



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: ☐ M ☐ 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			
ICD-10 Code (Required): _____		ICD-10 Description: _____	
Baseline Liver Enzymes, Including Bilirubin (Results): _____			
Date of Negative TB Test: _____ ☐ TB Test is Pending, Will Fax The Results			
OMVOH™ ORDERS			
Prescription type: ☐ New start ☐ Restart ☐ Continued therapy Total Doses Received: _____ Date of Last Injection/Infusion: _____			
Crohn's Disease Diagnosis			
Induction Dosing: ☐ 3 x 300 mg/15 mL single dose vial	☐ 900 mg by intravenous infusion at Weeks 0,4, and 8	Quantity: _____	
Maintenance Dosing: ☐ 1 x 100 mg/mL Prefilled Pen AND 1 x 200 mg/mL Prefilled Pen given as two consecutive subcutaneous injections ☐ 1 x 100 mg/mL Prefilled Syringe AND 1 x 200 mg/mL Prefilled Syringe given as two consecutive subcutaneous injections	☐ 300 mg given by subcutaneous injection, given as two consecutive injections of 100 mg and 200 mg in any order, at week 12 and every 4 weeks thereafter	Quantity: _____ Refills: _____	
Ulcerative Colitis Diagnosis			
Induction Dosing: ☐ 300 mg / 15 mL single dose vial	☐ 300 mg by intravenous infusion at Weeks 0, 4, and 8	Quantity: _____	
Maintenance Dosing: ☐ 2 x Prefilled Pen 100 mg/mL given as two consecutive subcutaneous injections ☐ 2 x Prefilled Syringe 100 mg/mL given as two consecutive subcutaneous injections	☐ 200 mg by subcutaneous injection, given as two consecutive injections of 100 mg each, at week 12 and every 4 weeks thereafter	Quantity: _____ Refills: _____	
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	
☐ Methylprednisolone	☐ 40mg ☐ 100mg	☐ Inject contents of 1 vial IV prior to infusion or as directed	

CONFIDENTIALITY STATEMENT: This facsimile and documents accompanying this transmission contain confidential health information that is legally privileged. This information is intended only for the use of the individual or entity named above. The authorized recipient of this information is prohibited from disclosing this information to any other party unless required to do so by law or regulation. If you are not the intended recipient, you are hereby notified that any disclosure, copying, distribution, or action taken in reliance on the contents of these documents is strictly prohibited. If you have received this information in error, please notify the sender at the address and telephone number set forth herein and arrange for return or destruction of the material. In no event should such material be read by anyone other than the named addressee, except by express authority of the sender to the named addressee.



☐	☐ 125mg	☐ Other: Inject 100mg IV 30 minutes prior to infusion
☐		

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 mid-antergluteal 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.

CONFIDENTIALITY STATEMENT: This facsimile and documents accompanying this transmission contain confidential health information that is legally privileged. This information is intended only for the use of the individual or entity named above. The authorized recipient of this information is prohibited from disclosing this information to any other party unless required to do so by law or regulation. If you are not the intended recipient, you are hereby notified that any disclosure, copying, distribution, or action taken in reliance on the contents of these documents is strictly prohibited. If you have received this information in error, please notify the sender at the address and telephone number set forth herein and arrange for return or destruction of the material. In no event should such material be read by anyone other than the named addressee, except by express authority of the sender to the named addressee.