



TREMFYA IV MEDICATION ORDER

This order is only for the infusion version of Tremfya, not the injectable version.

Patient's Name (Last, First, Middle) _____ DOB _____

Patient's height in feet and inches _____ Patient's weight in pounds _____

Diagnosis *These diagnoses are all without complications. For with complications please change last digit from 0 to 1.*

- K50.00 Crohn's disease of sm intestine w/o comp
- K50.10 Crohn's disease of lg intestine w/o comp
- K50.80 Crohn's disease of both int w/o comp
- K50.90 Crohn's disease unspecified w/o comp
- K51.00 UC pancolitis w/o comp
- K51.20 UC proctitis w/o comp
- K51.30 UC rectosigmoiditis w/o comp
- K51.50 Left-sided colitis w/o comp
- K51.80 Other UC w/o comp
- K51.90 UC unspecified w/o comp

Details Needed for Approval

- How is the patient's UC classified? Circle one: Mild Moderate Moderate-to-Severe Severe
- Provide evidence of no latent TB within 3 months or, if positive, document start of anti-TB therapy.
- Provide patient's vaccination history.
- Has the patient tried another systemic therapy that is FDA labeled for this condition? _____
- Has the patient had an inadequate response to a conventional agent (such as azathioprine or corticosteroids) after therapy lasting at least three (3) months? _____
- Does the patient have pouchitis? _____
- Has the patient tried an antibiotic, probiotic, corticosteroid enema or mesalamine enema? _____
- Is the patient planning to concurrently receive another biologic? _____

If you answered 'Yes' to any of the above questions, please provide us with comprehensive chart notes regarding those items.

IV Tremfya (guselkumab) Induction Order

- 200mg administered by IV over at least one hour three (3) times, at Weeks 0, 4 and 8

Administer according to manufacturer's instructions. Post infusion flush line. Check vitals and monitor for signs and symptoms of an infusion reaction at start, throughout infusion, and after completion.

Rescue Management in case of Infusion Therapy Reaction

These include fever, chills, rigors, headache, rash, itching, swelling, edema, nausea, vomiting, abdominal pain, hypotension, and respiratory distress.

- Stop medication infusion and start normal saline infusion at 50 ml/hr. Call ordering provider to report reaction.
- Follow standing reaction orders, including Diphenhydramine, Methylprednisolone, Albuterol and oxygen as needed. Famotidine 20mg IVP for minor cutaneous reactions which do not respond to diphenhydramine.
- For severe reactions, administer Epi-pen or equivalent and call 911. Repeat if severe symptoms persist.

Ordering Provider Authorization

Provider Signature: _____ Name: _____ Date: _____

Address: _____

Phone: _____ Fax: _____ Indiv. NPI #: _____ License: _____

Best Contact Person in Office: _____ Direct Phone to Contact Person: _____

FAX THIS ORDER AND SUPPORTING DOCUMENTATION TO 201-266-0437 OR UPLOAD USING YOUR SECURE DEDICATED WEBPAGE – TO GET A PERSONAL LINK PLEASE CONTACT THE INTAKE TEAM.