



PRIOR AUTHORIZATION REQUEST
Simponi

PATIENT: Name, Address, City, State, Zip, D.O.B., Member ID
Prescriber: Name, Address, City, State, Zip, Phone, Fax, NPI

Medication Requested: Qty Requested:

Your patient's prescription benefit requires that we review certain requests for coverage with the prescriber. You have prescribed a medication for your patient that requires Prior Authorization before benefit coverage or coverage of additional quantities can be provided.

SECTION A: Please answer the following questions

- 1. Yes No Is the patient currently receiving Simponi (Aria or SC)?
2. Is the medication to be used in combination with a BIOLOGIC disease-modifying antirheumatic drug (DMARD) or a targeted synthetic DMARD?
3. What is the patient's diagnosis?
4. Yes No Is Simponi SC being prescribed by, or in consultation with, a rheumatologist?
5. Yes No Is Simponi SC being prescribed by, or in consultation with, a rheumatologist or dermatologist?
6. Yes No Has the patient had a trial of one conventional systemic agent or a corticosteroid such as prednisone or methylprednisolone, or was intolerant to one of these agents for ulcerative colitis?
7. Yes No Has the patient tried one conventional synthetic disease-modifying antirheumatic drug

If you have any questions, call: 800-753-2851



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(DMARD) for at least 3 months?

- 8. Yes No Has the patient tried one biologic for at least 3 months?
- 9. Yes No Has the patient tried a biologic?
- 10. Yes No Does the patient have pouchitis?
- 11. Yes No Has the patient tried therapy with an antibiotic, probiotic, corticosteroid enema, or Rowasa (mesalamine) enema?
- 12. Yes No Is the requested medication prescribed by or in consultation with a gastroenterologist?
- 13. Yes No Does the patient have arthritis primarily in the knees, ankles, elbows, wrists, hands and/or feet?
- 14. Yes No Has the patient tried at least ONE conventional synthetic DMARD?
- 15. Yes No Does the patient have axial spondyloarthritis?
- 16. Yes No Does the patient have objective signs of inflammation, defined as: a C-reactive protein (CRP) elevated beyond the upper limit of normal for the reporting laboratory?
- 17. Yes No Does the patient have objective signs of inflammation, defined as: sacroiliitis reported on magnetic resonance imaging (MRI)?
- 18. Yes No Has the patient had a response, as determined by the prescriber?
- 19. Yes No Has the patient had a response, as determined by the prescriber?
- 20. Yes No Has the patient had a response, as determined by the prescriber?
- 21. Yes No Has the patient had a response, as determined by the prescriber?

Please document the diagnoses, symptoms, and/or any other information important to this review:

SECTION B

Physician Signature

PHYSICIAN SIGNATURE

DATE

FAX COMPLETED FORM TO: 877-251-5896

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