

Kaiser Permanente Health Plan of Mid-Atlantic States, Inc.
Kevzara (sarilumab) Prior Authorization (PA)
Pharmacy Benefits Prior Authorization Help Desk
Length of Authorizations: Initial- 12 months; Continuation- 12 months

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: <u>1-866-331-2104</u>]. If you have any questions or concerns, please call <u>1-866-331-2103</u>. **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

	5- Diagnosis/ Clinical Criteria
□ Initia 1.	Does the member have diagnosis of one of the following? AND Rheumatoid Arthritis (RA), moderate to severe
	□ Diagnosis of Polymyalgia Rheumatica (PMR)
	□ Polyarticular juvenile idiopathic arthritis (pJIA) in pts weighing ≥ 63 kg
	□ Other:
2.	,
Rheum	□ No □ Yes natoid Arthritis (RA):
1.	Is the patient ≥18 years old? AND □ No □ Yes
	Does the patient have a history of failure, contraindication, or intolerance to one non-biologic disease-modifying anti- eumatic drug (DMARD) [e.g., Rheumatrex /Trexall (methotrexate), Arava (leflunomide), Azulfidine (sulfasalazine)? AND □ No □ Yes If yes, list the products and the outcome of therapy:
Polym	yalgia Rheumatica (PMR):
2.	Is the patient ≥18 years old? AND
	□ No □ Yes
3.	2. Does the patient have a history of failure, contraindication, or intolerance to corticosteroids?
	□ No □ Yes
Polyar	ticular juvenile idiopathic arthritis (pJIA) in pts weighing ≥ 63 kg
1.	Is patient's weight ≥ 63 kg on the day of therapy being prescribed?
	No □ Yes
	Does the patient have a history of failure, contraindication, or intolerance to one non-biologic disease-modifying antieumatic drug (DMARD) [e.g., Rheumatrex /Trexall (methotrexate), Arava (leflunomide), Azulfidine (sulfasalazine)?
	No □ Yes
□ Rene	ew Criteria for:
Rheum	natoid Arthritis (RA) or Polymyalgia Rheumatica (PMR):
	 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) Does the member have a documented clinically significant benefit from medication? □ No □ Yes
Polyar	ticular juvenile idiopathic arthritis (pJIA) in pts weighing ≥ 63 kg
1.	Positive clinical response to Kevzara therapy □ No □ Yes

6 - Provider Sign-Off

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Additional Information –		
1. Please submit chart notes/medical records for the patient that are applicable to this request.		
If member has not tried preferred agent(s) please provide ration information that should be taken into consideration for the requ		
I certify that the information provided is accurate. Supporting	documentation is available for State audits.	
Provider Signature:	Date:	
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