

## Lucentis

## **Prior Authorization Request**

CVS Caremark administers the prescription benefit plan for the patient identified. This patient's benefit plan requires prior authorization for certain medications in order for the drug to be covered. To make an appropriate determination, providing the most accurate diagnosis for the use of the prescribed medication is necessary. **Please respond below and fax this form to CVS Caremark toll-free at 1-855-330-1720**. If you have questions regarding the prior authorization, please contact CVS Caremark at **1-888-877-0518**. For inquiries or questions related to the patient's eligibility, drug copay or medication delivery; please contact the Specialty Customer Care Team: CaremarkConnect® 1-800-237-2767.

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Patient's Name:		Date:
Patient's ID:	Patient's Date of Birth:	
Physician's Name:		
Specialty:		NPI#:
Physician Office Telephone:	Physician Office Fax:	
<b>Referring</b> Provider Info: ☐ Same as Re	equesting Provi	der
Name:		NPI#:
Fax:		Phone:
Rendering Provider Info: ☐ Same as Ro Name:	-	er  Same as Requesting Provider NPI#:
Fax:		Phone:
		in accordance with FDA-approved labeling, vidence-based practice guidelines.
Patient Weight:	kg	
Patient Height:	cm	
Please indicate the place of service for the	requested drug:	
☐ Ambulatory Surgical	☐ Home	Off Campus Outpatient Hospital
☐ On Campus Outpatient Hospital	$\square$ Office	□ Pharmacy

Exc	ception Criteria Questions: Skip to Criteria questions if the member is MD or VA Risk			
A.	The preferred product for your patient's health plan is Avastin.  Can the patient's treatment be switched to the preferred product?  Yes, Please obtain Form for preferred product and submit for corresponding PA.  No			
B.	Is this request for continuation of therapy with the requested product? $\square$ Yes $\square$ No, If No, skip to Question D			
C.	Is the patient currently receiving the requested product through samples or a manufacturer's patient assistance program? If unknown, answer Yes. $\square$ Yes $\square$ No If No, skip to Clinical Criteria Questions			
D.	Does the patient have a documented inadequate response or intolerable adverse event with the preferred product (Avastin)? <i>ACTION REQUIRED: If 'Yes'</i> , <i>attach supporting chart note(s)</i> . □ Yes □ No			
	What is the diagnosis?  Diabetic macular edema  Neovascular (wet) age-related macular degeneration  Macular edema due to retinal vein occlusion  Diabetic retinopathy  Myopic choroidal neovascularization  Other			
2.	What is the ICD-10 code?			
3.	Is this a request for continuation of therapy? $\square$ Yes $\square$ No If No, no further questions			
4.	Has the patient demonstrated a positive clinical response to therapy (e.g., improvement or maintenance in best corrected visual acuity [BCVA] or visual field, or a reduction in the rate of vision decline or the risk of more severe vision loss)? $\square$ Yes $\square$ No			

Step Therapy Override: Complete if Applicable for the state of Maryland.		Please Circle	
Is the requested drug being used to treat stage four advanced metastatic cancer?		No	
Is the requested drug's use consistent with the FDA-approved indication or the National Comprehensive Cancer Network Drugs & Biologics Compendium indication for the treatment of stage four advanced metastatic cancer and is supported by peer-reviewed medical literature?		No	
Is the requested drug being used for an FDA-approved indication OR an indication supported in the compendia of current literature (examples: AHFS, Lexicomp, Clinical Pharmacology, Micromedex, current accepted guidelines)?		No	
Does the prescribed quantity fall within the manufacturer's published dosing guidelines or within dosing guidelines found in the compendia of current literature (examples: package insert, AHFS, Lexicomp, Clinical Pharmacology, Micromedex, current accepted guidelines)?		No	
Do patient chart notes document the requested drug was ordered with a paid claim at the pharmacy, the pharmacy filled the prescription and delivered to the patient or other documentation that the requested drug was prescribed for the patient in the last 180 days?		No	
Has the prescriber provided proof documented in the patient chart notes that in their opinion the requested drug is effective for the patient's condition?		No	

Step Therapy Override: Complete if Applicable for the state of Virginia.		Please Circle	
Is the requested drug being used for an FDA-approved indication or an indication supported in the compendia of current literature (examples: AHFS, Micromedex, current accepted guidelines)?	Yes	No	
Does the prescribed dose and quantity fall within the FDA-approved labeling or within dosing guidelines found in the compendia of current literature?		No	
Is the request for a brand drug that has an AB-rated generic equivalent or interchangeable biological product available?		No	
Has the patient had a trial and failure of the AB-rated generic equivalent or interchangeable biological product due to an adverse event (examples: rash, nausea, vomiting, anaphylaxis) that is thought to be due to an inactive ingredient?		No	
Is the preferred drug contraindicated?		No	
Is the preferred drug expected to be ineffective based on the known clinical characteristics of the patient and the prescription drug regimen?		No	
Has the patient tried the preferred drug while on their current or previous health benefit plan and it was discontinued due to lack of efficacy or effectiveness, diminished effect, or an adverse event?		No	
Is the patient currently receiving a positive therapeutic outcome with the requested drug for their medical condition?	Yes	No	

I attest that this information is accurate and true, and that documentation supporting this information is available for review if requested by CVS Caremark or the benefit plan sponsor.

(	
Prescriber or Authorized Signature	Date (mm/dd/yy)

Send completed form to: Case Review Unit CVS Caremark Specialty Programs Fax: 1-855-330-1720

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