



PRIOR AUTHORIZATION REQUEST FORM
Cobimetinib (Cotellic®) / Unresectable or metastatic melanoma

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 policies			
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
Please enclose acceptance or refusal documents, if applicable			

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 to: 1-855-453-3942.
Telephone: 418-651-2588/1-800-380-2588 – 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 prescribing physician		
Name of physician	Specialty	License no.
Telephone	Fax	
I hereby certify that the information in this request is complete, true and accurate.		
Signature of physician _____		Date _____

Section 5 : Drug covered by the authorization			
Name of drug	Pharmaceutical form	Strength	Dosage Dose: _____ Frequency of administration: _____
Cobimetinib	Tablet		
Type of request	<input type="checkbox"/> First request Complete section 6	<input type="checkbox"/> Continuation of treatment Complete section 7 Also complete section 6 if this is the first authorization requested from SSQ	

Section 6 : Clinical information (First request)
Diagnosis <input type="checkbox"/> Unresectable melanoma <input type="checkbox"/> Metastatic melanoma <input type="checkbox"/> Other. Specify : _____
BRAF V600 Mutation <input type="checkbox"/> Presence <input type="checkbox"/> Absence
Administration of cobimetinib <input type="checkbox"/> In conjunction with vemurafenib <input type="checkbox"/> Other. Specify : _____



Section 6 : Clinical information (First request) (cont'd)

Pharmacological treatment

First-line treatment

Second-line treatment following failure of prior cytotoxic chemotherapy or immunotherapy targeting PD-1 or CTLA-4.

Other. Specify: _____

Actual value of the ECOG performance status

0 1 2 3 4

Summary of previous trials or contraindications

Cytotoxic chemotherapy		
<input type="checkbox"/> Dacarbazine	<input type="checkbox"/> Ineffectiveness <input type="checkbox"/> Intolerance <input type="checkbox"/> Contraindication <input type="checkbox"/> Other, specify: _____	from _____ to _____
<input type="checkbox"/> Carboplatin	<input type="checkbox"/> Ineffectiveness <input type="checkbox"/> Intolerance <input type="checkbox"/> Contraindication <input type="checkbox"/> Other, specify: _____	from _____ to _____
<input type="checkbox"/> Paclitaxel	<input type="checkbox"/> Ineffectiveness <input type="checkbox"/> Intolerance <input type="checkbox"/> Contraindication <input type="checkbox"/> Other, specify: _____	from _____ to _____
<input type="checkbox"/> Temozolomide	<input type="checkbox"/> Ineffectiveness <input type="checkbox"/> Intolerance <input type="checkbox"/> Contraindication <input type="checkbox"/> Other, specify: _____	from _____ to _____

Immunotherapy targeting :

<input type="checkbox"/> CTLA-4 :Ipilimumab	<input type="checkbox"/> Ineffectiveness <input type="checkbox"/> Intolerance <input type="checkbox"/> Contraindication <input type="checkbox"/> Other, specify: _____	from _____ to _____
<input type="checkbox"/> TPD-1 : Pembrolizumab	<input type="checkbox"/> Ineffectiveness <input type="checkbox"/> Intolerance <input type="checkbox"/> Contraindication <input type="checkbox"/> Other, specify: _____	from _____ to _____



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Section 6 : Clinical information (First request) (cont'd)

BRAF-inhibitors :		
<input type="checkbox"/> Dabrafenib	<input type="checkbox"/> Ineffectiveness <input type="checkbox"/> Intolerance <input type="checkbox"/> Contraindication <input type="checkbox"/> Other, specify: _____	from _____ to _____
<input type="checkbox"/> Vemurafenib	<input type="checkbox"/> Ineffectiveness <input type="checkbox"/> Intolerance <input type="checkbox"/> Contraindication <input type="checkbox"/> Other, specify: _____	from _____ to _____
MEK-inhibitors :		
<input type="checkbox"/> Cobimetinib	<input type="checkbox"/> Ineffectiveness <input type="checkbox"/> Intolerance <input type="checkbox"/> Contraindication <input type="checkbox"/> Other, specify: _____	from _____ to _____
<input type="checkbox"/> Trametinib	<input type="checkbox"/> Ineffectiveness <input type="checkbox"/> Intolerance <input type="checkbox"/> Contraindication <input type="checkbox"/> Other, specify: _____	from _____ to _____
Other agent Name : _____	<input type="checkbox"/> Ineffectiveness <input type="checkbox"/> Intolerance <input type="checkbox"/> Contraindication <input type="checkbox"/> Other, specify: _____	from _____ to _____



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Section 7 : Clinical information (Continuation of treatment)

Information necessary to evaluate the response to treatment

The drug covered by the present authorization request was first taken on: _____

Positive clinical effects observed

- Absence of disease progression
- Other, specify: _____

Positive clinical effects confirmation

- Response to treatment confirmed by medical imagery. Last imagery : _____
- Response to treatment confirmed by a physical exam
- Other. Specify : _____

Section 8 : Additional information
