



Drug Class Prior Authorization Criteria
Rheumatic and Inflammatory Diseases

Line of Business: Medicaid

P & T Approval Date: August 5, 2022

Effective Date: September 2, 2022

These criteria have been developed through review of medical literature, consideration of medical necessity, generally accepted medical practice standards, and approved by the IEHP Pharmacy and Therapeutics Subcommittee.

Drugs Requiring Prior Authorization Review: Rituxan (rituximab)

CRITERIA:

RITUXAN (RITUXIMAB)

Covered Uses: Rheumatoid arthritis

Exclusion Criteria: N/A

Required Medical Information:

Must meet all of the following requirements:

- a. Failure or clinically significant adverse effects to "1" of the following: azathioprine, cyclosporine, hydroxychloroquine, leflunomide, methotrexate, OR sulfasalazine
- b. Failure or clinically significant adverse effects to "1" of the following: **Enbrel** or **Humira**

Age Restrictions: N/A

Prescriber Restrictions: Dermatologist, Rheumatologist

Other Criteria:

Reauthorization Criteria: Must meet all of the following requirements:

- a. Documentation of meeting therapeutic goal (e.g., disease stability, improvement in daily activities and/or reduction in frequency of disease attacks)
 - b. Requested dosage and administration are consistent with the FDA recommendations
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Inland Empire Health Plan

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Change Control		
Date	Change	Author
06/23/2022	<ul style="list-style-type: none"> • Retired PA criteria for Medi-Cal Rx Benefits <ul style="list-style-type: none"> ○ Enbrel (etanercept) ○ Humira (adalimumab) ○ Otezla (apremilast) 	VM
07/15/2021	<ul style="list-style-type: none"> • Relocated Rituxan (rituximab) PA criteria for RA indication from the PA table 	SV
08/28/2020	<ul style="list-style-type: none"> • Renew with no changes 	RR
08/21/2019	<ul style="list-style-type: none"> • Added an indication table on page 1 for overview purpose • Retired PA criteria for non-preferred agents: <ul style="list-style-type: none"> ○ Actemra, Cimzia, Cosentyx, Entyvio, Kevzara, Kineret, Orencia, Renflexis, Simponi, Simponi Aria, Stelara, Taltz, Tremfya, Tysabri, Xeljanz • Retired PA criteria for low volume utilization <ul style="list-style-type: none"> ○ Humira: NIU (non-infectious uveitis) criteria retired • Created PA criteria for high PA volume indication: <ul style="list-style-type: none"> ○ Humira: HS (hidradenitis suppurativa) criteria added 	SV
05/15/2019	<ul style="list-style-type: none"> • Revise Xeljanz criteria: remove trial and failure requirement through Humira, Simponi and Renflexis per new updated 2019 ACG guidelines 	ND
08/17/2018	<ul style="list-style-type: none"> • Post P&T changes: Remove Inflectra and Remicade from document; Renflexis is the preferred infliximab agent 	HC
08/06/2018	<ul style="list-style-type: none"> • Added double lines to all drug names for consistency 	IK
07/15/2018	<ul style="list-style-type: none"> • Added new FDA indication of plaque psoriasis to Cimzia • Added new FDA indication of ulcerative colitis to Xeljanz • Remicade: Documented intolerance or contraindication to an inactive ingredient in Renflexis and Inflectra • Inflectra: Documented intolerance or contraindication to an inactive ingredient in Renflexis • Remove Ilaris criteria due to rare indications 	HC
07/11/2018	<ul style="list-style-type: none"> • Reformatted document 	IK