



PATIENT INFORMATION (Complete or Fax Existing Chart)	PRESCRIBER INFORMATION
Name: _____ DOB: _____	Prescriber Name: _____
Address: _____	State License: _____
City, State, Zip: _____	NPI #: _____ Tax ID: _____
Phone: _____ Alt. Phone: _____	Address: _____
Email: _____ SS#: _____	City, State, Zip: _____
Gender: <input type="checkbox"/> M <input type="checkbox"/> F Weight: _____ (lbs) Ht: _____	Phone: _____ Fax: _____
Allergies: _____	Office Contact: _____ Phone: _____
Is patient the main point of contact? <input type="checkbox"/> Yes <input type="checkbox"/> No	
If not, who? Name/Relationship: _____	
Phone Number: _____	

INSURANCE INFORMATION – AND – Send a copy of the patient's prescription/insurance cards (front & back)

Primary Insurance: _____	Secondary Insurance (If Applicable): _____
Plan #: _____	Plan #: _____
Group #: _____	Group #: _____
RX Card (PBM): _____	RX Card (PBM): _____
BIN: _____ PCN: _____	BIN: _____ PCN: _____
MEDICARE/MEDICARE ADVANTAGE PATIENTS ONLY:	
CMS Registry # (ALZH#): _____	CMS Registry Enrollment Date: _____

CLINICAL INFORMATION

Please Select Diagnosis:

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
 Other: _____

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:

Amyloid pathology confirmed via:
 Amyloid PET Scan CSF analysis Blood plasma Date: _____ Result: Amyloid Positive Amyloid Negative

Recent MRI obtained prior to initiating Leqembi® (including FLAIR and T2/GRE or SWI) to assess ARIA risk
 Prescriber has verified that this Patient does not have evidence of prior ARIA-H Date: _____

Completion of cognitive assessment type:
 MMSE MoCA CDR Other: _____ Date: _____

Completion of functional assessment type:
 FAQ FAST Other: _____ Date: _____

****Note: MRIs must be obtained prior to initial infusion to assess ARIA risk. During treatment, conduct an ARIA monitoring MRI before Infusions 3, 5, 7, and 14 and if symptoms consistent with ARIA occur.**

LEQEMBI® ORDERS

Prescription type: New start Restart Continued therapy Total Doses Received: _____ Date of Last Injection/Infusion: _____

Medication	Dose/Frequency	Refills
<input type="checkbox"/> Leqembi® (lecanemab-irmb) 500 mg/5 mL (100 mg/mL) <input type="checkbox"/> Leqembi® (lecanemab-irmb) 200 mg/2 mL (100 mg/mL)	<input type="checkbox"/> 10 mg/kg intravenous infusion over approximately one hour, once every two weeks. <input type="checkbox"/> Other: _____	_____
Pre-Medication	Dose/Strength	Directions
<input type="checkbox"/> Acetaminophen	<input type="checkbox"/> 500mg	<input type="checkbox"/> Take 1-2 tablets PO prior to infusion or post-infusion as directed

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<input type="checkbox"/> Diphenhydramine	<input type="checkbox"/> 25mg IV/PO <input type="checkbox"/> 50mg IV/PO	<input type="checkbox"/> Take 1 tablet PO prior to infusion or as directed OR Inject contents of 1 vial IV prior to infusion or as directed
<input type="checkbox"/> Methylprednisolone	<input type="checkbox"/> 40mg <input type="checkbox"/> 100mg <input type="checkbox"/> 125mg	<input type="checkbox"/> Inject contents of 1 vial IV prior to infusion or as directed <input type="checkbox"/> Other: Inject 100mg IV 30 minutes prior to infusion
<input type="checkbox"/> _____	_____	_____

INFUSION REACTION ORDERS

Mild reaction protocol:
 Diphenhydramine 25mg IV, one time, for pruritus.
If symptoms worsen, see orders for moderate to severe reactions.

Moderate reaction protocol:
 Acetaminophen 650mg PO, one time, for pyrexia or rigors
 Diphenhydramine 50mg IV, one time, for pruritus or urticaria
 Famotidine 20mg IV, one time, for, for pruritus or urticaria
 Methylprednisolone 125mg IV, one time, for respiratory or neurologic symptoms
If symptoms worsen, see interventions for severe reactions

Severe reaction protocol: (Call 911 if initiated):
 Titrate oxygen via continuous flow per nasal cannula or face mask to maintain spO2 of greater than ninety-five percent (>95%)
 Diphenhydramine 50mg IV, one time, for respiratory symptoms, edema, or anaphylaxis
 Methylprednisolone 125mg IV, one time, for respiratory symptoms, edema, or anaphylaxis
 Sodium Chloride 0.9% 500mL IV over 30-60 min, one time, for cardiovascular symptoms
 Epinephrine 0.3mg/0.3mL IM into mid-antrolateral aspect of thigh of anaphylaxis, may repeat x1 in 5-15 minutes if symptoms are not resolved or worsen

FLUSHING & LOCKING ORDERS

Flushing Protocol (>66lbs/33kg)

PIV and Midline: <input checked="" type="checkbox"/> 0.9% Sodium Chloride 2-5mL IV flush before and after each infusion	Implanted Port, PICC, Tunneled Catheter, and Non-tunneled Catheter: <input checked="" type="checkbox"/> 0.9% Sodium Chloride 5mL IV flush before infusion/lab draw and 10mL IV flush after infusion/lab draw
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Locking Protocol (>66lbs/33kg)

PIV and Midline: <input checked="" type="checkbox"/> Heparin Sodium 10 units/mL 1mL IV final flush post normal saline flush	PICC: <input checked="" type="checkbox"/> Heparin Sodium 10 units/mL 3mL IV final flush post normal saline flush	Implanted Port, Tunneled Catheter, and Non-tunneled Catheter: <input checked="" type="checkbox"/> Heparin Sodium 100 units/mL 3-5mL IV final flush post normal saline flush
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**** May substitute Dextrose 5% in Water, or alternative, for 0.9% Sodium Chloride, when indicated due to incompatibility with medications being infused**

SIGNATURE

We hereby authorize Talis Healthcare LLC to provide all supplies and additional services (nursing/patient training) required to provide and deliver the medicine as prescribed in this referral.

X _____ Date: _____
 Prescriber Signature

To ensure payment by insurance carrier, please include supporting clinical documentation for specified ICD 10 Code, demographic, and insurance information along with faxed order. Initial appointment will be verified upon insurance approval.

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