

# PRIOR AUTHORIZATION REQUEST FORM Ibrutinib (Imbruvica®) / Mantle-cell lymphoma

## **DECLARATION OF THE INSURED PERSON**

	1 ENSON						
Section 1: Information about the p	lan member and the p	atient					
Name of plan member	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 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		☐ Yes	□ No				
What is the status of the claim?		☐ Accepted	☐ Refused	☐ Under review			
Did this insurer ask you to complete a prio		☐ Yes	□ No				
If so, what is the status of the prior au	☐ 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 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 <b>patient</b> (parent/legal	guardian)		Dat	e			
IMPORTANT :							
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 PRESCRIBER**

Section 4: Information about the prescriber								
Name of prescriber		Specialty		Licence No.:				
Teleph	one				Fax			
I hereb	y certify that the	information in this req	uest is cor	nplete, true	and accura	te:		
Signatu	ure of <b>prescriber</b> _			Date				
Section	າ 5 : Drug covered	by the authorization						
Name	of drug	Pharmaceutical form	Strer	igth	Dosage			
Imbruv	uica.	Capsule	140 r	ma				
iiiibiuv	rica	Capsule	1401	ı ığ	Frequency o	f administration:		
Type o	f request	☐ First request	·		☐ Continua	ation of treatment		
		Complete section 6			Complete sec	tion 7		
						e section 6 if this is the first		
					authorization	requested from SSQ		
IMPOR	RTANT:							
Please	do not provide a	ny genetic test results						
		, 0						
Section	n 6 : Clinical inforr	nation (first request)						
Diagno	osis							
	☐ Recurrent or refractory mantle-cell lymphoma							
	Other. Specify:							
	· , -							
Imbruvica administration								
	As monotherapy	,						
	Other. Specify:							
Performance status ACTUAL value								
ECOG	□ 0	<b>1</b>	<b>□</b> 2	□ 3	ſ	<b>3</b> 4		



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Section 6 : Clinical information (first request) (cont'd)								
Summary of previous treatments or contraindications								
PROTOCOL ADMINISTERED	REASON FOR STOP OR NON-ADMINISTRATION							
☐ RITUXIMAB-based chemotherapy	☐ Failure Specify:	☐ Contraindication	☐ Other					
☐ No treatment received so far	Specify :							
☐ Other	Specify:							
Section 7 : Clinical information (continuation of Beneficial clinical effect observed with Imbruvi  Treatment was started on (YYYY-MM-DD):  Absence of disease progression  Other. Specify :	ca							
Section 8 : Additional information								