

DATE

FAX FORM TO: 1.866.233.8317 | PHONE: 1.855.492.0817 | EMAIL: contact@bluegrass-rx.com

Complete the following or include demographic sheet.

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DISPENSE AS WRITTEN

Complete the jollowin	ng <u>or include demographic sheel</u>	<u></u>			
1. PATIENT INFORMATION			2. PRESCRIBER INFORMATION		
Name:			Name:		
Address:			DEA #: NPI #:	State Lic. #:	
City, State, ZIP:			Group or Hospital:		
Primary Phone:	DOE	3: / /	Address:		
Alternate Phone:	Gen	der:	City, State, Zip:		
Email:			Phone:	Fax:	
Primary Language	e: Last	Four of SSN:	Contact Person:	Phone:	
3. INSURANCE	INFORMATION	Fax copy of pres c	cription and insurance cards with this f	orm. if available (front and	back)
		· ···· - - - · · · ·	Secondary Insurance Company Name:		
Primary Insurance Company Name:					
Primary Cardholder Name:			Secondary Cardholder Name:		
	Self Spouse/Partner	O Child/Dependent	Relationship: Self Spouse		endent
Phone: -	- Member ID:	Group #:	Phone: Membe	er ID: Group #:	
4. DIAGNOSIS A	AND CLINICAL INFORMATIO	N			
Needs by Date:	/ /		Ship to: Office Other:		
Date of Diagnosis:	/ /		Enterocutaneous/rectovaginal fistulas?	○ Ye	s O No
B16.9 Acute hepatitis B w/o delta-agent or hepatic coma			Has patient been diagnosed with heart fail	ure? Ye	s No
 ☐ B18.1 Chronic viral hepatitis B w/o delta-agent ☐ K5Ø.ØØ Crohn's disease of small intestine w/o complications 			Has patient been diagnosed with lyphoma		
K50.10 Crohn's disease of small intestine w/o complications K50.10 Crohn's disease of large intestine w/o complications			Does patients have serious/active infection? Yes No		
K5Ø.8Ø Crohn's disease of small and large intestine w/o complications			, , ,	No If yes, results:	
K50.90 Crohn's disease, unspecified, w/o complications			Is patient at risk for hepatitis B infection?	○ Ye	_
 ○ K51.8Ø Other ulcerative colitis without complications ○ Other: 			If yes, has hepatitis B been ruled out or tre		
Crohn's severity: Moderate Severe N/A			Does patient have a latex allergy? Or Yes No No No No No No No No No N		
Prior (failed) medications:			Are there any contraindications to previous treatments? Yes No If yes, drug: Reason:		
Allergies			If yes, drug: Reason: Does patient require injection training? Yes \(\) No		
Current Medications:			If no, reason: Patient is independent		
Height (in/cm): Weight (lb/kg):			Patient trained by MD or referred to alternate trainer		
5. PRESCRIPTION	ON INFORMATION				
Medication	Dose/Strength Directions			Quantity	Refills
○ Baraclude [™]	○ 0.5 mg ○ 1 mg				
Cimzia™	Cimzia starter kit	O Induction dose: Inject SQ 400 mg (2 vials) on day 1, and at weeks 2 and 4.		1 kit (6 vials)	
	200 mg/1 ml PFS	Maintenance dose: Inject SQ 400 mg (2 vials) every 4 weeks.		1 kit (2 vials)	
○ Epivir HBV™	200 vial 100 mg	Other:			
 Hepsera™ 	10 mg				
Humira [™]	Crohn's starter package	 Induction dose: Inject SQ 160 mg (4 pens) on day 1, then 80 mg (2 pens) on day 15, then maintenance dosing 		1 kit	
	0 40 mg self injectable pen	Maintenance dose: Inject SQ 40 mg (one pen) every other week.		1 kit	
O D : 1 TM	0 40 mg PFS		40 mg (one syringe) every other week.	" (100 · · ·	
○ Remicade [™]	○ 100 mg vialmg/kg	☐ Induction dose: IV at 5 mg/kg (Dose =mg) at 0, 2, and 6 weeks. ☐ Maintenance dose: IV at 5 mg/kg (Dose =mg every 8 weeks) ☐ Other:		# of 100 mg vials:	
○ Simponi [™]		 Induction dose: Inject 200 mg SQ initially, followed by 100 mg at week 2, and then 100 mg and then 100 mg every 4 weeks Maintenance dose: Inject 100 mg SQ every 4 weeks 			
Solesta™	Four 1ml PFS w/ needles				
Tysabri™					
Tyzeka™ (LD)*	(LD)* This is a limited distribution	drug that requires additional handlir	ng. Please call (1.855.492.0817) for more informat	ion.	
O Viread™					
Zorbtive [™]	8.8 mg vial				
Ancillary supplies and	kits will be provided as needed for	administration.			
6. PRESCRIBER	R SIGNATURE				

IMPORTANT NOTICE: This facsimile transmission is intended to be delivered only to the named addressee and may contain material that is confidential, privileged, proprietary or exempt from disclosure under applicable law. If it is received by anyone other than the named addressee, the recipient should immediately notify the sender at the address or telephone number set forth herein and obtain instructions as to disposal of the transmitted material. In no event should such material be read or retained by anyone other than the named addressee, except by express authority of the sender to the named addressee.

PRODUCT SUBSTITUTION PERMITTED

DATE