

Immunomodulators: Renflexis

Member Information

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

Request for Crohn's Disease (Adult)

1. Does the member have a diagnosis of moderate to severe Crohn's Disease? ☐ Yes ☐ No
2. Is the member not on another injectable biologic immunomodulator? ☐ Yes ☐ No
3. Has the member been considered and screened for the presence of latent tuberculosis infection? ☐ Yes ☐ No
4. Has the member been tested with Hep B SAG and Core Ab? ☐ Yes ☐ No
5. Has the member had a trial and failure of Humira or a clinical reason member cannot try Humira? ☐ Yes ☐ No

Request for Crohn's Disease (Pediatric)

1. Does the member have a diagnosis of moderate to severe Crohn's Disease? ☐ Yes ☐ No
2. Is the member not on another injectable biologic immunomodulator? ☐ Yes ☐ No
3. Has the member been considered and screened for the presence of latent tuberculosis infection? ☐ Yes ☐ No
4. Has the member been tested with Hep B SAG and Core Ab? ☐ Yes ☐ No
5. Has the member had a trial and failure of Humira or a clinical reason member cannot try Humira? ☐ Yes ☐ No

Request for Plaque Psoriasis (Adult)

1. Does the member have a documented definitive diagnosis of moderate-to-severe Chronic Plaque Psoriasis? ☐ Yes ☐ No
2. Is the member 18 years of age or older? ☐ Yes ☐ No
3. Is the member not on another injectable biologic immunomodulator? ☐ Yes ☐ No
4. Has the member been considered and screened for the presence of latent tuberculosis infection (not required for Otezla)? ☐ Yes ☐ No
5. Has the member been tested with Hep B SAG and Core Ab? ☐ Yes ☐ No
6. Does the member have a body surface area (BSA) involvement of at least 3%? ☐ Yes ☐ No
7. Does the member 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 member failed to respond to, or has been unable to tolerate phototherapy and **ONE** of the following medications or member has contraindications to these treatments: Soriatane (acitretin), Methotrexate, and/or Cyclosporine? ☐ Yes ☐ No
9. Has the member had a trial and failure of Cosentyx, Enbrel or Humira or a clinical reason member 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 member have a documented definitive diagnosis of Psoriatic Arthritis? ☐ Yes ☐ No
2. Is the member 18 years of age or older (OR 2 years or older for Simponi Aria)? ☐ Yes ☐ No
3. Is the member not on another injectable biologic immunomodulator? ☐ Yes ☐ No
4. Has the member been considered and screened for the presence of latent tuberculosis infection? ☐ Yes ☐ No
5. Has the member been tested with Hep B SAG and Core Ab? ☐ Yes ☐ No
6. Does the member have a documented inadequate response or inability to take methotrexate? ☐ Yes ☐ No
7. Has the member had a trial and failure of Cosentyx, Enbrel or Humira or a clinical reason member cannot try Cosentyx, Enbrel or Humira? ☐ Yes ☐ No

Request for Rheumatoid Arthritis

1. Does the member have a diagnosis of Rheumatoid Arthritis? ☐ Yes ☐ No
2. Is the member not on another injectable biologic immunomodulator? ☐ Yes ☐ No
3. Has the member been considered and screened for the presence of latent tuberculosis? ☐ Yes ☐ No
4. Has the member been tested with Hep B SAG and Core Ab? ☐ Yes ☐ No
5. Has the member 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 member unable to receive methotrexate or disease modifying antirheumatic drug due to contraindications or intolerabilities? ☐ Yes ☐ No
7. Does the member have clinical evidence of severe or rapidly progressing disease? ☐ Yes ☐ No
8. Has the member had a trial and failure of Enbrel or Humira or a clinical reason member cannot try Enbrel or Humira? ☐ Yes ☐ No

Request for Ulcerative Colitis (Adult)

1. Does the member have a diagnosis of ulcerative colitis? ☐ **Yes** ☐ **No**
2. Is the member not on another injectable biologic immunomodulator? ☐ **Yes** ☐ **No**
3. Has the member been considered and screened for the presence of latent tuberculosis? ☐ **Yes** ☐ **No**
4. Has the member been tested with Hep B SAG and Core Ab? ☐ **Yes** ☐ **No**
5. Has the member had a trial and failure of Humira or a clinical reason member 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.