

**AUTHORIZATION IS GRANTED TO DISPENSE AND ADMINISTER AN ALTERNATE DRUG PRODUCT ACCEPTABLE TO THE MEDICAL STAFF'S PHARMACY COMMITTEE UNLESS THE DRUG PRODUCT IS SPECIFICALLY CIRCLED.**

<input type="checkbox"/> MAINTENANCE every ____ weeks for one year.	Estimated treatment initiation date:	Allergies/Reactions:
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**Diagnosis (Complete One Box Below)**

<input type="checkbox"/> Treatment of <b>moderately to severely active Crohn's disease</b> (ICD 10 Code _____) (to reduce signs/symptoms and induce and maintain clinical remission) or to reduce the number of draining enterocutaneous and rectovaginal fistulas and maintain fistula closure <b>PLUS</b> (must complete below) <ul style="list-style-type: none"> <li><input type="checkbox"/> A. Previous inadequate response to conventional therapy                         <ul style="list-style-type: none"> <li>Drug Failed:</li> <li>Treatment Dates: From (mm/yy)_____ to (mm/yy)_____.</li> <li>Reason for Failure:</li> </ul> </li> </ul>	<input type="checkbox"/> Treatment of <b>moderately to severely active ulcerative colitis</b> (ICD 10 Code _____) (to reduce signs/symptoms and induce and maintain clinical remission) or to induce/maintain mucosal healing and eliminate corticosteroid use in adults <b>PLUS</b> (must complete below) <ul style="list-style-type: none"> <li><input type="checkbox"/> A. Previous inadequate response to conventional therapy                         <ul style="list-style-type: none"> <li>Drug Failed:</li> <li>Treatment Dates: From (mm/yy)_____ to (mm/yy)_____.</li> <li>Reason for Failure:</li> </ul> </li> </ul>
<input type="checkbox"/> Treatment of <b>moderately to severely active rheumatoid arthritis</b> (ICD 10 Code _____) (to reduce signs/symptoms of active arthritis and inhibit progression of structural damage and improve physical function) <b>PLUS</b> (must complete below) <ul style="list-style-type: none"> <li><input type="checkbox"/> A. On concurrent methotrexate therapy</li> </ul>	<input type="checkbox"/> Treatment of <b>active ankylosing spondylitis</b> (to reduce signs/symptoms) (ICD 10 Code _____)
<input type="checkbox"/> Treatment of <b>psoriatic arthritis</b> (to reduce signs/symptoms of active arthritis and inhibit progression of structural damage and improve physical function) (ICD 10 Code _____)	<input type="checkbox"/> Treatment of <b>chronic, severe (extensive and/or disabling) plaque psoriasis</b> as an alternative to other systemic therapy (ICD 10 Code _____)

**Lab orders:** CBC with differential and CMP per provider (provide separate order)

<b>Tuberculin Screening</b> complete on (Date): _____ Results: _____	<b>Weight:</b> _____ kg  <i>(Weigh patient prior to each dose, notify provider for fluctuations &gt; 10%)</i>	<b>HOLD treatment &amp; notify provider if:</b> <ul style="list-style-type: none"> <li>Fever or evidence of infection</li> <li>LFTs &gt; 5 x ULN</li> </ul>	<b>Emetic Risk: Minimal Monitor:</b> <ul style="list-style-type: none"> <li>For infection</li> <li>For infusion related reactions</li> <li>Liver function</li> </ul>
<b>REMS:</b> Medication Guide provided with each treatment			

**SAFETY LINE**

**0.9% NaCl** (gravity flow). Use as free-flow for IV push doses (if applicable) and post chemotherapy line flush.

TREATMENT	DOSAGE	ADMINISTRATION INSTRUCTIONS	FREQUENCY
<b>Choose one:</b> <b>Preferred:</b> <input type="checkbox"/> <b>InFLIXimab-abda</b> (Renflexis)  <b>Alternative:</b> <input type="checkbox"/> Infliximab-dyyb (Inflectra) <input type="checkbox"/> Infliximab (Remicade-generic)  <b>Reason for selecting alternative:</b> <input type="checkbox"/> Insurance requirements, including enrollment in patient assistance program <input type="checkbox"/> Intolerance  <input type="checkbox"/> <b>Unless checked</b> , pharmacy may select alternative biosimilar agent IF preferred by patient's insurance	_____ mg/kg _____ mg  <i>Dose rounded to the nearest 100mg</i>	In 0.9% NaCl 250 mL over at least 2 hours  Use in-line low protein binding filter (≤1.2 micron)  <i>Infusion should begin within 3 hours of preparation</i>	x 1 dose per maintenance schedule above

**IF PATIENT HAS A HYPERSENSITIVITY REACTION, BEGIN HYPERSENSITIVITY PROTOCOL**

Patient Name:

Date of Birth:

**The provider's full signature(s) is to follow the order**

_____ Provider Signature	_____ Date	_____ Time
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\_\_\_\_\_  
 Provider Printed Name

**InFLIXimab MAINTENANCE Dose  
 (Must sign both pages)**

**SYMPTOM MANAGEMENT**

DRUG	DOSE	ADMINISTRATION INSTRUCTIONS	FREQUENCY
<input type="checkbox"/> Acetaminophen	650 mg	Oral	x 1 dose if needed for headache during infusion
<input type="checkbox"/> DiphenhydrAMINE	25mg	IV Push over 1 minute	x 1 dose if needed for itching/rash during infusion

**ADDITIONAL ORDERS**

For symptomatic hypotension, slow infusion rate. If no improvement, stop infusion, bolus with 125-250 mL 0.9% NaCl and notify provider.

Discontinue IV upon completion of treatment, flush order per protocol.

**NURSING INSTRUCTIONS**
**For InFLIXimab:**

1. Monitor vital signs at baseline and every 30 minutes for the first hour or until stable, then every 60 minutes until completion of inFLIXimab infusion.
2. Pulse oximetry PRN dyspnea.

Reference: Arthritis Care Res (Hoboken)2012,64(5):625-39; Arthritis Rheum 2005,52(2):548-53

Am J Gastroenterol 2010, 105(3):501-23; Am J Gastroenterol 2009, 104(2): 465-83; Gastroenterology 2013,145(6):1459-1463

Patient Name:

Date of Birth:

**The provider's full signature(s) is to follow the order**

\_\_\_\_\_  
**Provider Signature**

\_\_\_\_\_  
**Date**

\_\_\_\_\_  
**Time**

\_\_\_\_\_  
**Provider Printed Name**

**InFLIXimab MAINTENANCE Dose  
 (Must sign both pages)**