

Pharmacy Prior Approval Request for Immunomodulators: Cimzia

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:

Drug Information

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

Phone #: Ext.

Clinical Information

Request for Ankylosing Spondylitis

- 1. Does the beneficiary have a diagnosis of Ankylosing Spondylitis? \Box Yes \Box No
- 2. Is the beneficiary not on another injectable biologic immunomodulator? \Box Yes \Box 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? \Box Yes \Box No
- 5. Has the beneficiary experienced inadequate symptom relief from treatment with at least two NSAIDS?
 Yes No
- 6. Is the 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
- 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 Crohn's Disease (Adult)

- 1. Does the beneficiary have a diagnosis of moderate to severe Crohn's Disease?
 Yes
 No
- 2. Is the beneficiary not on another injectable biologic immunomodulator? \Box Yes \Box 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 had a trial and failure of Humira or a clinical reason beneficiary cannot try Humira? 🗆 Yes 🗆 No

Request for Plaque Psoriasis (Adult)

1. Does the beneficiary have a documented definitive diagnosis of moderate-to-severe Chronic Plaque Psoriasis? **Yes No**

- 2. Is the beneficiary 18 years of age or older? \Box Yes \Box No
- 3. Is the beneficiary not on another injectable biologic immunomodulator? \Box Yes \Box No
- 4. Has the beneficiary been considered and screened for the presence of latent tuberculosis infection (not required

for Otezla)?

Yes
No

5. Has the beneficiary been tested with Hep B SAG and Core Ab? \Box Yes \Box No

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6. Does the beneficiary have a body surface area (BSA) involvement of at least 3%?
Yes No

7. Does the beneficiary have involvement of the palms, soles, head and neck, or genitalia, causing disruption in normal daily activities and/or employment? **Yes No**

8. Has the beneficiary failed to respond to, or has been unable to tolerate phototherapy and **ONE** of the following medications or beneficiary has contraindications to these treatments: Soriatane (acitretin), Methotrexate, and/or Cyclosporine?
Yes
No

9. 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

10. Are the beneficiaries, Providers, and Pharmacies utilizing Siliq registered appropriately in the Siliq Risk Evaluation and Mitigation Strategy Program (REMS program)?
Yes
No

Request for Psoriatic Arthritis

1. Does the beneficiary have a documented definitive diagnosis of Psoriatic Arthritis?

Yes
No

- 2. Is the beneficiary 18 years of age or older (OR 2 years or older for Simponi Aria)?

 Yes
 No
- 3. Is the beneficiary not on another injectable biologic immunomodulator? \Box Yes \Box No

4. Has the beneficiary been considered and screened for the presence of latent tuberculosis infection (not required

for Otezla? Yes No

- 5. Has the beneficiary been tested with Hep B SAG and Core Ab (not required for Otezla? \Box Yes \Box 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 I No

Request for Rheumatoid Arthritis

- 1. Does the beneficiary have a diagnosis of Rheumatoid Arthritis? \Box Yes \Box No
- 2. Is the beneficiary not on another injectable biologic immunomodulator? \Box Yes \Box 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?

8. Has the beneficiary had a trial and failure of Enbrel or Humira or a clinical reason beneficiary cannot try Enbrel or Humira?

I Yes I No

Request for Non-Radiographic Axial Spondyloarthritis

1. Does the beneficiary have a diagnosis of Non-Radiographic Axial Spondyloarthritis?

2. Is the beneficiary not on another injectable biologic immunomodulator? \Box Yes \Box No

3. Has the beneficiary failed an adequate trial of a Non-Steroidal Anti-Inflammatory Drug (NSAID) unless contraindicated?

Yes
No

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- 4. Has the beneficiary been considered and screened for the presence of latent tuberculosis? \Box Yes \Box No
- 5. Has the beneficiary been tested with Hep B SAG and Core Ab? \Box Yes \Box No
- 6. Has the beneficiary had a trial and failure of Cosentyx?
 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.