

Adalimumab (Abrilada[®], Amgevita[®], Hadlima[®], Hulio[®], Humira[®], Hyrimoz[®], Idacio[®], Simlandi[®], Yuflima[®])/ Hidradenitis

DECLARATION OF THE INSURED PERSON

Section 1: Information about the p	lan member and the	patient		
Name of plan member	Insurance policy / o	ertificate	Name of empl	oyer
Name of patient	Date of birth (YYYY/N	1M/DD)	Telephone	
Address	City/Taxwa		Dunasilana	Dontal and a
Address (house number and street name)	City/Town		Province	Postal code
Section 2: Other prescription drug	insurance policies			
Do you have other prescription drug insura	ance?		☐ Yes	□ No
If so, please answer the following:				
What type of plan is it?			□ Private	☐ Public
Have you ever submitted a claim for this d	rug to the other insurer?		☐ Yes	□ No
What is the status of the claim?		☐ Accepted	l □ Refused	Under review
Did this insurer ask you to complete a prio	r authorization request?		☐ Yes	□ No
If so, what is the status of the prior au	thorization request?	☐ Accepted	l □ Refused	☐ Under review
Please enclose acceptance or refu	sal documents, if app	olicable		
Section 3: Authorization to disclose	<u> </u>			
I certify that the information in this pr	ior authorization reque	est is complete	e, accurate and to	ue.
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 Régie de I 'assurance maladie du Québec, to disclose to SSQ, Life Insurance Company 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.				
Signature of patient (parent/legal	guardian)		Dat	e
IMPORTANT:				
All correspondence concerning this for	rm will be sent to the a	ddress indica	ted in the partici	pant's file.
Send us this duly completed form by mail	or by fax to: 1-855-453-3	3942.		
Telephone: 418-651-2588/1-800-380-2588 G1V 4H6	3 – Fax: 1-855-453-3942 A	ddress: 2525 L	aurier Blvd, P.O. Bo	ox 10500, Quebec City, QC



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DECLARATION OF THE PRESCRIBER

Section 4: Information a	bout the prescriber				
Name of prescriber		Specia	alty		Licence No.:
Telephone				Fax	
I hereby certify that the	information in this reques	t is complete,	true a	and accura	te:
				_	
Signature of prescriber			Date		
Section 5 : Drug covered	by the authorization				
Name of drug	Pharmaceutical form	Strength	[Oosage	
			F	requency o	f administration:
			-		
Type of request	☐ First request		ſ	J Continua	ation of treatment
	Complete section 6			Complete sec	
					e section 6 if this is the first requested from SSQ
Injection – administered	l at:				
☐ Home	☐ Outpatient clinic		☐ CI	HSLD	
☐ Doctor's office	☐ Hospital (patient is ad	mitted)	Пο	ther Specif	Y
Exact location's name and address:					
IMPORTANT:					
		as alone CCO			
To ensure sound management of its group insurance plans, SSQ gives preference to the use of biosimilar drugs. Eligibility for reference biologic products is subject to certain conditions.					
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IMPORTANT:					
	enetic test results				
Please do not provide genetic test results					



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Section 6 : Clinical information (first request)	
Diagnosis:		
☐ Active moderate to severe hid	dradenitis suppurativa	
☐ Other. Specify :		
Number of abscesses of inflamm	natory nodules :	
Lesions are in at least two distin	ct anatomical regions : 🗖 Yes 🛛 No	
At least one of the lesions is:		
Hurley stage II ☐ Yes □	J No	
Hurley stage III ☐ Yes ☐	J No	
Summary of previous trials or co	ontraindications	
Drug or other medical treatment	Reason for discontinuation	Duration of treatment
Name:	☐ Ineffectiveness ☐ Intolerance ☐ Contraindication ☐ Other, specify:	From To
Name:	☐ Ineffectiveness ☐ Intolerance ☐ Contraindication ☐ Other, specify:	From To
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Section 7 : Clinical information (co	<u> </u>			
Information necessary to evaluate the response to treatment				
	Evaluation prior to initiation of treatment	Last evaluation		
Number of inflammatory nodules	Date:	Date:		
	Number :	Number :		
Number of abscesses	Date:	Date:		
	Number:	Number:		
Number of draining fistulas	Date:	Date:		
	Number:	Number:		
Section 8 : Additional information				
Section 6 : Additional information				