

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: LYCLLBENDR

(Page 1 of 3)

DOCTOR'S ORDERS	Ht	cm	Wt	kg	BSA	m²
REMINDER: Please ensure drug allerg	ies and previou	s bleomyc	in are doc	umented	on the All	ergy & Alert Form
DATE:	To be given:			Сус	le #:	
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 10⁹/L 						
Dose modification for: Hematology	y 🗌 Othe	r Toxicity				
Proceed with treatment based on blood	d work from					
PREMEDICATIONS: Patient to take or	wn supply. RN/P	harmacist	to confirm _.			·
ondansetron 8 mg PO prior to bendamustine on Day 1 and Day 2 dexamethasone						
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 subcutaneous acetaminophen 650 mg to 975 mg PO prior to riTUXimab subcutaneous						
TREATMENT: Cycle 1 ONLY						
bendamustine 70 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	2.				
Continued on page 2						
DOCTOR'S SIGNATURE:					SIGN	IATURE:
					UC:	



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: LYCLLBENDR

(Page 2 of 3)

DOCTOR'S ORDERS						
DATE:						
** Have Hypersensitivity Reaction Tray and Protocol Available**						
TREATMENT: (Cycle 1 continued):						
riTUXimab (first dose) 375 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.						
Pharmacy to select	Pharmacy to select riTUXimab IV brand as per Provincial Systemic Therapy Policy III-190					
Drug	Brand (Pharmacist to complete. Please print.) Pharmacist	Initial and Date				
riTUXimab						
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 during all dos . Vital signs are not required, unless symptomatic.	e increases and for 30 minutes after				
TREATMENT: C	ycles 2, 3, 4, 5 and 6					
	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.					
	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 SIGN	NATURE:	SIGNATURE:				
		UC:				



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: LYCLLBENDR

(Page 3 of 3)

DOCTOR'S ORDERS

DATE:

** Have Hypersensitivity Reaction Tray and Protocol Available**

TREATMENT: (Cycles 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.

Pharmacy to select riTUXimab IV brand as per Provincial Systemic Therapy Policy III-190

Drug	Brand (Pharmacist to complete. Please print.)	Pharmacist Initial and Date
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 each cycle	
If clinically indicated:	
🗌 creatinine 🔲 ALT 🔄 total bilirubin	
HBV viral load every 3 months	
Other tests:	
Consults:	
See general orders sheet for additional requests.	
DOCTOR'S SIGNATURE:	SIGNATURE:
	UC: