

Kaiser Permanente Health Plan of Mid-Atlantic States, Inc.
Tyrvaya (varenicline) 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 **Tyrvaya (varenicline)** for **Commercial, Exchange, FEHB (Federal),** and **MD Medicaid** plans. <u>Please complete all sections, incomplete forms will delay processing.</u> 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: <u>Pharmacy</u>**

Community Provider Portal | Kaiser Permanente

	1 – Patient Information		
Patient Name:	Kaiser Medical ID#:	Date of Birth:	
2 – Provider Information			
Provider Name:	Specialty:	NPI:	
Provider Address:			
Provider Phone #:	Provider 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/Formulatio	n:		
Sig:			
Drug 2: Name/Strength/Formulatio	n:		
Sig:			
	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:			

Cli	nical Criteria:		
1.	Is this medication being prescribed by an Ophthalmologist or Optometrist? $\hfill\Box$ No $\hfill\Box$ Yes		
2.	Is the patient ≥18 years of age with a diagnosis of dry eye disease? □ No □ Yes		
3.	Has the patient tried and failed at least one month of one OTC ocular lubricant (e.g., artificial tears, lubricating gels/ointments)? \Box No \Box Yes		
4.	Has the patient had an adequate trial (3 months' treatment duration) and failure to ALL of the following? ○ Cyclosporine 0.05% ophthalmic solution (generic Restasis) ○ Cequa 0.09% ○ Xiidra 5% □ No □ Yes		
5.	 Is the patient concomitantly using an ophthalmic cyclosporine product (Cequa, Restasis, Vevye) OR Xiidra (lifitegrast ophthalmic solution) (therapy will not be approved if so)? □ No □ Yes 		
Foi	continuation of therapy, please respond to <u>additional questions</u> below. New members who were initiated on therapy		
	side of Kaiser, who have not been reviewed previously, must meet all above Clinical Criteria.		
	Is there documentation of positive clinical response to therapy (e.g., increased tear production or improvement in dry eye symptoms)? \Box No \Box Yes		
	6 – Provider Sign-Off		
	ditional 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:		
	certify that the information provided is accurate. Supporting documentation is available for State audits.		
	vider Signature: Date:		
Plea	se Note: This document contains confidential information, including protected health information, intended for a specific individual and purpose. The information is		

private and legally protected by law, including HIPAA. If you are not the intended recipient, you are hereby notified that any disclosure, copying, distribution or taking of any action in reliance on the contents of this telecopied information is strictly prohibited. Please notify sender if document was not intended for receipt by your facility