

SPECIAL AUTHORITY REQUEST INHALERS FOR CHRONIC OBSTRUCTIVE PULMONARY DISEASE (COPD)

HITH 5362 Rev 2022/10/1

LIMITED COVERAGE LONG ACTING MUSCARINIC ANTAGONIST (LAMA) INHALERS Complete sections 1, 2 and 3 LONG ACTING BETA-AGONIST (LABA) INHALERS Complete sections 1, 2 and 4 For up-to-date criteria and forms, please check: www.gov.bc.ca/pharmacs Fax requests to 1-800-609-4884 (toll free) OR mail requests to: PharmaCare, Box 9 This facsimile is Doctor privileged and contains confidential information intended only fo copying or disclosure is strictly prohibited. If PharmaCare approves this Special Authority request, approval is granted solely for the property of the property		Co INI Co ICS Co Carespeci 9652 Stn I for Pharmace	harmaCare. Any other distribution, If you have received this fax in error, please write MISDIRECTED across the front of the form and fax toll-free to 1-800-609-4884, then destroy the pages received in error.				
PharmaCare approval does not indicate that the requeste Forms with information missing will be returned		•	•	PharmaC	are will be unable to return a response.		
SECTION 1 – PRESCRIBER INFORMA	TION		SECTION 2 - PATIENT I	NFORM	IATION		
Name and Mailing Address			Patient (Family) Name				
			Patient (Given) Name(s)				
College ID (use ONLY College ID number)	hone Number (include area	code)	Date of Birth (YYYY / MM / DD)	Date of Application (YYYY / MM / DD)		
CRITICAL FOR A TIMELY RESPONSE Prescriber's Fax Number			CRITICAL FOR PROCESSING	Personal	ersonal Health Number (PHN)		
SECTION 3 – LIMITED COVERAGE LA	AMA INHALERS Sele	ct produ	uct below:		9901-0164		
○ Aclidinium (TUDORZA GENUAIR)	Tiotropium (SPIRIVA HA	ANDIHAL	ER) Glycopyrronium (SEEBRI B	REEZHALER)		
☐ Failure of ALL of the Regular Benefit long-a ☐ Tiotropium (SPIRIVA RESPIMAT)	<u> </u>	_	st (LAMA) devices after a mini m (INCRUSE ELLIPTA)	mum one	-month trial of EACH device:		
SECTION 4 – LABA INHALERS Select	product below:						
○ Salmeterol (SEREVENT) 9901-0341	Ondacaterol (ONBREZ	BREEZHA	LER) 9901-0229 (Maximum do	se of 75mc	g daily)		
☐ Diagnosis of COPD with a post-bronchodila	ator FEV1/FVC < 0.70. Plea	ise specify	FEV1/FVC ratio: 0.				
Contraindication or intolerance to a long-a	cting muscarinic receptor	antagoni	st (LAMAs)				
Please specify nature and severity of contra	aindication or intolerance:	:					
Please submit applications for LABA inhalers in a	asthma on the general spec	cial author	ity form.				
PHARMACARE USE ONLY							
STATUS		EFFECTIVE D	ATE (YYYY / MM / DD)	DURA	NTION OF APPROVAL		

PATIENT (FAMILY) NAME	PATIENT (GIVEN) NAME(S)		PERSONAL HEALTH NUMBER (PHN)	HLTH 5362 PAGE 2		
SECTION 5 - LAMA / LABA COMBINAT	ION INHALERS Selec	ERS Select product below:				
	rol/Glycopyrronium D BREEZHALER)	○ Tiotropium/Olodatero (INSPIOLTO RESPIMA	_	erol		
☐ Diagnosis of moderate to very severe COPD w	rith a post-bronchodilator F	EV1/FVC < 0.70 AND a post-	bronchodilator FEV1 < 80% predicted	d.		
Please specify FEV1/FVC ratio: 0.	AND FEV1:%					
AND						
☐ Inadequate response after minimum of 6 mo	onth trial of either LAMA or	LABA. Please specify LAMA	or LABA tried:			
SECTION 6 - INHALED CORTICOSTERC	DID (ICS) / LABA COM	BINATION INHALERS	1	9901-0063		
O Fluticasone Propionate/Salmeterol (ADVAI	R/GENERICS)	uticasone Furoate/Vilanter	ol (BREO ELLIPTA 100/25 only)			
☐ Diagnosis of moderate to very severe COPD w	vith a post-bronchodilator F	EV1/FVC < 0.70 AND a post-	bronchodilator FEV1 < 80% predicted	d.		
Please specify FEV1/FVC ratio: 0.	AND FEV1:%					
AND						
☐ Inadequate response after minimum of 6 mo	nth trial of either LAMA or	LABA. Please specify LAMA	or LABA tried:			
AND EITHER OF THE FOLLOWING						
	n the previous 12 months, o	lefined as requiring a prescri	bed antibiotic and/or using systemic	glucocorticoids.		
☐ History of ≥ 1 severe exacerbation in the	previous 12 months define	ed as requiring a hospital ad	mission or emergency department vi	sit.		
SECTION 7 - ICS / LAMA / LABA COMB	INATION INHALERS			9901-0340		
Fluticasone Furoate/Umeclidinium/Vilante (TRELEGY ELLIPTA 100/62.5/25 only)		de/Glycopyrronium/Formo	oterol			
☐ Diagnosis of moderate to very severe COPD w	vith a post-bronchodilator F	EV1/FVC < 0.70 AND a post-	bronchodilator FEV1 < 80% predicted	d.		
Please specify FEV1/FVC ratio: 0.	AND FEV1:%					
AND EITHER OF THE FOLLOWING:						
○ Inadequate response after minimum of	6 month trial of a LAMA/L	ABA combination inhaler. Ple	ease specify product:			
○ Inadequate response after minimum of	6 month trial of a ICS/LAB.	A combination inhaler. Pleas	e specify product:			
AND EITHER OF THE FOLLOWING			. ,,			
History of ≥ 2 moderate exacerbations in	n the previous 12 months, d	efined as requiring a prescri	bed antibiotic and/or using systemic	glucocorticoids.		
O History of ≥ 1 severe exacerbation in the	previous 12 months define	ed as requiring a hospital adr	mission or emergency department vi	sit.		
SECTION 8 - COMMENTS						
SECTION 9 - PRESCRIBER'S SIGNATUR	RE					
Personal information on this form is collected under the au	I have discussed with	the patient that the purpose of re	eleasing their			
with, the <i>British Columbia Pharmaceutical Services Act</i> 22(1) - <i>Protection of Privacy Act</i> 26 (a),(c),(e). The information is bein	ng collected for the purposes		naCare is to obtain Special Authori purposes set out here.	ty for prescription		
of (a) administering the PharmaCare program, (b) analyzing Special Authority and other Ministry programs and (c) to m		coverage and for the	pa.poses set out here.			
system generally. If you have any questions about the colle Health Insurance BC from Vancouver at 1-604-683-7151 or 1	ction of this information, call					
1-800-663-7100 and ask to consult a pharmacist concerning	Prescriber's Signature (Man	Prescriber's Signature (Mandatory)				