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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"/> Granulomatosis with polyangitis (GPA) (Wegener's granulomatosis) and microscopic polyangitis (MPA)					
<input type="checkbox"/> Immune or idiopathic thrombocytopenic purpura (ITP)					
<input type="checkbox"/> Moderately to severely active rheumatoid arthritis					
<input type="checkbox"/> Non-Hodgkin's lymphoma (NHL)					
<input type="checkbox"/> Pemphigus vulgaris					
<input type="checkbox"/> Other diagnosis: _____ ICD-10 Code(s): _____					
Prescriber's Specialty:					
Select if Rituxan is prescribed by or in consultation with the following, as appropriate for the patient's diagnosis:					
<input type="checkbox"/> Dermatologist					
<input type="checkbox"/> Hematologist/oncologist					
<input type="checkbox"/> Nephrologist					
<input type="checkbox"/> Pulmonologist					
<input type="checkbox"/> Rheumatologist					
For chronic lymphocytic leukemia (CLL), answer the following:					
Will the patient use Rituxan in combination with fludarabine? <input type="checkbox"/> Yes <input type="checkbox"/> No					
Will the patient use Rituxan in combination with cyclophosphamide? <input type="checkbox"/> Yes <input type="checkbox"/> No					
For granulomatosis with polyangitis (GPA) (Wegener's granulomatosis) and microscopic polyangitis (MPA), answer the following:					
Select the diagnosis that applies to the patient:					
<input type="checkbox"/> Granulomatosis with polyangitis (Wegener's granulomatosis)					
<input type="checkbox"/> Microscopic polyangitis					
Is the patient concurrently on glucocorticoids (e.g., prednisone)? <input type="checkbox"/> Yes <input type="checkbox"/> No					
If "no," does the patient have history of contraindication or intolerance to glucocorticoids (e.g., prednisone)? <input type="checkbox"/> Yes <input type="checkbox"/> No					
For immune or idiopathic thrombocytopenic purpura, answer the following:					
Select if the patient has had a trial and failure, contraindication, or intolerance to the following:					
<input type="checkbox"/> Corticosteroids					
<input type="checkbox"/> Immunoglobulins					
<input type="checkbox"/> Splenectomy					
Does the patient have a documented platelet count of less than $50 \times 10^9/L$? <input type="checkbox"/> Yes <input type="checkbox"/> No					

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Rituxan[®] Prior Authorization Request Form (Page 2 of 2)

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For moderately to severely active rheumatoid arthritis, answer the following:

Is the patient concurrently on methotrexate? Yes No

If "no" to the above question, does the patient have history of **contraindication or intolerance** to methotrexate? Yes No

Select if the patient has had a trial and failure, contraindication, or intolerance to the following, or attestation demonstrating a trial may be inappropriate:

- Cimzia (certolizumab)
- Humira (adalimumab)
- Simponi (golimumab) or Simponi Aria (golimumab IV)

Select if the patient has had a trial and failure, contraindication, or intolerance to the following:

- Kevzara (sarilumab)
- Xeljanz (tofacitinib) or Xeljanz XR (tofacitinib ER)

Is this request for continuation of prior Rituxan therapy? Yes No

Will the patient be receiving Rituxan in combination with a biologic DMARD [e.g., Enbrel (etanercept), Orenia (abatacept), Kineret (anakinra)]? Yes No

Reauthorization:

Is there documentation the patient has had a positive clinical response to Rituxan therapy? Yes No

Have at least 16 weeks elapsed since the last course of therapy? Yes No

Will the patient be receiving Rituxan in combination with a biologic DMARD [e.g., Enbrel (etanercept), Orenia (abatacept), Kineret (anakinra)]? Yes No

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

Select the diagnosis that applies to the patient:

- 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 a first-line treatment in combination with chemotherapy? Yes No
Will Rituxan be used as monotherapy for maintenance therapy? Yes No
Has the patient 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 chemotherapy? Yes No
- Relapsed or refractory, low-grade or follicular CD20-positive, B-cell NHL

For pemphigus vulgaris, answer the following:

Does the patient have moderate to severe disease? Yes No

Reauthorization:

Is there documentation the patient has had a positive clinical response to Rituxan therapy? Yes No

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.
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.

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