Trillium Health Resources
Pharmacy Prior Approval Request for

Trillium HEALTH RESOURCES
Transforming Lives. Building Community Well-Being.

Immunomodulators: Actemra

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 Polyarticular Juvenile Idiopathic Arthritis (PJIA):

1. Does the member have a diagnosis of Polyarticular Juvenile Idiopathic 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 infection? ☐ **Yes** ☐ **No**
4. Has the member been tested with Hep B SAG and Core Ab? ☐ **Yes** ☐ **No**
5. Has the member 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 member have PJIA subtype enthesitis related arthritis? ☐ **Yes** ☐ **No**
7. 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 Systemic Onset Juvenile Idiopathic Arthritis (SJIA)

1. Does the member have a diagnosis of Systemic Juvenile Idiopathic 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 infection? ☐ **Yes** ☐ **No**
4. Has the member been tested with Hep B SAG and Core Ab? ☐ **Yes** ☐ **No**
5. Does the member have systemic arthritis with active systemic features and features of poor prognosis, as determined by the prescribing physician (e.g. arthritis of the hip, radiographic damage)? ☐ **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 infection? ☐ **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 either Enbrel or Humira? ☐ **Yes** ☐ **No**

Request for Giant Cell Arteritis:

1. Does the member have a diagnosis of Giant Cell Arteritis? ☐ **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**

Request for Cytokine Release Syndrome:

1. Does the member have a diagnosis of Cytokine Release Syndrome? ☐ **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**

Request for Systemic Sclerosis-Associated Interstitial Lung Disease (SSc-ILD)

1. Does the member have a diagnosis of Systemic Sclerosis-Associated Interstitial Lung 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**

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.