

Information on this form is a guide only. User will be solely responsible for verifying its currency and accuracy with the corresponding BC Cancer treatment protocols located at <u>www.bccancer.bc.ca</u> and according to acceptable standards of care

PROTOCOL CODE: LYCLLFBR

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DOCTOR'S ORDERS	Ht	cm Wt	kg BSA_	m²	
REMINDER: Please ensure drug allergi	es and previous b	leomycin are do	cumented on th	e Allergy & Alert Form	
DATE: 1	o be given:		Cycle #:		
Date of Previous Cycle:					
 Delay treatment week(s) CBC & Diff Day 1 of treatment 					
Day 1: may proceed with doses as written, if within 96 hours ANC greater than or equal to 1.0 x 10⁹/L and platelets greater than or equal to 75 x 10 ⁹ /L					
Dose modification for: Image: Hematology Other Toxicity Proceed with treatment based on blood work from					
PREMEDICATIONS: Patient to take ow	/n supply. RN/Pha	rmacist to confirm	l		
ondansetron 8 mg PO prior to bendamustine on Day 1 and Day 2 dexamethasone 2 8 mg or 2 12 mg (select one) PO prior to bendamustine on Day 1 and Day 2					
For intravenous riTUXimab infusion: diphenhydrAMINE 50 mg PO prior to riTUXimab IV and then q 4 h if IV infusion exceeds 4 h acetaminophen 650 mg to 975 mg PO prior to riTUXimab IV and then q 4 h if IV infusion exceeds 4 h For subcutaneous riTUXimab injection: diphenhydrAMINE 50 mg PO prior to riTUXimab subcut acetaminophen 650 mg to 975 mg PO prior to riTUXimab subcutaneous					
** Have Hypers	ensitivity Reactio	n Tray and Prote	col Available**		
TREATMENT: Cycle 1 ONLY	,				
bendamustine 90 mg/m² x BSA = mg Dose Modification:% = mg/m² x BSA = mg IV in 250 to 500 mL NS over 1 hour on Day 1 and Day 2. Continued on page 2					
DOCTOR'S SIGNATURE:				SIGNATURE:	
				UC:	



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DOCTOR'S ORDERS							
DATE:							
** Have Hypersensitivity Reaction Tray and Protocol Available**							
TREATMENT: (0	TREATMENT: (Cycle 1 continued)						
riTUXimab (first dose) 375 mg/m ² x BSA = mg IV in 250 to 500 mL NS on Day 1 or 2 whenever possible, but not later than 72 hours after Day 1 of bendamustine.							
Pharmacy to select	riTUXimab IV brand as per Provincial Systemic The	erapy Policy III-190	_				
Drug	Brand (Pharmacist to complete. Please print.)	Pharmacist Initial and D	Date				
riTUXimab							
Start at 50 mg/hour occurs.	Start at 50 mg/hour. After 1 hour, increase rate by 50 mg/hour every 30 minutes until rate = 400 mg/hour unless toxicity occurs.						
	patients are to be under constant visual observation . Vital signs are not required, unless symptomatic.	during all dose increases	and for 30 minutes after				
Cycle 2, 3, 4, 5	and 6						
bendamustine 90	mg/m² x BSA = mg						
Dose Modific	cation:% = mg/m² x	BSA = mg					
IV in 250 to 500	mL NS over 1 hour on Day 1 and Day 2 .						
riTUXimab for Cycle 2 and subsequent treatments:							
Patient tolerated a full dose of IV riTUXimab (no severe reactions requiring early termination) and can proceed to subcutaneous riTUXimab:							
riTUXimab subcut (RITUXAN SC) 1600 mg (fixed dose in 13.4 mL) subcutaneously into abdomen over 7 minutes. Observe for 15 minutes after administration.							
NB: During treatment with subcutaneous riTUXimab, administer other subcutaneous drugs at alternative injection sites whenever possible.							
Continued on Page 3							
DOCTOR'S SIGI	NATURE:		SIGNATURE:				
			UC:				



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DOCTOR'S ORDERS						
DATE:						
** Have Hypersensitivity Reaction Tray and Protocol Available**						
TREATMENT: (Cycle 2, 3, 4, 5 and 6 continued)						
 Patient did not tolerate a full dose of IV riTUXimab (experienced severe reactions requiring early termination) in the previous treatment and will continue with IV riTUXimab for this cycle: riTUXimab 500 mg/m² x BSA = mg IV in 250 to 500 mL NS on Day 1 or 2 whenever possible, but not later than 72 hours after Day 1 of bendamustine. 						
	hab IV brand as per Provincial Systemic The		Note 1			
	(Pharmacist to complete. Please print.)	Pharmacist Initial and D	Jate			
riTUXimab						
Infuse 50 mL (or 100 mL of 500 mL bag) of the dose over 30 minutes, then infuse the remaining 200 mL (or 400 mL of 500 mL bag) over 1 hour. (total infusion time = 1 hour 30 min) If flushing, dyspnea, rigors, rash, pruritus, vomiting, chest pain, any other new acute discomfort or exacerbation of any existing symptoms occur, stop infusion and page physician. Constant visual observation is not required.						
	RETURN APPOINTMENT	ORDERS				
 Return in <u>four</u> weeks for Doctor and Cycle Book chemo on Day 1 and Day 2. Note: riTUXimab to be booked within 72 hours of bendamustine. Last Cycle. Return in week(s). 						
CBC & Diff prior to Day 1 of	of each cycle					
If clinically indicated:						
Creatinine ALT total bilirubin						
HBV viral load every 3 r						
Other tests:						
Consults:						
See general orders sheet for additional requests.						
DOCTOR'S SIGNATURE	E:		SIGNATURE:			

UC: