



# INFLIXIMAB

(Remicade or Biosimilars)

## Order Form

Outpatient Infusion

<b>Patient Name</b>	
<b>DOB</b>	
<b>Address</b>	
<b>Phone</b>	

**Order Status**       New Order                       Renewal                       Dose or Frequency Change

<b>Allergies:</b>		<b>Weight:</b>	
<b>Diagnosis</b>	<input type="checkbox"/> Rheumatoid arthritis <input type="checkbox"/> Ankylosing spondylitis <input type="checkbox"/> Psoriatic arthritis <input type="checkbox"/> Plaque Psoriasis <input type="checkbox"/> Ulcerative Colitis <input type="checkbox"/> Pustular Psoriasis <input type="checkbox"/> Crohn's Disease <input type="checkbox"/> Other:	<b>Height:</b>	
		<b>Diagnosis Code:</b>	

<b>Required Information</b>	Negative Quantiferon TB, T-spot or chest x-ray (no active disease)	Date:
	Diagnostic Hepatitis B panel	Date:
	CBC and CMP	Date:

**Please include labs attached and refer to page 2 for Required Documentation**

<b>Labs</b>	<input type="checkbox"/> Hepatic Function panel every 3 months	<input type="checkbox"/> Other:
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**Pre-Medications**

Diphenhydramine       PO                       25 mg       Acetaminophen                       325 mg  
 IV                       50 mg                       650 mg  
 Other:

**Medication Order**

<input type="checkbox"/> <b>Remicade (infliximab)</b> <input type="checkbox"/> <b>Inflectra (infliximab-dyyb)</b> <input type="checkbox"/> <b>Renflexis (infliximab-abda)</b> <input type="checkbox"/> <b>Avsola (infliximab-axxq)</b>	<input type="checkbox"/> 3 mg/kg IV <input type="checkbox"/> 5 mg/kg IV <input type="checkbox"/> 7.5 mg/kg IV <input type="checkbox"/> 10 mg/kg IV <input type="checkbox"/> Other: <b>Dose will be rounded up to nearest 100 mg</b>	<input type="checkbox"/> Induction: Week 0, 2, 6, then every _____ weeks thereafter <input type="checkbox"/> Maintenance: Every _____ weeks <input type="checkbox"/> Other:
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Remicade has several biosimilars. Certain payors may require use of a specific biosimilar. Please select allowed alternative if Remicade is not covered by payor. If more than one, note preference.

Alternative(s):

**Infusion Reaction Medications**

Hypersensitivity Reaction Protocol will be utilized unless otherwise specified

Provider (print name):	Date:
Provider Signature:	NPI:
Office Phone:	Office Fax:

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### Required Documentation

- H&P or progress note supporting diagnosis.
- Documented negative TB within 6 months of initiation.
- Chron's: Inadequate response to systemic corticosteroids
- UC: Inadequate response to systemic corticosteroids
- RA: documentation of methotrexate combination or intolerance or contraindication. Positive RF or anti-CCP
- AS: Inadequate response to two or more NSAIDs or intolerance or contraindication to two or more NSADs
- PsA: Inadequate response to another conventional synthetic drug or intolerance or enthesitis or predominately axial disease or severe disease
- PsO: Inadequate response or intolerance to UVB, PUVA or pharmacologic treatment with methotrexate, cyclosporin, acitretin or clinical reason to avoid pharmacologic treatment with methotrexate, cyclosporine and acitretin.
- Medication history
- Recent labs (as above) and/or diagnostic test results