ADULT Updated: February 3, 2022

Regimen Reference Order – LYMP – oBINutuzumab maintenance

Planned Course: Every 8 weeks for 12 doses (2 years)

Indication for Use: Non-Hodgkin Lymphoma

CVAD: At Provider's Discretion

Proceed with treatment if:

ANC equal to or greater than $1 \times 10^9/L$ AND Platelets equal to or greater than $75 \times 10^9/L$

Contact Hematologist if parameters not met

SEQUENCE OF MEDICATION ADMINISTRATION

	Pre-treatment Requirements					
	Drug	Dose	CCMB Administration Guideline			
None Required						

Establish primary solution 500 mL of: normal saline				
Drug	Dose	CCMB Administration Guideline		
dexamethasone	20 mg	<u>ONLY</u> to be given if patient had a grade 3 or 4 infusion-related reaction with their previous oBINutuzumab infusion or if their lymphocyte count is greater than $25 \times 10^9/L$		
		IV in normal saline 50 mL over 15 minutes <u>1 hour</u> prior to oBINutuzumab		
		*Nursing Alert: oBINutuzumab starts 1 hour after completion of dexamethasone infusion		
acetaminophen	650 mg	Orally 30 minutes prior to oBINutuzumab		
cetirizine	10 mg	Orally 30 minutes prior to oBINutuzumab		
oBINutuzumab	1000 mg	Rapid Infusion: IV in normal saline 250 mL following administration rates below:		
		0 to 30 minutes – 25 mL/hour		
		• 30 to 93 minutes – 225 mL/hour		
		*Alert: Pharmacy to ensure final volume in bag = 250 mL (4 mg/mL final concentration)		
		*Nursing Alert: IV tubing is primed with oBINutuzumab		
		OR		
		Slow Infusion: IV in normal saline 250 mL following administration rates below:		
		0 to 30 minutes – 25 mL/hour		
		• 30 to 60 minutes – 50 mL/hour		
		• 60 to 90 minutes – 75 mL/hour		



90 minutes onwards – 100 mL/hour
*Alert: Pharmacy to ensure final volume in bag = 250 mL (4 mg/mL final concentration) *Nursing Alert: IV tubing is primed with oBINutuzumab

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

REQUIRED MONITORING

All Cycles

- CBC as per Physician Orders
- Full vital signs (temperature, heart rate, respiratory rate, blood pressure and O₂ saturation) at baseline and as clinically indicated
- No observation period is required after oBINutuzumab administration. Patient can be discharged from treatment room if stable whether they had a reaction or not

	Recommended Support Medications					
	Drug	Dose	CCMB Administration Guideline			
None Required						

DISCHARGE INSTRUCTIONS

 Patients should be instructed to contact their cancer team immediately if symptoms of hypersensitivity reactions occur after discharge

ADDITIONAL INFORMATION

- · Administering nurse must document any infusion-related reactions with any dose of oBINutuzumab
- Ensure there were **no Grade 3 or 4** infusion-related reactions with the three preceding infusions and that lymphocyte count is less than 5 x 10⁹/L prior to administering oBlNutuzumab via rapid infusion
- Note: an entry called "Physician Reminder oBINutuzumab infusion time 1 Units Insert Miscellaneous once" will appear in the electronic drug order. No action is required. This prompt is to remind the prescriber to confirm that patient is eligible for oBINutuzumab rapid infusion

