

## **Prior Authorization Request**

ILARIS (canakinumab)

## **Instructions**

Please complete Part A and have your physician complete Part B. Completion and submission is not a guarantee of approval. Any fees related to the completion of this form are the responsibility of the plan member. Drugs in the Prior Authorization Program may be eligible for reimbursement if the patient does not qualify for coverage under a primary plan or a government program. Drugs used for indications not approved by Health Canada may be denied. For Quebec plan members, RAMQ exception drug criteria may apply. The decision for approval versus denial is based on pre-defined clinical criteria, primarily based on Health Canada approved indication(s) and on supporting evidence-based clinical protocols. The plan member will be notified whether their request has been approved or denied. Please note that you have the right to appeal the decision made by Express Scripts Canada.

## Part A - Patient

Patient information			T				
First Name:			Last Name:				
Insurance Carrier N	lame/Number:						
Group Number:			Client ID:				
Date of Birth (YYYY/MM/DD):			Relationship: Employee Spouse Dependent				
Language:	English Frenc	h	Gender: Male Female				
Address:							
City:		Province:		Postal Code:			
Email address:							
Telephone (home):		Telephone (cell):		Telephone (work):			
Coordination of ben	efits						
Patient Assistance	Is the patient enrolled in any patient assistance program? Yes No						
Program	Contact Name: Telephone:						
Provincial Coverage	Has the patient applied for reimbursement under a provincial plan? Yes No N/A						
	What is the coverage decision of the drug? Approved Denied *Attach decision letter*						
Primary Coverage	Has the patient applied for reimbursement under a primary plan? Yes No N/A						
	What is the coverage decision of the drug? Approved Denied *Attach decision letter*						
information containe administration and r	ed on this form. I give n management of my gro	ny consent on the un up benefit plan. This	derstanding that the in consent shall continue	er, and its agents, to exchange the personal formation will be used solely for purposes of so long as my dependents and I are covered val, or reinstatement thereof.			
Plan Member Signat	ure			Date			



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## Part B - Prescriber

Please see instructions on page 1 and complete all sections below. Incomplete forms may result in automatic denial. Please do not provide genetic test information or results.

SECTION 1 - DRUG REQUESTED									
ILARIS (canakinumab)		New request	Renewal request*						
Dose	Administration (ex: oral, IV, etc)	Frequency	Duration						
Site of drug administration:									
Home Physician	's office/Infusion clinic	Hospital (outpatient)	Hospital (inpatient)						
* Please submit proof of prior c	overage if available								
SECTION 2 – ELIGIBILITY CI	RITERIA								
Please indicate if the patient satisfies the below criteria:									
Cryopyrin-Associated Periodic Syndromes									
For the treatment of cryopyrin-associated periodic syndromes (CAPS), including familial cold autoinflammatory syndrome (FCAS) or familial cold urticaria (FCU), Muckle-Wells syndrome (MWS), neonatal-onset multisystem inflammatory disease (NOMID) or chronic infantile neurological, cutaneous, articular syndrome (CINCA), AND									
The patient is 2 years of	of age or older								
Systemic Juvenile Idiopathic Art	thritis								
For the treatment of sy	stemic juvenile idiopathic arthriti	is (sJIA), AND							
The patient is 2 years of	of age or older and weighs 9 kg o	r more, AND							
		locumented contraindication or in nic corticosteroids (Please list pric							
The patient has failed a	another biologic response modifi	er (Please list prior therapies in ti	he chart below)						
Adult-Onset Still's Disease									
For the treatment of ac	dult-onset Still's disease in an ad	ult, AND							
		locumented contraindication or in latory drugs (NSAIDs) <i>(Please list</i>							
Familial Mediterranean Fever									
For the treatment of fa	milial Mediterranean fever (FMF)	), AND							
The patient is 2 years of	of age or older, AND								
The patient has had ar list prior therapies in the		locumented contraindication or ir	ntolerance to colchicine ( <i>Please</i>						



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Tumor Necrosis Factor Receptor Asso	ciated Periodic Syndrome								
For the treatment of tumor necrosis factor receptor associated periodic syndrome (TRAPS), AND									
The patient is 2 years of age or older									
Hyperimmunoglobulin D Syndrome or Mevalonate Kinase Deficiency  For the treatment of hyperimmunoglobulin D syndrome (HIDS) or mevalonate kinase deficiency (MKD), AND  The patient is 2 years of age or older									
OR  None of the above criteria applies.									
Relevant additional information:									
Please list previously tried therap	ies								
		Duration of therapy		Reason for cessation					
<b>D</b> .	Dosage and	Duration	of therapy						
Drug	Dosage and administration	Duration From	of therapy To	Reason for Inadequate response	Allergy/ Intolerance				
Drug				Inadequate	Allergy/				
Drug				Inadequate	Allergy/				
Drug				Inadequate	Allergy/				
Drug				Inadequate	Allergy/				
Drug				Inadequate	Allergy/				
Drug				Inadequate	Allergy/				
Drug				Inadequate	Allergy/				
	administration			Inadequate	Allergy/				
SECTION 3 – PRESCRIBER INFOR	administration			Inadequate	Allergy/				
	administration			Inadequate	Allergy/				
SECTION 3 – PRESCRIBER INFOR	administration			Inadequate	Allergy/				
SECTION 3 – PRESCRIBER INFO	administration			Inadequate	Allergy/				
SECTION 3 – PRESCRIBER INFOR	administration	From		Inadequate	Allergy/				

Please fax or mail the completed form to Express Scripts Canada®

Fax:

Express Scripts Canada Clinical Services 1 (855) 712-6329

Mail:

Express Scripts Canada Clinical Services 5770 Hurontario Street, 10<sup>th</sup> Floor Mississauga, ON L5R 3G5