

Cystic Fibrosis Agents, Oral - Washington Prior Authorization Request Form

Please complete this **entire** form and fax it to: **866-940-7328**. If you have questions, please call **800-310-6826**.

**This form may contain multiple pages. Please complete all pages to avoid a delay in our decision.
Allow at least 24 hours for review.**

Section A – Member Information

First Name:	Last Name:	Member ID:
Address:		
City:	State:	ZIP Code:
Phone:	DOB:	Allergies:
Primary Insurance Information (if any):		
Is the requested medication: <input type="checkbox"/> New or <input type="checkbox"/> Continuation of Therapy? If continuation, list start date: _____		
Is this patient currently hospitalized? <input type="checkbox"/> Yes <input type="checkbox"/> No If recently discharged, list discharge date: _____		

Section B - Provider Information

First Name:	Last Name:	M.D./D.O.
Address:	City:	State: ZIP code:
Phone:	Fax:	NPI #: Specialty:
Office Contact Name / Fax attention to:		

Section C - Medical Information

Medication:	Strength:
Directions for use:	Quantity:
Diagnosis (Please be specific & provide as much information as possible):	ICD-10 CODE:
Is this member pregnant? <input type="checkbox"/> Yes <input type="checkbox"/> No If yes, what is this member's due date? _____	

Section D – Previous Medication Trials

Medication Name	Strength	Directions	Dates of Therapy	Reason for failure / discontinuation

Section E – Additional information and Explanation of why preferred medications would not meet the patient's needs: Please refer to the patient's PDL for a list of preferred alternatives

Member First name:	Member Last name:	Member DOB:
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Clinical and Drug Specific Information

ALL REQUESTS:

- Does the patient have a diagnosis of cystic fibrosis? Yes No

- Has the patient demonstrated failure or intolerance to a majority of the preferred oral Cystic Fibrosis agents?
 Yes No
 (If yes, complete Section D above with medication information, including dose, date of trial, and reason for discontinuation)

- Does the patient have confirmation of 2 copies of the F508del mutation in the CFTR gene (i.e. the patient is homozygous for the F508del mutation)? Yes No

- Does the patient have documentation of at least ONE mutation in the CFTR gene that is responsive to the requested medication potentiation? Yes No

<i>A1067T</i> <i>c.3199G>A</i>	<i>G1244E</i> <i>c.3731G>A</i>	<i>R352Q</i> <i>c.1055G>A</i>
<i>A455E</i> <i>c.1364C>A</i>	<i>G1349D</i> <i>c.4046G>A</i>	<i>R74W</i> <i>c.220C>T</i>
<i>D110E</i> <i>c.330C>A</i>	<i>G178R</i> <i>c.532G>A</i>	<i>S1251N</i> <i>c.3752G>A</i>
<i>D110H</i> <i>c.328G>C</i>	<i>G551D</i> <i>c.1652G>A</i>	<i>S1255P</i> <i>c.3763T>C</i>
<i>D1152H</i> <i>c.3454G>C</i>	<i>G551S</i> <i>c.1651G>A</i>	<i>S549N</i> <i>c.1646G>A</i>
<i>D1270N</i> <i>c.3808G>A</i>	<i>K1060T</i> <i>c.3179A>C</i>	<i>S549R</i> <i>c.1645A>C,</i> <i>c.1647T>G</i>
<i>D579G</i> <i>c.1736A>G</i>	<i>L206W</i> <i>c.617T>G</i>	<i>S945L</i> <i>c.2834C>T</i>
<i>E193K</i> <i>c.577G>A</i>	<i>P67L</i> <i>c.200C>T</i>	<i>S977F</i> <i>c.2930C>T</i>
<i>E56K</i> <i>c.166G>A</i>	<i>R1070Q</i> <i>c.3209G>A</i>	<i>2789+5G→A</i> <i>c.2657+5G>A</i>
<i>E831X</i> <i>c.2491G>T</i>	<i>R1070W</i> <i>c.3208C>T</i>	<i>3272-26A→G</i> <i>c.3140-26A>G</i>
<i>F1052V</i> <i>c.3154T>G</i>	<i>R117C</i> <i>c.349C>T</i>	<i>3849+10kbC→T</i> <i>c.3718-2477C>T</i>
<i>F1074L</i> <i>c.3222T>A</i>	<i>R117H</i> <i>c.350G>A</i>	<i>711+3A→G</i> <i>c.579+3A>G</i>
<i>G1069R</i> <i>c.3205G>A</i>	<i>R347H</i> <i>c.1040G>A</i>	<i>F508del/F508del</i> <i>c.1521_1523delCTT</i>

Requests for CONTINUATION OF THERAPY:

- Does the patient meet one of the following: Yes No (check which applies)
 - FEV₁ has significantly improved from baseline
 - Stabilization of disease

Provider Signature: _____ **Date:** _____

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