

Skyrizi (risankizumab-rzaa) Infusion/Injection Orders

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I	Patient Name:	DOB		🗖 Male 🗖 Female		
I	Diagnosis (please provide ICD10 code)					
	New Start Therapy Continuation of Therapy	Date of la	ast d	ose (if applicable):		
[NKDA Allergies:					
C	Ordering Provider:					
P	Provider NPI:	Phone:		Fax:		
F	Practice Address:	City:		State	:	Zip Code:
	PRE-MEDICATION			REQUIRED TESTING	5/LA	BS
	Acetaminophen1000mg PO 🔲 Solu-Medrol 125mg IVP	1	Ø	Clinical/Progress No	otes.	Labs, Tests supporting
	Diphenhydramine25mgI Solu-Cortef 100mg IVPPO Ceterizine 10mg POI Diphenhydramine 25mg		_	primary diagnosis at		
			V	TB status and date (
					produ	
			M	CMP (LFTs and Biliru	ubin	should be monitored
						tion, and periodically)
	SKYRIZI ORDERS			, J		
⊻	Initial Skyrizi Induction Infusion:	1				
	Dilute in 250ml D5W and administer intravenously over	i nour				
	600mg IV Infusion @ week 0, 4, and 8					
	□ Other:					
	*Maintenance dose: 360mg subcutaneously at week 12	, then every	8 we	eks thereafter to be adm	iniste	ered in home setting via OB
	Infuse One Standing Orders:					
	Provide treatment under Infuse One's Clinical Guidelines, Meand Action Plan for Infusion Reactions.	dication Saf	ety P	rotocol, Emergency Gui	deline	es,
	Provider Name					
	Provider Signature			Date		

Serious hypersensitivity reactions, including anaphylaxis, may occur. Skyrizki may increase the risk of infection. Instruct patients to seek medical advice if signs or symptoms of clinically important infection occur. If such an infection develops, do not administer Skyrizi until the infection resolves. Evaluate for TB prior to initiating treatment with Skyrizi. Hepatotoxicity in Treatment of Crohn's Disease: Drug-induced liver injury during induction has been reported. Monitor liver enzymes and bilirubin levels at baseline and, during induction, up to at least 12 weeks of treatment. Monitor thereafter according to routine patient management. Administration of Vaccines: Avoid use of live vaccines.