

Pharmacy Prior Authorization Form

Last Reviewed: Sept. 11

For Prior Authorization please fax to: (877)974-4411 toll free, or (616)942-8206

This form applies to: Commercial Plan Medicaid Plan Medicare Plan

Kineret[®] (anakinra) Urgent Non-urgent

Member Name:	Member #:
DOB:	Gender:
Provider Name:	Provider Phone:
Provider Office Address:	
Provider Office Contact Name:	Provider Fax:
Provider Signature:	Provider NPI:
Date:	Member's PCP:

Product:

Kineret single use Prefilled Syringe - 100 mg

Dose: _____ Start date: _____

Prescriber is a rheumatologist:

Yes

No

Priority Health Precertification Requirements:

Authorization of Kineret requires:

- Diagnosis of rheumatoid arthritis
 - Documented therapeutic trial of at least one DMARD
 - Documented therapeutic trial of Enbrel
- Negative TB test (must be done yearly)
- Patient must **not** have moderate to severe heart failure

Continuation of Kineret therapy requires:

- Patient must be compliant taking the medication as prescribed
- Patient must be tolerating the medication
- Patient must not be experiencing any severe adverse reactions while taking the medication
- Patient must be responding positively to the medication
- Patient must have a negative TB test within the past 12 months

Please Complete the Following Information:

Diagnosis:

Rheumatoid Arthritis

Other: _____ Please provide rationale for use: _____

Results of annual (within the past 12 months) TB test:

Positive - Rationale for use: _____

Negative

Test not done – Rationale for use: _____

Patient has moderate to severe heart failure:

- Yes – Rationale for use: _____
- No

New request or continuation of therapy:

- New (see section 1)
- Continuation (see section 2)

Section 1 – New requests:

Rheumatoid Arthritis

Patient has had a therapeutic trial of at least one of the following DMARDS:

- Yes

	Dose	Dates	Outcome
<input type="checkbox"/> azathioprine	_____	_____	_____
<input type="checkbox"/> cyclosporine	_____	_____	_____
<input type="checkbox"/> d-penicillamine	_____	_____	_____
<input type="checkbox"/> gold sodium thiomalate	_____	_____	_____
<input type="checkbox"/> auranofin	_____	_____	_____
<input type="checkbox"/> aurothioglucose	_____	_____	_____
<input type="checkbox"/> hydroxychloroquine	_____	_____	_____
<input type="checkbox"/> leflunomide	_____	_____	_____
<input type="checkbox"/> methotrexate	_____	_____	_____
<input type="checkbox"/> sulfasalazine	_____	_____	_____

- No – Rationale for use: _____

Patient has had a documented therapeutic trial and clinical failure of Enbrel:

- Yes Enbrel Dose: _____ Trial Dates: _____
- No – Rationale for use: _____

Section 2 – Requests for continuation of therapy:

- The patient is compliant in taking the medication as scheduled
- The patient tolerated the medication
- The patient did not experience any severe adverse reactions while taking the medication
- The patient has responded to treatment, as determined by the prescribing physician
- The patient has had a negative TB test result within the past 12 months Date of test: _____

*** All fields must be complete and legible for Prior Authorization Review***

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YOUR OFFICE WILL RECEIVE A RESPONSE VIA FAX