South Dakota Department of Social Services

Please note: All information below is required to process this request.
Fax to 1-844-403-1029
Mon-Sat: 7am to 7pm Central

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

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

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:
❑ Check if requesting **brand**	Directions for Use:	
❑ Check if request is for **continuation of therapy**		

Clinical Information (required)

Select the diagnosis below:

❑ Active psoriatic arthritis (PsA)
❑ Moderately to severely active polyarticular juvenile idiopathic arthritis (pJIA)
❑ Moderately to severely active rheumatoid arthritis (RA)
❑ Other diagnosis: _____ ICD-10 Code(s): _____

Clinical information:

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

Select if the requested medication is prescribed by or in consultation with one of the following specialists:
❑ Dermatologist ❑ Rheumatologist ❑ Other _____

Will the requested medication be used in combination with another biologic agent? ❑ **Yes** ❑ **No**

For active psoriatic arthritis (PsA), also answer the following:

Has the patient had an inadequate response to, intolerance to, or contraindication to methotrexate? ❑ **Yes** ❑ **No**

For moderately to severely active polyarticular juvenile idiopathic arthritis (pJIA), also answer the following:

Has the patient had an inadequate response to, intolerance to, or contraindication to one or more non-biologic disease modifying anti-rheumatic drugs (DMARDs)? ❑ **Yes** ❑ **No** List _____

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

Has the patient had an inadequate response to, intolerance to, or contraindication to one or more non-biologic disease modifying anti-rheumatic drugs (DMARDs)? ❑ **Yes** ❑ **No** List _____

Quantity limit requests:

What is the quantity requested per TREATMENT? _____ syringe every _____ weeks

What is the reason for exceeding the plan limitations?

❑ Titration or loading dose purposes
❑ Patient is on a dose-alternating schedule (e.g., one tablet in the morning and two tablets at night, one to two tablets at bedtime)
❑ Requested strength/dose is not commercially available
❑ Other: _____

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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 1-855-401-4262.
This form may be used for non-urgent requests and faxed to 1-844-403-1029.