

**NC Medicaid
Pharmacy Prior Approval
Immunomodulators: Actemra**

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 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

Request for Systemic Onset Juvenile Idiopathic Arthritis (SJIA)

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

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- 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 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 either Enbrel or Humira? Yes No

Request for Giant Cell Arteritis:

- 1. Does the beneficiary have a diagnosis of Giant Cell Arteritis? 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

Request for Cytokine Release Syndrome:

- 1. Does the beneficiary have a diagnosis of Cytokine Release Syndrome? 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

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

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

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.