



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: _____

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: _____

CLINICAL INFORMATION
Primary ICD-10 Code (Please Specify Diagnosis): _____
Secondary ICD-10 Code (Please Specify Diagnosis): _____
MG-ADL* score (if known): _____ Has the patient received Meningitis vaccination? <input type="checkbox"/> Yes <input type="checkbox"/> No Date of vaccination: _____
<input type="checkbox"/> Please check this box if the patient has declined vaccination Reason: _____
Adverse reactions with previous Ultomiris treatments? <input type="checkbox"/> No <input type="checkbox"/> Yes <i>If yes, Reason/Date:</i> _____
<input type="checkbox"/> Please check to confirm: The patient is enrolled in the ULTOMIRIS REMS program; The patient has been counseled about the risks of meningococcal infection; The patient has received information and a Patient Safety Card about the symptoms and risks of meningococcal infection.

ULTOMIRIS® ORDERS

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

Medication	Strength	Dose/Frequency	Refills
Intravenous Ultomiris® (ravulizumab)	<input type="checkbox"/> 1,100mg/11mL vial <input type="checkbox"/> 300mg/3mL vial <input type="checkbox"/> 300mg/30mL vial <input type="checkbox"/> Other: _____	<input type="checkbox"/> Loading dose: Begin _____ mg IV on day 1 Then 2 weeks later <input type="checkbox"/> Maintenance dose: Begin _____ mg IV every _____ weeks <input type="checkbox"/> Other: _____	_____
Subcutaneous Ultomiris® (ravulizumab)	<input type="checkbox"/> 245mg/3.5 mL prefilled cartridge with on body injector	<input type="checkbox"/> 490 mg once weekly in adult patients greater than or equal to 40 kg body weight with PNH or aHUS. <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
<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 <input type="checkbox"/> Inject contents of 1 vial IV prior to infusion or as directed
<input type="checkbox"/> Methylprednisolone	<input type="checkbox"/> 40mg <input type="checkbox"/> 125mg	<input type="checkbox"/> Inject contents of 1 vial IV prior to infusion or as directed
<input type="checkbox"/> _____	_____	_____

INFUSION REACTION ORDERS

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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:

0.9% Sodium Chloride 2-5mL IV flush before and after each infusion

Implanted Port, PICC, Tunneled Catheter, and Non-tunneled Catheter:

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:

Heparin Sodium 10 units/mL 1mL IV final flush post normal saline flush

PICC:

Heparin Sodium 10 units/mL 3mL IV final flush post normal saline flush

Implanted Port, Tunneled Catheter, and Non-tunneled Catheter:

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 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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