## Lecanemab-irmb (Leqembi)

**Provider Order Form rev.** 10/15/2025

Novella
Infusion

PATIENT INFORMATION	Referral Status:	☐ New Ref	ferral l	□ Updated Or	der	☐ Order Renewal		
Patient Name:		DOB:		Patient Ph				
Patient Address:	Patient Email:							
Allergies:		□ NKDA	Weight (	lbs/kg):	He	ight (in/cm):		
Sex: □ M / □ F Date of Last Infusion:	Next Due Date:			red Location:		.8 (, 0).		
			110101	rea Location.	-			
DIAGNOSIS (Please provide ICD-10 code in space	-							
☐ G30.0 Alzheimer's disease w/ early onset ☐		ner's disease	e 🗆 (	G30.9 Alzheim	er's dis	sease unspecified		
☐ G30.1 Alzheimer's disease w/ late onset ☐	Other:		Desc	ription:				
REQUIRED INFORMATION FOR MEDICARE  □ 200.6: Encounter for examination for normal compace control in clinical research program  Medicare Trial Registry Number:	mg IV  ment  all  most for this potain and to Novella icribed  p	PRE-MEDICATION ORDERS  □ Tylenol □ 500mg / □ 650mg PO □ Loratadine 10mg PO □ Pepcid 20mg □ PO / □ IVP □ Benadryl □ 25mg / □ 50mg □ PO / □ IVP □ Solumedrol □ 40mg / □ 125mg IVP □ Other: □ LABORATORY ORDERS □ Other: ■ Hold infusion and notify provider for: ■ Hold if amyloid beta pathology has not been confirmed. ■ Abnormal vital signs ■ No brain MRI results in chart (need MRI within one year of starting treatment, and prior to 3rd, 5th, 7th, and 14th infusion). ■ Signs of Amyloid Related Imaging Abnormalities (ARIA) as reported on MRI results. ■ New or worsening headache or altered mental status. □ Record vital signs before infusion and prior to patient discharge □ Provide nursing care per Nursing Procedure, including Hypersensitivity Reaction Management Protocol and post-procedure observation □ To report suspected adverse reactions, contact FDA at 1-800-FDA-1088 or www.fda.gov/medwatch						
PROVIDER INFORMATION Preferred Contact Name:		Preferred Contact Email:						
Ordering Provider:		Provider NPI:						
Referring Practice Name:	Pho			Fax:				
Practice Address:	City		Sta	ite:	7	Zip Code:		
REQUIRED DOCUMENTATION CHECKLIST (	Additional docume	ntation real	uired for	nrocessina an	d insu	rance annroval)		
Required Documentation: Patient demos, copy of f								
treatment failures or contraindications. Documenta	•	-	-					
initial and throughout treatment, PET or CSF analys						.0 1/ 2		
Provider Name (print) Pr	Provider Signature			 Date				

Order valid for one year unless otherwise indicated. IV solutions/diluents may be substituted as allowed per manufacturer's instructions as necessitated by product availability.