

Kaiser Permanente Health Plan of Mid-Atlantic States, Inc.
Olumiant (baricitinib) 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 **Olumiant (baricitinib).** 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 – Prescriber Information				
Is the prescriber a rheumatologist or dermatologist? □ No □ Yes				
If consulted with a specialist, specialist name and specialty:				
Prescriber Name:	Specialty:	NPI:		
Prescriber Address:				
Prescriber Phone #:	Prescriber Fax #:			
Do you have an approved provider referral number from Kaiser Permanente? □ Yes — please provide your provider referral number here:				
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:				

5- Diagnosis/Clinical Criteria

1.	. Is this request for initial or continuing therapy?		
	□ Initial therapy □ Continuing therapy, state start date:		
2.	Indicate the patient's diagnosis for the requested medication:		
Clir	nical Criteria:		
Rhe	eumatology:		
	Patient is ≥18 years old and has a diagnosis of moderate-to-severe rheumatoid arthritis,		
	□ No □ Yes		
2.	AND patient has a history of treatment failure after an adequate trial (≥3 months), intolerance or contraindication to at least one agent in each of the following categories:		
	 At least 1 non-biologic disease-modifying antirheumatic drug (DMARD) such as methotrexate, 		
	leflunomide, sulfasalazine or hydroxychloroquine		
	 At least 1 TNF-alpha biologic DMARD: e.g., Enbrel (etanercept), adalimumab biosimilars (Amjevita preferred) or 		
	Humira		
	O Xeljanz (tofacitinib)□ No □ Yes		
Der	rmatology (Alopecia Areata):		
 Patient is ≥ 18 years, 			
	□ No □ Yes		
2.	AND diagnosis of alopecia areata (with <50% scalp involvement, mild facial involvement, not rapidly progressive, not alopecia totalis/universalis), □ No □ Yes		
3.	AND has tried a 2-month trial of all of the following unless clinically significant adverse effects, contraindication, or clinical reason to avoid treatment: O Topical corticosteroid, O AND topical calcineurin inhibitor, O AND topical minoxidil, O AND intralesional Kenalog, O AND topical JAK inhibitor		
	□ No □ Yes		
4.	AND patient has tried a 3-month trial of at least one of the systemic immunosuppressants such as methotrexate or cyclosporine unless clinically significant adverse effects, contraindication, or clinical reason to avoid treatment No Yes		
O	R—		
1.	Patient is ≥ 18 years, □ No □ Yes		
2.	AND diagnosis of alopecia areata (with >50% scalp involvement, disfiguring facial involvement, rapidly progressive, alopecia totalis/universalis), □ No □ Yes		
3.	AND patient has tried a 3-month trial of at least one of the systemic immunosuppressants such as methotrexate or cyclosporine unless clinically significant adverse effects, contraindication, or clinical reason to avoid treatment \Box No \Box Yes		

For Continuation of Therapy, please respond to <u>additional questions</u> below:			
1. Member has documentation of positive clinical response, $\hfill\Box$ No $\hfill\Box$ Yes			
2. AND specialist follow-up has occurred in the past 12 months since last r □ No □ Yes	review		
6 – Prescriber Sign-Off			
 Additional Information – 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 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.			
Prescriber Signature:	Date:		
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