

**Orencia
Prior Authorization Request**

Send completed form to: Case Review Unit, CVS Caremark Prior Authorization Fax: 1-866-249-6155

CVS Caremark administers the prescription benefit plan for the patient identified. This patient's benefit plan requires prior authorization for certain medications in order for the drug to be covered. To make an appropriate determination, providing the most accurate diagnosis for the use of the prescribed medication is necessary. **Please respond below and fax this form to CVS Caremark toll-free at 1-866-249-6155.** If you have questions regarding the prior authorization, please contact CVS Caremark at **1-866-814-5506**. For inquiries or questions related to the patient's eligibility, drug copay or medication delivery; please contact the Specialty Customer Care Team: CaremarkConnect® 1-800-237-2767.

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Patient's Name: _____ **Date:** _____
Patient's ID: _____ **Patient's Date of Birth:** _____
Physician's Name: _____ **NPI#:** _____
Specialty: _____ **Physician Office Fax:** _____
Physician Office Telephone: _____
Request Initiated For: _____

1. What is the requested formulation? Orencia (SC)/Orencia ClickJect Orencia (IV)
2. Has the patient been diagnosed with any of the following?
 Moderately to severely active rheumatoid arthritis (RA)
 Moderately to severely active polyarticular juvenile idiopathic arthritis (pJIA)
 Active psoriatic arthritis (PsA)
 Systemic juvenile idiopathic arthritis (sJIA)
 Other _____
3. What is the ICD-10 code? _____

Section A: Preferred Product

4. These are the preferred products for which coverage is provided for treatment of the following conditions:
a) Rheumatoid arthritis: **Enbrel, Humira, Kevzara, Orencia (SC)/Orencia ClickJect, skip to Section B: All Requests if Orencia (SC)/Orencia ClickJect is being prescribed.**
b) Psoriatic arthritis: **Cosentyx, Enbrel, Humira, Otezla**
Can the patient's treatment be switched to a preferred product?
 Yes - Please specify: _____ **If Yes, please call 1-866-814-5506 to have the updated form faxed to your office OR you may complete the PA electronically (ePA). You may sign up online via CoverMyMeds at: www.covermymeds.com/epa/caremark/ or call 1-866-452-5017.**
 No
 Not applicable - Requested for condition not listed above, skip to Section B: All Requests
5. Is this request for continuation of therapy with the requested product? Yes No *If No, skip to #7*
6. Is the patient currently receiving the requested product through samples or a manufacturer's patient assistance program? If unknown, answer Yes. Yes No *If No, skip to Section B: All Requests*

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7. Has the patient had a documented inadequate response or intolerable adverse event with any of the following preferred products? List continues on following page. Please indicate ALL that apply.

ACTION REQUIRED: If Yes, attach supporting chart note(s).

- | | | |
|---|--|--|
| <input type="checkbox"/> Cosentyx: | <input type="checkbox"/> Inadequate response | <input type="checkbox"/> Intolerable adverse event |
| <input type="checkbox"/> Enbrel: | <input type="checkbox"/> Inadequate response | <input type="checkbox"/> Intolerable adverse event |
| <input type="checkbox"/> Humira: | <input type="checkbox"/> Inadequate response | <input type="checkbox"/> Intolerable adverse event |
| <input type="checkbox"/> Kevzara: | <input type="checkbox"/> Inadequate response | <input type="checkbox"/> Intolerable adverse event |
| <input type="checkbox"/> Orencia (SC)/Orencia Click Ject: | <input type="checkbox"/> Inadequate response | <input type="checkbox"/> Intolerable adverse event |
| <input type="checkbox"/> Otezla: | <input type="checkbox"/> Inadequate response | <input type="checkbox"/> Intolerable adverse event |
| <input type="checkbox"/> No - none of the above | | |

8. Does the patient have one of the following documented clinical reasons to avoid Enbrel and Humira?

ACTION REQUIRED: If Yes, attach supporting chart note(s).

- | | |
|---|---|
| <input type="checkbox"/> Yes - History of demyelinating disorder | <input type="checkbox"/> Yes - Autoantibody formation/lupus-like syndrome |
| <input type="checkbox"/> Yes - History of congestive heart failure | <input type="checkbox"/> Yes - Risk of lymphoma |
| <input type="checkbox"/> Yes - History of hepatitis B virus infection | <input type="checkbox"/> No - none of the above |

Section B: All Requests

9. Is this request for continuation of therapy? Yes No *If No, skip to #13*
10. Is the patient currently receiving Orencia through samples or a manufacturer's patient assistance program?
 Yes No Unknown *If Yes or Unknown, skip to #13*
11. How long has the patient been receiving the requested medication? _____ months
If less than 3 months, no further questions.
12. Has the patient achieved or maintained positive clinical response to treatment as evidenced by low disease activity or improvement in signs and symptoms? *If Yes, no further questions* Yes No
13. Has the patient received any of the following medications?
If Yes, please indicate the most recent medication and skip to diagnosis section.
 Actemra Cimzia Cosentyx Enbrel Humira Inflectra Kevzara Kineret Remicade
 Renflexis Rituxan Siliq Simponi Simponi Aria Stelara Taltz Tremfya Xeljanz
 Xeljanz XR No
14. Has the patient undergone pretreatment screening for latent tuberculosis (TB) infection with either a TB skin test or an interferon gamma release assay (e.g., QFT-GIT, T-SPOT.TB)? Yes No

Complete the following section based on the patient's diagnosis, if applicable.

Section A: Rheumatoid Arthritis

15. Has the patient experienced an inadequate response after at least 3 months of treatment with the methotrexate dose greater than or equal to 20 mg per week? *If Yes, no further questions* Yes No
16. Has the patient experienced intolerance to methotrexate? *If Yes, no further questions* Yes No
17. Does the patient have a contraindication to methotrexate? Yes No
If Yes, indicate the contraindication: _____

Section B: Polyarticular Juvenile Idiopathic Arthritis

18. Has the patient experienced an inadequate response to a tumor necrosis factor (TNF) inhibitor (e.g., Enbrel, Humira, or Remicade) after at least 3 months of treatment? *If Yes, no further questions.* Yes No
19. Has the patient experienced an intolerable adverse event to a tumor necrosis factor (TNF) inhibitor (e.g., Enbrel, Humira, or Remicade)? *If Yes, no further questions.* Yes No
20. Does the patient have contraindication to a tumor necrosis factor (TNF) inhibitor (e.g., Enbrel, Humira, or Remicade)? Yes No

I attest that this information is accurate and true, and that documentation supporting this information is available for review if requested by CVS Caremark or the benefit plan sponsor.

X _____
Prescriber or Authorized Signature Date (mm/dd/yy)