



Instructions:

This form is used by Kaiser Permanente and/or participating providers for coverage of **Kevzara (sarilumab)**. Please complete and fax this form back to Kaiser Permanente within 24 hours [fax: [1-866-331-2104](tel:1-866-331-2104)]. If you have any questions or concerns, please call [1-866-331-2103](tel:1-866-331-2103). **Requests will not be considered unless this form is complete. The KP-MAS Formulary can be found at: [Pharmacy | Community Provider Portal | Kaiser Permanente](#)**

1 – Patient Information

Patient Name: _____ Kaiser Medical ID#: _____ Date of Birth: _____

2 – Provider Information

Provider Name: _____ Specialty: _____ Provider NPI: _____

Provider Address: _____

Provider Phone #: _____ Provider Fax #: _____

Please check the boxes that apply:

- Initial Request Continuation of Therapy Request

3 – Pharmacy Information

Pharmacy Name: _____ Pharmacy NPI: _____

Pharmacy Phone # _____ Pharmacy Fax #: _____

4 – Drug Therapy Requested

Drug 1: Name/Strength/Formulation: _____

Sig: _____

Drug 2: Name/Strength/Formulation: _____

Sig: _____

5– Diagnosis/Clinical Criteria

Initial Therapy:

1. Does the member have diagnosis of one of the following? **AND**
 - Rheumatoid Arthritis (RA)
 - Diagnosis of Polymyalgia Rheumatica (PMR)
 - Other: _____
2. Is the patient ≥ 18 years old? **AND**
 - No Yes
3. Is this prescribed by or in consultation with a rheumatologist? **AND**
 - No Yes

If this is being used for Rheumatoid Arthritis (RA):

4. Does the patient have a history of failure, contraindication, or intolerance to one non-biologic disease-modifying anti-rheumatic drug (DMARD) [e.g., Rheumatrex /Trexall (methotrexate), Arava (leflunomide), Azulfidine (sulfasalazine)]? **AND**
 - No Yes If yes, list the products and the outcome of therapy: _____

If this is being used for Polymyalgia Rheumatica (PMR):

5. Does the patient have a history of failure, contraindication, or intolerance to corticosteroids?
 - No Yes

Renew Criteria:

1. Is the patient receiving Kevzara in combination with any of the following?
 - i. Biologic DMARD [e.g., Enbrel (etanercept), Humira (adalimumab), Cimzia (certolizumab), Simponi (golimumab)]
 - ii. Janus kinase inhibitor [e.g., Xeljanz (tofacitinib)]
 - iii. Phosphodiesterase 4 (PDE4) inhibitor [e.g. Otezla (apremilast)]
 - No Yes (if yes, PA will not be approved)
2. Does the member have a documented clinically significant benefit from medication?
 - No Yes

6 – Provider Sign-Off

Additional Information –

1. Please submit chart notes/medical records for the patient that are applicable to this request.
2. If member has not tried preferred agent(s) please provide rationale/explanation and any additional supporting information that should be taken into consideration for the requested medication:

I certify that the information provided is accurate. Supporting documentation is available for State audits.

Provider Signature:

Date:

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