DEPARTMENT OF HEALTH SERVICES
Division of Health Care Access and Accountability
F-11308 (12/12)

STATE OF WISCONSIN
DHS 107.10(2), Wis. Admin. Code

FORWARDHEALTH

PRIOR AUTHORIZATION / PREFERRED DRUG LIST (PA/PDL) FOR CYTOKINE AND CELL ADHESION MOLECULE (CAM) ANTAGONIST DRUGS FOR RHEUMATOID ARTHRITIS (RA) AND POLYARTICULAR JUVENILE RA

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 Rheumatoid Arthritis (RA) and Polyarticular Juvenile RA Completion Instructions, F-11308A. 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 Rheumatoid Arthritis (RA) and Polyarticular Juvenile Rheumatoid Arthritis (RA) 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 RHEUMATOID ARTHRITIS AND POLYARTICULAR JUVENILE RA (Required for all requests.)

12. Diagnosis Code and Description

13. Does the member have a diagnosis of polyarticular juvenile RA?	❑ Yes	❑ No
14. Does the member have a diagnosis of RA?	❑ Yes	❑ No
15. Does the member have moderate to severe symptoms of RA?	❑ Yes	❑ No
16. Is the prescription written by a rheumatologist or through a rheumatology consultation?	❑ Yes	❑ No

Continued

SECTION III — CLINICAL INFORMATION FOR RA AND POLYARTICULAR JUVENILE RA (Continued)

17. Has the member received **two** 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 boxes next to the drugs the member received. Indicate the dose of the drugs, specific details about the unsatisfactory therapeutic responses or clinically significant adverse drug reactions, and the approximate dates the drugs were taken in the space below.

1. ❑ azathioprine _____
2. ❑ cyclosporine _____
3. ❑ hydroxychloroquine _____
4. ❑ leflunomide _____
5. ❑ methotrexate _____
6. ❑ NSAIDs or COX-2 _____
7. ❑ oral corticosteroids _____
8. ❑ penicillamine _____
9. ❑ 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.)

18. 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 the preferred cytokine and CAM antagonist drug was taken in the space provided.

1. _____

2. _____

SECTION IIIB — ADDITIONAL CLINICAL INFORMATION FOR SIMPONI REQUESTS

19. Will the member continue to take methotrexate in combination with Simponi? ❑ Yes ❑ No

SECTION IV — AUTHORIZED SIGNATURE

20. **SIGNATURE** — Prescriber	21. Date Signed

SECTION V — FOR PHARMACY PROVIDERS USING STAT-PA

22. National Drug Code (11 digits)	23. Days' Supply Requested (Up to 365 Days)

24. NPI

25. 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.)

Continued

SECTION V — FOR PHARMACY PROVIDERS USING STAT-PA (Continued)

26. Place of Service

27. Assigned PA Number

28. Grant Date	29. Expiration Date	30. Number of Days Approved

SECTION VI — ADDITIONAL INFORMATION

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

Reset Form