PRIOR AUTHORIZATION REQUEST FORM
Aflibercept (Eylea®), ranibizumab (Byooviz®, Lucentis®) /
Visual impairment due to macular oedema following a central retinal vein occlusion (CRVO)

DECLARATION OF THE INSURED PERSON

Section 1 : Information about the plan 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 (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.

_____ YYYY-MM-DD

Signature of **patient** (parent/legal guardian) Date

IMPORTANT :

All correspondence concerning this form will be sent to the address indicated in the plan member'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	License no.
Telephone	Fax	
I hereby certify that the information in this request is complete, true and accurate. _____ YYYY-MM-DD Signature of **prescriber** Date		

Section 5 – Drug covered by the authorization			
Drug name	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:
To ensure sound management of its group insurance plans, SSQ gives preference to the use of biosimilar drugs. Eligibility for reference biologic products is subject to certain conditions.

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IMPORTANT:
Please do not provide any genetic test results

Section 6 : Clinical information (First request)	
Therapeutic indication ❒ Visual impairment due to macular oedema following a central retinal vein occlusion (CRVO) ❒ Other. Specify: _____	
Left eye	**Right eye**
Administration of the drug covered by the authorization	
❒ Monotherapy ❒ In conjunction: Specify agent: _____	❒ Monotherapy ❒ In conjunction: Specify agent: _____
Optimum visual acuity after correction	
❒ Between 6/12 and 6/96 ❒ Other. Specify: _____	❒ Between 6/12 and 6/96 ❒ Other. Specify: _____
Thickness of the central retina	
❒ ≥ 250µm ❒ Other. Specify: _____	❒ ≥ 250µm ❒ Other. Specify: _____
Afferent pupillary defect	
❒ Absence ❒ Presence	❒ Absence ❒ Presence

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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: YYYY-MM-DD	
Left eye	**Right eye**
Visual acuity measured by the Snellen test	
Date : YYYY-MM-DD ❒ Stabilization ❒ Improvement ❒ Deterioration	Date : YYYY-MM-DD ❒ Stabilization ❒ Improvement ❒ Deterioration
Macular oedema evaluated by optical coherence tomography	
Date : YYYY-MM-DD ❒ Stabilization ❒ Improvement ❒ Deterioration	Date : YYYY-MM-DD ❒ Stabilization ❒ Improvement ❒ Deterioration

Section 8 : Additional information