



PRIOR AUTHORIZATION REQUEST FORM
Omalizumab (Xolair®) / Moderate or severe chronic idiopathic urticaria

DECLARATION OF THE INSURED PERSON

Section 1: Information about the participant and the patient			
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?	<input type="checkbox"/> Yes	<input type="checkbox"/> No	
If so, please answer the following:			
What type of plan is it?	<input type="checkbox"/> Private	<input type="checkbox"/> Public	
Have you ever submitted a claim for this drug to the other insurer?	<input type="checkbox"/> Yes	<input type="checkbox"/> No	
What is the status of the claim?	<input type="checkbox"/> Accepted	<input type="checkbox"/> Refused	<input type="checkbox"/> Under review
Did this insurer ask you to complete a prior authorization request?	<input type="checkbox"/> Yes	<input type="checkbox"/> No	
If so, what is the status of the prior authorization request?	<input type="checkbox"/> Accepted	<input type="checkbox"/> Refused	<input type="checkbox"/> Under review
<i>Please enclose acceptance or refusal documents, if applicable</i>			

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 Régie de l'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) _____	Date: _____

IMPORTANT:
All correspondence concerning this form will be sent to the address indicated in the participant's file.

Send us this duly completed form by mail or by fax at: 1-855-453-3942.
Telephone: 418-651-2588 /1-866-332-2588 – Fax: 1-855-453-3942 Address: 2525 Laurier Blvd, P.O. Box 10500, Quebec City, QC G1V 4H6
ssq.ca



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DECLARATION OF THE PHYSICIAN

Section 4: Information about the physician		
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 Xolair	Pharmaceutical Form	Strength	Dosage Dose: _____ Frequency of administration: _____
Type of request			
<input type="checkbox"/> First request Complete Section 6	<input type="checkbox"/> Request for continuation of treatment after 24 weeks Complete Section 7 Also complete Section 6 if this is the first authorization requested from SSQ	<input type="checkbox"/> Subsequent request following a relapse after stopping treatment Complete Section 8	
For injection – Location where prescription drug is to be administered:			
<input type="checkbox"/> Home	<input type="checkbox"/> Outpatient	<input type="checkbox"/> CHSLD	
<input type="checkbox"/> Doctor's office	<input type="checkbox"/> Hospital	<input type="checkbox"/> Other. Specify _____	



Section 6: Clinical information (first request)

Therapeutic indication

For a person suffering from **moderate or severe chronic idiopathic urticaria (CIU)**

Other. Specify. _____

Urticaria Activity Score 7 (UAS7): _____

Summary of pervious tests

ANTIHISTAMINES	RESULTS	TEST PERIOD (IF APPLICABLE)
Name: _____ Dosage (optimized dose): _____	<input type="checkbox"/> Poor control <input type="checkbox"/> Other Specify: _____	From _____ To _____
Name: _____ Dosage (optimized dose): _____	<input type="checkbox"/> Poor control <input type="checkbox"/> Other Specify: _____	From _____ To _____
Other agent Name: _____ Dosage (optimized dose): _____	<input type="checkbox"/> Poor control <input type="checkbox"/> Other Specify: _____	From _____ To _____
Other agent Name: _____ Dosage (optimized dose): _____	<input type="checkbox"/> Poor control <input type="checkbox"/> Other Specify: _____	From _____ To _____

Section 7: Clinical information (continuation of treatment after 24 weeks)

Full response over a period of less than 12 weeks (UAS7 ≤ 6)

Partial response (drop in UAS7 score of at least 9.5 points since the start and UAS7 > 6)



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UAS7 scores over the 24 weeks of treatment

Evaluation date: _____	UAS7 score: _____
Evaluation date: _____	UAS7 score: _____
Evaluation date: _____	UAS7 score: _____
Evaluation date: _____	UAS7 score: _____
Evaluation date: _____	UAS7 score: _____
Evaluation date: _____	UAS7 score: _____

Section 8: Clinical information (request following a relapse after stopping treatment)

Previous treatment

Date of last injection: _____

Response:

Satisfactory

UAS7 score: _____

Other. Specify. _____

Current UAS7 score indicating a relapse: _____

Section 9: Additional information
