

Drug Prior Authorization Form Cinqair (reslizumab), Fasenra (benralizumab)

The purpose of this form is to obtain information required to assess your drug claim.

IMPORTANT: Please answer all questions. Your claim assessment will be delayed if this form is incomplete or contains errors.

Any costs incurred for the completion of this form are the responsibility of the plan member/patient.

Canada Life recognizes and respects the importance of privacy. Personal information collected is used for the purposes of assessing eligibility for this drug and for administering the group benefits plan. For a copy of our Privacy Guidelines, or if you have questions about Canada Life's personal information policies and practices (including with respect to service providers), refer to www.canadalife.com or write to Canada Life's Chief Compliance Officer.

I authorize Canada Life, any healthcare provider, my plan administrator, any insurance or reinsurance company, administrators of government benefits or patient support programs or other benefits programs, other organizations, or service providers working with Canada Life or any of the above, located inside or outside Canada, to exchange personal information when relevant and necessary for these purposes. I understand that personal information may be subject to disclosure to those authorized under applicable law within or outside Canada.

I acknowledge that the personal information is needed to assess eligibility for this drug and to administer the group benefits plan. I acknowledge that providing consent will help Canada Life to assess my claim and that refusing to consent may result in delay or denial of my claim. Canada Life reserves the right to audit the information provided on this form at any time and this consent extends to any audit of my claim. This consent may be revoked by me at any time by sending written instruction to that effect.

I also consent to the use of my personal information for Canada Life and its affiliates' internal data management and analytics purposes.

If the patient is a person other than myself, I confirm that the patient has given their consent to provide their personal information and for Canada Life to use and disclose it as set out above.

I certify that the information given below is true, correct, and complete to the best of my knowledge. Failure to provide true, correct and complete information on this form could result in revocation of any approval decision, a requirement to repay paid claims or other appropriate action.

Plan Member's signature:	Date:
ian Monibol o dignataro.	Date:

Form Completion Instructions:

- 1. Complete "Patient Information" sections.
- 2. Have the prescribing physician complete the "Physician Information" sections.
- 3. Send all pages of the completed form to us by mail, fax or email as noted below.

Note: As email is not a secure medium, any person with concerns about their prior authorization form/medical information being intercepted by an unauthorized party is encouraged to submit their form by other means.

Fax to:

Mail to: The Canada Life Assurance Company

Drug Claims Management

PO Box 6000

Winnipeg MB R3C 3A5

Email to: cldrug.services@canadalife.com

Attention: Drug Claims Management

For additional information regarding Prior Authorization and Health Case Management, please visit our Canada Life website at

<u>www.canadalife.com</u> or contact Group Customer Contact Services at 1-800-957-9777. Deaf or hard of hearing and require access to a telecommunications relay service? Please contact us at 711 for TTY to Voice or 1-800-855-0511 for Voice to TTY.

(Continued on next page)

Page 1 of 6

Fax 1-204-946-7664

The Canada Life Assurance Company

Attention: Drug Claims Management



Patient Information Cinqair (reslizumab), Fasenra (benralizumab)

Plan Member Information – Complete all sections of this page (please print)			
Plan Member:		Patient Name:	
Plan Name:	Plan Number:		Plan Member ID Number:
Patient Date of Birth (DD/MM/YYYY):	Address (number, street	, city, province, postal cod	de):
Please indicate preferred contact number and if the	 here are anv times when t	elephone contact with vo	u about vour claim would be most convenient.
	,	,	
May we contact you by email? (Note that some co	orrespondence may still r	eed to be sent by regular	mail).
\square Yes \square No $\:$ If yes, please provide email ad	dress:		
Tell us if you have been on this drug b	pefore		
Is the patient currently on, or previously been on	this drua? 🗆 Yes 🗆 1	No	
If Yes, a) indicate start date (DD/MM/YYYY):			
b) coverage provided by:			
(if coverage is not provided by Canada Lif	e please provide pharma	cy print-out showing purc	hase of this drug)
Tell us if you have coverage with any	other benefits plan		
	-		
Does the patient have drug coverage under any other group benefits plan? Yes No			
If Yes, name of other Insurance Company:			
Name of plan member:			
Relationship to patient:			
Provide details and attach documentation of			
Fromue details and attach documentation of acceptance of decime.			
Tell us about any Provincial or other of	coverage you may h	2210	
-			1
Does the patient have coverage under a provincial program or from any other source? Yes No			
If Yes, name of program or other source:			
Provide details and attach documentation of acceptance or decline:			
Is the patient currently receiving disability benefits for the condition for which this drug has been prescribed? Yes No			
Tell us about any Patient Support Pro	gram you might be	enrolled in	
Has the patient enrolled in the patient support pr	rogram for this drug? \Box	Yes □ No	
If Yes, please provide the following information:			
Patient support program patient ID Number:			
2. Patient support program contact person na	ame and phone number:		
Contact Name:		Phone Number	



Physician Information Cinqair (reslizumab), Fasenra (benralizumab)

Note to Physician: In order to assess a patient's claim for this drug, we require detailed information on the patient's prescription drug history as requested below.

Attach extra information if necessary. GENETIC TEST RESULTS ARE NOT REQUIRED

Physician's Information (please print)				
Name of prescribing physician:				
Specialty:				
Address (number, street, city, province, postal code):				
Telephone Number (including area code):	Fax Number (including area code):			
Prescribed dosage and regimen:				
☐ Cinqair (reslizumab) ☐ 3mg/kg every 4 weeks				
☐ Fasenra (benralizumab) ☐ 30mg every 4 weeks for 3 doses	☐ Fasenra (benralizumab) ☐ 30mg every 4 weeks for 3 doses then every 8 weeks thereafter ☐ RENEWAL: 30mg every 8 weeks			
☐ Other (please specify):				
Provide rationale if requested dose does not align with product monograph:				
Patients weight kg (required for weight-based dosing)				
Date determined (MM/YYYY):				
2. Health Canada Indication (include date of initial diagnosis)(MM/YYYY)	:			
☐ Severe Asthma				
Complete questions 1–6 and Physician's information				
Other (approved by Health Canada):				
Complete questions 1 – 6 and Other condition (Health Canada approved)				
☐ Other (prescribed use is not approved by Health Canada):				
Complete questions 1-6 and Off-label use				
3. What is the anticipated duration of treatment with this drug?				
4. Where will treatment be administered? \Box Home \Box Physician's Office \Box Private clinic \Box Hospital in-patient \Box Hospital out-patient				
5. Please provide medical rationale why this drug has been prescribed instead of an alternate drug in the same therapeutic class:				



Physician Information Cinqair (reslizumab), Fasenra (benralizumab)

Physician's Information (continued) (please print)

6. Drug and Treatment History - i	must be completed for	every request. If co	verage for these dru	gs was not provided by Canada Life, please
submit a pharmacy printout fo	r the last 12 months.			
Drug(s) and Treatment(s) past and present	Dosing Regimen	Start Date (DD/MM/YYYY)	End Date (DD/MM/YYYY)	Clinical Results/Outcome
				☐ Failure ☐ Intolerance ☐ Other Clinical details:
				☐ Failure ☐ Intolerance ☐ Other Clinical details:
				☐ Failure ☐ Intolerance ☐ Other Clinical details:
Severe Asthma				
Is the patient currently using a biologic therapy to treat their asthma? Yes No If yes, provide rationale for the change in therapy:				
Will the treatment be used as add-	on therapy to existing op	timized regimen? \Box	Yes ☐ No	
	corticosteroids (>250mcgicosteroids (>500mcg fluications (e.g. long-acting	g fluticasone propior ticasone propionate b-agonists, leukotri	nate or equivalent) or equivalent) ene receptor antago	onists, and/or theophylline)
Does the patient currently require of			J Yes ∟ No	
Please ensure the Drug and Trea				
Current blood eosinophil count:		Date (DD/MM/YYY	Y):	
If prior biologic therapy, was blood Value and date (DD/MM/YYYY):		ells/µL prior to initiati	on of biologic asthm	na therapy? ☐ Yes ☐ No
Recent clinically significant exacert Please provide the numb		exacerbations in the	e last 12 months:	

M6453(SEA)-4/24 Page 4 of 6



Physician Information Cinqair (reslizumab), Fasenra (benralizumab)

Severe Asthma (continued)
☐ Other. Please provide detail:
Date(s) (DD/MM/YYYY):
Will reslizumab or benralizumab be used in combination with another biologic medication (eg: omalizumab, mepolizumab, dupilumab, tezepelumab)?
☐ Yes ☐ No
Renewal Request
Date reslizumab or benralizumab was started:
Has the patient demonstrated a positive clinical response to treatment? ☐ Yes ☐ No
Please provide details:
□ Decrease or elimination of oral corticosteroid requirements
Dose at initiation: Current dose:
☐ Decrease in clinically significant exacerbations. Please provide details of any exacerbations since initiation of therapy:
☐ Improvement in asthma control questionnaire score from baseline
Result and date (MM/YYYY) prior to therapy:
Current result and date (MM/YYYY):
*Attach copy of completed questionnaires.
☐ Improvement in % predicted FEV1
FEV1 prior to therapy and date (MM/YYYYY):
Current FEV1 and date (MM/YYYY):
*Attach spirometry results. Other, Provide details:
Ottlei. Provide details.
Will reslizumab or benralizumab continue to be used as add-on maintenance asthma therapy? \square Yes \square No
Will reslizumab or benralizumab be used in combination with another biologic medication (eg: omalizumab, mepolizumab, dupilumab,
tezepelumab)? Yes No
Other condition (Health Canada approved)
Please provide any relevant information related to the disease and attach supporting documentation.

M6453(SEA)-4/24 Page 5 of 6



Email to: <u>cldrug.services@canadalife.com</u> Attention: Drug Claims Management

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Off-lab	el use		
Question	is 1 - 6 must be completed.		
Date of in	nitial diagnosis (DD/MM/YYYY):		
Is there of	clinical evidence supporting the off-label use of this drug? \Box Yes \Box	No	
Provide o	clinical literature/studies to support the request for off-label use, such	n as:	
•	At least two Phase II or two Phase III clinical trials showing consist	ent results	s of efficacy; and
•	Published recommendations in evidence-based guidelines support	ting its us	e.
Provide r	medical rationale why this drug has been prescribed off-label instead	of an alte	ernate drug with an approved indication for this condition.
Provide a	any pertinent medical history or information to support this off-label re	equest.	
If this is a	a renewal request, provide documentation showing treatment efficac	y since pr	evious request.
	Physician: To be eligible for reimbursement, Canada Life ation from a pharmacy designated by Canada Life. If applicant		
certify t	hat the information provided is true, correct, and complet	te.	
Physician	's Signature:		Date:
License N	lumber:		_
	rtant to provide the requested information in detail to help avoit to audit. The completed form can be returned to Canada Li		
	email is not a secure medium, any person with concerns abo ercepted by an unauthorized party is encouraged to submit th		
Mail to:	The Canada Life Assurance Company Drug Claims Management PO Box 6000 Winnipeg MB R3C 3A5	ax to:	The Canada Life Assurance Company Attention: Drug Claims Management Fax 1-204-946-7664

M6453(SEA)-4/24 Page 6 of 6