

PRIOR AUTHORIZATION REQUEST FORM

Abatacept (Orencia®), adalimumab (Abrilada®, Amgevita®, Hadlima®, Hulio®, Humira®, Hyrimoz®, Idacio®, Simlandi®, Yuflima®), etanercept (Brenzys®, Enbrel®, Erelzi®), infliximab (Remicade®)/ Juvenile idiopathic arthritis of polyarticular or systemic form

DECLARATION OF THE INSURED PERSON

Section 1: Information about the	plan member and the p	patient			
Name of Plan Member	Insurance Policy / Cer	tificate	Name of Employer		
Name of Patient	Date of Birth (yyyy/m	m/dd)	Telephone		
Address (house number and	City/Town		Province	Postal code	
street name)					
Section 2: Other prescription dru	g incurance				
Do you have other prescription drug insu			☐ Yes	□ No	
If so, please answer the following:	mance:		LJ Tes	□ NO	
What type of plan is it?			☐ Private	☐ Public	
Have you ever submitted a claim for this	drug to the other insurer?		☐ Yes	☐ No	
What is the status of the claim?	· ·	☐ Accepte	d 🗖 Refused	☐ Under review	
Did this insurer ask you to complete a pr	ior authorization request?	•	☐ Yes	□ No	
What is the status of the prior author	orization?	☐ Accepte	d 🗖 Refused	☐ Under review	
Please enclose acceptance or ref	usal documents, if app	licable			
Section 3: Authorization to disclo					
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 Pref		-	* *		
organization, including the Régie de				* *	
Inc. (SSQ) any of my relevant 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 relevant 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.					
Thotocopies of this document have the same value as the original.					
Signature of patient (parent/legal guardian)			Date		
IAADODTAAIT.					
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: 418-651-2588/1-800-380-2588 — Fax: 1-855-453-3942 Addresse: 2525 Laurier Blvd, P.O. Box 10500, Quebec City QC G1V 4H6					
	66 Tax. 1-655-455-5542				



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DECLARATION OF THE PRESCRIBER

Section 4: Information about the prescriber						
Name of prescriber		Spe	Specialty		Licence No.:	
				_		
Telephone:				Fax:		
I hereby certify that the	information in this reques	t is accurate	2.			
				_		
Signature of prescriber			Date			
Continu F. Duva anyond	hth.a. ath.a.vi-atia.va					
Section 5: Drug covered		Characth				
Name of Drug	Pharmaceutical Form	Strength		Dosage		
					 f administration:	
Tune of Doguest	T First Dogwood			Continue	ation of Treatment	
Type of Request	☐ First Request Complete section 6					
	complete section o			Complete sec	e section 6 if this is the first	
					requested from SSQ	
For Injection – Location	where the drug is to be ad	ministered:	:			
☐ 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.						
IMPORTANT:						
Please do not provide genetic test results						



Section 6: Clinical Information (first request)

Evaluation immediately before the start of treatment with the requested drug

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Date:	_	
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:	Intolerance	
Section 7: Clinical Information (continuatio	n of treatment)	
Information necessary to evaluate, after f	ive months or more, the respo	onse to treatment based on the
points evaluated initially 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 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)		



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Section 8: Additional information	