

TRICARE Prior Authorization Request Form for  
elexacaftor - tezacaftor - ivacaftor (**Trikafta**)



**JOHNS HOPKINS**  
HEALTH PLANS

7231 Parkway Drive, Suite 100, Hanover, MD 21076

## USFHP Pharmacy Prior Authorization Form

To be completed by requesting provider	
Drug Name:	Strength:
Dosage/Frequency (SIG):	Duration of Therapy:

**FAX Completed Form and  
Applicable Progress Notes to:**  
(410) 424-4037

**Questions?** Contact the Pharmacy Dept at: (888) 819-1043, option 4

Clinical Documentation must accompany form in order for a determination to be made.

**Step 1** Please complete patient and physician information (please print):

Patient Name: _____	Physician Name: _____
Address: _____	Address: _____
Sponsor ID #: _____	Phone #: _____
Date of Birth: _____	Secure Fax #: _____

**Step 2** Please complete the clinical assessment:

1. Is the requested medication being prescribed by or in consultation with a pulmonologist?	<input type="checkbox"/> Yes Proceed to question 2	<input type="checkbox"/> No <b>STOP</b> Coverage not approved
2. Is Trikafta being prescribed for the treatment of cystic fibrosis (CF)?	<input type="checkbox"/> Yes Proceed to question 3	<input type="checkbox"/> No <b>STOP</b> Coverage not approved
3. Is this drug being requested for an FDA approved age?	<input type="checkbox"/> Yes Proceed to question 4	<input type="checkbox"/> No <b>STOP</b> Coverage not approved
4. Does the patient have at least one F508del mutation in the cystic fibrosis transmembrane conductance regulator (CFTR) gene as detected by an FDA-approved CF mutation test?	<input type="checkbox"/> Yes Proceed to question 6	<input type="checkbox"/> No Proceed to question 5
5. Does the patient have a mutation in the CFTR gene that is responsive based on in vitro data?	<input type="checkbox"/> Yes Proceed to question 6	<input type="checkbox"/> No <b>STOP</b> Coverage not approved
6. Is the genotype known or unknown?	<input type="checkbox"/> Known - Proceed to question 8 <input type="checkbox"/> Unknown - Proceed to question 7	

7. Has an FDA-approved test been used to detect the presence of at least one F508del mutation or a mutation that is responsive based on in vitro data?	<input type="checkbox"/> Yes Proceed to question 8	<input type="checkbox"/> No <b>STOP</b> Coverage not approved
8. Is this agent being used in combination therapy with Symdeko, Orkambi or Kalydeco?	<input type="checkbox"/> Yes <b>STOP</b> Coverage not approved	<input type="checkbox"/> No <b>Sign and date below</b>

**Step 3** I certify the above is true to the best of my knowledge. Please sign and date:

**3**

\_\_\_\_\_

Prescriber Signature

\_\_\_\_\_

Date

[09 June 2021]

<b>For Internal Use Only</b>	
<input type="checkbox"/> Approved:	Duration of Approval: ____month(s)
<input type="checkbox"/> Denied:	Authorized By:
<input type="checkbox"/> Incomplete/Other:	PA#:
Date Faxed to MD:	Date Decision Rendered: