Skyrizi

Prior Authorization Request

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 1-855-330-1720**. If you have questions regarding the prior authorization, please contact CVS Caremark at **1-888-877-0518.** 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.

The recipient of this fax may make a request to opt-out of receiving telemarketing fax transmissions from CVS Caremark. There are numerous ways you may opt-out: The recipient may call the toll-free number at 877-265-2711, at any time, 24 hours a day/7 days a week. The recipient may also send an opt-out request via email to do_not_call@cvscaremark.com. An opt out request is only valid if it (1) identifies the number to which the request relates, and (2) if the person/entity making the request does not, subsequent to the request, provide express invitation or permission to CVS Caremark to send facsimile advertisements to such person/entity at that particular number. CVS Caremark is required by law to honor an opt-out request within thirty days of receipt.

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

Referring Provider Info: ❑ **Same as Requesting Provider**
Name: _____ **NPI#:** _____
Fax: _____ **Phone:** _____

Rendering Provider Info: ❑ **Same as Referring Provider** ❑ **Same as Requesting Provider**
Name: _____ **NPI#:** _____
Fax: _____ **Phone:** _____

Approvals may be subject to dosing limits in accordance with FDA-approved labeling, accepted compendia, and/or evidence-based practice guidelines.

Required Demographic Information:

Patient Weight: _____*kg*

Patient Height: _____*cm*

Please indicate the place of service for the requested drug:

❑ *Ambulatory Surgical*	❑ *Home*	❑ *Off Campus Outpatient Hospital*
❑ *On Campus Outpatient Hospital*	❑ *Office*	❑ *Pharmacy*

Criteria Questions:

What is the ICD-10 code? _____

1. Will the requested drug be used in combination with any other biologic (e.g., Humira) or targeted synthetic drug (e.g., Olumiant, Otezla, Xeljanz)?
❒ Yes, *Continue to 2*
❒ No, *Continue to 2*

2. Has the patient ever received (including current utilizers) a biologic (e.g., Humira) or targeted synthetic drug (e.g., Olumiant, Xeljanz) associated with an increased risk of tuberculosis?
❒ Yes, *Continue to 6*
❒ No, *Continue to 3*

Send completed form to: Case Review Unit CVS Caremark Specialty Programs Fax: 1-855-330-1720

3. Has the patient had a tuberculosis (TB) test (e.g., tuberculosis skin test [PPD], interferon-release assay [IGRA], chest x-ray) within 6 months of initiating therapy?
❒ Yes, *Continue to 4*
❒ No, *Continue to 4*

4. What were the results of the TB test?

❒ Positive for TB, *Continue to 5*

❒ Negative for TB, *Continue to 6*

❒ Unknown, *No further questions*

5. Which of the following applies to the patient?

❒ Patient has latent TB and treatment for latent TB has been initiated, *Continue to 6*

❒ Patient has latent TB and treatment for latent TB has been completed, *Continue to 6*

❒ Patient has latent TB and treatment for latent TB has not been initiated, *Continue to 6*

❒ Patient has active TB, *Continue to 6*

6. What is the diagnosis?

❒ Plaque psoriasis, *Continue to 8*

❒ Psoriatic arthritis WITH co-existent plaque psoriasis, *Continue to 7*

❒ Psoriatic arthritis, *Continue to 22*

❒ Crohn's disease, *Continue to 37*

❒ Other, please specify. _____, *No further questions*

7. What is the primary diagnosis being treated?

❒ Psoriatic arthritis, *Continue to 22*

❒ Plaque psoriasis, *Continue to 8*

8. Has the patient been diagnosed with moderate to severe plaque psoriasis?
❒ Yes, *Continue to 9*
❒ No, *Continue to 9*

9. Is the patient an adult (18 years of age or older)?
❒ Yes, *Continue to 10*
❒ No, *Continue to 10*

10. Is the requested drug being prescribed by or in consultation with a dermatologist?
❒ Yes, *Continue to 11*
❒ No, *Continue to 11*

11. Is this request for continuation of therapy with the requested drug?
❒ Yes, *Continue to 12*
❒ No, *Continue to 16*

12. Is the patient currently receiving the requested drug through samples or a manufacturer's patient assistance program?

❒ Yes, *Continue to 16*

❒ No, *Continue to 13*

❒ Unknown, *Continue to 16*

13. Has the patient achieved or maintained a positive clinical response as evidenced by low disease activity or improvement in signs and symptoms of the condition since starting treatment with the requested drug?
❒ Yes, *Continue to 14*
❒ No, *Continue to 14*

14. Has the patient experienced a reduction in body surface area (BSA) affected from baseline? ***ACTION REQUIRED***: If Yes, please attach chart notes or medical record documentation of decreased body surface area affected. ***ACTION REQUIRED***: Submit supporting documentation
❒ Yes, *Continue to 44*
❒ No, *Continue to 15*

15. Has the patient experienced an improvement in signs and symptoms of the condition from baseline (e.g., itching, redness, flaking, scaling, burning, cracking, pain)? ***ACTION REQUIRED***: If Yes, please attach chart notes or medical record documentation of improvement in signs and symptoms. ***ACTION REQUIRED***: Submit supporting documentation
❒ Yes, *Continue to 44*
❒ No, *Continue to 44*

16. Has the patient ever received or is currently receiving a biologic (e.g., Humira) or targeted synthetic drug (e.g., Sotyktu, Otezla) indicated for the treatment of moderate to severe plaque psoriasis (excluding receiving the drug via samples or a manufacturer's patient assistance program)? ***ACTION REQUIRED***: If Yes, please attach chart notes, medical record documentation, or claims history supporting previous medications tried. ***ACTION REQUIRED***: Submit supporting documentation
❒ Yes, *Continue to 44*
❒ No, *Continue to 17*

17. Are crucial body areas (e.g., hands, feet, face, neck, scalp, genitals/groin, intertriginous areas) affected? ***ACTION REQUIRED***: If Yes, please attach chart notes or medical record documentation of affected areas. ***ACTION REQUIRED***: Submit supporting documentation
❒ Yes, *Continue to 44*
❒ No, *Continue to 18*

18. What is the percentage of body surface area (BSA) affected (prior to starting the requested medication)? Indicate percentage. ***ACTION REQUIRED***: Please attach chart notes or medical record documentation of body surface area affected.
❒ Greater than or equal to 3% to less than 10% of BSA _____ ***ACTION REQUIRED****: Submit supporting documentation, Continue to 19*
❒ Greater than or equal to 10% of BSA _____ ***ACTION REQUIRED****: Submit supporting documentation, Continue to 44*
❒ Less than 3% of BSA _____, *No further questions*

19. 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? ***ACTION REQUIRED***: If Yes, please attach chart notes, medical record documentation, or claims history supporting previous medications tried, including response to therapy. ***ACTION REQUIRED***: Submit supporting documentation
❒ Yes, *Continue to 44*
❒ No, *Continue to 20*

20. Does the patient have a clinical reason to avoid pharmacologic treatment with methotrexate, cyclosporine, and acitretin? ***ACTION REQUIRED***: If Yes, please attach documentation of clinical reason to avoid therapy. ***ACTION REQUIRED***: Submit supporting documentation
❒ Yes, *Continue to 21*

❒ No, *Continue to 21*

21. Please indicate the clinical reason to avoid pharmacologic treatment with methotrexate, cyclosporine, and acitretin.

❒ Clinical diagnosis of alcohol use disorder, alcoholic liver disease, or other chronic liver disease, *Continue to 44*

❒ Drug interaction, *Continue to 44*

❒ Risk of treatment-related toxicity, *Continue to 44*

❒ Pregnancy or currently planning pregnancy, *Continue to 44*

❒ Breastfeeding, *Continue to 44*

❒ Significant comorbidity prohibits use of systemic agents (e.g., liver or kidney disease, blood dyscrasias, uncontrolled hypertension), *Continue to 44*

❒ Hypersensitivity, *Continue to 44*

❒ History of intolerance or adverse event, *Continue to 44*

❒ Other, please specify. _____, *Continue to 44*

22. Is the patient an adult (18 years of age or older)?
❒ Yes, *Continue to 23*
❒ No, *Continue to 23*

23. Is the requested drug being prescribed by or in consultation with a rheumatologist or dermatologist?
❒ Yes, *Continue to 24*
❒ No, *Continue to 24*

24. Is this request for continuation of therapy with the requested drug?
❒ Yes, *Continue to 25*
❒ No, *Continue to 28*

25. Is the patient currently receiving the requested drug through samples or a manufacturer's patient assistance program?

❒ Yes, *Continue to 28*

❒ No, *Continue to 26*

❒ Unknown, *Continue to 28*

26. Has the patient achieved or maintained a positive clinical response as evidenced by low disease activity or improvement in signs and symptoms of the condition since starting treatment with the requested drug?
❒ Yes, *Continue to 27*
❒ No, *Continue to 27*

27. Which of the following has the patient experienced an improvement in from baseline? ***ACTION REQUIRED***: Please attach chart notes or medical record documentation supporting positive clinical response.

❒ Number of swollen joints ***ACTION REQUIRED****: Submit supporting documentation, Continue to 44*

❒ Number of tender joints ***ACTION REQUIRED****: Submit supporting documentation, Continue to 44*

❒ Dactylitis ***ACTION REQUIRED****: Submit supporting documentation, Continue to 44*

❒ Enthesitis ***ACTION REQUIRED****: Submit supporting documentation, Continue to 44*

❒ Axial disease ***ACTION REQUIRED****: Submit supporting documentation, Continue to 44*

❒ Skin and/or nail involvement ***ACTION REQUIRED****: Submit supporting documentation, Continue to 44*

❒ None of the above, *Continue to 44*

28. Has the patient been diagnosed with active psoriatic arthritis (PsA)?

❒ Yes, *Continue to 29*
❒ No, *Continue to 29*

29. Has the patient ever received or is currently receiving a biologic (e.g., Humira) or targeted synthetic drug (e.g., Rinvoq, Otezla) that is indicated for the treatment of active psoriatic arthritis (excluding receiving the drug via samples or a manufacturer's patient assistance program)? ***ACTION REQUIRED***: If Yes, please attach chart notes, medical record documentation, or claims history supporting previous medications tried. ***ACTION REQUIRED***: Submit supporting documentation
❒ Yes, *Continue to 44*
❒ No, *Continue to 30*

30. What is the patient's disease severity?

❒ Mild to moderate, *Continue to 31*

❒ Severe, *Continue to 44*

31. Does the patient have enthesitis or predominantly axial disease?
❒ Yes, *Continue to 44*
❒ No, *Continue to 32*

32. Has the patient had an inadequate response to methotrexate, leflunomide, or another conventional synthetic drug (e.g., sulfasalazine) administered at an adequate dose and duration? ***ACTION REQUIRED***: If Yes, please attach chart notes, medical record documentation, or claims history supporting previous medications tried, including response to therapy. ***ACTION REQUIRED***: Submit supporting documentation
❒ Yes, *Continue to 44*
❒ No, *Continue to 33*

33. Has the patient had an intolerance to methotrexate, leflunomide, or another conventional synthetic drug (e.g., sulfasalazine)? ***ACTION REQUIRED***: If Yes, please attach chart notes, medical record documentation, or claims history supporting previous medications tried, including response to therapy. ***ACTION REQUIRED***: Submit supporting documentation
❒ Yes, *Continue to 44*
❒ No, *Continue to 34*

34. Does the patient have a contraindication to methotrexate or leflunomide? ***ACTION REQUIRED***: If Yes, please attach documentation of clinical reason to avoid therapy. ***ACTION REQUIRED***: Submit supporting documentation
❒ Yes, *Continue to 36*
❒ No, *Continue to 35*

35. Does the patient have a contraindication to another conventional synthetic drug (e.g., sulfasalazine)? ***ACTION REQUIRED***: If Yes, please attach documentation of clinical reason to avoid therapy. ***ACTION REQUIRED***: Submit supporting documentation
❒ Yes, *Continue to 44*
❒ No, *Continue to 44*

36. Please indicate the contraindication methotrexate or leflunomide.

❒ Clinical diagnosis of alcohol use disorder, alcoholic liver disease, or other chronic liver disease, *Continue to 44*

❒ Drug interaction, *Continue to 44*

❒ Risk of treatment-related toxicity, *Continue to 44*

❒ Pregnancy or currently planning pregnancy, *Continue to 44*

❒ Breastfeeding, *Continue to 44*

❒ Significant comorbidity prohibits use of systemic agents (e.g., liver or kidney disease, blood dyscrasias,

uncontrolled hypertension), *Continue to 44*

❒ Hypersensitivity, *Continue to 44*

❒ History of intolerance or adverse event, *Continue to 44*

❒ Other, please specify. _____, *Continue to 44*

37. Has the patient been diagnosed with moderately to severely active Crohn's disease (CD)?
❒ Yes, *Continue to 38*
❒ No, *Continue to 38*

38. Is the patient an adult (18 years of age or older)?
❒ Yes, *Continue to 39*
❒ No, *Continue to 39*

39. Is the requested drug being prescribed by or in consultation with a gastroenterologist?
❒ Yes, *Continue to 40*
❒ No, *Continue to 40*

40. Which of the following applies to this request for the requested drug?

❒ Initiation of the intravenous (IV) loading dose, *Continue to 44*

❒ Initiation of the subcutaneous (SQ) maintenance dose, *Continue to 44*

❒ Continuation of the subcutaneous (SQ) maintenance dose, *Continue to 41*

41. Has the patient achieved or maintained remission? ***ACTION REQUIRED***: If Yes, please attach chart notes or medical record documentation of remission. ***ACTION REQUIRED***: Submit supporting documentation
❒ Yes, *Continue to 44*
❒ No, *Continue to 42*

42. Has the patient achieved or maintained a positive clinical response as evidenced by low disease activity or improvement in signs and symptoms of the condition since starting treatment with the requested drug?
❒ Yes, *Continue to 43*
❒ No, *Continue to 43*

43. Which of the following has the patient experienced improvement in from baseline? ***ACTION REQUIRED***: Please attach chart notes or medical record documentation supporting positive clinical response.

❒ Abdominal pain or tenderness ***ACTION REQUIRED****: Submit supporting documentation, Continue to 44*

❒ Diarrhea ***ACTION REQUIRED****: Submit supporting documentation, Continue to 44*

❒ Body weight ***ACTION REQUIRED****: Submit supporting documentation, Continue to 44*

❒ Abdominal mass ***ACTION REQUIRED****: Submit supporting documentation, Continue to 44*

❒ Hematocrit ***ACTION REQUIRED****: Submit supporting documentation, Continue to 44*
❒ Appearance of the mucosa on endoscopy, computed tomography enterography (CTE), magnetic resonance enterography (MRE), or intestinal ultrasound ***ACTION REQUIRED****: Submit supporting documentation, Continue to 44*
❒ Improvement on a disease activity scoring tool (e.g., Crohn's Disease Activity Index [CDAI] score) ***ACTION REQUIRED****: Submit supporting documentation, Continue to 44*

❒ None of the above, *Continue to 44*

44. What is the diagnosis?

❒ Plaque Psoriasis, *Continue to 45*

❒ Psoriatic arthritis WITH co-existent plaque psoriasis, *Continue to 45*

❒ Psoriatic arthritis, *Continue to 45*

❒ Crohn's disease, *Continue to 50*

45. Is the patient currently receiving the requested drug?
❒ Yes, *Continue to 46*
❒ No, *Continue to 48*

46. Does the prescribed dose exceed 150 mg?
❒ Yes, *Continue to 47*
❒ No, *Continue to 47*

47. Is the prescribed frequency for the maintenance dose more frequent than one dose every 12 weeks?
❒ Yes, *No Further Questions*
❒ No, *No Further Questions*

48. Does the prescribed dose exceed a loading dose of 150 mg at weeks 0 and 4, and a maintenance dose of 150 mg thereafter?
❒ Yes, *Continue to 49*
❒ No, *Continue to 49*

49. Is the prescribed frequency for the maintenance dose more frequent than one dose every 12 weeks?
❒ Yes, *No Further Questions*
❒ No, *No Further Questions*

50. Which of the following applies to this request for the requested drug?

❒ Initiation of the intravenous (IV) loading dose, *Continue to 51*

❒ Initiation of the subcutaneous (SQ) maintenance dose, *Continue to 53*

❒ Continuation of the subcutaneous (SQ) maintenance dose, *Continue to 53*

51. Does the prescribed dose exceed a loading dose of 600 mg at weeks 0, 4, and 8, and a maintenance dose of 360 mg at week 12 and thereafter?
❒ Yes, *Continue to 52*
❒ No, *Continue to 52*

52. Is the prescribed frequency for the maintenance dose more frequent than one dose every 8 weeks?
❒ Yes, *No Further Questions*
❒ No, *No Further Questions*

53. Does the prescribed dose exceed 360 mg?
❒ Yes, *Continue to 54*
❒ No, *Continue to 54*

54. Is the prescribed frequency for the maintenance dose more frequent than one dose every 8 weeks?
❒ Yes, *No Further Questions*
❒ No, *No Further Questions*

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)**

Send completed form to: Case Review Unit CVS Caremark Specialty Programs Fax: 1-855-330-1720