

OMVOH (MIRIKIZUMAB-MRKZ) ORDERS

P: 877.365.5566 | **F:** 855.889.2946

A Carelon Company	1107710	03.3300 11.033.003.23 10
PATIENT INFORMATION: Fax completed form, insura	nce information,	and clinical documentation to 855.889.2946
Patient Name: Patient Status: □ New to Therapy □ Continuing Therapy	Next Treatr	nent Date:
MEDICAL INFORMATION		
Patient Weight: lbs. (required) Allergies: Diagnosis: Ulcerative Colitis Crohn's Disease Oth ICD-10 Code:		
THERAPY ORDER		
Omvoh (Ulcerative Colitis dosing) IV induction dose: 300mg IV at week 0, 4, and 8 Maintenance dose: 200mg subcutaneously at week 0, 4, and 8 year (to be evaluated by Paragon Specialty Phase)	eek 12, then	every 4 weeks thereafter x 1
Omvoh (Crohn's Disease dosing) IV induction dose: 900mg IV at week 0, 4, and 8	3	
☐ Maintenance dose: 300mg subcutaneously at w year (to be evaluated by Paragon Specialty Pha		every 4 weeks thereafter x 1
Lab Orders: **LFTs and Bilirubin should be monitored at baseline, o	during first 24	 weeks of treatment, and periodically**
Lab frequency: ☐ Prior to 4 and 8 week dose ☐ Other	r:	
Required labs to be drawn by: 🗆 Paragon 🗆 Referring	g Provider	
Other orders:		
Home IV Biologic Ana-kit Orders (adult): • Epinephrine: >30kg (>66lbs): EpiPen 0.3mg or compounded s • Diphenhydramine: Administer 25-50mg orally OR IV (adult) • NS 0.9% 1000mL IV bolus per protocol PRN (adult) Home biologic injection Ana-kit (adult): • Dispense per protocol EpiPen 0.3mg IM (2-pack) Flush orders: NS 1-20mL pre/post infusion PRN and Heparin 10U/mL Supply IV Infusion Pump (E0781) as needed		
PROVIDER INFORMATION		
By signing this form and utilizing our services, you are authorizing Paragon Healthcare, Inc. and its enagent in dealing with medical and prescription insurance companies, and to select the preferred site of Provider Name: Provider NPI: Phone: Fax: Opt out of Paragon selecting site of care (if checked, pleas	of care for the patient	
Provider NP1Priorie rax		

PREFERRED LOCATION

______ State: _____ View our locations here:







COMPREHENSIVE SUPPORT FOR OMVOH THERAPY

A Carelon Company

PATIENT INFORMATION:
Patient Name: DOB:
REQUIRED DOCUMENTATION FOR REFERRAL PROCESSING & INSURANCE APPROVAL
\square Include <u>signed</u> and <u>completed</u> order (MD/prescriber to complete page 1)
\square Include patient demographic information and insurance information
☐ Include patient's medication list
☐ Supporting clinical notes to include any past tried and/or failed therapies, intolerance, benefits, or contraindications to conventional therapy
☐ Does the patient have a contraindication/intolerance or failed trial to corticosterioids or immunomodulators (i.e., 6-MP, azathioprine, budesonide)? ☐ Yes ☐ No If yes, which drug(s)?
□ Does the patient have a contraindication/intolerance or failed trial to any biologic (i.e., Humira, Remicade, Stelara, Cimzia)? □ Yes □ No If yes, which drug(s)?
☐ Include labs and/or test results to support diagnosis
If applicable - Last known biological therapy: and last date received: If patient is switching to biologic therapies, please perform a washout period of weeks prior to starting Omvoh.
Other medical necessity:
REQUIRED PRE-SCREENING
☐ TB screening test completed - attach results ☐ Positive ☐ Negative
☐ Baseline liver function tests and bilirubin - attach results
If TB results are positive - please provide documentation of treatment or medical clearance, and a negative CXR

Paragon Healthcare will complete insurance verification and submit all required documentation for approval to the patient's insurance company for eligibility. Our team will notify you if any additional information is required. We will review financial responsibility with the patient and refer him/her to any available co-pay assistance as needed. Thank you for the referral.

Please fax all information to (855) 889-2946 or call (877) 365-5566 for assistance