

Regimen Reference Order – riTUXimab desensitization

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

To order this therapy in ARIA, refer to Additional Information below

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	<p>IV in normal saline made up to a final concentration of 1 mg/mL following the administration rates below:</p> <p>Step 1: 2 mL/hour for 15 minutes, then</p> <p>Step 2: 4 mL/hour 15 minutes, then</p> <p>Step 3: 6 mL/hour for 15 minutes, then</p> <p>Step 4: 8 mL/hour for 15 minutes, then</p> <p>Step 5: 10 mL/hour 15 minutes, then</p> <p>Step 6: 15 mL/hour for 15 minutes, then</p> <p>Step 7: 30 mL/hour for 15 minutes, then</p> <p>Step 8: 60 mL/hour for 15 minutes, then</p> <p>Step 9: 80 mL/hour for 15 minutes, then</p> <p>Step 10: 100 mL/hour for 15 minutes, then</p> <p>Step 11: 120 mL/hour for 15 minutes, then</p> <p>Step 12: 140 mL/hour for 15 minutes, then</p> <p>Step 13: 160 mL/hour for 15 minutes, then</p> <p>Step 14: 180 mL/hour for 15 minutes, then</p> <p>Step 15: 200 mL/hour until infusion is complete</p> <p><i>*Alert: Pharmacy to ensure final volume on label</i></p> <p><i>*Alert: riTUXimab must be the first agent administered when given in combination with another chemotherapy agent</i></p> <p><i>*Nursing Alert: IV tubing is primed with riTUXimab</i></p> <p><i>*Alert: Ensure brand name on prescription label (indicated in brackets on prescription label) matches prescribed order</i></p> <p><i>*Nursing Alert: There is no interruption in riTUXimab infusion unless patient is experiencing an infusion-related reaction</i></p>

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**
- **ARIA ordering:** Support protocols are available under **ritUX desens. q7d, ritUX desens. q21d, ritUX desens. q28d** and **ritUX desens. maint.** in the “Lymphoma” folder