

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



JOHNS HOPKINS
HEALTH PLANS

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

USFHP Pharmacy Prior Authorization Form

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

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 Symdeko being prescribed for the treatment of cystic fibrosis?	<input type="checkbox"/> Yes Proceed to question 2	<input type="checkbox"/> No STOP Coverage not approved
2. Is this drug being requested for an FDA approved age?	<input type="checkbox"/> Yes Proceed to question 3	<input type="checkbox"/> No STOP Coverage not approved
3. Is the requested medication prescribed by or in consultation with a pulmonologist?	<input type="checkbox"/> Yes Proceed to question 4	<input type="checkbox"/> No STOP Coverage not approved
4. Is the patient homozygous for the F508del mutation in the cystic fibrosis transmembrane conductance regulator (CFTR) gene?	<input type="checkbox"/> Yes Proceed to question 6	<input type="checkbox"/> No Proceed to question 5
5. Does the patient have at least one mutation in the cystic fibrosis transmembrane conductance regulator (CFTR) gene that is responsive to Symdeko potentiation based on in vitro data and/or clinical evidence?	<input type="checkbox"/> Yes Proceed to question 6	<input type="checkbox"/> No STOP 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 a CFTR mutation followed by verification with bi-directional sequencing when recommended by the mutation test instructions for use?	<input type="checkbox"/> Yes Proceed to question 8	<input type="checkbox"/> No STOP Coverage not approved
8. Is this agent being used in combination therapy with Orkambi, Kalydeco, or Trikafta?	<input type="checkbox"/> Yes STOP Coverage not approved	<input type="checkbox"/> No Sign and date below

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

_____ Date
Prescriber Signature

[09 June 2021]

For Internal Use Only	
<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: