

Renflexis Order Form (infliximab-abda)

FAX TO: 972.499.9210

PATIENT INFORMATION				
Patient Name:	DOB:	Phone:	Sex: M / F Ht: \	Nt: lbs / kg
Primary Language:	Allergies:			
Patient Preferred Location:		_		
<icd 10="" code="" required=""> DIAGNOSIS & CLINICAL INFORMATION</icd>				
ICD 10 Code (PROVIDE COMPLETE CODE) DERMATOLOGY L40.5 Psoriatic Arthritis/Arthropathy L40. Psoriasis GASTROENTEROLOGY K50.0 Crohn's Disease, Small Intestine K50.1 Crohn's Disease, Large Intestine	☐ K50.9 Crohn's Dis ☐ K51.8 Other Ulce ☐ K51.5 Left Sided ☐ K51.0 Universal U ☐ K51.9 Ulcerative ☐ K60.3 Anal Fistula	rative Colitis, Chronic Ulcerative Colitis, Chronic Ulcerative Pancolitis, Chronic Colitis, Unspecified	RHEUMATOLOGY M05 Rheumatoid Arthritis, w/ R M06 Rheumatoid Arthritis, w/o L40.5 Psoriatic Arthritis/Arthropa M45 Ankylosing Spondylitis D86.0 Sarcoidosis of the Lung OTHER:	Rheumatoid Factor athy
<u>REQUIRED:</u> Demographics & Most Recent: H&P, clinical notes, & medication list. Supporting clinical notes to include any past tried and/or failed therapies, intolerance, outcomes, or contraindications to conventional therapy. <u>LAB RESULTS:</u> Include Negative Hepatitis B within 3 years and Negative TB within 12 months.				
PRESCRIPTION				
Pre-Medications ☐ Acetaminophen: 650 mg PO ☐ Cetirizine: 10 mg PO ☐ Diphenhydramine: 25 mg PO ☐ Diphenhydramine: 25 mg IVP ☐ Famotidine: 20 mg PO ☐ Methylprednisolone: 125 mg SIVP Other:		ative TB, annually on will draw maintenand	e labs unless otherwise directed by Re	ferring Provider.
Renflexis (infliximab-abda)				
Infuse in 250 mL of 0.9% NS over at least 2 hours via pump with 0.2-micron filter. Doses > 1000 mg need total volume of 500 mL.				
Loading Dose: (SELECT ONE) □ IV: Infuse 3 mg/kg at weeks 0, 2, and 6 □ IV: Infuse 5 mg/kg at weeks 0, 2, and 6 □ IV: Infuse mg or mg/kg at weeks 0, 2, and 6 Maintenance: (SELECT ONE) □ IV: Infuse 3 mg/kg every 8 weeks for one year □ IV: Infuse 5 mg/kg every 8 weeks for one year □ IV: Infuse mg or mg/kg every weeks for one year				
In the event of an adverse reaction occurring at a Medix Infusion suite, utilize the Medix Infusion adverse reaction protocol.				
Post Treatment Observations: The patient is observed for 30 minutes following the first infusion.				
Comments:				
PRESCRIBER INFORMATION				
Prescriber Name:				
Date:NPI#:				
Supervising Physician:				(If Applicable)
Address:		_ City:	State: 2	<u>Z</u> ip:
Contact Name:	Phone:	Fax:	Email:	