



PATIENT INFORMATION (Complete or Fax Existing Chart)	PRESCRIBER INFORMATION
Name: _____ DOB: _____	Prescriber Name: _____
Address: _____	State License: _____
City, State, Zip: _____	NPI #: _____ DEA: _____
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 #: _____	Secondary Insurance: _____
Group #: _____	Plan #: _____
RX Card (PBM): _____	Group #: _____
BIN: _____ PCN: _____	BIN: _____ PCN: _____

CLINICAL INFORMATION			
Primary ICD-10 Code: _____	Diagnosis Description: _____		
Secondary ICD-10 Code: _____	Diagnosis Description: _____		
Hepatitis B Vaccination: <input type="checkbox"/> Yes <input type="checkbox"/> No	Patient on Methotrexate: <input type="checkbox"/> Yes <input type="checkbox"/> No	Line Access: <input type="checkbox"/> PIV <input type="checkbox"/> Port <input type="checkbox"/> PICC <input type="checkbox"/> Midline	

### RITUXIMAB ORDERS

Prescription type:  New start  Restart  Continued therapy Total Doses Received: \_\_\_\_\_ Date of Last Dose: \_\_\_\_\_

Medication	Dose/Frequency	Refills
<input type="checkbox"/> Rituxan® (Rituximab) 100mg/10ml Vial <input type="checkbox"/> Rituxan® (Rituximab) 500mg/50ml Vial	<input type="checkbox"/> 1000mg IV x 2 Doses separated by 14 days, repeat every 24 weeks <input type="checkbox"/> Other: _____ Frequency: _____	_____
<input type="checkbox"/> Riabni™ (rituximab-arrx) 100mg Vial <input type="checkbox"/> Riabni™ (rituximab-arrx) 500mg Vial	<input type="checkbox"/> 375mg/m2 once weekly for 4 weeks <input type="checkbox"/> 500mg IV infusion separated by 2 weeks, followed by a 500mg IV infusion every 6 months <input type="checkbox"/> Other: _____ Frequency: _____	_____
<input type="checkbox"/> Ruxience® (Rituximab-pvvr) 100mg Vial <input type="checkbox"/> Ruxience® (Rituximab-pvvr) 500mg Vial	<input type="checkbox"/> 1000mg IV x 2 Doses separated by 14 days, repeat every 24 weeks <input type="checkbox"/> Other: _____ Frequency: _____	_____

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"/> 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

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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

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

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:**

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 \_\_\_\_\_

Prescriber Signature

Date: \_\_\_\_\_

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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