

**OPTIMA HEALTH COMMUNITY CARE  
AND  
OPTIMA FAMILY CARE  
(MEDICAID)**

**PHARMACY PRIOR AUTHORIZATION/STEP-EDIT REQUEST**

**Directions:** The prescribing physician must sign and clearly print name (preprinted stamps not valid) on this request. All other information may be filled in by office staff; **fax to 1-800-750-9692.** No additional phone calls will be necessary if all information (including phone and fax #s) on this form is correct. **Incomplete form will delay authorization process.**

**Drug Requested:** **Symdeko®** (tezacaftor/ivacaftor)

**DRUG INFORMATION:** Complete all information below or authorization process will be delayed.

**Drug Form/Strength:** \_\_\_\_\_

**Dosing Schedule:** \_\_\_\_\_ **Length of Therapy:** \_\_\_\_\_

**Diagnosis:** \_\_\_\_\_ **ICD Code, if applicable:** \_\_\_\_\_

**Recommended dose:** 1 tablet (tezacaftor 100mg/ivacaftor 150mg) in the morning and 1 tablet (ivacaftor 150mg) in the evening (approximately 12 hours apart).

**Authorization approval – 1 year**

**CLINICAL CRITERIA:** **ALL** boxes below **MUST** be checked to qualify. Lab results and chart notes **MUST** be attached to ensure authorization will **NOT** be delayed.

1. Is the member 12 ≥ years of age?  Yes  No

**AND**

2. Does the member have a diagnosis of Cystic Fibrosis.  Yes  No

**AND**

3. Is member homozygous for the **F508del** mutation in the CFTR gene as confirmed by an FDA-cleared CF mutation Test? **(Document required; include a copy of the test with this request)**  Yes  No

**OR**

4. Does member have one (1) of the following mutations (see chart below) in the CFTR gene as confirmed by an FDA-cleared CF mutation Test? **(Lab documentation required; include a copy of the test with this request)**  Yes  No

**\*CFTR Mutations Responsive to Symdeko:**

E56K	R352Q	A1067T
P67L	A455E	R1070W
R74W	D579G	F1074L
D110E	711+3A→G	D1152H
D110H	E831X	D1270N
R117C	S945L	2789+5G→A
E193K	S977F	3272-26A→G
L206W	F1052V	3849+10kbC→T
R347H	K1060T	

(continued on next page)

5. Has baseline ALT and AST testing been done? (**Documentation required; include a copy of the test with this request**)  Yes  No

**For Re-Authorization approval  
liver function testing (LFT) documentation is required**

**Medication being provided by a Specialty Pharmacy  
Sentara Norfolk General CM Pharmacy**

**\*\*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.\***

Patient Name: \_\_\_\_\_

Member Optima #: \_\_\_\_\_ Date of Birth: \_\_\_\_\_

Prescriber Name: \_\_\_\_\_

Prescriber Signature: \_\_\_\_\_ Date: \_\_\_\_\_

Office Contact Name: \_\_\_\_\_

Phone Number: \_\_\_\_\_ Fax Number: \_\_\_\_\_

**DEA OR NPI #:** \_\_\_\_\_

\*Approved by Pharmacy and Therapeutics Committee: 4/19/2018

REVISED/UPDATED ~~6/21/2018~~; 8/30/2018