

Federal Employee Program.

## SYMDEKO PRIOR APPROVAL REQUEST

Send completed form to: Service Benefit Plan Prior Approval P.O. Box 52080 MC 139 Phoenix, AZ 85072-2080 Attn. Clinical Services Fax: 1-877-378-4727

Additional information is required to process your claim for prescription drugs. Please complete the patient portion, and have the prescribing physician complete the physician portion and submit this completed form.

	atient Inform	ation (required)			Information (1	required)	L
Date:				Provider Name:			
Patient Name:				Specialty:	NPI:		
Date of Birth:		Sex:  Male	Female	Office Phone:	Office Fax:		
Street Address:		l		Office Street Address:			
City:		State:	Zip:	City:	State:	Zip:	
Patient ID: <b>R</b>	I I	1 1 1		Physician Signature:	-		
		P	HYSICIAN (	COMPLETES			
			Symo	deko			
			(tezacaftor ar				
	**Check	www.fepblue.org/fori	nulary to confirm	which medication is part of the pa	itient's benefit		
		NOTE: Form m	ust be complete	ed in its entirety for processing	<u>ng</u>		
Is this request for	r brand or generic	? □Brand □G	eneric				
How many tablet	s will the patient	need for an 84-day	y supply?	tablet(s) per 84 days			
Dosing direction	ç. -						
Dosing directions	s						-
-	atient's diagnosis?	)					
•	brosis (CF)	•••					
☐ Other dia	agnosis ( <i>please spe</i>	cify):					_
	o be used in comb □Yes* □No	ination with anoth	ner *cystic fibro	sis transmembrane conductar	nce regulator (CFT	ľR)	
		nedication:					_
*CFTR P	otentiators: Kalyde	co (ivacaftor), Orka	ımbi (ivacaftor/lu	macaftor), and Trikafta (ivacaj	ftor/tezacaftor/elexa	caftor)	
3. Has the patien	nt been on Symdel	ko continuously fo	or the last <b>4 mor</b>	nths, excluding samples? Plea	ase select answer be	low:	
		of therapy, please		• •			
_				e CFTR gene? □Yes □No			
	_	*CFTR gene mutati CFTR gene mutati	_	to Symdeko?  Yes  No onsive to Symdeko	0		
c. What i	s the pretreatment	t percent predicted	l forced expirate	ory volume (ppFEV <sub>1</sub> )?	<b>U</b> nk	cnown	
d. Have b	paseline levels of	ALT, AST, and bi	lirubin been obt	tained? □Yes* □No			
* <i>If</i> Y	<b>ES</b> , will the patie	ent's ALT, AST, a	and bilirubin lev	els be tested every three mon	iths for the first ye	ar? □Yes □No	)
e. Has Sy	mdeko been pres	cribed by a pulmo	nologist or gast	roenterologist?	<b>1</b> 0		
$\Box$ <b>YES</b> – this	is a PA renewal f	for <b>CONTINUAT</b>	TON of therapy	, please the following question	ons:		
	e patient been stab seline? TYes		een an improver	ment of percent predicted for	ced expiratory volu	ume (ppFEV <sub>1</sub> ) from	m

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 $\square$ No

b. Will the patient have annual testing of their ALT, AST, and bilirubin levels after the first year of therapy?  $\Box$ Yes



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## CFTR Gene Mutations that are Responsive to Symdeko

546insCTA	E92K	G576A	L346P	R117G	S589N
711+3A → G	E116K	G576A; R668C †	L967S	R117H	S737F
2789+5G→A	E193K	G622D	L997F	R117L	S912L
3272-26A → G	E403D	G970D	L1324P	R117P	S945L
3849+10kbC→T	E588V	G1069R	L1335P	R170H	S977F
A120T	E822K	G1244E	L1480P	R258G	S1159F
A234D	E831X	G1249R	M152V	R334L	S1159P
A349V	F191V	G1349D	M265R	R334Q	S1251N
A455E	F311del	H939R	M952I	R347H	S1255P
A554E	F311L	H1054D	M952T	R347L	T338I
A1006E	F508C	H1375P	P5L	R347P	T1036N
A1067T	F508C; S1251N†	I148T	P67L	R352Q	T1053I
D110E	F508del ^	1175V	P205S	R352W	V201M
D110H	F575Y	1336K	Q98R	R553Q	V232D
D192G	F1016S	I601F	Q237E	R668C	V562I
D443Y	F1052V	I618T	Q237H	R751L	V754M
D443Y; G576A;R668C†	F1074L	1807M	Q359R	R792G	V1153E
D579G	F1099L	1980K	Q1291R	R933G	V1240G
D614G	G126D	11027T	R31L	R1066H	V1293G
D836Y	G178E	11139V	R74Q	R1070Q	W1282R
D924N	G178R	11269N	R74W	R1070W	Y109N
D979V	G194R	11366N	R74W; D1270N †	R1162L	Y161S
D1152H	G194V	K1060T	R74W; V201M †	R1283M	Y1014C
D1270N	G314E	L15P	R74W; V201M;D1270N †	R1283S	Y1032C
E56K	G551D	L206W	R75Q	S549N	
E60K	G551S	L320V	R117C	S549R	

<sup>^</sup> A patient must have two copies of the F508del mutation or at least one copy of a responsive mutation presented above to be indicated.

<sup>†</sup> Complex/compound mutations where a single allele of the CFTR gene has multiple mutations; these exist independent of the presence of mutations on the other allele.



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Message:

Attached is a Prior Authorization request form.

For your convenience, there are 3 ways to complete a Prior Authorization request:

Electronically Online (ePA) Results in 2-3 minutes FASTEST AND EASIEST	Now you can get responses to drug Prior Authorization requests <b>securely</b> online. <b>Online</b> submissions may receive <b>instant</b> responses and do not require faxing or phone calls.  Requests can be made 24 hours a day, 7 days a week. For more information on electronic prior authorization (ePA) and to register, go to <b>Caremark.com/ePA.</b>
Phone (4-5 minutes for response)	The FEP Clinical Call Center can be reached at (877)-727-3784 between the hours of 7AM-9PM Eastern Time. A live representative will assist with the Prior Authorization, asking for the same information contained on the attached form. Please review the form and have your answers ready for faster service.  The process over the phone takes on average between 4 and 5 minutes.
Fax (3-5 days for response)	Fax the attached form to (877)-378-4727. Requests sent via fax will be processed and responded to within 5 business days. The form must be filled out completely, if there is any missing information the Prior Authorization request cannot be processed.  Please only fax the completed form once as duplicate submissions may delay processing times.

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