

Symdeko[®] Prior Authorization Request Form (Page 1 of 2)

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Member Information (required)			Provider Information (required)		
Member Name:			Provider Name:		
Insurance ID#:			NPI#:		Specialty:
Date of Birth:			Office Phone:		
Street Address:			Office Fax:		
City:	State:	Zip:	Office Street Address:		
Phone:			City:	State:	Zip:
Medication Information (required)					
Medication Name:			Strength:		Dosage Form:
<input type="checkbox"/> Check if requesting brand			Directions for Use:		
<input type="checkbox"/> Check if request is for continuation of therapy					
Clinical Information (required)					
Select the diagnosis below: <input type="checkbox"/> Cystic fibrosis (CF) <input type="checkbox"/> Other diagnosis: _____ ICD-10 Code(s): _____					
Clinical Information: Is the patient homozygous for the F508del mutation in the cystic fibrosis transmembrane conductance regulator (CFTR) gene? <input type="checkbox"/> Yes <input type="checkbox"/> No <i>Lab documentation must be submitted</i> Select if the patient has one of the following mutations in the CFTR gene that is responsive to Symdeko based on in vitro data and/or clinical evidence: <i>Lab documentation must be submitted</i>					
<input type="checkbox"/> A1067T		<input type="checkbox"/> E193K		<input type="checkbox"/> L206W	
<input type="checkbox"/> A455E		<input type="checkbox"/> E56K		<input type="checkbox"/> P67L	
<input type="checkbox"/> D110E		<input type="checkbox"/> E831X		<input type="checkbox"/> R1070W	
<input type="checkbox"/> D110H		<input type="checkbox"/> F1052V		<input type="checkbox"/> R117C	
<input type="checkbox"/> D1152H		<input type="checkbox"/> F1074L		<input type="checkbox"/> R347H	
<input type="checkbox"/> D1270N		<input type="checkbox"/> F508del		<input type="checkbox"/> R352Q	
<input type="checkbox"/> D579G		<input type="checkbox"/> K1060T		<input type="checkbox"/> R74W	
<input type="checkbox"/> S945L		<input type="checkbox"/> 711+3A ->G			
<input type="checkbox"/> S977F		<input type="checkbox"/> 2789+5G ->A			
<input type="checkbox"/> 3272-26A ->G		<input type="checkbox"/> 3849+10kbC ->T			
Select the following that applies to the patient: <ul style="list-style-type: none"> <input type="checkbox"/> Baseline FEV1 within the last 30 days is $\leq 90\%$ (<i>Lab documentation must be submitted</i>) <input type="checkbox"/> Baseline eGFR or SCr within the last 3 months (<i>Lab documentation must be submitted</i>) <input type="checkbox"/> Baseline LFTs prior to initiating therapy, then every 3 months the first year, then annually (<i>Lab documentation must be submitted</i>) <input type="checkbox"/> Documentation of the number of pulmonary exacerbations in preceding 6 months <input type="checkbox"/> Documentation of baseline body mass index <input type="checkbox"/> Documentation of baseline ophthalmic examination to monitor lens opacities/cataracts (<i>Lab documentation must be submitted</i>) <input type="checkbox"/> Patient does not have positive cultures for <i>Burkholderia cenocepacia</i>, <i>Burkholderia dolosa</i>, or <i>Mycobacterium abscessus</i> (<i>Lab documentation within last six (6) months of THIS request must be submitted</i>) 					
Select if the patient is currently COMPLIANT on the following: <ul style="list-style-type: none"> <input type="checkbox"/> CFTR modulator therapy [Orkambi (lumacaftor/ivacaftor)] (Must be compliant on Orkambi if switching to Symdeko) <input type="checkbox"/> Dornase alfa <input type="checkbox"/> Hypertonic saline <input type="checkbox"/> Inhaled or oral antibiotics within the last three months 					

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Reauthorization:

If this is a reauthorization request, answer the following questions:

Select the following that applies to the patient:

- FEV1 \geq 5%
- Hospitalization decreased since prior to Symdeko therapy
- Patient's body weight increased at least 1.5kg

Select the following the patient has lab results documenting:

Lab documentation must be submitted

- Follow up ophthalmic examination
- Patient does not have positive cultures for *Burkholderia cenocepacia*, *Burkholderia dolosa*, or *Mycobacterium abscessus*
(*Lab documentation within last six (6) months of THIS request must be submitted*)
- Recent LFTs (within the last month)

Select if the patient is currently COMPLIANT on the following:

- Dornase alfa
- Hypertonic saline
- Inhaled or oral antibiotics within the last three months

Chart notes and any lab results MUST be submitted with this request

****Use of samples to initiate therapy does not meet step edit/ preauthorization criteria.****

Previous therapies will be verified through pharmacy paid claims or submitted chart notes.

Prescriber Signature: _____ **Date:** _____

Are there any other comments, diagnoses, symptoms, medications tried or failed, and/or any other information the physician feels is important to this review?

Please note: This request may be denied unless all required information is received.
For urgent or expedited requests please call 1-800-711-4555.
This form may be used for non-urgent requests and faxed to 1-800-527-0531.