

**Riabni™ (Rituximab-arrx), Ruxience™ (Rituximab-pvvr),  
and Truxima® (Rituximab-abbs)  
Prior Authorization Form**

Member Name: \_\_\_\_\_ Date of Birth: \_\_\_\_\_ Member ID#: \_\_\_\_\_

**Drug Information**

Physician billing (HCPCS code: \_\_\_\_\_) Start Date (or date of next dose): \_\_\_\_\_

Dose: \_\_\_\_\_ Regimen: \_\_\_\_\_

**Billing Provider Information**

Provider NPI: \_\_\_\_\_ Provider Name: \_\_\_\_\_

Provider Phone: \_\_\_\_\_ Provider Fax: \_\_\_\_\_

**Prescriber Information**

Prescriber NPI: \_\_\_\_\_ Prescriber Name: \_\_\_\_\_

Prescriber Phone: \_\_\_\_\_ Prescriber Fax: \_\_\_\_\_ Specialty: \_\_\_\_\_

**Criteria**

**For Initial Authorization:**

1. Please provide all of the following:

A. Diagnosis: \_\_\_\_\_

B. A patient-specific, clinically significant reason why the member cannot use Rituxan® (rituximab):

\_\_\_\_\_  
\_\_\_\_\_  
\_\_\_\_\_

Additional Information: \_\_\_\_\_

\_\_\_\_\_

**For Continued Authorization:**

1. Date of last dose: \_\_\_\_\_

2. Does patient have any evidence of progressive disease while on rituximab therapy? Yes \_\_\_ No \_\_\_

3. Has the member experienced any adverse drug reactions related to rituximab therapy? Yes \_\_\_ No \_\_\_

If yes, please specify adverse reactions: \_\_\_\_\_

Additional Information: \_\_\_\_\_

\_\_\_\_\_

**Prescriber Signature:** \_\_\_\_\_ **Date:** \_\_\_\_\_

*I certify that the indicated treatment is medically necessary and all information is true and correct to the best of my knowledge.*

*Please do not send in chart notes. Specific information will be requested if necessary. Failure to complete this form in full will result in processing delays.*

PLEASE PROVIDE THE INFORMATION REQUESTED AND RETURN TO:

University of Oklahoma College of Pharmacy  
Pharmacy Management Consultants  
Product Based Prior Authorization Unit

Fax: 1-800-224-4014  
Phone: 1-800-522-0114 Option 4

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