Trillium Health Resources
Pharmacy Prior Approval Request for

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

Immunomodulators: Kineret

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 Neonatal Onset Multisystem Inflammatory Disease (NOMID)
1. Does the member have a diagnosis of neonatal-onset multisystem inflammatory 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**

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 Enbrel or Humira? ☐ **Yes** ☐ **No**

Request for Deficiency of Interleukin-1 Receptor Antagonist (DIRA)
1. Does the member have a diagnosis of Deficiency of Interleukin-1 Receptor Antagonist (DIRA)? ☐ **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** ☐ **N0**

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.