

ICS/LABA Combo - Arizona Prior Authorization Request Form

Please complete this <u>entire</u> 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 Infor	mation								
First Name:	Last Name:				Member ID:				
Address:					•				
City:	State:				ZIP Code:				
Phone:	DOB:				Allergies:				
Primary Insurance Information	(if any):	l							
Is the requested medicati	ion: New or	Continua	tion of Ther	apy? If continuation	n, list sta	rt date:			
Is this patient currently h									
Section B - Provider Infor	mation								
First Name:			Last Name:				M.D./D.O.		
Address:			City:		State:	State: ZIP code:			
Phone:	Phone: Fax:			NPI #:			Specialty:		
Office Contact Name / Fax atte	ention to:				ı				
Section C - Medical Inforn	mation								
Medication:						Strength:			
Directions for use:						Quantity:			
Diagnosis (Please be specific		ICD-10 CODE:							
Is this member pregnant? □	 Yes □ No	If yes	, w hat is this	member's due date?					
Section D - Previous Med	ication Trials	_							
Medication Name	Strength	Dire	ctions	Dates of Thera	ару	Reason for failure / discontinuation			
						41300	Milliaution		
	+								
Section E – Additional info	ormation and Ex	nlanation	of why pref	erred medications	would no	ot meet th	ne natient's needs		
Please refer	to the patient's	PDL at wv	vw.uhcprov	ider.com for a list o	f preferr	ed alterna	atives		



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Date: _____

Member First name:		Member Last name:	Member DOB:	Member DOB:				
		Clinical and Drug Spec	cific Information					
ALL REQUESTS								
□ Yes □ No	Does the patient have o □ Asthma □ COPD	ne of the following diagnos	ses? (If yes, check which applies)					
□ Yes □ No	Is the patient already established on ICS/LABA combination therapy (e.g. Advair, Symbicort, Dulera)? (If yes, complete Section D above)							
□ Yes □ No	Does the patient have a history of failure, contraindication, or intolerance to treatment with at least a 30-day trial of any of the following? (If yes, check which applies and complete Section D above) □ Beclomethasone dipropionate (Qvar) □ Budesonide (Pulmicort) □ Fluticasone propionate (Flovent HFA) □ Mometasone (Asmanex Twisthaler)							
□ Yes □ No	30 day trial of any of the ☐ A long-acting beta-ag ☐ An orally inhaled antic	e following? (If yes, check wonist (e.g. Foradil, Serevent, cholinergic agent (e.g. Spiriva	dication, or intolerance to treatment which applies and complete Section D a Striverdi, Arcapta) a, Atrovent, Combivent, Tudorza, Incru beta-agonist combination agent (e.g.	above) se Ellipta)				
□ Yes □ No	Does the patient have a history of failure, contraindication, or intolerance to treatment with any of the following preferred products? (If yes, check which applies and complete Section D above) □ Advair HFA (fluticasone/salmeterol) □ Dulera (mometasone/formoterol) □ Symbicort (budesonide/formoterol)							

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Provider Signature: _____

error, please notify the sender immediately.