

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 <a href="https://www.bccancer.bc.ca">www.bccancer.bc.ca</a> and according to acceptable standards of care

## PROTOCOL CODE: LYVENOB

(Ramp-up phase: Low or Medium TLS Risk venetoclax PLUS oBINutuzumab combination therapy - Cycle 2)

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DOCTOR'S ORDERS W	tkg	
REMINDER: Please ensure drug allergies and previous bleomycin are documented on the Allergy & Alert Form		
DATE: Start date of dose ramp-up (must be on a Thursday):	Cycle # 2	
Date of previous cycle:		
Weeks 1 to 5 - Outpatient		
☐ Delay treatment week(s)		
May proceed with doses as written if lab work is within 72 h of venetoclax initiation: ANC <u>greater than or equal to</u> 1.0 x 10 <sup>9</sup> /L, platelets <u>greater than or equal to</u> 25 x 10 <sup>9</sup> /L, total bilirubin <u>less than or equal to</u> 3 x ULN		
Day 8: May proceed with oBINutuzumab as written if within 72 hours <b>ANC</b> greater than or equal to 1.0 x 10 <sup>9</sup> /L, platelets greater than or equal to 25 x 10 <sup>9</sup> /L		
Dose modification for:		
Proceed with treatment based on blood work from		
Tumor Lysis Prophylaxis: Patient to take own supply. RN/Pharmacist to confirm	 Day 1)	
TREATMENT:		
Note: Week 1 starts on Day 1 of Cycle 2 (on a Thursday)		
Week 1: venetoclax 20 mg (2 x 10 mg) PO once daily for 7 days Week 2: venetoclax 50 mg (1 x 50 mg) PO once daily for 7 days Week 3: venetoclax 100 mg (1 x 100 mg) PO once daily for 7 days Week 4: venetoclax 200 mg (2 x 100 mg) PO once daily for 7 days **DO NOT take Day 2 dose on weeks 1 and 2, until approval received**  **DO NOT start weekly dose increase, until approval received**  AND		
Week 5: <b>venetoclax 400 mg</b> (4 x 100 mg) PO once daily for 7 days		
**DO NOT start dose increase, until approval received**		
venetoclax mg PO once daily for days (to last until next dose ramp up to	start on a Thursday)	
OR		
☐ Dose modifications: venetoclax mg PO once daily. Start on	(enter date)	
Mitte: weeks		
DOCTOR'S SIGNATURE:	SIGNATURE: UC:	



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DATE:	
**Have Hypersensitivity Reaction Tray and Protocol Available**	
PREMEDICATIONS FOR oBINutuzumab INFUSION:  Patient to take own supply of oral medications. RN/Pharmacist to confirm:  ☐ If previous reaction was Grade 3, or if lymphocyte count greater than 25 x 10 <sup>9</sup> /L in labs of 60 minutes prior to infusion: dexamethasone 20 mg IV  30 minutes prior to infusion: acetaminophen to 650 to 975 mg PO and diphenhydrAMINE	
TREATMENT:	<u> </u>
Note: Day 8 must be on a Thursday	
oBINutuzumab 1000 mg IV in 250 mL NS on Day 8.	
If no infusion reaction or only Grade 1 infusion reaction in the previous infusion and final infusion at 100 mg/h. Increase by 100 mg/h every 30 minutes until rate = 400 mg/h unless toxi appendix for oBINutuzumab infusion rate titration table.	
RETURN APPOINTMENT ORDERS	
Return in <u>five</u> weeks for Doctor and Cycle 3. Book treatment on Day 1.	
**ALL LABS FROM WEEKS 1 TO 5 MUST BE ORDERED AS <u>STAT</u> AT A LABORATOR' TURNAROUND TIME (e.g. BC Cancer or hospital laboratory)**	Y WITH RAPID
CBC & Diff on Day 7 of weeks 1, 2, 3, and 4	
Ramp up labs: potassium, calcium, phosphate, uric acid, creatinine, LDH, albumin on the following days and times:	
Note: Day 7 labs must be on a Wednesday  Week 1 Day 1 at 12 noon  Week 1 Day 2 at 8 am  Week 1 Day 7 before 12 noon  Week 2 Day 1 at 12 noon (drawn during oBINutuzumab infusion if applicable)  Week 2 Day 2 at 8 am  Week 2 Day 7 before 12 noon  Week 3 Day 7 before 12 noon  Week 4 Day 7 before 12 noon	
Telephone nursing assessment on Day 6 of weeks 1, 2, 3 and 4	
Pharmacy booking as per centre specific standard on the following days: Week 1 and Week 2: Days 1, 2 and 7 Week 3 and Week 4: Day 7	
Prior to next cycle: CBC & Diff, creatinine, total bilirubin, ALT	
If clinically indicated:    HBV viral load     Other tests:   Consults:   See general orders sheet for additional requests	
DOCTOR'S SIGNATURE:	SIGNATURE:
	UC: