# Regimen Reference Order – LYMP – R-CHOP

ARIA: LYMP - [R-CHOP]

LYMP – [R-CHOP (Split Day SLOW on Cycle 1)]

Planned Course: Every 21 days for 6 cycles Indication for Use: Non-Hodgkin Lymphoma,

OR

Hodgkin Lymphoma, Nodular lymphocytic predominant

**CVAD: At Provider's Discretion (VESICANT INVOLVED)** 

## Proceed with treatment if:

ANC equal to or greater than  $0.8 \times 10^9/L$  AND Platelets equal to or greater than  $100 \times 10^9/L$ 

Contact Hematologist if parameters not met

**Note:** Hepatitis B serology results must be reviewed in accordance with CCMB Policy Hepatitis B Monitoring for Oncology and Hematology Patients

## **SEQUENCE OF MEDICATION ADMINISTRATION**

Pre-treatment Requirements				
Drug	Dose	CCMB Administration Guideline		
Instruct patient to start vigorous oral pre-hydration (600-900 mL) the morning of cyclophosphamide treatment (Self-administered at home)				
allopurinol	300 mg	Orally once daily for 10 days to begin 3 days prior to Cycle 1 (Self-administered at home)		
		Only patients at risk of tumor lysis syndrome will be prescribed allopurinol		
		Note: allopurinol should not be prescribed beyond 10 days unless under the direction of the hematologist. See Additional Information		

Treatment Regimen – LYMP – R-CHOP  Establish primary solution 500 mL of: normal saline				
Cycle 1				
Day 1				
predniSONE	100 mg	Orally once in the morning with food (Self-administered at home) predniSONE is started on Day 1 regardless if R-CHOP is given over one day or split over two days		
cetirizine	10 mg	Orally 30 minutes prior to riTUXimab		
acetaminophen	650 mg	Orally 30 minutes prior to riTUXimab		
dexamethasone*	20 mg	IV in normal saline 50 mL over 15 minutes		

ADULT LYMP – R-CHOP

riTUXimab (IV brand	375 mg/m <sup>2</sup>	Slow infusion (if greater than 6 months since last riTUXimab dose
name specific)	O.	or no previous riTUXimab): IV made up to a final concentration of
		mg/mL in normal saline. Start at 50 mg/hr for 60 minutes and
		escalate infusion rate in 50 mg/hr increments every 30 minutes to
		maximum of 400 mg/hr
		*Nursing Alert: IV tubing is primed with riTUXimab
		*Alert: Ensure brand name on prescription label (indicated in brackets on prescription label) matches prescribed order
		*Alert: Pharmacy to ensure final volume on label
		OR
		Slow infusion (if equal to or less than 6 months since last
		riTUXimab dose): IV made up to a final concentration of 1 mg/mL
		in normal saline. Start at 100 mg/hr for 30 minutes and escalate
		infusion rate in 100 mg/hr increments every 30 minutes to a
		maximum of 400 mg/hr
		*Nursing Alert: IV tubing is primed with riTUXimab
		*Alert: Ensure brand name on prescription label (indicated in brackets on prescription label) matches prescribed order
		*Alert: Pharmacy to ensure final volume on label
ondansetron*	16 mg	
	16 mg	Orally 30 minutes pre-chemotherapy
DOXOrubicin	50 mg/m <sup>2</sup>	IV Push over 10 to 15 minutes
vinCRIStine	1.4 mg/m²; maximum dose 2 mg	IV in normal saline 25 mL over 2 to 3 minutes by gravity infusion
cyclophosphamide	750 mg/m <sup>2</sup>	IV in normal saline 250 mL over 1 hour
Days 2, 3, 4 and 5		
predniSONE	100 mg	Orally once daily in the morning with food
		(Self-administered at home)
*If R-CHOP is split ove	r two days, give dexamet	hasone 12 mg and ondansetron prior to CHOP on Day 2
Cycle 2 and Onward	S	
Day 1		
predniSONE	100 mg	Orally once in the morning with food
		(Self-administered at home)
cetirizine	10 mg	Orally 30 minutes prior to riTUXimab
acetaminophen	650 mg	Orally 30 minutes prior to riTUXimab
dexamethasone	12 mg	IV in normal saline 50 mL over 15 minutes
Wait 30 minutes after	completion of IV pre-me	dication(s) before starting riTUXimab
riTUXimab	1400 mg	<u>Subcutaneous</u> : Administer over 5 minutes into abdomen
	(1400 mg = 11.7 mL)	Syringe should be held in hand for 5 minutes to warm up and
(Subcutaneous)	(1400 mg = 11.7 ml)	1
(Subcutaneous)	(1400 mg – 11.7 mc)	decrease viscosity Use 25G needle
(Subcutaneous)	(1400 ing – 11.7 inc)	decrease viscosity



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		OR
riTUXimab (IV brand name specific)	375 mg/m <sup>2</sup>	Rapid infusion: IV in normal saline over 90 minutes: Infuse 50 mL of a 250 mL bag (or 100 mL of a 500 mL bag) over 30 minutes, then infuse the remaining 200 mL (or 400 mL of a 500 mL bag) over 60 minutes  *Alert: Ensure brand name on prescription label (indicated in brackets on prescription label) matches prescribed order  *Alert: Pharmacy to ensure final volume on label
		OR
		Slow infusion: IV made up to a final concentration of 1 mg/mL in normal saline. Start at 100 mg/hr for 30 minutes and escalate infusion rate in 100 mg/hr increments every 30 minutes to a maximum of 400 mg/hr  *Nursing Alert: IV tubing is primed with riTUXimab  *Alert: Ensure brand name on prescription label (indicated in brackets on prescription label) matches prescribed order  *Alert: Pharmacy to ensure final volume on label
ondansetron	16 mg	Orally 30 minutes pre-chemotherapy
DOXOrubicin	50 mg/m <sup>2</sup>	IV Push over 10 to 15 minutes
vinCRIStine	1.4 mg/m²; maximum dose 2 mg	IV in normal saline 25 mL over 2 to 3 minutes by gravity infusion
cyclophosphamide	750 mg/m <sup>2</sup>	IV in normal saline 250 mL over 1 hour
Days 2, 3, 4 and 5		
predniSONE	100 mg	Orally once daily in the morning with food (Self-administered at home)

In the event of an infusion-related hypersensitivity reaction, refer to the 'Hypersensitivity Reaction Standing Order'

## **REQUIRED MONITORING**

#### Hepatitis B serology

• Hepatitis B surface antigen and Hepatitis B core antibody (drawn within preceding 5 years)

### Cardiac Monitoring

• Left Ventricular Ejection Fraction (LVEF) monitoring is recommended at baseline and as clinically indicated

## All Cycles

• CBC, serum creatinine, urea, electrolytes, liver enzymes, LDH, total bilirubin, uric acid and albumin as per Physician Orders

#### **INTRAVENOUS** riTUXimab

- Full vital signs (temperature, heart rate, respiratory rate, blood pressure and O<sub>2</sub> saturation) at baseline and as clinically indicated
- No observation period is required. Patient can be discharged from treatment room if stable whether they had a reaction or not



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#### SUBCUTANEOUS riTUXimab

• Full vital signs (temperature, heart rate, respiratory rate, blood pressure and O₂ saturation) at baseline, <u>at discharge</u> and as clinically indicated

• 15-minute observation period required after each dose

Recommended Support Medications				
Drug	Dose	CCMB Administration Guideline		
filgrastim (brand name specific)	5 mcg/kg (round to nearest 300 mcg or 480 mcg)	ONLY to be given if patient eligible for Growth Factor Support (refer to CCMB Drug Formulary Web App for Primary Prophylaxis eligibility criteria) Subcutaneous once daily for 5 days to start on Day 3		
metoclopramide	10 – 20 mg	Orally every 4 hours as needed for nausea and vomiting		

### **DISCHARGE INSTRUCTIONS**

- Patients should be instructed to contact their cancer team immediately if symptoms of hypersensitivity reactions occur after discharge
- · For eligible patients, ensure patient receives filgrastim supply if patient is self-administering at home
- Instruct patient to:
  - Continue taking anti-emetic(s) at home
  - o Maintain oral intake of 2000 mL (8 glasses) of fluid daily at home
  - Empty bladder every 2 hours while awake and at bedtime for 24 hours after each dose of cyclophosphamide
  - o Obtain immediate assistance as per your clinic's contact instructions if:
    - Symptoms of hemorrhagic cystitis (e.g. dysuria, hematuria)
    - Unable to drink recommended amount of fluid
- predniSONE is a cancer therapy in this treatment regimen. Remind patient to take predniSONE at home
- Reinforce applicable safe handling precautions of medications, blood and body fluids for 48 hours after completion of chemotherapy

## **ADDITIONAL INFORMATION**

- Cumulative DOXOrubicin dose should be calculated and should not exceed 450 mg/m<sup>2</sup>
- Due to the risk of reactivation of Hepatitis B virus (HBV) while on this treatment regimen, prescriber must adhere to CCMB Policy *Hepatitis B Monitoring for Oncology and Hematology Patients* for ordering and interpreting HBV serology and prescribing antiviral prophylaxis
- Unless patient was taking allopurinol for gout or other reasons unrelated to the patient's underlying lymphoma, allopurinol should not be prescribed with cycle 2 and onwards unless directed by hematologist
- Administering nurse must document any infusion-related reactions with any dose of riTUXimab
- Ensure there were **no Grade 3 or 4** infusion-related reactions with any previous dose prior to administering riTUXimab via subcutaneous injection or rapid infusion
- Intravenous riTUXimab formulation is available from more than one manufacturer and uses several different brand names. Brand name will be indicated in brackets after riTUXimab. Ensure prescription label matches the brand name on prescribed order for intravenous riTUXimab
- Note: At Cycle 1, an entry called "Physician Reminder- Growth Factor 60 y.o." will appear in the electronic drug order. This prompt is to remind the prescriber to order filgrastim for eligible patients

