

Tremfya
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 diagnosis?
 Moderate to severe plaque psoriasis
 Other _____
2. What is the ICD-10 code? _____
3. These are the primary preferred products for which coverage is provided for treatment of the following condition:
 Plaque psoriasis: **Humira (primary); Stelara/Taltz (secondary)***
**Note: Secondary preferred products for plaque psoriasis are Stelara and Taltz. These preferred product options only apply to members who have had a documented inadequate response or intolerable adverse event with Humira.*
 Can the patient's treatment be switched to a preferred drug?
 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 #8
4. Is this request for continuation of therapy with the requested product? Yes No *If No, skip to #6*
5. 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 #8*
6. Has the patient had a documented inadequate response or intolerable adverse event with any of the following preferred products? Please indicate ALL that apply. **ACTION REQUIRED: If Yes, attach supporting chart note(s).**

<input type="checkbox"/> Humira:	<input type="checkbox"/> Inadequate response	<input type="checkbox"/> Intolerable adverse event
<input type="checkbox"/> Stelara:	<input type="checkbox"/> Inadequate response	<input type="checkbox"/> Intolerable adverse event
<input type="checkbox"/> Taltz:	<input type="checkbox"/> Inadequate response	<input type="checkbox"/> Intolerable adverse event

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No - none of the above

7. Does the patient have one of the following documented clinical reasons to avoid Humira?

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

- Yes - History of demyelinating disorder
- Yes - History of congestive heart failure
- Yes - History of hepatitis B virus infection
- Yes - Autoantibody formation/lupus-like syndrome
- Yes - Risk of lymphoma
- No - none of the above

8. Is this request for continuation of therapy? Yes No *If No, skip to #12*

9. Is the patient currently receiving Tremfya through samples or a manufacturer's patient assistance program? Yes No Unknown *If Yes or Unknown, skip to #12*

10. How long has the patient been receiving the requested medication? _____ months
If less than 4 months, no further questions.

11. 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

12. Has the patient received any of the following medications? *If Yes, please indicate the most recent medication.*

- Actemra Cimzia Cosentyx Enbrel Humira Inflectra Kevzara Orencia
- Otezla Remicade Renflexis Siliq Simponi Simponi Aria Stelara Taltz
- Xeljanz Xeljanz XR No

13. 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

14. What is the percentage of body surface area (BSA) affected? _____ % of BSA

15. *If less than 5% BSA affected, are crucial body areas (e.g., hands, feet, face, neck, scalp, genitals/groin, intertriginous areas) affected?* Yes No

16. Has the patient experienced an inadequate response, or has an intolerance to phototherapy (e.g., UVB, PUVA) or pharmacologic treatment with methotrexate, cyclosporine or acitretin?
If Yes, no further questions Yes No

17. Does the patient have a clinical reason to avoid pharmacologic treatment with methotrexate, cyclosporine, or acitretin? Yes No

If Yes, indicate clinical reason: _____

18. Does the patient have severe psoriasis that warrants a biologic DMARD as first-line therapy?
 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)