Kisunla (donanemab-azbt)

Office Phone



REFERRAL STATUS: □ New Referral □ Dose or Frequency Change □ Order Renewal Infusion Office Preference:			
PATIENT INFORMATION			
Date: Patient Name:		DOB:	
□ NKDA Allergies:		Weight (lbs / kg):	Height:
Patient Status: ☐ New to Therapy ☐ Continuing Therapy - Last Treatment Date: Next Due Date:			
DIAGNOSIS AND ICD 10 CODE			
 □ G30.0 Alzheimer's disease with early onset □ G30.1 Alzheimer's disease with late onset □ G30.8 Other Alzheimer's disease 		☐ G30.9 Alzheimer's disease, unspecified ☐ G31.84 Mild cognitive impairment, so stated	
REQUIRED DOCUMENTATION			
☐ This signed order form by the provider ☐ Patient demographics AND insurance info ☐ 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)			
□ Amyloid pathology confirmed via: (Kisunla is not a treatment option for this Pt, if checked) □ Amyloid PET Scan OR □ CSF analysis OR □ Blood plasma Date:Result: □ Amyloid Positive □ Amyloid Negative			
MEDICATION ORDERS			
Initial dosing	☐ Kisunla 350mg IV day 0, 700mg IV @ week 4, 1050mg IV @ week 8		
Maintenance dosing ☐ Kisunla 1400 mg IV every 4 weeks thereafter			
Refills*: None X1 year Other: *(if not indicated order will expire one year from date signed)			
SPECIAL INSTRUCTIONS			
PRESCRIBER INFORMATION			
Provider Name (print)	Provider Signa	ture*:	Date:

Fax referral to 866-507-1164 or email to bionurses@metroinfusioncenter.com

Office Fax