# Lucentis® (ranibizumab)



## **Pharmacy Coverage Policy**

Effective Date: January 01, 2024 Revision Date: February 26, 2025 Review Date: February 19, 2025 Line of Business: Medicaid - Louisiana

Policy Type: Prior Authorization

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#### **Products Affected**

Lucentis intravitreal solution for injection

Lucentis intravitreal syringe

#### **Listed Indications**

Neovascular (Wet) Age-Related Exudative Macular Degeneration (AMD)

<u>Diabetic Macular Edema (DME)</u> Diabetic Retinopathy (DR)

Macular Edema Following Retinal Vein Occlusion (RVO)

Myopic Choroidal Neovascularization (mCNV)

Neovascular (Wet) Age-Related Exudative Macular Degeneration (AMD)			
Does the member meet all of the following criteria?			
Criteria #1	Diagnosed with neovascular (wet) age-related macular degeneration		
Criteria #2	Has a contraindication, or intolerance to bevacizumab. OR Has had prior therapy with bevacizumab and provider attests that the member has NOT demonstrated a positive clinical response to bevacizumab (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).		
Approval Duration			
Initial	Lucentis (ranibizumab) will be approved in plan year duration or as determined through clinical review.		
Renewal	Lucentis (ranibizumab) will be approved in plan year duration or as determined through clinical review.		
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Does the member meet all of the following criteria?  Criteria #1  Diagnosed with Diabetic Macular Edema	Diabetic Macular Edema (DME)			
	Does the member meet all of the following criteria?			
Criteria #2  Has a contraindication, or intolerance to bevacizumab. OR Has had prior therapy was provider attests that the member has NOT demonstrated a positive clinical responsing improvement or maintenance in best corrected visual acuity [BCVA] or visual field, wision decline or the risk of more severe vision loss).	nse to bevacizumab (e.g.,			
Approval Duration				
<u>Lucentis (ranibizumab) will be approved in plan year duration or as determined through the plan year duration of the plan year </u>	ough clinical review.			
Renewal Lucentis (ranibizumab) will be approved in plan year duration or as determined through the control of t	ough clinical review.			

Diabetic Retinopathy (DR)		
Does the member meet all of the following criteria?		
Criteria #1	Diagnosed with Diabetic Retinopathy	
	Has a contraindication, or intolerance to bevacizumab. OR Has had prior therapy with bevacizumab and provider attests that the member has NOT demonstrated a positive clinical response to bevacizumab (e.g.,	

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	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).	
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Macular Edema Following Retinal Vein Occlusion (RVO)			
Does the member meet all of the following criteria?			
Criteria #1	Diagnosed with macular edema following Retinal Vein Occlusion		
Criteria #2	Has a contraindication, or intolerance to bevacizumab. OR Has had prior therapy with bevacizumab and provider attests that the member has NOT demonstrated a positive clinical response to bevacizumab (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).		
Approval Duration			
<u>Initial</u>	Lucentis (ranibizumab) will be approved in plan year duration or as determined through clinical review.		
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Myopic Choroidal Neovascularization (mCNV)			
Does the member meet all of the following criteria?			
Criteria #1	Diagnosed with Myopic Choroidal Neovascularization (mCNV)		
<u>Criteria #2</u>	Has a contraindication, or intolerance to bevacizumab. OR Has had prior therapy with bevacizumab and provider attests that the member has NOT demonstrated a positive clinical response to bevacizumab (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).		
Approval Duration			
<u>Initial</u>	Lucentis (ranibizumab) will be approved in plan year duration or as determined through clinical review.		
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#### **Background**

This is a prior authorization policy about Lucentis (ranibizumab).

Lucentis (ranibizumab) is a recombinant monoclonal antibody, ophthalmic Vascular Endothelial Growth Factor (VEGF) Inhibitor.

Lucentis (ranibizumab) binds to and inhibits vascular endothelial growth factor (VEGF-A) from promoting growth of new blood vessels beneath the retina, by intravitreal injection.

Ranibizumab is indicated for the treatment of Exudative (wet) Age-related Macular Degeneration (AMD), Macular Edema following Retinal Vein Occlusion (RVO), Diabetic Macular Edema (DME), Diabetic Retinopathy (DR), and Myopic Choroidal Neovascularization (mCNV).

Ranibizumab is available as Lucentis:

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- Single-use prefilled syringe designed to provide 0.05 mL for intravitreal injection
  - 10 mg/mL solution (Lucentis 0.5 mg)
  - 0.3 mg/0.05mL solution (Lucentis 0.3mg)

Lucentis (ranibizumab) is contraindicated in patients with ocular or periocular infections.

Lucentis (ranibizumab) should not be used concurrently with other VEGF inhibitors for intraocular use in the absence of documentation indicating that individual products are to be used in different eyes.

See product prescribing information for complete list of warnings and precautions.

#### **Provider Claim Codes**

For medically billed requests, please visit www.humana.com/PAL. Select applicable Preauthorization and Notification List(s) for medical and procedural coding information.

## **Medical Terms**

Lucentis; ranibizumab; Neovascular (wet) Age Related Macular Degeneration; AMD; Diabetic Macular Edema; DME; Diabetic Retinopathy; DR; Macular Edema, Retinal Vein Occlusion; RVO; Myopic Choroidal Neovascularization; mCNV; Intravitreal; Pharmacy

#### References

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- 4. Flaxel CJ, Adelman RA, et al. Age-Related Macular Degeneration Preferred Practice Pattern. Ophthalmology. 2020; 127;1:p1-65.
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