

## Regimen Reference Order – LYMP – mogamulizumab

ARIA: LYMP - [mogamulizumab]

**Planned Course:** Until disease progression or unacceptable toxicity (1 cycle = 28 days)

**Indication for Use:** Mycosis fungoides and Sezary syndrome

**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  $50 \times 10^9/L$**

❖ **Contact Hematologist if parameters not met**

### SEQUENCE OF MEDICATION ADMINISTRATION

#### Pre-treatment Requirements

Drug	Dose	CCMB Administration Guideline
Not Applicable		

### Treatment Regimen – LYMP – mogamulizumab

Establish primary solution 500 mL of: normal saline

Drug	Dose	CCMB Administration Guideline
<b>Cycle 1</b>		
<b>Days 1, 8, 15 and 22</b>		
cetirizine	20 mg	Orally 30 minutes prior to mogamulizumab
acetaminophen	650 mg	Orally 30 minutes prior to mogamulizumab
dexamethasone	20 mg	<p><b><u>Days 1 and 8 only:</u></b>                      IV in normal saline 50 mL over 15 minutes <b>1 hour</b> prior to mogamulizumab  <i>*Nursing Alert: mogamulizumab starts 1 hour after completion of dexamethasone infusion</i></p> <p><b><u>Days 15 and 22:</u></b>  <b><i>ONLY</i></b> to be given if patient had a grade 3 or 4 infusion-related reaction with their previous mogamulizumab infusion                      IV in normal saline 50 mL over 15 minutes <b>1 hour</b> prior to mogamulizumab  <i>*Nursing Alert: mogamulizumab starts 1 hour after completion of dexamethasone infusion</i></p>

**If applicable, wait 1 hour after completion of IV pre-medication(s) before starting mogamulizumab**

mogamulizumab	1 mg/kg	<p><b>Day 1:</b> IV in normal saline 250 mL over 2 hours Use 0.2 or 0.22 micron filter</p> <p><b>Days 8, 15 and 22:</b> IV in normal saline 250 mL over 1 to 2 hours Use 0.2 or 0.22 micron filter <i>*Nursing Alert: mogamulizumab should be infused over 2 hours if patient experienced an infusion-related reaction with their previous mogamulizumab infusion</i></p>
<b>Cycle 2 and Onwards</b>		
<b>Days 1 and 15</b>		
cetirizine	20 mg	Orally 30 minutes prior to mogamulizumab
acetaminophen	650 mg	Orally 30 minutes prior to mogamulizumab
mogamulizumab	1 mg/kg	<p>IV in normal saline 250 mL over 1 to 2 hours Use 0.2 or 0.22 micron filter <i>*Nursing Alert: mogamulizumab should be infused over 2 hours if patient experienced an infusion-related reaction with their previous mogamulizumab infusion</i></p>
All doses will be automatically rounded that fall within CCMB Approved Dose Bands. See Dose Banding document for more information		

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

## REQUIRED MONITORING

### Cycle 1

#### Days 1 and 15

- CBC, serum creatinine, urea, electrolytes, liver enzymes, LDH, total bilirubin, uric acid, albumin, glucose and TSH 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
- Cycle 1, Day 1 ONLY: Observe patient for 30 minutes after mogamulizumab infusion (first dose). Full vital signs prior to discharge

#### Days 8 and 22

- No blood work required
- 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 required. Patient can be discharged from treatment room if stable whether they had a reaction or not

## Cycle 2 and Onwards

## Days 1 and 15

- CBC, serum creatinine, urea, electrolytes, liver enzymes, LDH, total bilirubin, uric acid, albumin, glucose and TSH 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 required. 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
- Patients should report rash to clinic

### ADDITIONAL INFORMATION

- mogamulizumab can cause severe skin rash
- Serious immune-mediated complications and infections have been reported with mogamulizumab