

**Pharmacy Prior Approval Request for  
Immunomodulators: Simponi Aria**

**Beneficiary Information**

1. Beneficiary Last Name: \_\_\_\_\_ 2. First Name: \_\_\_\_\_  
3. Beneficiary ID #: \_\_\_\_\_ 4. Beneficiary Date of Birth: \_\_\_\_\_ 5. Beneficiary Gender: \_\_\_\_\_

**Prescriber Information**

6. Prescribing Provider NPI #: \_\_\_\_\_  
7. Requester Contact Information - Name: \_\_\_\_\_ Phone #: \_\_\_\_\_ Ext. \_\_\_\_\_

**Drug Information**

8. Drug Name: \_\_\_\_\_ 9. Strength: \_\_\_\_\_ 10. Quantity Per 30 Days: \_\_\_\_\_  
11. Length of Therapy (in days):  up to 30 Days  60 Days  90 Days  120 Days  180 Days  365 Days  Other \_\_\_\_\_

**Clinical Information****Request for Ankylosing Spondylitis**

1. Does the beneficiary have a diagnosis of Ankylosing Spondylitis?  **Yes**  **No**
2. Is the beneficiary not on another injectable biologic immunomodulator?  **Yes**  **No**
3. Has the beneficiary been considered and screened for the presence of latent tuberculosis infection?  **Yes**  **No**
4. Has the beneficiary been tested with Hep B SAG and Core Ab?  **Yes**  **No**
5. Has the beneficiary experienced inadequate symptom relief from treatment with at least two NSAIDS?  
 **Yes**  **No**
6. Is beneficiary unable to receive treatment with NSAIDS due to contraindications?  **Yes**  **No**
7. Does the beneficiary have clinical evidence of severe or rapidly progressing disease?  **Yes**  **No**
8. Has the beneficiary had a trial and failure of Cosentyx, Enbrel or Humira or a clinical reason beneficiary cannot try Cosentyx, Enbrel or Humira?  **Yes**  **No**

**Request for Polyarticular Juvenile Idiopathic Arthritis (PJIA)**

1. Does the beneficiary have a diagnosis of Polyarticular Juvenile Idiopathic Arthritis?  **Yes**  **No**
2. Is the beneficiary not on another injectable biologic immunomodulator?  **Yes**  **No**
3. Has the beneficiary been considered and screened for the presence of latent tuberculosis infection?  **Yes**  **No**
4. Has the beneficiary been tested with Hep B SAG and Core Ab?  **Yes**  **No**
5. Has the beneficiary tried one systemic corticosteroid (e.g. prednisone, methylprednisolone) or methotrexate, leflunomide or sulfasalazine with inadequate response or is unable to take these therapies due to contraindications?  
 **Yes**  **No**
6. Does the beneficiary have PJIA subtype enthesitis related arthritis?  **Yes**  **No**
7. Has the beneficiary had a trial and failure of Enbrel or Humira or a clinical reason beneficiary cannot try Enbrel or Humira?  **Yes**  **No**

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**Request for Psoriatic Arthritis**

1. Does the beneficiary have a documented definitive diagnosis of Psoriatic Arthritis?  **Yes**  **No**
2. Is the beneficiary 2 years of age or older?  **Yes**  **No**
3. Is the beneficiary not on another injectable biologic immunomodulator?  **Yes**  **No**
4. Has the beneficiary been considered and screened for the presence of latent tuberculosis infection?  **Yes**  **No**
5. Has the beneficiary been tested with Hep B SAG and Core Ab?  **Yes**  **No**
6. Does the beneficiary have a documented inadequate response or inability to take methotrexate  **Yes**  **No**
7. Has the beneficiary had a trial and failure of Cosentyx, Enbrel or Humira or a clinical reason beneficiary cannot try Cosentyx, Enbrel or Humira?  **Yes**  **No**

**Request for Rheumatoid Arthritis**

1. Does the beneficiary have a diagnosis of Rheumatoid Arthritis?  **Yes**  **No**
2. Is the beneficiary not on another injectable biologic immunomodulator?  **Yes**  **No**
3. Has the beneficiary been considered and screened for the presence of latent tuberculosis?  **Yes**  **No**
4. Has the beneficiary been tested with Hep B SAG and Core Ab?  **Yes**  **No**
5. Has the beneficiary experienced a therapeutic failure/inadequate response with methotrexate or at least one disease modifying antirheumatic drug (e.g. leflunomide, hydroxychloroquine, minocycline, sulfasalazine)?  
 **Yes**  **No**
6. Is the beneficiary unable to receive methotrexate or disease modifying antirheumatic drug due to contraindications or intolerabilities?  **Yes**  **No**
7. Does the beneficiary have clinical evidence of severe or rapidly progressing disease?  **Yes**  **No**
8. Has the beneficiary had a trial and failure of Enbrel or Humira or a clinical reason beneficiary cannot try Enbrel or Humira?  **Yes**  **No**

Signature of Prescriber: \_\_\_\_\_ Date: \_\_\_\_\_

**(Prescriber Signature Mandatory)**

I certify that the information provided is accurate and complete to the best of my knowledge, and I understand that any falsification, omission, or concealment of material fact may subject me to civil or criminal liability.