

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"/> LOADING dose at 0, 2, and 6 weeks	Estimated treatment initiation date: _____	Allergies/Reactions: _____
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Diagnosis (Complete One Box Below)

<input type="checkbox"/> Treatment of moderately to severely active Crohn's disease (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 PLUS (must complete below) <input type="checkbox"/> A. Previous inadequate response to conventional therapy Drug Failed: Treatment Dates: From (mm/yy) _____ to (mm/yy) _____. Reason for Failure: _____	<input type="checkbox"/> Treatment of moderately to severely active ulcerative colitis (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 PLUS (must complete below) <input type="checkbox"/> A. Previous inadequate response to conventional therapy Drug Failed: Treatment Dates: From (mm/yy) _____ to (mm/yy) _____. Reason for Failure: _____
<input type="checkbox"/> Treatment of moderately to severely active rheumatoid arthritis (ICD 10 Code _____) (to reduce signs/symptoms of active arthritis and inhibit progression of structural damage and improve physical function) PLUS (must complete below) <input type="checkbox"/> A. On concurrent methotrexate therapy	<input type="checkbox"/> Treatment of active ankylosing spondylitis (to reduce signs/symptoms) (ICD 10 Code _____)
<input type="checkbox"/> Treatment of psoriatic arthritis (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 chronic, severe (extensive and/or disabling) plaque psoriasis as an alternative to other systemic therapy (ICD 10 Code _____)

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

Tuberculin Screening complete on (Date): _____ Results: _____	Weight: _____ kg <i>(Weigh patient prior to each dose, notify provider for fluctuations > 10%)</i>	HOLD treatment & notify provider if: <ul style="list-style-type: none"> Fever or evidence of infection LFTs > 5 x ULN 	Emetic Risk: Minimal Monitor: <ul style="list-style-type: none"> For infection For infusion related reactions Liver function
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REMS: 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
Choose one: Preferred: <input type="checkbox"/> InFLIXimab-abda (Renflexis) Alternative: <input type="checkbox"/> Infliximab-dyyb (Inflixtra) <input type="checkbox"/> Infliximab (Remicade-generic) Reason for selecting alternative: <input type="checkbox"/> Insurance requirements, including enrollment in patient assistance program <input type="checkbox"/> Intolerance <input type="checkbox"/> Unless checked , 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 loading dose schedule above

IF PATIENT HAS A HYPERSENSITIVITY REACTION, BEGIN HYPERSENSITIVITY PROTOCOL, Policy 061.060

Patient Name: _____ Date of Birth: _____	<div style="background-color: black; color: white; padding: 5px; text-align: center;"> The provider's full signature(s) is to follow the order </div> <div style="border: 1px solid black; height: 40px; margin-top: 5px;"></div> <table border="1" style="width: 100%; border-collapse: collapse; margin-top: 5px;"> <tr> <td style="width: 70%; padding: 5px;">Provider Signature</td> <td style="width: 15%; padding: 5px;">Date</td> <td style="width: 15%; padding: 5px;">Time</td> </tr> </table> <div style="border: 1px solid black; padding: 5px; margin-top: 5px;"> Provider Printed Name </div>	Provider Signature	Date	Time
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InFLIXimab LOADING Dose

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.

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:

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_____ Provider Signature	_____ Date	_____ Time
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InFLIXimab LOADING Dose