Kaiser Permanente Health Plan of Mid-Atlantic States, Inc.
TRIKAFTA (Elexacaftor-Tezacaftor-Ivacaftor) 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 **TRIKAFTA (Elexacaftor-Tezacaftor-Ivacaftor) .** 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: http://www.providers.kaiserpermanente.org/mas/formulary.html

1 – Patient Information

Patient Name: ______________________________ Kaiser Medical ID#: _________________ Date of Birth: _____________

2 – Prescriber Information

Prescriber Name: ______________________________ Specialty: ____________________________ NPI: _________________

Prescriber Address: ____

Prescriber Phone #: __________________________Prescriber Fax #: ___________________________

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 date: ________________________
2. Indicate the Member's diagnosis for the requested medication: ____
3. Is the member ≥12 years of age? **AND**
   □ No □ Yes
4. Was the member diagnosis of CF confirmed by a clinician with expertise in providing CF care? **AND**
   □ No □ Yes
5. At least one F508del mutation in the CFTR gene detected using either an FDA-cleared CF mutation test or testing was completed by a CLIA certified laboratory? **AND**
   □ No □ Yes
6. Member does not have either of the following:
   a. Severe liver impairment (Child-Pugh Class C), **OR**
   b. Prior solid organ or hematological transplantation, unless use of the medication is approved by the transplant center

   □ No □ Yes

For Continuation of Therapy, Please Respond to Additional Questions Below:

1. Was there documentation of positive clinical response? **AND**
   □ No □ Yes
2. Did the specialist follow-up occur in the past 12 months? **AND**
   AST, ALT, bilirubin and ophthalmic changes (patients up to 17 years) are monitored at least annually
   □ No □ Yes

6 – Prescriber Sign-Off

Additional Information – Please submit chart notes/medical records for the patient that are applicable to this request. Provide any additional supporting information that should be taken into consideration:

____

I certify that the information provided is accurate. Supporting documentation is available for State audits.

Prescriber Signature:	**Date:**

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