Leqembi[™] (Lecanemab-irmb)

Provider Order Form rev. 8/10/2023

PATIENT INFORMATION	Referral Status (check one): New Referral Updated Order Order Renewal
Patient Name:	DOB:
NKDA 🗆 Allergies:	Weight Please specify: Dbs bkg Height:
Patient Status <i>(check one):</i> New to Therapy Continuing Therapy	Last Treatment Date: Next Due Date:
ICD-10 code (required): ICD-10 c	description:
	ical notes, & medication list. Supporting clinical notes to include any ce, outcomes, or contraindications to conventional therapy.
PRESCRIPTION	
 Provide nursing care per AdaptIV Infusion Nursing Procedures, incl reaction management and post-procedure observation Medicare Registry # 	Iuding PRE-INFUSION: Confirm baseline MRI results prior to initiation of treatment. Confirm MRI completed and reviewed by prescriber prior to the 5th, 7th, and 14th treatment.
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	 Measure and record weight prior to each treatment to determine dose. Hold infusion and notify provider if patient reports: Headache Vision changes Dizziness New or worsening confusion Nausea
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 BE MET (PLEASE PROVIDE DOCUMENTATION): Beta Amyloid Pathology Confirmed Via: Amyloid PET Scan Date:	 MEDICATION: Administer LEQEMBI 10 mg/kg intravenously over at least 60 minutes. Dilute required volume of lecanemab-irmb in 250 ml 0.9% sodium chloride and infuse using a terminal low-protein binding 0.2-micron in-line filter. If infusion-related reaction occurs, stop infusion and treat per orders/ protocol as clinically indicated.
OR Result: CSF Analysis Date: Result: Cognitive Assessment Used: Date: Date: Result: ApoE E4 Genetic Test: Date: Result: Homozygote Heterozygote Noncarrier	TREATMENT FREQUENCY: Schedule treatments every two weeks (at least 14 days apart). POST-INFUSION: Educate patient/care partner to report headache, dizziness, nausea, vision changes, or new/worsening confusion. Fax treatment notes to provider at number below.

adaptIV infusion

SPECIAL INSTRUCTIONS

PROVIDER INFORMATION

Referral Coordinator Name: Referral Coordinator Email: Ordering Provider: Provider NPI: Referring Practice Name: Phone: Practice Address: City: State: Zip Code: