

Regorafenib (Stivarga®) / Hepatocellular carcinoma

DECLARATION OF THE INSURED PERSON

Section 1: Information about the pla	n member and the	patient				
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 in	curance					
Section 2: Other prescription drug insurance						
Do you have other prescription drug insurance?			☐ Yes	□ No		
If so, please answer the following:			- - · ·	7 5 1 11		
What type of plan is it?			☐ Private	☐ Public		
Have you ever submitted a claim for this dru	g to the other insurer?	. .	☐ Yes	□ No		
What is the status of the claim?	☐ Accepted			☐ Under review		
	Did this insurer ask you to complete a prior authorization request? If so, what is the status of the prior authorization request? Accepted		☐ Yes	□ No		
If so, what is the status of the prior auth	d □ Refused	☐ Under review				
Please enclose acceptance or refusa	il documents, if app	olicable				
Section 3: Authorization to disclose p	acreanal informatio	n				
I certify that the information in this prior			e, accurate and ti	rue.		
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 ori	ginal.				
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: 418-651-2588/1-800-380-2588 – G1V 4H6	Fax: 1-855-453-3942 A	ddress: 2525 L	aurier Blvd, P.O. Bo	ox 10500, Quebec City, QC		

PRIOR AUTHORIZATION REQUEST FORM



Regorafenib (Stivarga®) / Hepatocellular carcinoma

DECLARATION OF THE PRESCRIBER

Section 4: Information a	bout the prescriber						
Name of Prescriber			Specialty		Licence No.:		
Telephone				Fax			
I hereby certify that the information in this request is accurate:							
Signature of Prescriber				Date			
Signature of Frescriber							
Section 5: Drug covered	by the authorization						
Drug name	Pharmaceutical form	Strength		Dosage			
Regorafenib				Dose:			
Regulateriib				Frequency o	f administration:		
Type of request	☐ First request	Tirst request		☐ Continuation of treatment			
	Complete Section 6			Complete Sec	Complete Section 7		
				Also complete Section 6 if this is the first authorization requested from SSQ			
				authorization	requested from SSQ		
IMPORTANT:							
Please do not provide any genetic test results							
Section 6: Clinical inform	lation (first request)						
Therapeutic Indication							
Hepatocellular carcinoma resistant to sorafenib							
☐ Other. Specify:							
About sorafenib							
☐ The patient tolerated a previous sorefenib treatment, defined by the administration of a dose equal to							
400 mg or more per day for less than 20 of the 28 days before ceasing sorafenib							
☐ Other. Specify:							
Hepatic damage stage							
Child-Pugh ☐ A ☐ B ☐	C Other. Specify:						





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Section 6: Clinical information (first request) (Cont'd)
Value of the <u>ACTUAL</u> performance status
ECOG 🗆 0 🗆 1 🗔 2 🗔 3 🗔 4
Administration of regorafenib
☐ In monotherapy
☐ Other. Specify:
Section 7: Clinical information (Continuation of treatment)
Observed beneficial clinical effect Start date of treatment:
Start date of treatment.
☐ Absence of disease progression
☐ Other. Specify:
Confirmation by imaging
☐ Response to treatment confirmed by imaging
Date of last imaging:
☐ Response to treatment NOT confirmed by imaging
Date of next imaging:
Reason that prevented proceeding with imaging:
Section 8: Additional information