# **Regimen Reference Order** – LYMP – brentuximab vedotin + CHP

ARIA: LYMP - [brentuximab + CHP]

Planned Course:Every 21 days for 6 cyclesIndication for Use:T Cell Lymphoma

CVAD: At Provider's Discretion (VESICANT INVOLVED)

## Proceed with treatment if:

ANC equal to or greater than  $1 \times 10^9/L$  AND Platelets equal to or greater than  $50 \times 10^9/L$ 

Contact Hematologist if parameters not met

# SEQUENCE OF MEDICATION ADMINISTRATION

Pre-treatment Requirements				
Drug	Dose	CCMB Administration Guideline		
Instruct patient to st (Self-administered a	• • •	n (600-900 mL) the morning of cyclophosphamide treatment		
allopurinol*	300 mg	Orally once daily for 10 days to begin 3 days prior to Cycle 1 and at provider's discretion for subsequent cycles		
		(Self-administered at home)		
		* Only patients at risk of tumor lysis syndrome will be prescribed allopurinol		

Treatment Regimen – LYMP – brentuximab vedotin + CHP					
Establish primary solution 500 mL of: normal saline					
Drug	Dose	CCMB Administration Guideline			
Day 1					
predniSONE	100 mg	Orally once in the morning with food (Self-administered at home)			
ondansetron	16 mg	Orally 30 minutes pre-chemotherapy			
dexamethasone	12 mg	Orally 30 minutes pre-chemotherapy			
DOXOrubicin	50 mg/m <sup>2</sup>	IV Push over 10 to 15 minutes			
cyclophosphamide	750 mg/m <sup>2</sup>	IV in normal saline 250 mL over 1 hour			
cetirizine	10 mg	Orally 30 minutes prior to brentuximab vedotin			
acetaminophen	650 mg	Orally 30 minutes prior to brentuximab vedotin			
brentuximab vedotin	1.8 mg/kg; maximum dose 180 mg	IV in normal saline 100 mL over 30 minutes			



Days 2, 3, 4 and 5				
predniSONE	100 mg	Orally once daily in the morning with food (Self-administered at home)		
All doses will be automation	tically rounded that fall with	in CCMB Approved Dose Bands. See Dose Banding document for		

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

### **REQUIRED MONITORING**

#### 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
- Assess patient for neuropathy prior to every cycle
- Full vital signs (temperature, heart rate, respiratory rate, blood pressure and O<sub>2</sub> saturation) prior to each dose of brentuximab vedotin and as clinically indicated
- No observation period is required after brentuximab vedotin 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		
filgrastim (brand name specific) (See Filgrastim Clinical Guide)	5 mcg/kg (rounded to nearest 300 mcg or 480 mcg)	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
- Ensure patient receives filgrastim supply if patient is self-administering at home
- Instruct patient to:

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- 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
  - 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
- Avoid grapefruit and grapefruit juice, Seville oranges (i.e. orange marmalade), and starfruit
- Reinforce applicable safe handling precautions of medications, blood and body fluids for 48 hours after completion of chemotherapy



# **ADDITIONAL INFORMATION**

- brentuximab vedotin must be the last medication administered on Day 1
- Cumulative DOXOrubicin dose should be calculated and should not exceed 450 mg/m<sup>2</sup>
- brentuximab vedotin can cause peripheral neuropathy

