

Patient Name
 DOB
 MRN
 Physician
 CSN

Physician's Orders

USTEKINUMAB (STELARA) FOR CROHN'S DISEASE - ADULT, OUTPATIENT, COREWELL HEALTH INFUSION CENTER

Page 1 of 3

Defaults for orders not otherwise specified below:

- Interval: **INDUCTION** – Once x 1 treatment (begin maintenance dose on day 56)
- Interval: **MAINTENANCE** – Every 56 days

Duration:

- Until date: _____
- 1 year
- _____ # of Treatments

Anticipated Infusion Date _____ ICD 10 Code with Description _____

Height _____ (cm) Weight _____ (kg) Allergies _____

Site of Service

- CH Gerber
- CH Lemmen Holton (GR)
- CH Pennock
- CH Greenville
- CH Helen DeVos (GR)
- CH Ludington
- CH Reed City
- CH Zeeland
- CH Blodgett (GR)

Provider Specialty

- Allergy/Immunology
- Infectious Disease
- OB/GYN
- Rheumatology
- Cardiology
- Internal Med/Family Practice
- Other
- Surgery
- Gastroenterology
- Nephrology
- Otolaryngology
- Urology

Appointment Requests

- Infusion Appointment Request**
 Status: Future, Expected: S, Expires: S+365, Sched. Tolerance: Schedule appointment at most 3 days before or at most 3 days after, Infusion and possible labs. Verify that all INDUCTION/LOADING DOSES have been scheduled and offset appropriately when scheduling MAINTENANCE DOSES.

Provider Ordering Guidelines

- ONC PROVIDER REMINDER 15**
 USTEKINUMAB (STELERA) Crohn disease:

 Tuberculosis surveillance and management: Screen prior to starting therapy. Treat latent infection prior to starting therapy.

 Induction: IV:

 Less than or equal to 55 kg: 260 mg as single dose

 Greater than 55 kg to 85 kg: 390 mg as single dose

 Greater than 85 kg: 520 mg as single dose

 Maintenance: SubQ: 90 mg every 8 weeks; begin maintenance dosing 8 weeks after the IV induction dose.

Safety Parameters and Special Instructions

- ONC SAFETY PARAMETERS AND SPECIAL INSTRUCTIONS 6**
 Verify all INDUCTION/LOADING DOSES given prior to start of MAINTENANCE DOSES
- ONC SAFETY PARAMETERS AND SPECIAL INSTRUCTIONS 5**
 TUBERCULOSIS SURVEILLANCE AND MANAGEMENT RECOMMENDATIONS: Screen prior to treatment. Treat latent infection prior to starting therapy.

CONTINUED ON PAGE 2 →

NOTE: Epic Treatment/Therapy Plan Orders. To be scanned/attached to the appropriate Infusion Referral Order in Epic.

Confidentiality of this medical record shall be maintained except when use or disclosure is required or permitted by law, regulation, or written authorization by the patient.

Patient Name

DOB

MRN

Physician

CSN

USTEKINUMAB (STELARA) FOR CROHN'S DISEASE - ADULT, OUTPATIENT, COREWELL HEALTH INFUSION CENTER (CONTINUED)

Page 2 of 3

Labs

Arrange For Patient To Have Id Tb Skin Test Administered And Read Or Serum Tb Screening Lab Prior To Therapy Or Annually

ONC PROVIDER REMINDER 28

Arrange for patient to have intradermal TB skin test (tuberculin PPD) screening performed and read prior to initiating therapy and annually.

TB Screen (Quantiferon Gold)

Status: Future, Expected: S, Expires: S+365, URGENT, Clinic Collect, Blood, Blood, Venous

Additional Lab Orders

	Interval	Duration
<input type="checkbox"/> Labs: _____	<input type="checkbox"/> Every ___ days	<input type="checkbox"/> Until date: _____
	<input type="checkbox"/> Once	<input type="checkbox"/> 1 year
		<input type="checkbox"/> _____ # of Treatments

Nursing Orders

ONC NURSING COMMUNICATION 15

USTEKINUMAB (STELERA):

Hypersensitivity, including anaphylaxis and angioedema, has been reported. Discontinue immediately with signs/symptoms of hypersensitivity reaction and treat appropriately as indicated.

Monitor for signs/symptoms of infection, reversible posterior leukoencephalopathy syndrome (RPLS), and squamous cell skin carcinoma.

ONC NURSING COMMUNICATION 100

May Initiate IV Catheter Patency Adult Protocol

Hypersensitivity Reaction Adult Oncology Protocol

S

Until discont'd

Routine, Until discontinued Starting when released for 24 hours

HYPERSENSITIVITY REACTIONS:

Discontinue the medication infusion immediately.

Activate emergency response for severe or rapidly progressing symptoms. Where available consider calling RAP and have crash cart available. Call 911 or code team (if applicable) as needed for an absence of pulse and respirations. Refer to site specific emergency response policy.

Stay with patient until symptoms have resolved.

Initiate/Continue Oxygen to maintain SpO2 greater than 90% and discontinue Oxygen Therapy to maintain SpO2 above 90%

For severe or rapidly progressing hypersensitivity reaction symptoms, monitor vital signs and pulse oximeter readings every 2 to 5 minutes until the patient is stable and symptoms resolve.

Document medication infusing and approximate dose received at time of reaction in the patient medical record. Document allergy to medication attributed with causing reaction in patient medical record. Complete Adverse Drug Reaction form per Pharmacy Clinical Policy.

Treatment Parameters

ONC MONITORING AND HOLD PARAMETERS 4

May proceed with treatment if tuberculosis screening test with either TB Screen blood test (QuantiFERON® Gold Plus) or TB skin test have been resulted prior to first dose and the results are negative.

Confidentiality of this medical record shall be maintained except when use or disclosure is required or permitted by law, regulation, or written authorization by the patient.

