

**PATIENT INFORMATION**

Fax completed form, insurance information, and clinical documentation to 855-889-2946

Name:		DOB:	Gender: <input type="checkbox"/> Female <input type="checkbox"/> Male	
Address:		City:	State:	ZIP:
Phone:	Email:	Height:	<input type="checkbox"/> inches <input type="checkbox"/> cm	Weight: <input type="checkbox"/> lbs. <input type="checkbox"/> kg
Allergies:				

**Diagnosis Code ICD-10 (required):**
**Diagnosis Description:**

 Patient Status:  New to Therapy  Continuing Therapy

Next Treatment Date:

**PHYSICIAN INFORMATION**

Prescriber Name:		Phone:	Fax:
Office Contact:		Email:	
Address:		City:	State: ZIP:
NPI #:	DEA#:	Tax ID:	

**INSURANCE INFORMATION (or attach copy of cards)**

Primary Insurance:	Policy #:	Group #:
Secondary Insurance:	Policy #:	Group #:

**PRESCRIPTION INFORMATION (or attach a copy of the prescription)**

Drug	Medication Orders	Refills
<b>Leqembi</b> (lecanemab)	<input type="checkbox"/> 10mg/kg IV every 2 weeks <input type="checkbox"/> 10mg/kg IV every 4 weeks (after 18 months of treatment, patient can transition to q 4 weeks*)  <ul style="list-style-type: none"> <li>Patients may transition to every 4 weeks <u>after 18 months</u> or remain on every 2 weeks</li> <li>MRIs should be performed at baseline &amp; prior to the 3<sup>rd</sup>, 5<sup>th</sup>, 7<sup>th</sup>, and 14<sup>th</sup> infusion</li> <li>HOLD infusion if MRI is not performed at indicated interval</li> </ul>	<input type="checkbox"/> x 1 year <input type="checkbox"/> _____
<b>Kisunla</b> (donanemab)	<input type="checkbox"/> Initial start: Infusion 1: 350mg IV at week 0 Infusion 2: 700mg IV at week 4 Infusion 3: 1,050mg IV at week 8 Infusion 4 and beyond: 1,400mg at week 12 and every 4 weeks thereafter  <input type="checkbox"/> Maintenance: 1400mg IV every 4 weeks <input type="checkbox"/> Other: _____  <ul style="list-style-type: none"> <li>Protocol pre-medication (if no contraindications): acetaminophen 500mg PO &amp; loratadine 10mg PO</li> <li>MRIs should be performed at baseline &amp; prior to the 2<sup>nd</sup>, 3<sup>rd</sup>, 4<sup>th</sup>, and 7<sup>th</sup> infusion</li> <li>HOLD infusion if MRI is not performed at indicated interval</li> </ul>	<input type="checkbox"/> x 1 year <input type="checkbox"/> _____

Additional orders: \_\_\_\_\_

Lab orders: \_\_\_\_\_ Frequency: \_\_\_\_\_

 Required labs to be drawn by  Paragon  Referring Provider

As required by your state, Prescriber to check "Dispense as written" or handwritten "Brand Medically Necessary" and sign to prevent generic substitution.

 Dispense as written

**PRESCRIBER SIGNATURE**

By signing this form and utilizing our services, you are authorizing Paragon Healthcare, Inc. and its employees to serve as your prior authorization and specialty pharmacy designated agent in dealing with medical and prescription insurance companies, and to select the preferred site of care for the patient.

Prescriber Signature: X

Date:

**PATIENT INFORMATION**

Name: \_\_\_\_\_

DOB: \_\_\_\_\_

**REQUIRED DOCUMENTATION FOR REFERRAL PROCESSING & INSURANCE APPROVAL**

- Include signed and completed order (MD/prescriber to complete page 1)
- Include patient demographic information and insurance information
- Include patient's current medication list
- Supporting clinical notes (H&P) to support primary diagnosis - Including tried/failed medications.
- Other medical necessity: \_\_\_\_\_

**REQUIRED PRE-SCREENING**

- Patient enrolled in the CMS National Patient Registry (Medicare & Medicare Advantage required)**  
Issue number: \_\_\_\_\_ Date of registry enrollment: \_\_\_\_\_
  - Provide copy of CMS national patient registry confirmation  
<https://qualitynet.cms.gov/alzheimers-ced-registry/submission>
- Confirmed presence of amyloid pathology**  
Attach results: Amyloid PET scan OR +CSF (cerebrospinal fluid)
- MRI of the brain (within 1 year) - attach results**
- Cognitive assessment scores (list all available, attach results):**
  - MMSE:** Score \_\_\_\_\_ Date of assessment \_\_\_\_\_
  - MoCA:** Score \_\_\_\_\_ Date of assessment \_\_\_\_\_
  - CDR:** Score \_\_\_\_\_ Memory box: Score \_\_\_\_\_ Date of assessment \_\_\_\_\_
  - Other: \_\_\_\_\_ Score \_\_\_\_\_ Date of assessment \_\_\_\_\_
- Functional assessment score: \_\_\_\_\_ (attach results)**  
Assessment Name:  FAQ  FAST  Other: \_\_\_\_\_ Assessment date: \_\_\_\_\_
- Include labs and/or test results for the following:**
  - Genotype testing for ApoE4
  - OR -
  - ApoE4 genetic testing has NOT been completed. Provider has counselled the patient on how testing for ApoE4 status informs the risk of developing ARIA and the patient has shared decision-making to initiate Leqembi
- Does the patient have objective impairment in episodic memory as evidenced by a memory test** (i.e., Free and Cued, Wechsler, etc.)? (BCBS required)  Yes  No
- Is the patient on therapeutic anticoagulation/antiplatelet therapy?**  Yes  No  
If yes, please note therapy and dose: \_\_\_\_\_