

Regimen Reference Order – riTUXimab desensitization

riTUXimab desensitization protocol is prescribed in combination with a riTUXimab-based protocol

- ARIA: LYMP - [riTUX desens. q7d]
- LYMP - [riTUX desens. q21d]
- LYMP - [riTUX desens. q28d]
- LYMP - [riTUX desens. maintenance]

Planned Course: Refer to prescribed riTUXimab-based protocol
Indication for Use: Eligible patients with previous hypersensitivity reactions to riTUXimab
Alert: Desensitization protocol

riTUXimab:

- *riTUXimab is prepared to a final concentration of 1 mg/mL by Pharmacy*
- *riTUXimab must be the first chemotherapy agent administered when given in combination with another chemotherapy agent*
- *IV tubing is primed with riTUXimab*
- *riTUXimab is administered slowly following specified rate increases*
- *riTUXimab administration duration depends upon total dose. riTUXimab infusion can take up to 7 hours to complete*

CVAD: At Provider’s Discretion

Proceed with treatment if:

- ❖ *Refer to prescribed riTUXimab-based protocol*

SEQUENCE OF MEDICATION ADMINISTRATION

| Pre-treatment Requirements | | |
|---|------|-------------------------------|
| Drug | Dose | CCMB Administration Guideline |
| <i>Refer to prescribed riTUXimab-based protocol</i> | | |

| Treatment Regimen – riTUXimab desensitization | | |
|---|--------|---|
| Establish primary solution 500 mL of: normal saline | | |
| Drug | Dose | CCMB Administration Guideline |
| cetirizine | 20 mg | Orally 1 hour prior to riTUXimab |
| acetaminophen | 650 mg | Orally 1 hour prior to riTUXimab |

| | | |
|---|-------------------------------|--|
| dexamethasone | 20 to 40 mg | IV in normal saline 50 mL over 15 minutes 1 hour prior to rituximab <i>*Nursing Alert: ritUXimab starts 1 hour after completion of dexamethasone infusion</i> |
| famotidine | 20 mg | IV in normal saline 50 mL over 15 minutes 45 minutes prior to ritUXimab |
| Wait 45 minutes after completion of IV pre-medication(s) before starting ritUXimab | | |
| riTUXimab (brand name specific) | Dose as specified in protocol | IV in normal saline made up to a final concentration of 1 mg/mL following the administration rates below: Step 1: 2 mL/hour for 15 minutes, then Step 2: 4 mL/hour 15 minutes, then Step 3: 6 mL/hour for 15 minutes, then Step 4: 8 mL/hour for 15 minutes, then Step 5: 10 mL/hour 15 minutes, then Step 6: 15 mL/hour for 15 minutes, then Step 7: 30 mL/hour for 15 minutes, then Step 8: 60 mL/hour for 15 minutes, then Step 9: 80 mL/hour for 15 minutes, then Step 10: 100 mL/hour for 15 minutes, then Step 11: 120 mL/hour for 15 minutes, then Step 12: 140 mL/hour for 15 minutes, then Step 13: 160 mL/hour for 15 minutes, then Step 14: 180 mL/hour for 15 minutes, then Step 15: 200 mL/hour until infusion is complete <i>*Alert: Pharmacy to ensure final volume on label</i> <i>*Alert: ritUXimab must be the first agent administered when given in combination with another chemotherapy agent</i> <i>*Nursing Alert: IV tubing is primed with ritUXimab</i> <i>*Alert: Ensure brand name on prescription label (indicated in brackets on prescription label) matches prescribed order</i> <i>*Nursing Alert: There is no interruption in ritUXimab infusion unless patient is experiencing an infusion-related reaction</i> |

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

REQUIRED MONITORING

- Full vital signs (temperature, heart rate, respiratory rate, blood pressure and O₂ saturation) prior to each ritUXimab infusion and as clinically indicated
- No observation period is required. Patient can be discharged from treatment room if stable whether they have had a reaction or not
- Refer to the prescribed ritUXimab-based protocol for additional monitoring

Recommended Support Medications

| Drug | Dose | CCMB Administration Guideline |
|---|------|-------------------------------|
| <i>Refer to the prescribed ritUXimab-based protocol</i> | | |

DISCHARGE INSTRUCTIONS

- Patients should be instructed to contact their cancer team immediately if symptoms of hypersensitivity reactions occur after discharge
- Refer to the prescribed ritUXimab-based protocol for additional discharge instructions

ADDITIONAL INFORMATION

- Once the patient requires ritUXimab desensitization protocol, all subsequent **ritUXimab doses must be given using ritUXimab desensitization protocol**
- Hematologist must write first prescription of ritUXimab desensitization protocol
- Administering nurse must document any infusion-related reactions with any dose of ritUXimab
- Refer to the prescribed ritUXimab-based protocol for additional ritUXimab information
- Due to the duration of treatment, administration site restrictions may be in place
- Intravenous ritUXimab is available from more than one manufacturer and uses several different brand names. Brand name will be indicated in brackets after ritUXimab. **Ensure prescription label matches the brand name on prescribed order**
- Support protocols are available under **ritUX desens. q7d, ritUX desens. q21d, ritUX desens. q28d and ritUX desens. maint.** in the “Lymphoma” folder