



## Please Fax Completed Form To: 888-898-9113

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

PATIENT INFORMATION (Com	plete or Fax Existing Cha	rt) PRESCRIBER INFORMATIO	N				
Name: DOB:			Prescriber Name:				
Address:			State License:				
City, State, Zip:		NB1 #	NPI #: Tax ID:				
Phone: Al			Address:				
Email:		6': 6: - 7'	City, State, Zip:				
Gender:     M   F   Weight:(lbs)   Ht:		BI SI	Phone: Fax:				
Allergies:		Office Contact:	Office Contact: Phone:				
INSURANCE INFORMATION – AND – Send a copy of the patient's prescription/insurance cards (front & back)							
Primary Insurance:		Secondary Insurance (If Applicable	Secondary Insurance (If Applicable):				
Plan #:							
Group #:							
RX Card (PBM):			RX Card (PBM):				
BIN: PCN:			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			:				
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	• .	et to commin diagnosis and that Fatient na	s evidence of AD neuro	pathology and has			
☐ Amyloid pathology confirmed via:							
$\square$ Amyloid PET Scan $\square$ CSF analysis	☐ Blood plasma	Date: Result	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 Pa		or ARIA-H Date:					
☐ Completion of cognitive assessment type:         ☐ MMSE         ☐ MoCA         ☐ 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.	musion to assess ARIA risk. During tre	athent, conduct an ANA monitoring with before initia	sions 2, 3, 4 and 7 and 11 sympt	onis consistent with			
ORDERS							
Prescription type: ☐ New start ☐	Restart   Continued therapy	Total Doses Received: Da	te of Last Injection/Infu	sion:			
Dose/Frequency			Quantity	Refills			
$\square$ Starting Dose: Infuse 700 mg intravend	ously over approximately 30 minute	es once every 4 weeks for Infusions 1, 2, and 3	☐ 1 Vial ☐ 2 Vials	□ 2			
$\square$ Alternate Starting Dose: Infuse 350mg			□ 3 Vials □ 4 Vials	□ 0			
☐ Maintenance Dose: Infuse 1400 mg intravenously over approximately 30 minutes once every 4 weeks thereafter							
Pre-Medication	Dose/Strength	Directions					
☐ Acetaminophen	□ 500mg	☐ Take 1-2 tablets PO prior to infusion or post-infusion as directed					
☐ Diphenhydramine	☐ 25mg IV/PO	☐ Take 1 tablet PO prior to infusion or as directed OR					
	☐ 50mg IV/PO	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	d after each infusion						
		IV flush after infusion/lab draw					
Locking Protocol (>66lbs/33kg)							
PIV and Midline:	PICC:		Implanted Port, Tunneled Catheter, and Non-				
⋈ Heparin Sodium 10 units/mL 1mL IV final			tunneled Catheter:				
flush post normal saline flush	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.							

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.

**Prescriber Signature** 

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