

**Beovu (brolucizumab-dbb)**  
**Effective 07/01/2020**

<b>Plan</b>	<input type="checkbox"/> MassHealth UPPL <input checked="" type="checkbox"/> Commercial/Exchange	<b>Program Type</b>	<input checked="" type="checkbox"/> Prior Authorization <input type="checkbox"/> Quantity Limit <input type="checkbox"/> Step Therapy
<b>Benefit</b>	<input checked="" type="checkbox"/> Pharmacy Benefit <input checked="" type="checkbox"/> Medical Benefit		
<b>Specialty Limitations</b>	This medication has been designated specialty and must be filled at a contracted specialty pharmacy.		
<b>Contact Information</b>	<b>Medical and Specialty Medications</b>		
	All Plans	Phone: 877-519-1908	Fax: 855-540-3693
<b>Exceptions</b>	<b>Non-Specialty Medications</b>		
	All Plans	Phone: 800-711-4555	Fax: 844-403-1029
<b>Exceptions</b>	N/A		

### Overview

Brolucizumab is a recombinant humanized monoclonal antibody vascular endothelial growth factor (VEGF) inhibitor that binds to the 3 major isoforms of VEGF-A, thereby suppressing endothelial cell proliferation, neovascularization, and vascular permeability to slow vision loss.

### Coverage Guidelines

Authorization may be granted for members who are currently receiving treatment with Beovu excluding when the product is obtained as samples or via manufacturer's patient assistance programs.

#### OR

Authorization may be granted for members when all the following criteria are met, and documentation is provided:

1. Member has a diagnosis of neovascular (wet) age-related macular degeneration
2. Member has had an inadequate response to previous trials with and/or contraindication to Eylea, Lucentis, AND Avastin

### Continuation of Therapy

Authorization of 24 months may be granted for members who have 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).

### Limitations

Initial authorization will be approved for a duration of 24 months

### References

1. Beovu [package insert]. East Hanover, NJ: Novartis Pharmaceuticals Corporation; October 2019.

### Review History

05/20/2020 – reviewed and approved by P&T. Effective 7/1/20.

