

QC G1V 4H6

Prior authorization request form Abatacept (Orencia[®]), etanercept (Enbrel[®], Erelzi[®]), infliximab (Remicade[®]), adalimumab (Humira[®])/ Juvenile idiopathic arthritis of

polyarticular or systemic form

DECLARATION OF THE INSURED PERSON

Section 1: Information about the	participant and the pati	ent				
Name of Participant	Insurance Policy / Certificate		Name of Employer			
Name of Patient	Date of Birth (yyyy/mm/dd)		Telephone			
Address (house number and street name)	City/Town		Province	Postal code		
Section 2: Other prescription drug	insurance			,		
Do you have other prescription drug insurance? If so, please answer the following:			☐ Yes	□ No		
What type of plan is it?	What type of plan is it?		☐ Private	☐ Public		
Have you ever submitted a claim for this drug to the other insurer?		7 At-	☐ Yes	□ No		
What is the status of the claim? Did this insurer ask you to complete a prior authorization request?		☐ Accepte	ed	☐ Under review☐ No		
What is the status of the prior autho		□ Accepte	ed	☐ Under review		
Please enclose acceptance or refu		•	a B Neruseu	_ onder review		
Section 3: Authorization to disclose personal information I certify that the information in this prior authorization request is complete, accurate and true.						
I authorize physicians and other health care professionals, medical, paramedical or clinical institutions, care coordinators, members of SSQ's Preferred Pharmacy Network (outside Quebec only) and any public or parapublic organization, including the Régie de I 'assurance maladie du Québec, to disclose to SSQ, Life Insurance Company Inc. (SSQ) any of my personal information including and without limitation, any medical information and medical evaluations in connection with the processing of this request. I hereby waive their confidentiality obligation and authorize them to disclose the requested information to SSQ. In addition, I authorize SSQ to disclose to the previously named third parties any of my personal information including and without limitation any medical information and medical evaluations in connection with the processing of this request.						
Photocopies of this document have the same value as the original.						
Signature of patient (parent/legal guardian)		Dat	e			
IMPORTANT: All correspondence concerning this form will be sent to the address indicated in the participant's file.						
Send us the completed form by email or by fax at: 1-855-453-3942. Telephone 418, 651, 2588/4, 800, 380, 3588 — Fey: 1, 855, 453, 3043, Addresses, 3535, Lourier Blad, B.O., Boy 10500, Overhea City.						
Telephone: 418-651-2588/1-800-380-2588 — Fax: 1-855-453-3942 Addresse: 2525 Laurier Blvd, P.O. Box 10500, Quebec City						



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DECLARATION OF THE PHYSICIAN

Section 1: Information a	bout the prescribing physic					
Name of Physician			Specialty		Licence No.:	
Telephone:		•		Fax:		
I hereby certify that the information in this request is accurate.						
Signature of physician				Date		
Section 5: Drug covered	by the authorization					
Name of Drug	Pharmaceutical Form	Stren	ath	Dosage		
Name of Drug	Filarinaceuticarronni	301611	gui	Dosage Dose:		
					f administration:	
				l requestoy o	administration.	
Type of Request	☐ First Request			☐ Continua	ation of Treatment	
	Complete section 6			Complete sec	tion 7	
					e section 6 if this is the first	
				authorization	requested from SSQ	
For Injection – Location where the drug is to be administered:						
☐ Home	☐ Outpatient			☐ CHSLD		
☐ Doctor's office	☐ Patient is hospitalized			☐ Other. Specify		
Exact name and address:						

Important:

To ensure sound management of its group insurance plan, SSQ gives preference to the use of biosimilar drugs. The eligibility of claims for brand-name drugs is subject to certain conditions.



Section 6: Clinical Information (first request)

Date: _____

Evaluation immediately before the start of treatment with the requested drug

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Number of joints with active synovitis: _		-
Provide at least one of the following		
Value of the C-reactive protein	mg/l	
Value of the sedimentation rate	mm/h	
Summary of trials with methotrexate		
Methotrexate		
Dosage: Ineffectiveness	☐ Intolerance ☐ Contrai	ndication
Specify:		to
Section 6: Clinical Information (continua	ation of treatment)	
Information necessary to evaluate, after	er five months or more, the	response to treatment based on the
points evaluated initially	T	T
Information related to the evaluation	First evaluation	Most recent evaluation
	Date:	Date:
Number of joints with active synovitis:		
Number of joints affected by limitation of movement:		
Value of the C-reactive protein	mg/l	mg/l
value of the c-reactive protein		
Value of sedimentation rate	mm/h	mm/h
Score on the pediatric health		
questionnaire (CHAQ) or a return to school		
Overall evaluation of the physician, the		
individual or the parent (visual analogue		
scale)		
		EV70264 (2010 02)



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Section 8: Additional information	