

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 chronic obstructive pulmonary disease (COPD), including chronic bronchitis and/or emphysema? Yes No
If no, list diagnosis: _____

- Does the patient have a history of failure, contraindication, or intolerance to treatment with at least a 30 day trial of **both** of the following used in combination: Yes No
 - Stiolto Respimat (tiotropium/olodaterol)
 - Arnuity Ellipta (fluticasone)(If yes, complete Section D above with medication information, including dose, date of trial, and reason for discontinuation)

Provider Signature: _____ **Date:** _____

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