

FORWARDHEALTH
**PRIOR AUTHORIZATION / PREFERRED DRUG LIST (PA/PDL) 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/Preferred Drug List (PA/PDL) for Cytokine and Cell Adhesion Molecule (CAM) Antagonist Drugs for Ankylosing Spondylitis Completion Instructions, F-11304A. Providers may refer to the Forms page of the ForwardHealth Portal at www.forwardhealth.wi.gov/WIPortal/Content/provider/forms/index.htm.spage for the completion instructions.

Pharmacy providers are required to have a completed Prior Authorization/Preferred Drug List (PA/PDL) for Cytokine and Cell Adhesion Molecule (CAM) Antagonist Drugs for Ankylosing Spondylitis form signed by the prescriber before calling the Specialized Transmission Approval Technology-Prior Authorization (STAT-PA) system or submitting a PA request on the Portal or on paper. Providers may call Provider Services at (800) 947-9627 with questions.

SECTION I — MEMBER INFORMATION

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

2. Member Identification Number

3. Date of Birth — Member

SECTION II — PRESCRIPTION INFORMATION

4. Drug Name

5. Drug Strength

6. Date Prescription Written

7. Directions for Use

8. Name — Prescriber

9. National Provider Identifier (NPI) — Prescriber

10. Address — Prescriber (Street, City, State, ZIP+4 Code)

11. Telephone Number — Prescriber

SECTION III — CLINICAL INFORMATION FOR ANKYLOSING SPONDYLITIS

12. Diagnosis Code and Description

13. Does the member have a diagnosis of ankylosing spondylitis? Yes No

14. Is the prescription written by a rheumatologist or through a rheumatology consultation? Yes No

15. Does the member have moderate to severe axial symptoms of ankylosing spondylitis? Yes No

Continued



DT-PA072-072

SECTION III — CLINICAL INFORMATION FOR ANKYLOSING SPONDYLITIS (Continued)

16. Has the member received **one** or more of the drugs listed below and taken each drug for at least **three** consecutive months and experienced an unsatisfactory therapeutic response or experienced a clinically significant adverse drug reaction? Yes No

If yes, check the box next to the drug(s) the member received. Indicate the dose of the drug(s), specific details about the unsatisfactory therapeutic response(s) or clinically significant adverse drug reaction(s), and the approximate dates the drug(s) was taken in the space provided.

- 1. leflunomide _____
- 2. methotrexate _____
- 3. NSAID or COX-2 _____
- 4. oral corticosteroids _____
- 5. sulfasalazine _____

SECTION IIIA — ADDITIONAL CLINICAL INFORMATION FOR NON-PREFERRED CYTOKINE AND CAM ANTAGONIST DRUG REQUESTS (Prior authorization requests for non-preferred cytokine and CAM antagonist drugs must be submitted on paper.)

17. Has the member taken **two** preferred cytokine and CAM antagonist drugs for at least **three** consecutive months and experienced an unsatisfactory therapeutic response or experienced a clinically significant adverse drug reaction? Yes No

If yes, indicate the two preferred cytokine and CAM antagonist drugs and doses, specific details about the unsatisfactory therapeutic responses or clinically significant adverse drug reactions, and the approximate dates each preferred cytokine and CAM antagonist drug was taken in the space provided.

- 1. _____
- 2. _____

SECTION IV — AUTHORIZED SIGNATURE

18. SIGNATURE — Prescriber	19. Date Signed
----------------------------	-----------------

SECTION V — FOR PHARMACY PROVIDERS USING STAT-PA

20. National Drug Code (11 digits)	21. Days' Supply Requested (Up to 365 Days)
------------------------------------	---

22. NPI _____

23. Date of Service (MM/DD/CCYY) (For STAT-PA requests, the date of service may be up to 31 days in the future or up to 14 days in the past.) _____

24. Place of Service _____

25. Assigned PA Number _____

26. Grant Date	27. Expiration Date	28. Number of Days Approved
----------------	---------------------	-----------------------------

SECTION VI — ADDITIONAL INFORMATION

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