

Request for Prior Authorization for Lucentis (ranibizumab) Website Form – www.highmarkhealthoptions.com Submit request via: Fax - 1-855-476-4158

All requests for Lucentis (ranibizumab) require a prior authorization and will be screened for medical necessity and appropriateness using the criteria listed below.

<u>Lucentis (ranibizumab) Prior Authorization Criteria:</u>

For all requests for Lucentis (ranibizumab) all of the following criteria must be met:

- The member is 18 years of age or older
- The treatment is prescribed by, or in consultation with, an ophthalmologist
- The requested dose and frequency is in accordance with FDA-approved labeling, nationally recognized compendia, and/or evidence-based practice guidelines
- The member does not have an active ocular or periocular infection
- The member has tried and failed or had an intolerance to Avastin

Coverage may be provided with a <u>diagnosis</u> of Neovascular (Wet) Age-Related Macular Degeneration (AMD).

Coverage may be provided with a <u>diagnosis</u> of Macular Edema Following Retinal Vein Occlusion.

Coverage may be provided with a <u>diagnosis</u> of Diabetic Macular Edema and the following criteria is met:

- Must provide documentation of clinically significant macular edema (CSME) defined as any of the following:
 - o Retinal thickening within 500 µm of the macular center
 - $\circ~$ Hard exudates within 500 μm of the macular center with adjacent retinal thickening
 - One or more disc diameters of retinal thickening, part of which is within one disc diameter of the macular center

Coverage may be provided with a <u>diagnosis</u> of Diabetic Retinopathy.

Coverage may be provided with a <u>diagnosis</u> of Myopic Choroidal Neovascularization.

Initial Duration of Approval: 12 months

Reauthorization criteria:

o Member continues to meet initial criteria for medical necessity

Reauthorization Duration of Approval: 12 months



Coverage may be provided for any non-FDA labeled indication if it is determined that the use is a medically accepted indication supported by nationally recognized pharmacy compendia or peer-reviewed medical literature for treatment of the diagnosis(es) for which it is prescribed. These requests will be reviewed on a case by case basis to determine medical necessity.



LUCENTIS (ranibizumab) PRIOR AUTHORIZATION FORM

Please complete and fax all requested information below including any progress notes, laboratory test results, or chart documentation as applicable to Highmark Health Options Pharmacy Services. **FAX:** (855) 476-4158

If needed, you may call to speak to a Pharmacy Services Representative.

PHONE : (844) 325-6253 Monday tl	nrough Friday 8:30am to 5:00pm		
PROVIDER INF	ORMATION		
Requesting Provider:	NPI:		
Provider Specialty:	Office Contact:		
Office Address:	Office Phone:		
	Office Fax:		
MEMBER INFO	DRMATION		
Member Name:	DOB:		
Health Options ID:	Member weight:pounds orkg		
REQUESTED DRUG	INFORMATION		
Medication:	Strength:		
Frequency:	Duration:		
Is the member currently receiving requested medication? Yes	No Date Medication Initiated:		
Is this medication being used for a chronic or long-term condition			
the patient? Yes No	•		
Billing Info	rmation		
This medication will be billed: at a pharmacy OR			
medically (if medically please	provide a JCODE:		
	per's home Other		
Place of Service	Information		
Name:	NPI:		
Address:	Phone:		
MEDICAL HISTORY (Con	nplete for ALL requests)		
Diagnosis:			
Neovascular (Wet) Age-Related Macular Degeneration (AMD	D) ICD-10 Code:		
Macular Edema following Retinal Vein Occlusion	ICD-10 Code:		
Diabetic Macular Edema	ICD-10 Code:		
Diabetic Retinopathy	ICD-10 Code:		
Myopic Choroidal Neovascularization	ICD-10 Code:		
Other:	ICD-10 Code:		
Guiei.	105 To Code.		
Does the member have an active ocular or periocular infection?	Yes No		
Has the member tried and failed Avastin? Yes No			
For the diagnosis of Diabetic Macular Edema:			
Must provide documentation of clinically significant macular ede	ma (CSME) defined as any of the following:		
 Retinal thickening within 500 μm of the macular center 			
 Hard exudates within 500 μm of the macular center with 			
 One or more disc diameters of retinal thickening, part of which is within one disc diameter of the macular center 			



LUCENTIS (ranibizumab) PRIOR AUTHORIZATION FORM (CONTINUED) – PAGE 2 OF 2

Please complete and fax all requested information below including any progress notes, laboratory test results, or chart documentation as applicable to Highmark Health Options Pharmacy Services. **FAX:** (855) 476-4158

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	ONE: (844) 325-6253 Mond	•	•	
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Member Name:		DOB:		
Health Options ID:		Member weight: _	pounds or	kg
	CURRENT or PR	REVIOUS THERAPY	Y	
Medication Name	Strength/ Frequency	Dates of Therapy	Status (Discontinued & Why/Current)	
Has the member experienced as		ORIZATION	□ No	
Has the member experienced a significant improvement with treatment? Yes No SUPPORTING INFORMATION or CLINICAL RATIONALE				
	II OKIINO INFORMATI	ON OF CLINICAL R	AHONABE	
Prescribing Prov	ider Signature		Date	