

FORWARDHEALTH  
**PRIOR AUTHORIZATION DRUG ATTACHMENT FOR CYTOKINE AND CELL ADHESION  
MOLECULE (CAM) ANTAGONIST DRUGS FOR ANKYLOSING SPONDYLITIS**

**INSTRUCTIONS:** Type or print clearly. Before completing this form, read the Prior Authorization Drug Attachment for Cytokine and Cell Adhesion Molecule (CAM) Antagonist Drugs for Ankylosing Spondylitis Instructions, F-11304A. Prescribers may refer to the Forms page of the ForwardHealth Portal at <https://www.forwardhealth.wi.gov/WIPortal/Subsystem/Publications/ForwardHealthCommunications.aspx?panel=Forms> for the completion instructions.

Pharmacy providers are required to have a completed Prior Authorization Drug Attachment for Cytokine and CAM Antagonist Drugs for Ankylosing Spondylitis form signed and dated by the prescriber before submitting a PA request on the Portal, by fax, or by mail. Prescribers and pharmacy providers may call Provider Services at 800-947-9627 with questions.

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**SECTION I – MEMBER INFORMATION**

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1. Name – Member (Last, First, Middle Initial)

2. Member ID Number

3. Date of Birth – Member

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**SECTION II – PRESCRIPTION INFORMATION**

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4. Drug Name

5. Drug Strength

6. Date Prescription Written

7. Directions for Use

8. Name – Prescriber

9. Address – Prescriber (Street, City, State, Zip+4 Code)

10. Phone Number – Prescriber

11. National Provider Identifier – Prescriber

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**SECTION III – CLINICAL INFORMATION FOR ANKYLOSING SPONDYLITIS**

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12. Diagnosis Code and Description

**Note: Supporting clinical information and a copy of the member's current medical records must be submitted with all PA requests.**

13. Does the member have ankylosing spondylitis?

Yes  No

14. Is the prescription written by a rheumatologist or through a rheumatology consultation?

Yes  No



DT-PA072-072

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15. Is the member currently using the requested non-preferred cytokine and CAM antagonist drug?  Yes  No

If yes, indicate the approximate date therapy was started.

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16. Indicate the preferred cytokine and CAM antagonist drugs the member has taken and provide specific details regarding member's response to treatment and the reason(s) for discontinuing. If additional space is needed, continue documentation in Section V of this form.

1. Drug Name \_\_\_\_\_ Dose \_\_\_\_\_ Dates Taken \_\_\_\_\_

Description of Treatment Response and Reason(s) for Discontinuing

\_\_\_\_\_  
\_\_\_\_\_

2. Drug Name \_\_\_\_\_ Dose \_\_\_\_\_ Dates Taken \_\_\_\_\_

Description of Treatment Response and Reason(s) for Discontinuing

\_\_\_\_\_  
\_\_\_\_\_

3. Drug Name \_\_\_\_\_ Dose \_\_\_\_\_ Dates Taken \_\_\_\_\_

Description of Treatment Response and Reason(s) for Discontinuing

\_\_\_\_\_  
\_\_\_\_\_

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17. Indicate the clinical reason(s) why the prescriber is requesting a non-preferred cytokine and CAM antagonist drug.

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**SECTION IV – AUTHORIZED SIGNATURE**

18. **SIGNATURE** – Prescriber

19. Date Signed

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**SECTION V – ADDITIONAL INFORMATION**

20. Include any additional information in the space below. Additional diagnostic and clinical information explaining the need for the drug requested may be included here.

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