

DECLARATION OF THE INSURED PERSON

Section 1: Information about the plan member and the patient			
Name of plan member	Insurance policy / certificate	Name of empl	oyer
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?		🗖 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 other insurer?		🗖 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 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 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) ____

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



DECLARATION OF THE PRESCRIBER

Section 4: Information about the prescriber			
Name of prescriber	Specialty		Licence No.:
Telephone		Fax	
I hereby certify that the information in this request is complete, true and accurate:			
		_	
Signature of prescriber		D	Date

Section 5 : Drug covered by the authorization			
Name of drug	Pharmaceutical form	Strength	Dosage Dose: Frequency of administration:
Type of request	First request Complete section 6		Continuation of treatment Complete section 7 Also complete section 6 if this is the first authorization requested from SSQ

IMPORTANT:

Please do not provide any genetic test results

Section 6 : Clinical information (first request)

Diagnosis

Non-small cell lung cancer (NSCLC) non resectable locally advanced or metastatic in compliance with Health Canada indication

For informational purposes only:

Alecensaro, Alunbrig, and Zycadia are Health Canada indicated for:

- the first line treatment of adult patients with anaplastic lymphoma kinase (ALK)-positive locally advanced (not amenable to curative therapy) or metastatic non-small cell lung cancer (NSCLC).
- monotherapy treatment of adult patients with anaplastic lymphoma kinase (ALK)-positive metastatic nonsmall cell lung cancer (NSCLC) who have progressed on or who were intolerant to an ALK inhibitor (crizotinib).



Alectinib (Alecensaro[®]), Brigatinib (Alunbrig[®]), Ceritinib (Zykadia[®]) / Locally advanced or metastatic non-small cell lung cancer (NSCLC)

Section 6 : Clinical information (first request) (cont'd)							
Adr	nin	istration					
	١n ı	monotherapy					
	Otl	her. Specify :					
ACT	ΓUA	L value of performa	ance status				
ECC)G	□ 0	□ 1	□ 2	□ 3	□ 4	
Pha	rm	acologic treatment					
	Firs	st-line treatment					
	Otl	her. Specify :					
Crizotinib previous trial							
	Yes	5					
Discontinuation reason:							
	0	Cancer has progres	ssed despite the	e administration			
	0	Intolerance. Specif	y:				
	0	Other. Specify :					
	No						

Section 7 : Clinical information (continuation of treatment)

Administration

	In monotherapy	
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□ Other. Specify : _____

Beneficial clinical effect observed

Treatment start date	•
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Absence of disease progress	ion
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Other. Specify : ______



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Section 7 : Clinical information (continuation of treatment) (cont'd)

Imaging confirmation (if crizotinib previous trials EXCLUSIVELY) :

- **T**reatment response **confirmed** by imaging :
 - Last imaging date : ______

□ Treatment response **not confirmed** by imaging :

- Last imaging date : ______
- Reasons preventing to proceed with imaging :

Section 8 : Additional information	