Skyrizi IV (Risankizumab-rzaa)

Provider Order Form rev. 8/29/2025

PATIENT INFORMATION



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PATIENT INFORMATION	Referr	al Statu	s: □ New Referr	al 🗆 Updat	ed Order	☐ Order Renewal
Date: Patient Name:				D	OB:	
CD-10 code (required): ICD-10 descrip	tion:					
□ NKDA Allergies:			V	Veight (lbs/kg	<u>;</u>):	Height:
Patient Status: ☐ New to Therapy ☐ Continuing Therapy	Last Treatment Date: Next Due Date:				ate:	
PROVIDER INFORMATION						
Referral Coordinator Name:	Referral Coordinator Email:					
Ordering Provider:	Pro	ovider NI	PI:			
Referring Practice Name:	Phone:		Fax:			
Practice Address:	City:		State:	Zip C	ode:	
NURSING ☐ TB status & date (list results here & attach clinicals) ☐ Baseline Liver Enzymes, including bilirubin (results) ☐ Provide nursing care per IVCare Infusion's ☐ Nursing Procedures, including reaction management and ☐ post- procedure observation PRE-MEDICATION ORDERS ☐ acetaminophen (Tylenol) ☐ 500mg / ☐ 650mg / ☐ 1000mg PO ☐ cetirizine (Zyrtec) 10mg PO ☐ loratadine (Claritin) 10mg PO ☐ diphenhydramine (Benadryl) ☐ 25mg / ☐ 50mg ☐ PO / ☐ IV ☐ methylprednisolone (Solu-Medrol) ☐ 40mg / ☐ 125mg IV ☐ hydrocortisone (Solu-Cortef) ☐ 100mg IV ☐ Other: ☐ Dose: ☐ Route: ☐ Frequency: ☐ Frequency:	□ □ TH I	CMP □ at each dose □ every □ Other: THERAPY ADMINISTRATION Risankizumab-rzaa (Skyrizi) induction IV dose Dose: □ 600 mg / □ 1200 mg Frequency: week 0, week 4, and week 8 Route: Intravenous Infuse over 60 minutes Flush with 0.9% sodium chloride at infusion completion				
Evaluate for TB prior to initiating treatment with SKYRIZI. Hepatotoxicity in Treatment of Crohn's disease: Drug-induced liver injury during in induction, up to at least 12 weeks of treatment. Monitor thereafter according to responsible to the provider Name (Print)	outine pati	ent manag		nzymes and bili	rubin levels a	

Email info@ivcareinfusion.com or fax this Form, Insurance card (both sides), Demographics, Recent H&P, Labs & supporting Clinicals to:

FAX NUMBER FOR SKYRIZI REFERRALS: (615) 471-8674