

Kaiser Permanente Health Plan of Mid-Atlantic States, Inc.
Takhzyro (lanadelumab-flyo) 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 **Takhzyro (lanadelumab-flyo)** for **Commercial, Exchange, FEHB (Federal),** and **MD Medicaid** plans. Please complete all sections, incomplete forms will delay processing. Fax this form back to Kaiser Permanente within 24 hours (fax: 1-866-331-2104). If you have any questions or concerns, please call 1-866-331-2103. **Requests will not be considered unless all sections are complete. KP-MAS Formulary can be found at:** Pharmacy | Community Provider Portal | Kaiser Permanente

1 – Patient Information 2 – Prescriber Information 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: 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:								
Clinical Critoria									
	Clinical Criteria: 1. Is the prescriber an Allergist, Immunologist, or Physician specializing in the treatment of hereditary angioedema (HAE)? □ No □ Yes								
2.	Is the patient ≥2 years of age? □ No □ Yes								
3.	 Does the patient have a diagnosis of HAE type I or type II as confirmed by ONE of the following? Mutation known to cause HAE in either the SERPING1 or F12 gene, OR A C4 level below the lower limit of normal and/or a C1 inhibitor (C1-INH) antigenic level or functional level below the lower limit of normal No Yes 								
4.	Does provider attest that Takhzyro is NOT being used in combination with other products indicated for prophylaxis against HAE attacks? □ No □ Yes								
For continuation of therapy, please respond to <u>additional questions</u> below. New members who were initiated on therapy outside of Kaiser, who have not been reviewed previously, must meet all above Clinical Criteria. 1. Has specialist follow-up occurred in the past 12 months with documented improvement while on Takhzyro? □ No □ Yes									
2.	Does provider attest that Takhzyro is NOT being used in combination with other products indicated for prophylaxis against HAE attacks? \Box No \Box Yes								
3.	If patient has been stable (i.e. attack free) for >6 months: Is there documentation that extending the dosing interval to 300 mg every 4 weeks has been considered, OR is patient's current dosing already 300 mg every 4 weeks? □ No □ Yes								
6 – Prescriber 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.									
	scriber Signature: Date:								

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