TALIS HEALTHCARE
AN INFUSION MANAGEMENT COMPANY

KISUNLA™

Please Fax Completed Form To: 888-898-9113

Please Send a Copy of The Patient's Insurance Cards (Front & Back)

PATIENT INFORMATION (Complete or Fax Existing Chart)

Name: _____ DOB: _____
Address: _____
City, State, Zip: _____
Phone: _____ Alt. Phone: _____
Email: _____ SS#: _____
Gender: ☐ M ☐ F Weight: _____(lbs) Ht: _____
Allergies: _____

PRESCRIBER INFORMATION

Prescriber Name: _____
State License: _____
NPI #: _____ Tax ID: _____
Address: _____
City, State, Zip: _____
Phone: _____ Fax: _____
Office Contact: _____ Phone: _____

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

Primary Insurance: _____
Plan #: _____
Group #: _____
RX Card (PBM): _____
BIN: _____ PCN: _____

Secondary Insurance (If Applicable): _____
Plan #: _____
Group #: _____
RX Card (PBM): _____
BIN: _____ PCN: _____

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 Kisunla™ (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: _____

☐ **Completion of CMS approved CED registry (only required for Patients with Medicare) ClinicalTrials.gov Registry Number: NCT _____**
CED Submission Date: _____ Submission Number (if applicable): _____

****Note: MRIs must be obtained prior to initial infusion to assess ARIA risk. During treatment, conduct an ARIA monitoring MRI before Infusions 2, 3, 4 and 7 and if symptoms consistent with ARIA occur.**

ORDERS

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

Dose/Frequency	Quantity	Refills
☐ Starting Dose: Infuse 700 mg intravenously over approximately 30 minutes once every 4 weeks for Infusions 1, 2, and 3	2 Vials	2
☐ Maintenance Dose: Infuse 1400 mg intravenously over approximately 30 minutes once every 4 weeks thereafter	4 Vials	_____

Pre-Medication	Dose/Strength	Directions
☐ Acetaminophen	☐ 500mg	☐ Take 1-2 tablets PO prior to infusion or post-infusion as directed
☐ Diphenhydramine	☐ 25mg IV/PO ☐ 50mg IV/PO	☐ Take 1 tablet PO prior to infusion or as directed OR ☐ Inject contents of 1 vial IV prior to infusion or as directed
☐ _____	_____	_____

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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

☒ 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 mis-anterolateral 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:	**Implanted Port, PICC, Tunneled Catheter, and Non-tunneled Catheter:**
☒ 0.9% Sodium Chloride 2-5mL IV flush before and after each infusion	☒ 0.9% Sodium Chloride 5mL IV flush before infusion/lab draw and 10mL IV flush after infusion/lab draw

Locking Protocol (>66lbs/33kg)

PIV and Midline:	**PICC:**	**Implanted Port, Tunneled Catheter, and Non-tunneled Catheter:**
☒ Heparin Sodium 10 units/mL 1mL IV final flush post normal saline flush	☒ Heparin Sodium 10 units/mL 3mL IV final flush post normal saline flush	☒ Heparin Sodium 100 units/mL 3-5mL IV final flush post normal saline flush

**** May substitute Dextrose 5% in Water, or alternative, for 0.9& Sodium Chloride, when indicated due to incompatibility with medications bring 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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