



Cytokine and CAM Antagonists: T-Lymphocyte Inhibitors

Please fax this completed form to (833) 645-2734 OR mail to: Centene Pharmacy Services | 5 River Park Place East, Suite 210 | Fresno, CA 93720. You can also complete online at [CoverMyMeds.com](https://www.covermymeds.com).

Coordinated Care of Washington, Inc. (Apple Health) Preferred Drug list:

https://www.coordinatedcarehealth.com/content/dam/centene/centene-pharmacy/pdl/FORMULARY-CoordinatedCare_Washington.pdf

For policy criteria, see: https://www.coordinatedcarehealth.com/content/coordinatedcare/en_us/providers/resources/clinical-payment-policies.html/

Date of request:	Reference #:	MAS:	
Patient	Date of birth	ProviderOne ID or Coordinated Care ID	
Pharmacy name	Pharmacy NPI	Telephone number	Fax number
Prescriber	Prescriber NPI	Telephone number	Fax number
Medication and strength	Directions for use	Qty/Days supply	

1. Is this request for a continuation of therapy? Yes No

If yes, does patient have clinical documentation demonstrating disease stability or a positive clinical response? Yes No

2. Is this prescribed by, or in consultation with, any of the following? Check all that apply:

Dermatologist Hematologist Oncologist
 Rheumatologist Other. Specify: _____

3. Will the requested medication be used in combination with another Cytokine and CAM medication?

Yes No

4. If request is non-preferred, has patient had treatment with one or more preferred Cytokine and CAM medications on the Apple Health Preferred Drug List (AHPDL) that was ineffective, contraindicated or not tolerated?

Yes. List each medication and duration of trial:

Medication Name: _____ Duration: _____

Medication Name: _____ Duration: _____

Medication Name: _____ Duration: _____

No. Explain why a preferred product(s) have not been tried: _____

5. What is patient current weight: _____ kg Date taken: _____

6. Indicate patient's diagnosis and answer the associated questions as indicated:

Graft Versus Host Disease (questions 7-9)

Polyarticular Juvenile Idiopathic Arthritis (questions 10 – 11)

Psoriatic Arthritis (PsA) (questions 12 - 15)

Rheumatoid Arthritis (questions 16 - 18)

For diagnosis of Graft Versus Host Disease:

7. **If patient has received a hematopoietic stem cell transplant (HSCT):** Indicate the following for patient. Check all that apply:

- Requested drug will be used as additional therapy in combination with corticosteroids for chronic GVHD
 Patient has no response (e.g., steroid-refractory disease) to first-line therapy options

8. **If patient is undergoing a hematopoietic stem cell transplant (HSCT) from a matched or 1 allele-mismatched unrelated-donor:** Indicate the following for patient. Check all that apply:

- Requested drug will be used for prophylaxis of acute graft versus host disease (aGVHD)
 Requested drug will be used in combination with a calcineurin inhibitor and methotrexate
 Patient will receive antiviral prophylactic treatment for Epstein-Barr Virus (EBV) reactivation and prophylaxis will continue for 6 months post-transplantation

9. **If patient received the requested medication previously,** indicate the dates and duration of treatment:

Date(s) received: _____ Duration of treatment: _____

For diagnosis of Polyarticular Juvenile Idiopathic Arthritis

10. Has patient had treatment with at least one non-Cytokine and CAM DMARD (e.g., methotrexate, sulfasalazine, leflunomide, hydroxychloroquine, azathioprine, cyclosporine) that has been ineffective, unless all are contraindicated, or not tolerated [minimum trial of 3 months]? Yes No

11. **For continuation of therapy:** Has documentation been submitted demonstrating disease stability or a positive clinical response (e.g., improvement in joint pain, swelling, activities of daily living, reduction in diseases flares, etc.)? Yes No

For diagnosis of Psoriatic Arthritis

12. Has patient had treatment with at least one non-Cytokine and CAM disease-modifying antirheumatic drug (DMARD) that has been ineffective, contraindicated or not tolerated [minimum trial of 3 months]?
 Yes No

13. Does patient have presence of active, severe disease indicated by provider assessment?
 Yes No

14. Does patient have presence of any of the following? Check all that apply:

- Erosive disease
 Elevated C-reactive protein (CRP) or erythrocyte sedimentation rate (ESR)
 Long-term damage interfering with function (e.g., joint deformities, vision loss)
 Major impairment of quality of life due to high disease activity at many sites (including dactylitis, enthesitis) or functionally limiting arthritis at a few sites.

15. **For continuation of therapy:** Has documentation been submitted demonstrating disease stability or a positive clinical response (e.g., improvement in joint pain, swelling, activities of daily living, reduction in diseases flares, etc.)? Yes No

For diagnosis of Rheumatoid Arthritis (RA)

16. Have baseline assessments been submitted (e.g., Disease Activity Score for 28 joints (DAS28) with the CRP, DAS28 with ESR, Simplified Disease Activity Index (SDAI), Clinical Disease Activity Index (CDAI), Routine Assessment of Patient Index Data 3 (RAPID3), Patient Activity Scale (PAS) II)? Yes No
17. Has patient had treatment with at least one non-Cytokine and CAM DMARD (e.g., methotrexate, sulfasalazine, hydroxychloroquine, leflunomide, cyclosporine, azathioprine) that has been ineffective, unless all are contraindicated, or not tolerated [minimum trial of 3 months]? Yes No
18. **For continuation of therapy:** Has documentation been submitted demonstrating disease stability or a positive clinical response (e.g. improvement in DAS28 with CRP/ESR, SDAI, CDAI, RAPID3, PAS II scores)?
 Yes No

CHART NOTES ARE REQUIRED WITH THIS REQUEST

Prescriber signature	Prescriber specialty	Date
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Centene Pharmacy Services will respond via fax or phone within 24 hours of receipt of the request. Requests for prior authorization must include member name, ID#, and drug name. Please include lab reports with requests when appropriate (e.g., Culture and Sensitivity; Hemoglobin A1C; Serum Creatinine; CD4; Hematocrit; WBC, etc.)