

## **Prior Authorization Request Form**

Adalimumab (Abrilada®, Amgevita®, Hadlima®, Hulio®, Humira®, Hyrimoz®, Idacio®, Simlandi®, Yuflima®), apremilast (Otezla®), brodulamab (Siliq™), certolizumab pegol (Cimzia®), deucravacitininb (Sotyktu®), etanercept (Brenzys®, Enbrel®, Erelzi®), guselkumab (Tremfya™), infliximab (Avsola®, Inflectra®, Remicade®, Renflexis®), ixekizumab (Taltz®), risankizumab (Skyrizi®), secukinumab (Cosentyx®), tildrakizumab (Ilumya®), ustekinumab (Stelara®) Severe to moderate chronic plaque psoriasis

DECLARATION OF THE INSURED PERSON				
Section 1: Information about the plan member and the	patient			
Name of plan member	Policy Cer	rtificate Name of	employer	
Name of patient		Telephone		
Address (number and street name)	Town/Ci	ity	Province	Postal code
Section 2: Other prescription drug insurance policies				
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 <b>FOR THIS DRUG</b> to the other insur	rer?	□Yes	□No	
What is the status of the claim?		☐ Accepted	☐ Refused	☐ Under review
Did this insurer ask you to complete a prior authorization request?		□Yes	□No	
If so, what is the status of the prior authorization request?		☐ Accepted	Refused	☐ Under review
Please enclose acceptance or refusal documents, if applicable				
Section 3: Authorization to disclose personal informat	ion			
I certify that the information in this prior authorization request is comp				
I authorize physicians and other health care professionals, medical, Quebec only) and any public or parapublic organization, including Ré- personal information including and without limitation, any medical confidentiality obligation and authorize them to disclose the requested personal information including and without limitation, any medical info	gie de l'assurance maladie information and medical e l information to SSQ. In addi	du Québec, to disclose to SSQ, Li evaluations in connection with th ition, I authorize SSQ to disclose to	ife Insurance Company ne processing of this ro the previously named	r Inc. (SSQ) any of my relevant request. I hereby waive the
Photocopies of this document have the same value as the original.				
O' to				
Signature of patient (parent/legal guardian)		Date		
IMPORTANT:				
All correspondence concerning this form will be sent to the	e address indicated in the	he participant's file.		
Send us this duly completed form by mail or by fax to: 1-855-4	53-3942.			
Telephone: 418-651-2588/1-800-380-2588 – Fax: 1-855-453-3942				

DECLARATION OF THE PRESCRIBER			
Section 4: Information about the pre	escriber		
Name of prescriber	Specialty		License no.
Telephone	Fax		
I hereby certify that the information in this requ	uest is complete, true, and accurate.		
Signature of prescriber		Date	
Section 5: Drug covered by the auth	orization		
Drug name	Pharmaceutical form	Strength	Dosage
			Dose:
			Frequency of administration:
Type of request	tion 6 Complete section 7	the first authorization requested from SSQ use of biosimilar drugs. Eligibility for reference	ce biologic products is subjectto certain
IMPORTANT: Please do not provide genetic test result:	s.		
Section 6: Clinical information (Findagnosis  ☐ Severe to moderate chronic plaque pso ☐ Other, specify:	rst request) riasis		
Administration of the prescription drug  ☐ Monotherapy (without standard systemi ☐ Other, specify:			
Evaluation before the start of treatment  Date of evaluation:  DLQI:  PASI:  OU		_kg	
☐ Presence of large lesions on the face, p	alms, soles or the genital area.		

Specify: \_\_\_

Section 6: Clinical information (First request) (cont'd)			
Summary of previous trials or contraindications			
Drug or other medical treatment	Reason for discontinuation	Duration of treatment	
Phototherapy  Number of treatments:	☐ Inaccessibility ☐ Ineffectiveness ☐ Intolerance ☐ Contraindication ☐ Other, specify:	From To	
Methotrexate	☐ Ineffectiveness	F	
Dose:	☐ Intolerance ☐ Contraindication ☐ Other, specify:	From To	
Cyclosporine	☐ Ineffectiveness☐ Intolerance	From	
Dose:	Contraindication Other, specify:	To	
Acitretin	☐ Ineffectiveness☐ Intolerance	From	
Dose:	☐ Contraindication ☐ Other, specify:	To	
Was a biologic agent previously used to treat psoriasis?			
☐ Yes (please complete information below) ☐ No			
Drug or other medical treatment	Reason for discontinuation	Duration of treatment	
Biologic agent <sup>(1)</sup>	☐ Ineffectiveness☐ Intolerance	From	
Name:	☐ Contraindication	From To	
Dose:	☐ Other, specify:		
Biologic agent <sup>(2)</sup>	☐ Ineffectiveness	F	
Name:	☐ Intolerance☐ Contraindication	From To	
Dose:	☐ Other, specify:		
Biologic agent <sup>(3)</sup>	☐ Ineffectiveness☐ Intolerance		
Name:	☐ Contraindication	From To	
Dose:	Other, specify:		
Other	☐ Ineffectiveness☐ Intolerance		
Name:	☐ Contraindication	From To	
Dose:	Other, specify:		
	0		
Section 7: Clinical information (Continuation of treatm	ent)		
Information necessary to evaluate the response to treatment			
The drug covered by the present authorization request was first take	en on:		
	First evaluation	The most recent follow-up evaluation	
Date of evaluation			
PASI			
DLQI			
Patient's weight	kg	kg	
Was there a significant improvement in the lesions on the face, palms, soles or the genital area?	N/A	☐ Yes ☐ No	

ection 8: Additional information	