

## Oregon Health & Science University Hospital and Clinics Provider's Orders



ADULT AMBULATORY INFUSION ORDER
Ustekinumab (STELARA) for
Inflammatory Bowel Disease
(Crohn's Disease and Ulcerative
Colitis)

ACCOUNT NO. MED. REC. NO. NAME BIRTHDATE

Patient Identification

Weight:kg	Page 1 of 3
Altergies:	ALL ORDERS MUST BE MARKED IN INK WITH A CHECKMARK ( ✓ ) TO BE ACTIVE.
Diagnosis Code:	Weight:kg Height:cm
Treatment Start Date: Patient to follow up with provider on date:	
1. Send FACE SHEET and H&P or most recent chart note. 2. A Tuberculin test must have been placed and read as negative prior to initiation of treatment (PPD or QuantiFERON Gold blood test). Please send results with order. If result is indeterminate, a follow up chest X-ray must be performed to rule out TB. Please send results with order. Patients should not have an active ongoing infection at the onset of ustekinumab therapy. 3. Monitor patients for signs / symptoms of active TB, infection, reversible posterior leukoencephalopathy syndrome (RPLS), and malignancy throughout therapy.  PRE-SCREENING: (Results must be available prior to initiation of therapy):  Tuberculin skin test or QuantiFERON Gold blood test results scanned with orders Chest X-Ray result scanned with orders if TB test result is indeterminate.  LABS:  CBC+DIFF, Routine, ONCE  COMPLETE METABOLIC PANEL, Routine, ONCE  NURSING ORDERS 1. TREATMENT PARAMETER – Hold treatment and contact provider if TB test result is positive or if screening has not been performed. 2. Follow facility policies and/or protocols for vascular access maintenance with appropriate flush solution, declotting (alteplase), and/or dressing changes. 3. For signs and symptoms of active infection contact provider prior to administering.  MEDICATIONS:  Initial Dose:  Ustekinumab (STELARA) in sodium chloride 0.9 %, intravenous, ONCE, over 1 hour Less than or equal to 55 kg  Greater than 55-85 kg  300 mg (three 130 mg vials)	Treatment Start Date: Patient to follow up with provider on date:
1. Send FACE SHEET and H&P or most recent chart note.  2. A Tuberculin test must have been placed and read as negative prior to initiation of treatment (PPD or QuantiFERON Gold blood test). Please send results with order. If result is indeterminate, a follow up chest X-ray must be performed to rule out TB. Please send results with order. Patients should not have an active ongoing infection at the onset of ustekinumab therapy.  3. Monitor patients for signs / symptoms of active TB, infection, reversible posterior leukoencephalopathy syndrome (RPLS), and malignancy throughout therapy.  PRE-SCREENING: (Results must be available prior to initiation of therapy):  Tuberculin skin test or QuantiFERON Gold blood test results scanned with orders  Chest X-Ray result scanned with orders if TB test result is indeterminate.  LABS:  CBC+DIFF, Routine, ONCE  COMPLETE METABOLIC PANEL, Routine, ONCE  NURSING ORDERS  1. TREATMENT PARAMETER – Hold treatment and contact provider if TB test result is positive or if screening has not been performed.  2. Follow facility policies and/or protocols for vascular access maintenance with appropriate flush solution, declotting (alteplase), and/or dressing changes.  3. For signs and symptoms of active infection contact provider prior to administering.  MEDICATIONS:  Initial Dose:  Ustekinumab (STELARA) in sodium chloride 0.9 %, intravenous, ONCE, over 1 hour Less than or equal to 55 kg  390 mg (three 130 mg vials)	**This plan will expire after 365 days at which time a new order will need to be placed**
<ul> <li>□ Tuberculin skin test or QuantiFERON Gold blood test results scanned with orders</li> <li>□ Chest X-Ray result scanned with orders if TB test result is indeterminate.</li> <li>LABS:</li> <li>□ CBC+DIFF, Routine, ONCE</li> <li>□ COMPLETE METABOLIC PANEL, Routine, ONCE</li> <li>NURSING ORDERS</li> <li>1. TREATMENT PARAMETER – Hold treatment and contact provider if TB test result is positive or if screening has not been performed.</li> <li>2. Follow facility policies and/or protocols for vascular access maintenance with appropriate flush solution, declotting (alteplase), and/or dressing changes.</li> <li>3. For signs and symptoms of active infection contact provider prior to administering.</li> <li>MEDICATIONS:</li> <li>Initial Dose:</li> <li>□ ustekinumab (STELARA) in sodium chloride 0.9 %, intravenous, ONCE, over 1 hour Less than or equal to 55 kg</li> <li>□ 260 mg (two 130 mg vials)</li> <li>□ 390 mg (three 130 mg vials)</li> </ul>	<ol> <li>A Tuberculin test must have been placed and read as negative prior to initiation of treatment (PPD or QuantiFERON Gold blood test). Please send results with order. If result is indeterminate, a follow up chest X-ray must be performed to rule out TB. Please send results with order. Patients should not have an active ongoing infection at the onset of ustekinumab therapy.</li> <li>Monitor patients for signs / symptoms of active TB, infection, reversible posterior leukoencephalopathy</li> </ol>
<ul> <li>□ CBC+DIFF, Routine, ONCE</li> <li>□ COMPLETE METABOLIC PANEL, Routine, ONCE</li> <li>NURSING ORDERS</li> <li>1. TREATMENT PARAMETER – Hold treatment and contact provider if TB test result is positive or if screening has not been performed.</li> <li>2. Follow facility policies and/or protocols for vascular access maintenance with appropriate flush solution, declotting (alteplase), and/or dressing changes.</li> <li>3. For signs and symptoms of active infection contact provider prior to administering.</li> <li>MEDICATIONS:</li></ul>	
<ol> <li>TREATMENT PARAMETER – Hold treatment and contact provider if TB test result is positive or if screening has not been performed.</li> <li>Follow facility policies and/or protocols for vascular access maintenance with appropriate flush solution, declotting (alteplase), and/or dressing changes.</li> <li>For signs and symptoms of active infection contact provider prior to administering.</li> </ol> MEDICATIONS: Initial Dose: <ul> <li>ustekinumab (STELARA) in sodium chloride 0.9 %, intravenous, ONCE, over 1 hour</li> <li>Less than or equal to 55 kg</li> <li>Greater than 55-85 kg</li> <li>390 mg (three 130 mg vials)</li> </ul>	
Initial Dose:  ☐ ustekinumab (STELARA) in sodium chloride 0.9 %, intravenous, ONCE, over 1 hour  Less than or equal to 55 kg ☐ 260 mg (two 130 mg vials)  Greater than 55-85 kg ☐ 390 mg (three 130 mg vials)	screening has not been performed.  2. Follow facility policies and/or protocols for vascular access maintenance with appropriate flush solution, declotting (alteplase), and/or dressing changes.
	<ul> <li>□ ustekinumab (STELARA) in sodium chloride 0.9 %, intravenous, ONCE, over 1 hour</li> <li>Less than or equal to 55 kg</li> <li>□ 260 mg (two 130 mg vials)</li> <li>□ 390 mg (three 130 mg vials)</li> </ul>

☐ ustekinumab (STELARA) 90 mg, subcutaneous, ONCE, every 8 weeks

**Maintenance Doses:** (starting 8 weeks after initial dose)



## Oregon Health & Science University Hospital and Clinics Provider's Orders

ADULT AMBULATORY INFUSION ORDER

# Ustekinumab (STELARA) for Inflammatory Bowel Disease (Crohn's Disease and Ulcerative Colitis)

ACCOUNT NO.
MED. REC. NO.
NAME
BIRTHDATE

Patient Identification

Page 2 of 3

ALL ORDERS MUST BE MARKED IN INK WITH A CHECKMARK ( ✓ ) TO BE ACTIVE.

#### AS NEEDED MEDICATIONS:

- 1. acetaminophen (TYLENOL) tablet, 650 mg, oral, EVERY 4 HOURS AS NEEDED for fever
- 2. diphenhydrAMINE (BENADRYL) capsule, 25 mg, oral, EVERY 4 HOURS AS NEEDED for itching

#### **HYPERSENSITIVITY MEDICATIONS:**

- NURSING COMMUNICATION If hypersensitivity or infusion reactions develop, temporarily hold the
  infusion and notify provider immediately. Administer emergency medications per the Treatment
  Algorithm for Acute Infusion Reaction (OHSU HC-PAT-133-GUD, HMC C-132). Refer to algorithm for
  symptom monitoring and continuously assess as grade of severity may progress.
- 2. diphenhydrAMINE (BENADRYL) injection, 25-50 mg, intravenous, AS NEEDED x 1 dose for hypersensitivity or infusion reaction
- 3. EPINEPHrine HCI (ADRENALIN) injection, 0.3 mg, intramuscular, AS NEEDED x 1 dose for hypersensitivity or infusion reaction
- 4. hydrocortisone sodium succinate (SOLU-CORTEF) injection, 100 mg, intravenous, AS NEEDED x 1 dose for hypersensitivity or infusion reaction
- 5. famotidine (PEPCID) injection, 20 mg, intravenous, AS NEEDED x 1 dose for hypersensitivity or infusion reaction

By signing below, I represent the following: I am responsible for the care of the patient (who is identified at the top of this form); I hold an active, unrestricted license to practice medicine in:   Oregon   (check box that corresponds with state where you provide care to patient and where you are currently licensed. Specify state if not Oregon);					
My physician license Number is #	ope of practice and aut	ECOMPLETED TO BE thorized by law to order	A VALID Infusion of the		
Provider signature:	Date	e/Time:			
Printed Name:	Phone:	Fax:			



## Oregon Health & Science University Hospital and Clinics Provider's Orders

ADULT AMBULATORY INFUSION ORDER
Ustekinumab (STELARA) for
Inflammatory Bowel Disease
(Crohn's Disease and Ulcerative
Colitis)

ACCOUNT NO.
MED. REC. NO.
NAME
BIRTHDATE

Patient Identification

Page 3 of 3

ALL ORDERS MUST BE MARKED IN INK WITH A CHECKMARK ( ✓ ) TO BE ACTIVE.

### Please check the appropriate box for the patient's preferred clinic location:

☐ Hillsboro Medical Center

Infusion Services 364 SE 8th Ave, Medical Plaza Suite 108B Hillsboro, OR 97123

Phone number: (503) 681-4124 Fax number: (503) 681-4120

☐ Mid-Columbia Medical Center

Celilo Cancer Center 1800 E 19th St The Dalles, OR 97058

Phone number: (541) 296-7585 Fax number: (541) 296-7610 ☐ Adventist Health Portland

Infusion Services 10123 SE Market St Portland, OR 97216

Phone number: (503) 261-6631 Fax number: (503) 261-6756