one year from date signed)</i>		
REQUIRED MRIs		
Note: MRIs must be obtained prior to initial infusion to assess ARIA risk. During treatment, conduct an ARIA monitoring MRI before Infusions 2, 3, 4 and 7 infusion (this is the responsibility of the ordering provider).		
PRESCRIBER INFORMATION		
Provider Name (print)	Provider Signature*:	Date:
Office Phone	Office Fax	

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 part of the patient's medical record.