

(OmvoH IV)

# mirikizumab-mrkz

Infusion orders

Date: \_\_\_\_\_

## PATIENT INFORMATION

Name:	DOB:	SEX: M <input type="checkbox"/> F <input type="checkbox"/>
ICD-10 code (required):	ICD-10 description:	
<input type="checkbox"/> NKDA Allergies:	Weight lbs/kg:	

## REFERRAL STATUS

☐ New Referral ☐ Referral Renewal ☐ Medication/Order Change ☐ Benefits Verification Only ☐ Discontinuation Order

## PHYSICIAN INFORMATION

Referral Coordinator Name:	Referral Coordinator Email:		
Ordering Provider:	Provider NPI:		
Referring Practice Name:	Phone:	Fax:	
Practice Address:	City:	State:	Zip Code:

### DIAGNOSIS (and ICD 10 code)

- ☐ Ulcerative Colitis ICD-10 Code: K51.90  
☐ Other Diagnosis: ICD-10 Code: \_\_\_\_\_

### NOTE

#### List Tried & Failed Therapies, including duration of treatment:

- 1)  
2)

\*\*Referring physician is responsible for monitoring and reviewing the following labs prior to treatment:

- Fasting phosphorus level prior to each dose for first 3 doses and administer only if below ULN
- Fasting phosphorus level 2-4 weeks after dose modifications  
If dose adjustments are needed, new order must be sent by provider based on PI dose calculations

## MIRIKIZUMAB-MRKZ (OmvoH IV) ORDERS

### Medication ordered

OmvoH 300 mg IV at weeks 0 , 4 , 8

### SPECIAL INSTRUCTIONS

### PATIENT WEIGHT

\_\_\_\_\_ lbs.  
\_\_\_\_\_ kg

\*\*Hepatotoxicity in treatment of Crohn's disease. Drug induced liver injury during induction has been reported. Monitor LFT's and bilirubin at baseline and during induction, up to at least 24 weeks of treatment. Monitor thereafter according to routine patient management.

## REQUIRED DOCUMENTATION:

- ☐ This signed order form by the provider  
☐ Patient demographics AND insurance information  
☐ Clinical/Progress notes supporting primary dx  
☐ Confirmed negative TB testing  
☐ LFT and Bilirubin lab results

## ORDERING PROVIDER

Signature X Date \_\_\_\_\_

Provider \_\_\_\_\_ Phone \_\_\_\_\_ Fax \_\_\_\_\_