## FORWARDHEALTH

# PRIOR AUTHORIZATION DRUG ATTACHMENT FOR CYTOKINE AND CELL ADHESION MOLECULE (CAM) ANTAGONIST DRUGS FOR GIANT CELL ARTERITIS AND NON-RADIOGRAPHIC AXIAL SPONDYLOARTHRITIS (NR-AXSPA)

**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 Giant Cell Arteritis and Non-Radiographic Axial Spondyloarthritis (nr-axSpA) Instructions, F-01952A. Prescribers may refer to the Forms page of the ForwardHealth Portal at <u>forwardhealth.wi.gov/WIPortal/Subsystem/Publications/ForwardHealthCommunications.aspx?panel=Forms</u> for the completion instructions.

Pharmacy providers are required to have a completed Prior Authorization Drug Attachment for Cytokine and CAM Antagonist Drugs for Giant Cell Arteritis and nr-axSpA 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.

## **SECTION I – MEMBER INFORMATION**

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

2. Member ID 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.	Address -	Prescriber	(Street,	City,	State,	Zip+4	Code)
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10. Phone Number – Prescriber	11. National Provider Identifier – Prescriber
SECTION III – CLINICAL INFORMATION (Required for Al	I PA Requests)

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.



DT-PA120-120

SECTION III A – CLINICAL INFORMATION FOR GIANT CELL ARTERITIS ONLY				
13. Does the member have giant cell arteritis?		Yes		No
14. Is the prescription written by a rheumatologist or through a rheumatology consultation?		Yes		No
15. Is the member currently using the requested non-preferred drug?		Yes		No
If yes, indicate the approximate date therapy was started.				
16. Has the member taken Tyenne subQ for <b>at least three</b> consecutive months and experienced an unsatisfactory therapeutic response or a clinically significant adverse drug reaction?		Yes		No
If yes, list the Tyenne subQ dose and dates taken and describe the unsatisfactory theraped significant adverse drug reaction. If additional space is needed, continue documentation in				
Dose: Dates Taken:				
	tion.			
Describe the unsatisfactory therapeutic response or clinically significant adverse drug react		onton	onist	drug
Describe the unsatisfactory therapeutic response or clinically significant adverse drug react 17. Indicate the clinical reason(s) why the prescriber is requesting a non-preferred cytokine and SECTION III B – CLINICAL INFORMATION FOR NR-AXSPA ONLY		antag	onist o	drug.
17. Indicate the clinical reason(s) why the prescriber is requesting a non-preferred cytokine and	d CAM	antag Yes	onist o	drug. No
17. Indicate the clinical reason(s) why the prescriber is requesting a non-preferred cytokine and SECTION III B – CLINICAL INFORMATION FOR NR-AXSPA ONLY	d CAM			
17. Indicate the clinical reason(s) why the prescriber is requesting a non-preferred cytokine and         SECTION III B – CLINICAL INFORMATION FOR NR-AXSPA ONLY         18. Does the member have nr-axSpA?		Yes		No
<ul> <li>17. Indicate the clinical reason(s) why the prescriber is requesting a non-preferred cytokine and</li> <li>SECTION III B – CLINICAL INFORMATION FOR NR-AXSPA ONLY</li> <li>18. Does the member have nr-axSpA?</li> <li>19. Is the prescription written by a rheumatologist or through a rheumatology consultation?</li> </ul>		Yes Yes		No
<ul> <li>17. Indicate the clinical reason(s) why the prescriber is requesting a non-preferred cytokine and</li> <li>SECTION III B – CLINICAL INFORMATION FOR NR-AXSPA ONLY</li> <li>18. Does the member have nr-axSpA?</li> <li>19. Is the prescription written by a rheumatologist or through a rheumatology consultation?</li> <li>20. Is the member currently using the requested non-preferred drug?</li> </ul>		Yes Yes Yes		No No No
<ul> <li>17. Indicate the clinical reason(s) why the prescriber is requesting a non-preferred cytokine and</li> <li>SECTION III B – CLINICAL INFORMATION FOR NR-AXSPA ONLY</li> <li>18. Does the member have nr-axSpA?</li> <li>19. Is the prescription written by a rheumatologist or through a rheumatology consultation?</li> <li>20. Is the member currently using the requested non-preferred drug?</li> <li>If yes, indicate the approximate date therapy was started.</li> <li>21. Has the member taken Cimzia for at least three consecutive months and experienced an unsatisfactory therapeutic response or a clinically significant adverse drug reaction?</li> </ul>	CAM	Yes Yes Yes Yes or clir		No No No

Describe the unsatisfactory therapeutic response or clinically significant adverse drug reaction.

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

#### SECTION IV – AUTHORIZED SIGNATURE

23. SIGNATURE - Prescriber

24. Date Signed

## SECTION V – ADDITIONAL INFORMATION

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