

Patient Information

Patient Name: _____ DOB: _____ Sex: ☐ Male ☐ Female SSN: _____ Wt (kg/lbs): _____ Ht (cm/in): _____
 Address: _____ Phone: _____ Alternate: _____
 Caregiver Name: _____ Relation to Patient: _____ Phone: _____
 Insurance Plan: _____ Plan ID: _____ BIN #: _____ PCN #: _____ GRP #: _____

Please fax a copy of the front and back of the insurance card(s).

Prescriber + Shipping Information

Prescriber Name: _____ DEA: _____ NPI: _____
 Address: _____
 Phone: _____ Alternate: _____ Fax: _____ Email: _____
 If shipping to prescriber: ☐ First Fill ☐ Always ☐ Never

Clinical Information (Please fax all pertinent clinical and lab information)

Crohn's Disease: ☐ K50.0 (Crohn's Disease of the **Small** Intestine) ☐ K50.1 (Crohn's Disease of the **Large** Intestine)
☐ K50.8 (Crohn's Disease of **Both** Intestines) ☐ K50.9 (Crohn's Disease, unspecified)
Ulcerative Colitis: ☐ K51.0 (Ulcerative Pancolitis) ☐ K51.2 (Ulcerative Procolitis) ☐ K51.3 (Ulcerative Rectosigmoiditis)
☐ K51.5 (Left Sided Colitis) ☐ K51.8 (Other Ulcerative Colitis) ☐ K51.9 (Ulcerative Colitis, unspecified)
Other: ☐ _____
 Diagnosis Date: _____ TB Test: ☐ Yes ☐ No Neg. Test Date: _____

Prior Therapy ☐ Yes ☐ No _____
 Reason for Discontinuation of Therapy _____
 Approximate Start Date _____ Approximate End Date _____

Comorbidities: _____
 Concomitant Medications: _____
 Allergies: ☐ NKDA ☐ Other: _____

Prescription

☐ Cimzia® (certolizumab)	☐ Inject 400 mg subq at weeks 0, 2 and 4	☐ 6 x 200 mg/mL	☐ PFS ☐ Vials	0
	☐ Inject 400 mg subq every 4 weeks	☐ 2 x 200 mg/mL	☐ PFS ☐ Vials	
☐ Humira® (adalimumab) <i>Adults</i>	☐ Inject 160 mg subq on day 1, then 80 mg on day 15	☐ 6 x 40 mg/0.8mL	☐ Pens ☐ PFS	0
	☐ Inject 40 mg subq on day 29 and every other week thereafter	☐ 2 x 40 mg/0.8mL	☐ Pens ☐ PFS	
☐ Humira® (adalimumab) <i>Pediatrics ≥ 6 years</i>	☐ Inject 80 mg subq day 1, then 40 mg on day 15 (17 to <40 kg)	☐ 3 x 40 mg/0.8mL	☐ Pens ☐ PFS	0
	☐ Inject 160 mg subq day 1, then 80 mg on day 15 (≥40 kg)	☐ 6 x 40 mg/0.8mL		
	☐ Inject 20 mg subq on day 29 and every other week thereafter (17 to <40 kg)	☐ 2 x 20 mg/0.4mL	☐ PFS	
	☐ Inject 40 mg subq on day 29 and every other week thereafter (≥40 kg)	☐ 2 x 40 mg/0.8mL	☐ Pens ☐ PFS	
☐ Simponi® (golimumab)	☐ Inject 200 mg subq at week 0, then 100 mg at week 2	☐ 3 x 100 mg/mL	☐ SmartJect® Autoinjector ☐ PFS	0
	☐ Inject 100 mg subq every 4 weeks	☐ 1 x 100 mg/mL	☐ SmartJect® Autoinjector ☐ PFS	
☐ Stelara® (ustekinumab)	☐ Infuse 260 mg intravenously over no less than one hour (≤55kg)	☐ 2 x 130 mg/26 mL	☐ Vials	0
	☐ Infuse 390 mg intravenously over no less than one hour (>55 kg to <85 kg)	☐ 3 x 130 mg/26 mL		
	☐ Infuse 520 mg intravenously over no less than one hour (>85 kg)	☐ 4 x 130 mg/26mL		
	☐ Inject 90 mg subq 8 weeks following initial intravenous dose, then every 8 weeks thereafter	☐ 1 x 90 mg/mL	☐ PFS	
Patient eligible for self-administration? ☐ Yes ☐ No				
Date of last infusion: _____				
☐ Cosentyx® (secukinumab)	Initial Dose: ☐ 150 mg subq at weeks 0,1,2,3, and 4	☐ 5 x 150 mg	☐ PFS	0
	☐ 300 mg subq at weeks 0,1,2,3, and 4	☐ 10 x 150 mg	☐ Sensoready PEN	0
	Maintenance Dose: ☐ 150 mg every 4 weeks	☐ Qty:	☐ PFS ☐ Sensoready PEN	
☐ 300 mg every 4 weeks				

Injection training provided by: ☐ Physician's Office ☐ Pharmacy ☐ Other:

Per state-specific law, prescriptions will be dispensed as generic, if applicable, unless notated otherwise:

Prescriber's Signature: _____ Date: _____

I authorize Apollo Pharmacy and its representatives to act as an agent to initiate and execute the insurance prior authorization process for this prescription and any future fills of the same prescription for the patient listed above. I understand that I can revoke this designation at any time by providing written notice to Apollo Pharmacy.

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