



**PRIOR AUTHORIZATION REQUEST FORM**  
**Aflibercept (Eylea®), Ranibizumab (Byooviz®, Lucentis®) / Neovascular (wet)**  
**age-related macular degeneration (AMD)**

**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 (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
<b>Please enclose acceptance or refusal documents, if applicable</b>			

Section 3: Authorization to disclose personal information
<p>I certify that the information in this prior authorization request is complete, accurate and true.</p> <p>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.</p> <p>Photocopies of this document have the same value as the original.</p> <p>Signature of <b>patient</b> (parent/legal guardian) _____ Date _____</p>

**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



**PRIOR AUTHORIZATION REQUEST FORM**  
**Aflibercept (Eylea®), Ranibizumab (Byooviz®, Lucentis®) / Neovascular (wet) age-related macular degeneration (AMD)**

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 <b>prescriber</b> _____		Date _____

Section 5: Drug covered by the authorization			
Name of drug	Pharmaceutical form	Strength	<b>Dosage</b> Dose: _____ Frequency of administration: _____
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

**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.



**PRIOR AUTHORIZATION REQUEST FORM**  
**Aflibercept (Eylea®), Ranibizumab (Byooviz®, Lucentis®) / Neovascular (wet) age-related macular degeneration (AMD)**

**IMPORTANT:**  
Please do not provide any genetic test results

**Section 6: Clinical information (first request)**

<b>Therapeutic indication</b>	
<input type="checkbox"/> Neovascular (wet) age-related macular degeneration (AMD)	
<input type="checkbox"/> Other. Specify: _____	
<b>Left eye</b>	<b>Right eye</b>
<b>Administration of requested prescription drug</b>	
<input type="checkbox"/> Monotherapy	<input type="checkbox"/> Monotherapy
<input type="checkbox"/> In conjunction with Specify the agent: _____	<input type="checkbox"/> In conjunction with Specify the agent: _____
<b>Optimum visual acuity after correction</b>	
<input type="checkbox"/> Between 6/12 and 6/96	<input type="checkbox"/> Between 6/12 and 6/96
<input type="checkbox"/> Other. Specify: _____	<input type="checkbox"/> Other. Specify: _____
<b>Linear dimension of the lesion</b>	
<input type="checkbox"/> ≤ 12 disc surfaces	<input type="checkbox"/> ≤ 12 disc surfaces
<input type="checkbox"/> Other. Specify: _____	<input type="checkbox"/> Other. Specify: _____
<b>State of the centre of the macula</b>	
<input type="checkbox"/> No significant permanent structural damage*	<input type="checkbox"/> No significant permanent structural damage*
<input type="checkbox"/> Other. Specify: _____	<input type="checkbox"/> Other. Specify: _____
<small>*The damage is defined as fibrosis, atrophy or chronic disciform scarring, the seriousness of which prevents obtaining a functional benefit according to the attending physician</small>	
<b>Evolution of the illness over the past 3 months confirmed by:</b>	
<input type="checkbox"/> Retinal angiography	<input type="checkbox"/> Retinal angiography
<input type="checkbox"/> Optical coherence tomography	<input type="checkbox"/> Optical coherence tomography
<input type="checkbox"/> Recent visual acuity change	<input type="checkbox"/> Recent visual acuity change
<input type="checkbox"/> Other. Specify: _____	<input type="checkbox"/> Other. Specify: _____



**PRIOR AUTHORIZATION REQUEST FORM**  
**Aflibercept (Eylea®), Ranibizumab (Byooviz®, Lucentis®) / Neovascular (wet) age-related macular degeneration (AMD)**

**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
----------	-----------

**Response to treatment**

<input type="checkbox"/> Stabilization <input type="checkbox"/> Improvement <input type="checkbox"/> Deterioration	<input type="checkbox"/> Stabilization <input type="checkbox"/> Improvement <input type="checkbox"/> Deterioration
--	--

**Medical exam used**

Date: _____ <input type="checkbox"/> Retinal angiography <input type="checkbox"/> Optical coherence tomography <input type="checkbox"/> Other. Specify: _____	Date: _____ <input type="checkbox"/> Retinal angiography <input type="checkbox"/> Optical coherence tomography <input type="checkbox"/> Other. Specify: _____
--	--

**Section 8: Additional information**
