# Regimen Reference Order – LYMP – oBINutuzumab maintenance

To order this therapy in ARIA, refer to ADDITIONAL INFORMATION

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

**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		
Not Applicable				

Treatment Regimen – LYMP – oBINutuzumab maintenance				
Establish primary solution 500 mL of: normal saline				
Drug	Dose	CCMB Administration Guideline		
cetirizine	10 mg	Orally 30 minutes prior to oBINutuzumab		
acetaminophen	650 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**

#### Hepatitis B serology

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

#### All Cycles

- CBC as per Physician Orders
- 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 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**

- 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
- · 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  $5 \times 10^9$ /L prior to administering oBINutuzumab 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
- ARIA ordering: Support protocols are available under oBINutuzumab maint. in the "Lymphoma" folder

