

Kaiser Permanente Health Plan of Mid-Atlantic States, Inc.
Tremfya (guselkumab) Prior Authorization (PA)
Pharmacy Benefits Prior Authorization Help Desk
Length of Authorizations: Initial- 6 months; Continuation- 12 months

Instructions:

This form is used by Kaiser Permanente and/or participating providers for coverage of **Tremfya (guselkumab)**. 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.**

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, speci	alist name and specialty:	
Prescriber Name:	Specialty:	NPI:
Prescriber Address:		
Prescriber Phone #:	Prescriber Fax #:	
, , , , , , , , , , , , , , , , , , , ,	referral number from Kaiser Permanente? er referral number here:	
	3 – Pharmacy Information	
Pharmacy Name:	Pharmacy NPI:	
Pharmacy Phone #	Pharmacy Fax #:	
	4 – Drug Therapy Requested	
	on:	
Sig:		
Drug 2: Name/Strength/Formulation	on:	

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:
Cli	nical Criteria:
Pso	oriatic Arthritis
1.	Does the member have a diagnosis of active psoriatic arthritis? □ No □ Yes
2.	Does the member have documented inadequate response (of at least a 3 month trial), intolerance, or contraindication to BOTH of the following? a. ONE or more tumor necrosis factor (TNF alpha) inhibitors: Inflectra or Remicade (infliximab), Enbrel*PA
	 (etanercept), adalimumab biosimilars (Amjevita preferred) or Humira*PA b. Cosentyx*PA (secukinumab) □ No □ Yes
Pla	ique Psoriasis
	Does the member have a diagnosis of moderate-to-severe plaque psoriasis? □ No □ Yes
2.	Has the member had inadequate response or contraindication to at least a 3-month trial of phototherapy unless involvement in sensitive areas (e.g., face, body folds, etc.)? \Box No \Box Yes
3.	Has the member failed at least a 3-month trial of one of the following unless clinically significant adverse effects, contraindication or clinical reason to avoid treatment (i.e. pregnancy/breastfeeding, history of alcoholism or alcoholic liver disease, chronic liver disease, immunodeficiency syndrome, pre-existing blood dyscrasia, hemodialysis, or end-stage renal disease)? a. Methotrexate b. Acitretin □ No □ Yes
4.	Is there documentation of inadequate response (at least 3-month trial), intolerance, or contraindication to ALL of the following?
	 a. At least one TNF inhibitor [i.e. adalimumab (Amjevita preferred) or infliximab product (Inflectra preferred)] b. Secukinumab (Cosentyx)*PA No Yes
*PA	This medication is also subject to PA review
Fo	r continuation of therapy, please respond to <u>additional questions</u> below:
1.	Has the member had a positive clinical response to medication? $\hfill\Box$ No $\hfill\Box$ Yes
2.	Has specialist follow-up occurred in the past 12 months since last review? □ No □ Yes

6 - Prescriber Sign-Off

dditional 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 information that should be taken into consideration for the	
I certify that the information provided is accurate. Supporting doc	
I certify that the information provided is accurate. Supporting doce Prescriber Signature:	umentation is available for State audits. Date: