Rituximab (Rituxan, Ruxience)





PATIENT INFORMATION	Referral Sta	tus: □ New R	eferral 🔲 Updated O	rder 🔲 Order Renewal	
Patient Name:		DOB:	Patient Ph		
Patient Address:	Patient Email:				
Allergies:		□ NKDA	Weight (lbs/kg):	Height (in/cm):	
Sex: ☐ M / ☐ F Date of Last Infu	sion: Next Due		Preferred Location:		
Sex. Li Mi / Li Date of East mile	JIOII. NEXT DUC	Date.	Treferred Education.		
DIAGNOSIS (Please provide ICD-					
Non-Hodgkin's Lymphoma:	Chronic Lymphocytic Le	eukemia:	Rheumatoid Arthri	itis:	
Other:	Description:				
THERAPY ADMINISTRATION ☐ Infuse rituximab (Rituxan) OR ritux by patient's insurance. ☐ Infuse this rituximab product (sub	kimab biosimilar as required	☐ CBC ☐ CMP ☐ CRP ☐ Other:	DRY ORDERS ☐ at each dose ☐ at each dose ☐ at each dose ☐ at each dose ☐ CATION ORDERS	□ every: □ every: □ every:	
 □ Rituximab mg IV □ Rituximab mg/m2 x (Current BSA) m2 = mg (Dose will be rounded up to 10% to nearest 100 mg per protocol unless specified below). □ Dose rounding prohibited. ☑ Doses less than 500mg will go in final volume 250ml ml NS. Doses greater than 500mg will go in final volume 500 ml NS. FREQUENCY □ Infuse on Day 0 and Day 14 □ Infuse on Day 0, Day 7, Day 14, and Day 21 □ Other: □ Repeat dosing in weeks. □ Repeat dosing in months. 		□ Loratadine 10mg PO □ Required Tylenol 500mg PO □ Solumedrol 125mg IV (Required for diagnosis of RA) □ Required Benadryl 25 mg PO □ Other: NURSING □ Hold infusion and notify provider for: • Signs/symptoms of infection, surgical procedures, recent live vaccines, neurological or mood changes. □ Provide nursing care per Nursing Procedure, including Hypersensitivity Reaction Management Protocol and post-procedure observation ADDITIONAL ORDERS			
PROVIDER INFORMATION					
Preferred Contact Name:		Preferred Contact Email:			
Ordering Provider:		Provider NPI:			
Referring Practice Name:		Phone:	Fax:	7in Cada:	
Practice Address:		City:	State:	Zip Code:	
REQUIRED DOCUMENTATION	I CHECKLIST (Additional doc	cumentation req	uired for processing an	d insurance approval)	
Required Documentation: Patient of treatment failures or contraindication Required Labs: Include negative Health Programme Pro	demos, copy of front and back ons, biologic agent and steroid	of primary and s	secondary insurance, 2 red skin (by indication)	most recent OVN including	
Provider Name (print)	Provider Signa	Provider Signature			