Fax: 832-895-4040 Phone: 832-895-5000

E-mail: intake@adaptivinfusion.com



$\mathsf{Kisunla}^\mathsf{TM} \left(\mathsf{donanemab-azbt} \right)$

Provider Order Form rev. 8/10/2024

PATIENT INFORMATION	Dafaw	al Chahara (I I I I)	□ Novy Deferral	□ I Indated Orde	or Order Deneural
PATIENT INFORMATION	Referr	al Status (check one):	☐ New Referral	·	er Order Renewal
Patient Name:				DOB:	
NKDA Allergies:		Weight	Please specify	r: □lbs □kg	Height:
Patient Status (check one): \square New to Therapy \square Continuing Therap	ру	Last Treatment Date:		Next Due Date	::
ICD-10 code (required): ICD-1	0 descrip	tion:			
REQUIRED: Demographics & Most Recent: H&P, cli	inical not	es, & medication list.	Supporting clinical	al notes to include	any
past tried and/or failed therapies, intolera PRESCRIPTION	ance, out	comes, or contraindi	cations to conven	tional therapy.	
	a a la callaca	PRE-INFUSION:			
 □ Provide nursing care per AdaptIV Infusion Nursing Procedures, including reaction management and post-procedure observation □ Medicare Registry # □ DIAGNOSIS: □ G30.0 Alzheimer's Disease, Early Onset □ G30.1 Alzheimer's Disease, Late Onset □ G30.8 Other Alzheimer's disease □ G30.9 Alzheimer's disease, unspecified □ G31.84 Mild Cognitive Impairment, So Stated G30.X codes require secondary F02.8X code BELOW: □ F02.80 Dementia without behavioral disturbance □ F02.81 Dementia with behavioral disturbance □ PRESCRIBER MUST INDICATE THE FOLLOWING REQUIREMENTS HAVE BEEN MET (PLEASE PROVIDE DOCUMENTATION): □ Obtain a recent baseline brain magnetic resonance imaging (MRI) prior to initiating treatment with KISUNLA. □ Beta Amyloid Pathology Confirmed Via: □ Amyloid PET Scan Date: □ OR 		 □ Confirm baseline MRI results prior to initiation of treatment. □ Confirm MRI completed and reviewed by prescriber prior to 2nd, 3rd, 4th, and 7th infusions. □ Hold infusion and notify provider if patient reports: • Headache • Vision changes • Dizziness • New or worsening confusion • Nausea TREATMENT FREQUENCY: □ Schedule treatments every 28 days/monthly/every 4 weeks. DOSE □ 700mg infusions 1,2, and 3 □ 1400mg infusion 4 and beyond MEDICATION: □ Administer KISUNLA over approximately 30 minutes □ If infusion-related reaction occurs, stop infusion and treat per orders/ protocol as clinically indicated. 			
CSF Analysis Date: Result:		POST-INFUSION:			
□ ApoE E4 Genetic Test: Date: Result: □ Homozygote □ Heterozygote □ Noncarrier		Educate patiervision changes	nt/care partner to r s, or new/worsenir notes to provider a		zziness, nausea,
SPECIAL INSTRUCTIONS					
PROVIDER INFORMATION					
Referral Coordinator Name:		Referral Coordinator Email:			
Ordering Provider:		Provider NPI:			
Referring Practice Name:		Phone:	Fax		
Practice Address:		City:	Sta	te: Zip Co	de:
Provider Name (Print) Provider Sign	ature			Date	