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Please note: All information below is required to process this request.

Mon-Fri: 5am to 10pm Pacific / Sat: 6am to 3pm Pacific

Rituxan® Prior Authorization Request Form (Page 1 of 2)

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Member Information (required)			Provider Information (required)		
Member Name:			Provider Name:		
Insurance ID#:			NPI#:		Specialty:
Date of Birth:			Office Phone:		
Street Address:			Office Fax:		
City:	State:	Zip:	Office Street Address:		
Phone:			City:	State:	Zip:
Medication Information (required)					
Medication Name:			Strength:		Dosage Form:
<input type="checkbox"/> Check if requesting brand			Directions for Use:		
<input type="checkbox"/> Check if request is for continuation of therapy					
Clinical Information (required)					
Select the diagnosis below:					
<input type="checkbox"/> Chronic lymphocytic leukemia (CLL)					
<input type="checkbox"/> Immune or idiopathic thrombocytopenic purpura (ITP)					
<input type="checkbox"/> Microscopic polyangiitis (MPA)					
<input type="checkbox"/> Moderately to severely active rheumatoid arthritis (RA)					
<input type="checkbox"/> Non-Hodgkin's lymphoma (NHL)					
<input type="checkbox"/> Wegener's granulomatosis					
<input type="checkbox"/> Other diagnosis: _____ ICD-10 Code(s): _____					
Clinical Information:					
Is this request for continuation of prior Rituxan therapy? <input type="checkbox"/> Yes <input type="checkbox"/> No					
Has the patient used Rituxan within the past 120 days? <input type="checkbox"/> Yes <input type="checkbox"/> No					
Select if Rituxan is prescribed by or in consultation with one of the following specialists, as appropriate for the patient's diagnosis:					
<input type="checkbox"/> Hematologist/oncologist					
<input type="checkbox"/> Nephrologist					
<input type="checkbox"/> Pulmonologist					
<input type="checkbox"/> Rheumatologist					
For immune or idiopathic thrombocytopenic purpura (ITP), also answer the following:					
Has the patient had trial and failure, contraindication, or intolerance to corticosteroids, immunoglobulins, or splenectomy? <input type="checkbox"/> Yes <input type="checkbox"/> No					
Does the patient have a documented platelet count of less than 50 x 10 ⁹ /L? <input type="checkbox"/> Yes <input type="checkbox"/> No					
For moderately to severely active rheumatoid arthritis, also answer the following:					
Is the patient concurrently on methotrexate? <input type="checkbox"/> Yes <input type="checkbox"/> No					
If "no" to the above question, does the patient have history of contraindication or intolerance to methotrexate? <input type="checkbox"/> Yes <input type="checkbox"/> No					
Has the patient had trial and failure, contraindication, or intolerance to one TNF antagonist [e.g., Humira (adalimumab), Enbrel (etanercept), Remicade (Infliximab)]? <input type="checkbox"/> Yes <input type="checkbox"/> No					
Will the patient be receiving Rituxan in combination with a biologic DMARD [e.g., Enbrel (etanercept), Orencia (abatacept), Kineret (anakinra)]? <input type="checkbox"/> Yes <input type="checkbox"/> No					
For Wegener's granulomatosis or microscopic polyangiitis, also answer the following:					
Is the patient concurrently on glucocorticoids (e.g., prednisone)? <input type="checkbox"/> Yes <input type="checkbox"/> No					
If "no" to the above question, does the patient have history of contraindication or intolerance to glucocorticoids (e.g., prednisone)? <input type="checkbox"/> Yes <input type="checkbox"/> No					

This document and others if attached contain information that is privileged, confidential and/or may contain protected health information (PHI). The Provider named above is required to safeguard PHI by applicable law. The information in this document is for the sole use of OptumRx. Proper consent to disclose PHI between these parties has been obtained. If you received this document by mistake, please know that sharing, copying, distributing or using information in this document is against the law. **If you are not the intended recipient, please notify the sender immediately.**

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For non-Hodgkin's lymphoma (NHL), also answer the following:

Select if the patient has one of the following forms of non-Hodgkin's lymphoma (NHL):

- Diffuse large B-cell, CD20-positive, NHL
Will Rituxan be used as first-line treatment in combination with CHOP (cyclophosphamide, doxorubicin, vincristine, prednisone) or other anthracycline-based chemotherapy regimens? Yes No
- Follicular, CD20-positive, B-cell NHL
Will Rituxan be used as first-line treatment in combination with chemotherapy? Yes No
Will Rituxan be used as a single-agent maintenance therapy in a patient who has achieved a complete or partial response to Rituxan in combination with chemotherapy? Yes No
- Low-grade, CD20-positive, B-cell NHL
Does the patient have stable disease following first-line treatment with CVP (cyclophosphamide, vincristine, prednisolone/prednisone) chemotherapy? Yes No
Has the patient achieved a partial or complete response following first-line treatment with CVP (cyclophosphamide, vincristine, prednisolone/prednisone) chemotherapy? Yes No
- Relapse or refractory, low grade or follicular CD20-positive, B-cell NHL

Are there any other comments, diagnoses, symptoms, medications tried or failed, and/or any other information the physician feels is important to this review?

Please note:

This request may be denied unless all required information is received.
If the patient is not able to meet the above standard prior authorization requirements, please call 1-800-711-4555.
For urgent or expedited requests please call 1-800-711-4555.
This form may be used for non-urgent requests and faxed to 1-800-527-0531.