

## Ustekinumab (Stelara) Treatment Plan for Gastroenterology

Patient Name: \_\_\_\_\_ DOB: \_\_\_\_\_

Height (cm): \_\_\_\_\_ Weight (kg): \_\_\_\_\_ Allergies: \_\_\_\_\_

### **Diagnosis (select one and complete the 2<sup>nd</sup> and 3<sup>rd</sup> digits to complete the ICD-10 code):**

- K50.0\_\_\_ Crohn's Disease (small intestine)                       K50.8\_\_\_ Crohn's Disease (small and large intestine)
- K50.1\_\_\_ Crohn's Disease (large intestine)                       K50.9\_\_\_ Crohn's Disease, Unspecified
- Other: ICD 10 Code: \_\_\_\_\_ Diagnosis Description: \_\_\_\_\_

### **Pre-Medications: \*\*administered 30 minutes prior to infusion\*\***

- None
- Acetaminophen 650 mg PO
- Diphenhydramine:    Dose:  25 mg    50 mg                      Route:  PO or  IVP
- Methylprednisolone: Dose:  40 mg or  125                      Route: IVP
- Famotidine:                      Dose: 20 mg                      Route:  PO or  IVPB
- Other (include drug, dose, and route): \_\_\_\_\_

### **Drug Orders:**

- Induction: Ustekinumab (Stelara) (J3358) per 250 mL Sodium Chloride 0.9% IV to infuse over 1 hour for 1 dose

Dose:  Weight < 55 kg: 260 mg

Weight= 55-85 kg: 390 mg

Weight > 85 kg: 520 mg

- Subcutaneous maintenance dosing to be initiated 8 weeks following loading dose and coordinated by physician office

### **Lab Orders:**

\_\_\_\_\_

### **Standing Orders:**

- Infusion Reaction Protocol (CPOE-1396) will be activated if any hypersensitivity reaction occurs, including anaphylaxis. Infusion will be stopped and physician notified.

Physician Signature: \_\_\_\_\_ Date: \_\_\_\_\_

Physician Name: \_\_\_\_\_ Phone: \_\_\_\_\_

**Pre-Screening Requirements:**

- Provide TB screening results (PPD or QuantiFERON Gold Test) prior to start of therapy and within last 12 months
- Provide Hepatitis screening (Hepatitis B Surface Antigen) prior to start of therapy and within last 12 months

**Previous Therapies:**

- For new patient referrals, please send history and physical and most recent physician note with completed plan
- If patient has previously received ustekinumab at another facility, please provide last date received: \_\_\_\_\_
- If patient has previously received another biologic therapy, please provide the name: \_\_\_\_\_  
and the last date received: \_\_\_\_\_

**Insurance/Authorization Information:**

Insurance Type: \_\_\_\_\_

Insurance Authorization Reference Number: \_\_\_\_\_

Date Obtained: \_\_\_\_\_ Authorization Valid Until: \_\_\_\_\_

Additional Notes: \_\_\_\_\_

Fax completed Treatment Plan with authorization information to McLeod Infusion Services at the number below or call with any questions.

Florence: 843-777-6001 (Fax)

843-777-4655 (Phone)