

Medical drug benefit
Prior authorization request



Orencia

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Phone: 877-417-1839 (NH Medicaid)

Fax: 866-539-7185

* Some plans might not accept this form for Medicare or Medicaid requests

This form is being used for:

Check if Expedited Review/Urgent Request:

☐ (In checking this box, I attest to the fact that this request meets the definition and criteria for expedited review and is an urgent request.)

1. Patient Information

Patient Name:

DOB:

Member ID #:

2. Prescriber Information

Prescribing Clinician:

Phone #:

Specialty:

Secure Fax #:

NPI #:

DEA/xDEA:

Prescriber Point of Contact Name (POC) (if different than provider):

POC Phone #:

POC Secure Fax #:

POC Email (not required):

Prescribing Clinician or Authorized Representative Signature:

Date:

3. Requested dosing

Please document the requested dosing:

4. Member's weight in kg

Please document the member's weight in kg:

5. Concurrent use

Will the member be using the requested medication concurrently with a biologic or a targeted synthetic DMARD?

☐ Yes

☐ No

6. Other conditions

Does the member have any of the following conditions? Please choose any of the following that apply

☐ Ankylosing spondylitis

☐ Inflammatory bowel disease (i.e. Crohn's disease, ulcerative colitis)

☐ Psoriasis

☐ None of the above

7. Diagnosis

Please choose the appropriate diagnosis

☐ Juvenile Idiopathic Arthritis/Juvenile Rheumatoid Arthritis (proceed to Q8)

☐ Psoriatic Arthritis (proceed to Q9)

☐ Rheumatoid Arthritis (proceed to Q9)

☐ Other, please indicate diagnosis below and include supporting clinical documentation for use in this indication and attached applicable chart notes in faxed request

8. Juvenile idiopathic arthritis/Juvenile rheumatoid arthritis

Please choose **all** of the following that apply: (proceed to Q10)

- ☐ Member has tried Simponi Aria
- ☐ Member has heart failure, previously treated lymphoproliferative disorder, or a previous serious infection
- ☐ Member has been established on Orencia Intravenous or Orencia Subcutaneous for at least 90 days
- ☐ Other (please specify):

9. Psoriatic arthritis or rheumatoid arthritis

Please choose **all** of the following that apply (proceed to Q11 for rheumatoid arthritis or to Q12 for psoriatic arthritis)

- ☐ Member has tried Simponi Aria or Cimzia or both
- ☐ Member has heart failure, previously treated lymphoproliferative disorder, or a previous serious infection
- ☐ Member has been established on Orencia Intravenous or Orencia Subcutaneous for at least 90 days
- ☐ Other (please specify):

10. Juvenile idiopathic arthritis/juvenile rheumatoid arthritis

Please choose **one** of the following that apply (proceed to Q12):

- ☐ Member has tried one other agent for this condition(e.g. methotrexate, sulfasalazine, or leflunomide, and an NSAID)
- ☐ Member will be starting on therapy concurrently with methotrexate, sulfasalazine, or leflunomide
- ☐ Member has an aggressive disease
- ☐ Member has an absolute contraindication to methotrexate, sulfasalazine, or leflunomide (please specify the drug and associated contraindication)
- ☐ Other (please specify):

11. For rheumatoid arthritis

Please choose **all** of the following that apply (proceed to Q12)

- ☐ Member has tried one conventional synthetic DMARD for at least 3 months (e.g. methotrexate, leflunomide, hydroxychloroquine, and sulfasalazine)
- ☐ Other (please specify)

12. Specialty of the Prescriber

Please indicate what specialty the prescriber is (select any that apply):

- ☐ Dermatologist
- ☐ Rheumatologist
- ☐ Other (please indicate what specialty below):

13. Initial or Continuing Therapy

Is the request for initial or continuing therapy?

- ☐ Initial (Proceed to Q16)
- ☐ Continuation (Proceed to Q14)

14. Response

Did the member have a response to therapy?

- ☐ Yes (proceed to Q15)
- ☐ No

15. Appropriate Response to Therapy

Please choose the appropriate response to therapy

- ☐ Decrease soft tissue swelling in joints or tendon sheaths
- ☐ Improvement in limitation of movement
- ☐ Improved function or activities of daily living
- ☐ Improved laboratory values
- ☐ Less joint pain, tenderness, morning stiffness or fatigue
- ☐ Improvements in C-reactive protein
- ☐ Reduced dosage of corticosteroids
- ☐ Other, please specify:

16. HCPCS Codes

Please document the applicable HCPCS codes (e.g. J codes or Q codes) being requested, including the number of units and number of visits (*using the space below*):

- ☐ HCPCS / Qcodes:
- ☐ Number of units:
- ☐ Number of visits:

Providers should consult the health plan's coverage policies, member benefits, and medical necessity guidelines to complete this form. Providers may attach any additional data relevant to medical necessity criteria.

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