FUTURE SCRIPTS Secure

Please complete ALL information below and fax your request to 1-888-671-5285

Orkambi® Coverage Determination Request Form (Page 1 of 2)

DO NOT COPY FOR FUTURE USE. FORMS ARE UPDATED FREQUENTLY AND MAY BE BARCODED

Member Information (required)

Member Name:		
Insurance ID#:		
Date of Birth:		
Street Address:		
City:	State:	Zip:
Phone:		

Provider Information (required)

Provider Name:		
NPI#:	Specialty:	
Office Phone:		
Office Fax:	Office Contact:	
Office Street Address:		
City:	State:	Zip:

Medication Information (required)

Medication Name: Select one of the following: ❑ Request is for **GENERIC** ❑ Request is for **BRAND** (unable to take the generic)	Strength:	Dosage Form:
❑ Check if request is for **continuation of therapy**	Directions for Use:	

Clinical Information (required)

Select the Type(s) of Coverage Determination Requested:

- ❑ **Non-Formulary**- Request is for a drug not on the plan's list of covered drugs OR was previously included on the plan's list is being/was removed from this list during the plan year.
- ❑ **Prior Authorization**- Request is for a drug that requires prior authorization under the plan.
- ❑ **Quantity Limit**- Request is for an exception to the plan's quantity limit.

  Quantity per DAY requested? _____

Select the diagnosis below:

- ❑ Cystic fibrosis (CF)
- ❑ Other diagnosis: _____ ICD-10 Code(s): _____

Medication History:

List the medication(s) (in the **same class**) the patient has a history of trial and failure, or intolerance to: _____

_____

List the medication(s) (that are **therapeutic equivalent alternatives**) the patient has a history of trial and failure, or intolerance to: _____

_____

Clinical Information:

Is the patient homozygous for the F508del mutation in the cystic fibrosis transmembrane conductance regulator (CFTR) gene? ❑ **Yes** ❑ **No**

If the patient's genotype is unknown, has an FDA-cleared CF mutation test been used to detect the presence of the F508del mutation on both alleles of the CFTR gene? ❑ **Yes** ❑ **No**

Does the patient have diagnosis of CF other than those homozygous for the F508del mutation? ❑ **Yes** ❑ **No**

Quantity Limit Requests:

Is there a high risk of significant adverse clinical outcome with medication change or dosage change? ❑ **Yes** ❑ **No**

Is the requested quantity and dose within FDA approved maximum dosing limits or supported by peer-reviewed medical literature, accepted standards of medical practice and/or medical compendia? ❑ **Yes** ❑ **No**

If **yes,** please specify:_____

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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.