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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

Simponi® & Simponi Aria® Prior Authorization Request Form

DO NOT COPY FOR FUTURE USE. FORMS ARE UPDATED FREQUENTLY AND MAY BE BARCODED

Member Information (required)	Provider Information (required)
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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"/> Active ankylosing spondylitis	<input type="checkbox"/> Moderately to severely active rheumatoid arthritis
<input type="checkbox"/> Active psoriatic arthritis	<input type="checkbox"/> Moderately to severely active ulcerative colitis
<input type="checkbox"/> Other diagnosis: _____ ICD-10 Code(s): _____	

Clinical Information:

Is this request for continuation of prior therapy? Yes No

Will Simponi/Simponi Aria be used in combination with a biologic DMARD [e.g., Enbrel (etanercept), Humira (adalimumab), Cimzia (certolizumab), Orenzia (abatacept)]? Yes No

Select if the requested medication is prescribed by or in consultation with one of the following specialists:

Dermatologist Gastroenterologist Rheumatologist

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

Cosentyx (secukinumab) Enbrel (etanercept) Humira (adalimumab)

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

Is the patient concurrently on Rheumatrex/Trexall (methotrexate)? Yes No

Has the patient had trial and failure, contraindication, or intolerance to Rheumatrex/Trexall (methotrexate)? Yes No

For moderately to severely active ulcerative colitis, also answer the following:

Is the patient corticosteroid dependent (e.g., an inability to successfully taper corticosteroids without a return of the symptoms of ulcerative colitis)? Yes No

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

Oral aminosalicylates Oral corticosteroid Azathioprine 6-mercaptopurine

Reauthorization:

If this is a reauthorization request, answer the following questions:

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

Will Simponi/Simponi Aria be used in combination with a biologic DMARD [e.g., Enbrel (etanercept), Humira (adalimumab), Cimzia (certolizumab), Orenzia (abatacept)]? 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.
 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.

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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