

ssq.ca

Prior Authorization Request Form

Abatacept (Orencia^{MD}), adalimumab (Humira®),apremilast (Otezla®), certrolizumab pegol (Cimzia®), etanercet (Enbrel®, Erelzi®), golimumab (Simponi®), infliximab (Inflectra®, Remicade®, Renflexis™), ixekizumab (Taltz®), secukinumab (Cosentyx®), Tofacitinib (Xeljanz®) ustekinumab (Stelara®) /
Moderate or severe psoriatic arthritis of rheumatoid or non-rheumatoid form

DECLARATION OF THE INSURED PERSON					
Section 1: Information about the participant and	the patient				
	Policy Certif				
Name of participant	icate Name o	of employer			
Name of patient	Telephone	Telephone			
Address (number and street name)		Postal code			
Section 2: Other prescription drug insurance police	cies				
Do you have other prescription drug insurance?	□Yes	□No			
If so, please answer the following:					
What type of plan is it?	☐ Private	☐ Public			
Have you ever submitted a claim FOR THIS DRUG to the oth	□Yes	□ No			
What is the status of the claim?		☐ Accepted	☐ Refused	☐ Under review	
Did this insurer ask you to complete a prior authorization req	□Yes	□ No			
If so, what is the status of the prior authorization request?		□Accepted	Refused	☐ Under review	
Please enclose acceptance or refusal documents, if app	licable				
Section 3: Authorization to disclose personal info	ormation				
I certify that the information in this prior authorization request i	s complete, accurate and true.				
I authorize physicians and other health care professionals, medica Quebec only) and any public or parapublic organization, includi personal information including and without limitation, any medi confidentiality obligation and authorize them to disclose the req my personal information including and without limitation, any n	ng Régie de l'assurance mala ical information and medical ex juested information to SSQ. In	die du Québec, to disclose to valuations in connection with addition, I authorize SSQ to d	SSQ, Life Insurance C the processing of this isclose to the previous	ompany Inc. (SSQ) any of my request. I hereby waive their ly named third parties any of	
Photocopies of this document have the same value as the origin	nal.				
Signature of patient (parent/legal guardian)		Date			
IMPORTANT: All correspondence concerning this form will be sent	to the address indicated in	n the narticinant's file			
		in the participant's me.			
Send us this duly completed form by mail or by fax to: 1- Telephone: 418-651-2588/1-800-380-2588 – Fax: 1-855-453-3					
Address: 2525 Laurier Blvd, P.O. Box 10500, Quebec City, QC (

DECLARATION OF THE	PHYSICIAN					
Section 4: Informati	on about the p	rescribing physician				
Name of physician Specialty				License no.		
Telephone		Fax				
I hereby certify that the in	formation in this re	equest is complete, true and	accurate.			
Signature of physician				Dat	e	
Section 5: Drug cove	ered by the aut	horization				
Drug name		Pharmaceutical form Streng		th	Dosage	
						Dose: Frequency of administration:
						——————————————————————————————————————
Type of request	☐ First reques Complete sectio	on 6 Complete sectio		first authorization requ	ested from SSQ	
IMPORTANT:						
	gement of its ara	oup insurance plans. SSO c	gives preference t	to the use of biosi	milar drugs. Eligil	bility for reference biologic products is
subject to certain cond						
First evaluation Date of evaluation Patient's weight: Number of joints with a Drug administration Monotherapy In conjunction with Specify the associar	tic Arthritis soriatic Arthritis active synovitis:					
Rheumatoid Psoriatic Arthritis		Non-Rheumato	id Psoriatic Arthritis			
Please provide at least o	ne of the following:	:			_	
Erosions are visible on x-	rays		□Yes	□No	llaalul A	ocement Questiennaine (UAQ)
Health Assessment Ques	tionnaire (HAQ) scc	ore			Health Ass	essment Questionnaire (HAQ) score:
C reactive protein (CRP)	value			mg/L		
Sedimentation rate value	<u> </u>			mm/hr		

Section 6: Clinical information (First request) (con	t'd)	
Summary of previous trials or contraindications		
Drug or other medical treatment	Reason for discontinuation	
Methotrexate Dose:	☐ Ineffectiveness ☐ Intolerance ☐ Contraindication ☐ Other, specify:	From To
Azathioprine Dose:	☐ Ineffectiveness ☐ Intolerance ☐ Contraindication ☐ Other, specify:	From To
Hydroxychloroquine Dose:	☐ Ineffectiveness ☐ Intolerance ☐ Contraindication ☐ Other, specify:	From To
Leflunomide Dose:	☐ Ineffectiveness ☐ Intolerance ☐ Contraindication ☐ Other, specify:	From To
Sulfasalazine Dose:	☐ Ineffectiveness ☐ Intolerance ☐ Contraindication ☐ Other, specify:	From To
Anti-TNF Name: Dose:	☐ Ineffectiveness ☐ Intolerance ☐ Contraindication ☐ Other, specify:	From To
Other biological agent Name: Dose:	☐ Ineffectiveness ☐ Intolerance ☐ Contraindication ☐ Other, specify:	From To
Other Name: Dose:	☐ Ineffectiveness ☐ Intolerance ☐ Contraindication ☐ Other, specify:	From To
Section 7 : Clinical information (Continuation of t	was the suit	
Information necessary to evaluate the response to treate The drug covered by the present authorization request was firs	nent	
Evaluation information	Initial evaluation	The most recent subsequent evaluation
Date of evaluation		·
Number of joints with active synovitis		
Health Assessment Questionnaire (HAQ) score		
C reactive protein (CRP) value	mg/L	mg/L
Sedimentation rate value	mm/hr	mm/r
Patient's weight	kg	kg
Return to work, where applicable	N/A	□Yes □No □N/A

Section 8: Additional information	