



Your Lifetime Pharmacy Solution

GASTROENTEROLOGY ENROLLMENT FORM

Phone: (813) 871-5161 ext. 34993

Fax: (813) 877-2479

PATIENT INFORMATION (OR ATTACH PATIENT DEMOGRAPHIC SHEET)					
Patient Name:		<input type="checkbox"/> Male <input type="checkbox"/> Female	Allergies:		<input type="checkbox"/> NKDA
Date of Birth:	SSN:	Weight:	<input type="checkbox"/> kg <input type="checkbox"/> lb	Date:	
Address:		City:	State:	Zip:	
Phone # (Home):		Work #:	Email (Optional):		
INSURANCE INFORMATION (PLEASE PROVIDE COPIES OF MEDICAL AND PRESCRIPTION CARDS, IF AVAILABLE)					
Primary Insurance:			RX Bin:	RX PCN:	
RX Group:		RX ID:	RX Phone:		
Policy Holder's Name:		Policy Holder's DOB:	Policy Holder's SSN:		
DIAGNOSIS/MEDICAL INFORMATION (COMPLETE CLINICAL INFO BELOW OR ATTACH PATIENTS LABS)					
<input type="checkbox"/> K50.90 Crohn's Disease NOS		<input type="checkbox"/> K50.91 Crohn's Disease with complications		<input type="checkbox"/> Other:	
<input type="checkbox"/> K50.00 Crohn's Small Intestine without complications		<input type="checkbox"/> K51.90 Ulcerative Colitis without complications			
<input type="checkbox"/> K50.10 Crohn's Large Intestine without complications		<input type="checkbox"/> K51.91 Ulcerative Colitis with complications			
Severity type:		Has the patient had a NEGATIVE tuberculin skin test, or if positive, has treatment for latent TB been initiated prior to anti-TNF therapy?		Does the patient have NYHA Class III/IV CHF?	
<input type="checkbox"/> Mild		<input type="checkbox"/> YES <input type="checkbox"/> NO Test date: _____		<input type="checkbox"/> YES <input type="checkbox"/> NO	
<input type="checkbox"/> Moderate to severe				Does the patient have a clinically important active infection?	
<input type="checkbox"/> Fistulizing				<input type="checkbox"/> YES <input type="checkbox"/> NO If yes: _____	
For moderate to severe Crohn's disease: Has patient failed/contraindicated to optimal dosing/adequate duration of at least one of the following therapies: <input type="checkbox"/> YES <input type="checkbox"/> NO <i>Corticosteroids, Mesalamine or Sulfasalazine, Immunomodulator (methotrexate, 6-mercaptopurine, azathioprine)</i>					
Is the patient currently on any therapy? <input type="checkbox"/> YES <input type="checkbox"/> NO					

PRESCRIPTION INFORMATION				
MEDICATION	DOSE	DIRECTIONS	QTY	REFILLS
<input type="checkbox"/> Cimzia® (certolizumab pegol)	<input type="checkbox"/> 200mg Pre-filled Syringes Starter Kit <input type="checkbox"/> 200mg Pre-filled Syringes	<input type="checkbox"/> Initial Dose: 400mg subcutaneously at weeks 0,2 and 4 <input type="checkbox"/> Maintenance Dose: 400mg subcutaneously every four weeks		
<input type="checkbox"/> Entyvio® (vedolizumab)	<input type="checkbox"/> 300mg Powdered Vial	<input type="checkbox"/> Initial Dose: 300mg IV infused over 30 minutes at week 0, 2 and 6 <input type="checkbox"/> Maintenance Dose: 300mg IV infused over 30 minutes every 8 weeks after initial dosing		
<input type="checkbox"/> Humira CF® (adalimumab)	<input type="checkbox"/> 80mg / 0.8mL Pens	<input type="checkbox"/> Initial Dose: Inject 2 pens (160mg) subcutaneously on day 1 and inject 1 pen (80mg) on day 15		
	<input type="checkbox"/> 40mg / 0.4mL Pens	<input type="checkbox"/> Maintenance Dose: 40mg subcutaneously every other week <input type="checkbox"/> Maintenance Dose: 40mg subcutaneously every _____ weeks		
<input type="checkbox"/> Humira® (adalimumab)	<input type="checkbox"/> Crohn's / UC Starter Kit	<input type="checkbox"/> Initial Dose: Inject 2 pens (80mg) subcutaneously on day 1, then inject 2 pens (80mg) on day 2, then inject 2 pens (80mg) at week 2 <input type="checkbox"/> Initial Dose: Inject 4 pens (160mg) subcutaneously on day 1 then inject 2 pens (80mg) at week 2		
	<input type="checkbox"/> 40mg / 0.8mL Pens	<input type="checkbox"/> Maintenance Dose: 40mg subcutaneously every other week <input type="checkbox"/> Maintenance Dose: 40mg subcutaneously every _____ weeks		
<input type="checkbox"/> Remicade® Auto injector (infliximab)	<input type="checkbox"/> 100mg Vial	<input type="checkbox"/> Initial Dose: 5mg/kg at week 0, 2 and 6. Patients weight: _____ x 5mg = _____ mg <input type="checkbox"/> Maintenance Dose: 5mg/kg every 8 weeks. Patients weight: _____ x 5mg = _____ mg		

DELIVERY INSTRUCTIONS		
<input type="checkbox"/> Physician's Office	<input type="checkbox"/> Patient's Home	<input type="checkbox"/> 1st dose MD office, Refills to patient's home
PHYSICIANS CONTACT INFORMATION & AUTHORIZATION		
Physician's Name:	Office Contact:	Institution:
Phone #:	Fax #:	Specialty:
Address:	City/State/Zip:	
Tax ID:	DEA #:	NPI #:
Physician's Signature:		Date:
<small>*Prescriber Authorization: I authorize this pharmacy and its representatives to act as my authorized agent to secure coverage and initiate the insurance prior authorization process for my patient(s), and to sign any necessary forms on my behalf as my authorized agent, including the receipt of any required prior authorization forms and the receipt and submission of patient lab values and other patient data. In the event that this pharmacy determines that it is unable to fulfill this prescription, I further authorize this pharmacy to forward this information and any related materials related to coverage of the product to another pharmacy of the patient's choice or in the patient's insurer's provider network. The information contained in this transmission may contain privileged and confidential information, including patient information protected by federal and state privacy laws. It is intended only for the use of the person(s) named above. If you are not the intended recipient, you are hereby notified that any review, dissemination, distribution, or duplication of this communication is strictly prohibited. If you are not the intended recipient, please contact the sender and destroy all copies of the original document. Created: 08/28/18 Revised: 05/01/19, 08/22/19, 02/04/20</small>		



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<input type="checkbox"/> Moderate to severe				Does the patient have a clinically important active infection?	
<input type="checkbox"/> Fistulizing				<input type="checkbox"/> YES <input type="checkbox"/> NO If yes: _____	
For moderate to severe Crohn's disease: Has patient failed/contraindicated to optimal dosing/adequate duration of at least one of the following therapies: <input type="checkbox"/> YES <input type="checkbox"/> NO <i>Corticosteroids, Mesalamine or Sulfasalazine, Immunomodulator (methotrexate, 6-mercaptopurine, azathioprine)</i>					
Is the patient currently on any therapy? <input type="checkbox"/> YES <input type="checkbox"/> NO					

PRESCRIPTION INFORMATION												
MEDICATION	DOSE	DIRECTIONS	QTY	REFILLS								
<input type="checkbox"/> Simponi® UC (golimumab)	<input type="checkbox"/> 100mg SmartJect® Auto Injector <input type="checkbox"/> 100mg Pre-filled Syringe	<input type="checkbox"/> Initial Dose: 200mg subcutaneously at week 0, followed by 100mg at week 2 <input type="checkbox"/> Maintenance Dose: 100mg subcutaneously once every four weeks										
<input type="checkbox"/> Stelara® (ustekinumab)	<input type="checkbox"/> 90mg / mL Syringe <input type="checkbox"/> 45mg / mL Syringe <input type="checkbox"/> 130mg / 26mL Vial	<input type="checkbox"/> Initial Dose: A single IV infusion using weight-based dosing (patient weight in kg): <table border="1" style="margin-left: 20px;"> <thead> <tr> <th>Weight Range (kg)</th> <th>Recommended Dose</th> </tr> </thead> <tbody> <tr> <td><input type="checkbox"/> Up to 55 kg or less</td> <td>260 mg (2 vials)</td> </tr> <tr> <td><input type="checkbox"/> Greater than 55 kg to 85 kg</td> <td>390 mg (3 vials)</td> </tr> <tr> <td><input type="checkbox"/> Greater than 85 kg</td> <td>520 mg (4 vials)</td> </tr> </tbody> </table> <input type="checkbox"/> Maintenance Dose: 90mg SQ 8 weeks after initial IV dose, then every 8 weeks thereafter	Weight Range (kg)	Recommended Dose	<input type="checkbox"/> Up to 55 kg or less	260 mg (2 vials)	<input type="checkbox"/> Greater than 55 kg to 85 kg	390 mg (3 vials)	<input type="checkbox"/> Greater than 85 kg	520 mg (4 vials)		
Weight Range (kg)	Recommended Dose											
<input type="checkbox"/> Up to 55 kg or less	260 mg (2 vials)											
<input type="checkbox"/> Greater than 55 kg to 85 kg	390 mg (3 vials)											
<input type="checkbox"/> Greater than 85 kg	520 mg (4 vials)											
<input type="checkbox"/> Xeljanz® (tofacitinib)	<input type="checkbox"/> 5mg Tablet <input type="checkbox"/> 10mg Tablet	<u>Ulcerative Colitis:</u> <input type="checkbox"/> Initial Dose: 10mg twice daily for at least 8 weeks <input type="checkbox"/> Maintenance Dose: <input type="checkbox"/> 5mg twice daily <input type="checkbox"/> 10mg twice daily *Discontinue after 16 weeks of 10mg twice daily if adequate response not achieved. Use the lowest effective dose to maintain response.										

DELIVERY INSTRUCTIONS		
<input type="checkbox"/> Physician's Office	<input type="checkbox"/> Patient's Home	<input type="checkbox"/> 1 st dose MD office, Refills to patient's home
PHYSICIANS CONTACT INFORMATION & AUTHORIZATION		
Physician's Name:	Office Contact:	Institution:
Phone #:	Fax #:	Specialty:
Address:	City/State/Zip:	
Tax ID:	DEA #:	NPI #:
Physician's Signature:		Date:
<small>*Prescriber Authorization: I authorize this pharmacy and its representatives to act as my authorized agent to secure coverage and initiate the insurance prior authorization process for my patient(s), and to sign any necessary forms on my behalf as my authorized agent, including the receipt of any required prior authorization forms and the receipt and submission of patient lab values and other patient data. In the event that this pharmacy determines that it is unable to fulfill this prescription, I further authorize this pharmacy to forward this information and any related materials related to coverage of the product to another pharmacy of the patient's choice or in the patient's insurer's provider network. The information contained in this transmission may contain privileged and confidential information, including patient information protected by federal and state privacy laws. It is intended only for the use of the person(s) named above. If you are not the intended recipient, you are hereby notified that any review, dissemination, distribution, or duplication of this communication is strictly prohibited. If you are not the intended recipient, please contact the sender and destroy all copies of the original document. Created: 08/28/18 Revised: 05/01/19, 08/22/19, 02/04/20</small>		