

Pharmacy Prior Approval Request for Immunomodulators: Remicade and Infliximab

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 or is unable to receive treatment with NSAIDs due to contraindications or has clinical evidence of severe or rapidly progressing disease? Yes No
6. 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? 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 had a trial and failure of Humira or a clinical reason beneficiary cannot try Humira? Yes No

Request for Crohn's Disease (Pediatric)

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? 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 had a trial and failure of Humira or a clinical reason beneficiary cannot try Humira? Yes No

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

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? Yes 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)? 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

Request for Ulcerative Colitis (Adult)

1. Does the beneficiary have a diagnosis of ulcerative colitis? 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 had a trial and failure of Humira or a clinical reason beneficiary cannot try Humira? Yes No

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Request for Ulcerative Colitis (Pediatric)

1. Does the beneficiary have a diagnosis of ulcerative colitis? **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 had a trial and failure of Humira or a clinical reason beneficiary cannot try 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.