

# **GASTROENTEROLOGY REFERRAL FORM**

P: (877) 778-0318 F: (877) 778-0399

Revised: 08/18/2025

Patient Information				
Patient Name:	DOB: Phone:			
	Therapy   Continuing Therapy   Next Treatment Date:			
Patient Weight: lk				
Diagnosis	ICD-10			
☐ Crohn's Disease	K50			
<ul><li>Ulcerative Colitis</li><li>Prescription</li></ul>	Directions, Quantity, Form	Refill		
Tremfya	Induction:			
yu	<ul> <li>IV: **Anaphylaxis kit and epinephrine pen to be dispensed**</li> <li>Tremfya 200mg/20mL Vial: 200mg IV on weeks 0, 4, and 8. Qty: 3 Sodium Chloride 0.9% 250mL Bag - UAD for Tremfya infusion. Qty: 3</li> <li>SubQ:</li> <li>Tremfya 200mg/2mL Pen: 400mg SubQ on weeks 0, 4, and 8. Qty: 6</li> <li>Maintenance:</li> <li>SubQ:</li> <li>Tremfya 100mg/mL One Press or Pen (as covered by insurance): 100mg SubQ every 8 weeks (beginning at week 16). Qty: 1</li> </ul>	□ x1 year □ Other:		
Humira (or Humira biosimilar as covered by insurance)	Induction:  Humira 80mg/0.8mL Pen Citrate Free: 160mg SubQ on day 1, then inject 80mg SubQ 2 weeks later (day 15). Qty: 3  Maintenance:  Humira 40mg/0.4mL Pen Citrate Free: 40mg SubQ every other week (starting on day 29). Qty: 1	□ x1 year □ Other:		
<b>Skyrizi</b> (for Crohn's disease)	Induction:  • IV: **Anaphylaxis kit and epinephrine pen to be dispensed**  □ Skyrizi 600mg/10mL Vial: 600mg IV on weeks 0, 4, and 8. Qty: 3  □ Sodium Chloride 0.9% 250mL Bag - UAD for Skyrizi infusion. Qty: 3  Maintenance:  • SubQ: (select one of the following doses)  □ Skyrizi 180mg/1.2mL Cartridge: 180mg SubQ at week 12, then 8 weeks thereafter. Qty: 1  □ Skyrizi 360mg/2.4mL Cartridge: 360mg SubQ at week 12, then 8 weeks thereafter. Qty: 1	□ x1 year □ Other:		
<b>Skyrizi</b> (for Ulcerative colitis)	Induction:  IV: **Anaphylaxis kit and epinephrine pen to be dispensed**  Skyrizi 600mg/10mL Vial: 1200mg IV on weeks 0, 4, and 8. Qty: 6  Sodium Chloride 0.9% 250mL Bag - UAD for Skyrizi infusion. Qty: 3  Maintenance:  SubQ: (select one of the following doses)  Skyrizi 180mg/1.2mL Cartridge: 180mg SubQ at week 12, then 8 weeks thereafter. Qty: 1  Skyrizi 360mg/2.4mL Cartridge: 360mg SubQ at week 12, then 8 weeks thereafter. Qty: 1	□ x1 year □ Other:		
Stelara	Induction:  Iv: **Anaphylaxis kit and epinephrine pen to be dispensed**    ≤55kg: Stelara 130mg/26mL Vial: 260mg IV as a single dose. Qty: 2   Sodium Chloride 0.9% 250mL Bag - UAD for Stelara infusion. Qty: 1    55kg to 85kg: Stelara 130mg/26mL Vial: 390mg IV as a single dose. Qty: 3   Sodium Chloride 0.9% 250mL Bag - UAD for Stelara infusion. Qty: 1    >85kg: Stelara 130mg/26mL Vial: 520mg IV as a single dose. Qty: 4   Sodium Chloride 0.9% 250mL Bag - UAD for Stelara infusion. Qty: 1    Maintenance:   SubQ:   Stelara 90mg/mL PFS: 90mg SubQ after every 8 weeks from initial infusion, then 8 weeks thereafter. Qty: 1	□ x1 year □ Other:		

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Prescription	Directions, Quantity, Form	Refill
Entyvio	OPTION 1: IV Only	□ x1 year
	Induction:	□ Other:
	IV: **Anaphylaxis kit and epinephrine pen to be dispensed**	
	☐ Entyvio 300mg Vial: 300mg IV at weeks 0, 2, and 6. Qty: 3	
	Sodium Chloride 0.9% 250mL Bag - Use as directed for Entyvio infusion. Qty: 3	
	Sodium Chloride 0.9% 50mL Bag - Flush IV line with 30mL after Entyvio infusion. Qty: 3  Sterile Water for Injection 10mL Vial - Use as directed to reconstitute Entyvio. Qty: 3	
	Maintenance:	
	☐ Entyvio 300mg Vial: 300mg IV every 8 weeks. Qty: 1	
	Sodium Chloride 0.9% 250mL Bag - UAD for Entyvio infusion. Qty: 1	
	Sodium Chloride 0.9% 50mL Bag - Flush IV line with 30mL after Entyvio infusion. Qty: 1 Sterile Water for Injection 10mL Vial - UAD to reconstitute Entyvio. Qty: 1	
	OPTION 2: IV to SubQ	
	Induction:	
	IV: **Anaphylaxis kit and epinephrine pen to be dispensed**	
	☐ Entyvio 300mg Vial: 300mg IV at weeks 0 and 2. Qty: 2	
	Sodium Chloride 0.9% 250mL Bag - UAD for Entyvio infusion. Qty: 2 Sodium Chloride 0.9% 50mL Bag - Flush IV line with 30mL after Entyvio infusion. Qty: 2	
	Sterile Water for Injection 10mL Vial - UAD to reconstitute Entyvio. Qty: 2	
	Maintenance:	
	☐ Entyvio 108mg/0.68mL Pen: 108mg SubQ at week 6, then once every 2 weeks thereafter	
	(beginning after at least 2 IV infusions). Qty: 2	
Infliximab	Induction:	□ x1 year
(or Infliximab biosimilar as	IV: **Anaphylaxis kit and epinephrine pen to be dispensed**	□ Other:
covered by insurance)	☐ Infliximab 100mg Vial (or Infliximab biosimilar) - Infusemg (dose 5mg/kg)	
	intravenously on weeks 0, 2, and 6. Qty: QS Sterile Water for Injection 10mL Vial - UAD to reconstitute Infliximab. Qty: QS	
	Sodium Chloride 0.9% 250mL OR 500mL Bag- UAD for Infliximab infusion. Qty: 1/dose	
	For doses < 999 mg= 250mL bag. For doses > 1000 mg= 500 mL bag	
	Premedications (15-30 minutes prior to infusion):  • Acetaminophen 325mg Tablet - 2 tablets by mouth. Qty: 2/dose	
	Acetaminophen 325mg Tablet - 2 tablets by mouth, Qty. 2/dose     Diphenhydramine HCL 25mg Capsule - 1-2 capsules by mouth, Qty: 2/dose	
	OR	
	☐ Loratadine 10mg Tablet - 1 tablet by mouth. Qty: 1/dose (Diphenhydramine PO will not be dispensed if selected)	
	☐ Certirizine 10mg Tablet - 1 tablet by mouth. Qty: 1/dose (Diphenhydramine PO will not be dispensed if selected)	
	Maintenance: **Anaphylaxis kit and epinephrine pen to be dispensed**	
	☐ Infliximab 100mg Vial (or Infliximab biosimilar) - Infusemg (dose 5mg/kg)	
	intravenously every 8 weeks (starting at week 14). Qty: QS Sterile Water for Injection 10mL Vial - UAD to reconstitute Infliximab. Qty: QS	
	Sodium Chloride 0.9% 250mL OR 500mL Bag- UAD for Infliximab infusion. Qty: 1/dose	
	For doses < 999 mg= 250mL bag. For doses > 1000 mg= 500 mL bag	
	Premedications (15-30 minutes prior to infusion):  • Acetaminophen 325mg Tablet - 2 tablets by mouth. Qty: 2/dose	
	Diphenhydramine HCL 25mg Capsule - 1-2 capsules by mouth. Qty: 2/dose	
	OR	
	☐ Loratadine 10mg Tablet - 1 tablet by mouth. Qty: 1/dose (Diphenhydramine PO will not be dispensed if selected) ☐ Certirizine 10mg Tablet - 1 tablet by mouth. Qty: 1/dose (Diphenhydramine PO will not be dispensed if selected)	
	Certifizine formy fablet - Flablet by mouth, xty. Indose (bipinelinyuranime FO with not be dispensed it selected)	
Tysabri	REQUIRED: Tysabri TOUCH Patient Enrollment #:	□ x1 year
	IV: **Anaphylaxis kit and epinephrine pen to be dispensed**	□ Other:
	☐ Tysabri DS 300mg/15mL Vial: 300mg IV every 4 weeks. Qty: 1  Sodium Chloride 0.9% 100mL Bag - UAD for Tysabri infusion. Qty: 1	
Cimzia	Induction:	□ x1 year
	☐ Cimzia PFS Start Kit 6 X 200mg/mL: 400mg SubQ on week 0, 2, and 4. Qty: 3	□ Other:
	Maintenance:	_ 001.
	☐ Cimzia PFS 2 x 200mg/mL Kit: 400mg SubQ every 4 weeks. Qty: 1	

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# **COMPREHENSIVE SUPPORT FOR GASTROENTEROLOGY THERAPY**

Revised: 08/18/2025

Additional Pre-Medication (15-30 minutes prior to infusion):  Solu-Medrol #1 vial/infusion + bacteriostatic water #1 vial:	
Flushing Protocol (for IV Infusions only) (Dispense quantity sufficient per protocol)  Sodium Chloride 0.9% 5-10mL per SASH protocol  #3 per infusion day  +1 per additional IV infusion  #1 per infusion day	SASH protocol If Port/PICC (500u/mL)
Anaphylactic Reaction Orders (for infusions only: Entyvio IV, Tremfya IV, Skyrizi IV, Stelara IV,	Tvsabri IV. Infliximab IV)
■ Epinephrine (dosed by patient weight) #2 pens  •>30kg (>66lbs): EpiPen 0.3mg use as directed for anaphylaxis; may repeat in 5-10 minutes x1  •15-30kg (33-66lbs): EpiPen 0.15mg use as directed for anaphylaxis; may repeat in 5-10 minutes x1  •#2 Diphenhydramine HCL Injection 56mg/mL Vial (IV repeats to the control of t	aphylactic reaction per protocol.
Provider Information	
Provider Name: Provider's Signature:	Date:
Address, City, State, Zip:	
Provider NPI: Phone: Fax: Co	
Please kindly sign above to grant authorization for Harper's Pharmacy, Inc. operating as AmeriPharma, to undertake all necessary and reasonable actions, including submitting any neces the prompt and accurate dispensing of the medication(s) prescribed for the patient mentioned above. Pharmacy to dispense stock vials and IV supplies for administration. Prescriber	
*For AmeriPharma Use Only	
Notes:	
Required Documentation for Referral Processing & Insurance Approval	
☐ Include signed and completed order (MD/prescriber to complete page 1)	
☐ Include patient demographic information and insurance information	
☐ Include patient's medication list	
<ul> <li>Supporting clinical notes to include any past tried and/or failed therapies, intoleration</li> </ul>	ance, benefits, or
contraindications to conventional therapy	,,
☐ For biologic orders, has the patient had a documented contraindication/intolera	nce or failed trial
of a conventional therapy (i.e., 6MP, Azathioprine)?	
If yes, which drug(s)?	
☐ For biologic orders, does the patient have a documented contraindication/intole trial to any other biologic (i.e., Humira, Stelara, Cimzia)? ☐ Yes ☐ No If yes, which drug(s)?	erance or failed
☐ Include labs and/or test results to support diagnosis	
☐ If applicable - Last known biological therapy: and last date rec	eived:
If patient is switching to biologic therapies, please perform a wa	
period ofweeks prior to starting ordered biologic therapy.	
□ Other medical necessity:	

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Required F	ro-Scroonin	n (Based n	n drug therapy	201
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<b>TB Screening test completed within 12 months - attach results</b> Required for: Cimzia, Infliximab, Stelara, Entyvio, Skyrizi
□ Positive □ Negative
Hepatitis B screening test completed (Hepatitis B surface antigen) - attach results Required for: Cimzia, Infliximab  □ Positive □ Negative
JCV antibody & TOUCH authorization Required for: Tysabri  Positive Negative

AmeriPharma Specialty Care will complete insurance verification and submit all required documentation for approval to the patient's insurance company for eligibility. Our team will notify you if any additional information is required. We will review financial responsibility with the patient and refer them to any available co-pay assistance as needed. Thank you for the referral.

## **Nursing Orders:**

### **Assessment and Infusion Administration:**

- RN to complete patient assessment prior to infusion.
- Administer via ambulatory rateflow regulator administration set.
- Notify the pharmacy immediately of any pump or supply issues.
- Pharmacy to provide all required infusion supplies, including: Catheter Care Kit [A4221], Supply Kit [A222], and ancillary medications/equipment [E0779 or E0781]

#### Vascular Access:

- RN to insert, maintain, and remove peripheral IV catheter (PIVC) or access central venous catheter using aseptic technique.
- Rotate PIVC as clinically indicated for signs of infiltration or irritation.
- . Monitor patients throughout infusion. Discontinue PIVC upon completion of the infusion or infusion cycle.

#### **Catheter Maintenance:**

- Flush PIVC with 3-5 mL of 0.9% Sodium Chloride before and after each dose, with lab draws, and as needed to assess catheter patency.
- Flush Central Line (PICC, Hickman, etc.) with 10mL of 0.9% Sodium Chloride before and after infusion.
- Lock Central Line (PICC, Hickman, etc) with 100units/mL, 5ml to maintain patency on treatment days, daily and PRN.
- For port access, use sterile technique with a non-coring needle. Flush with 10 mL of 0.9% Sodium Chloride before and after infusion.
- Lock port with Heparin 100 units/mL, 5 mL to maintain patency. Flush port on treatment days, at least once monthly, and PRN.
- De-access port post-infusion and apply sterile pressure gauze and transparent dressing.
- Discontinue port maintenance upon discontinuation of pharmacy services.
- RN to prepare and administer ordered medication, infusing full contents of infusion bag/vials per prescribed dose.
- Infusions may be administered within a +/- 3-day window to accommodate patient scheduling.

#### **Patient and Caregiver Education:**

• RN to provide education on: Monitoring for signs/symptoms of complications, Infection prevention and IV access safety, Emergency preparedness, 24-hour on-call availability, Safe medication handling and administration.

#### **Coordination of Care:**

- Pharmacy and nursing services may be discontinued if continuation orders are not received.
- · Signed discharge orders must be on file prior to termination of services.

Please fax all information to (877) 778-0399 or call (877) 778-0318 for assistance.

Concernancy statement. This resissage is increased in the interest of the concernancy statement of the

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