

This document applies to the following:

| Formulary                                      | Applies                             |
|--|-------------------------------------|
| Standard Control (SF)                          | <input checked="" type="checkbox"/> |
| Standard Control – Choice (SCCF)               | <input checked="" type="checkbox"/> |
| Preferred Drug Plan Design (PDPD)              | <input type="checkbox"/>            |
| Advanced Control Specialty (ACSF)              | <input checked="" type="checkbox"/> |
| Advanced Control Specialty – Choice (ACSCF)    | <input checked="" type="checkbox"/> |
| Managed Medicaid Template (MMT)                | <input type="checkbox"/>            |
| Marketplace (MF)                               | <input type="checkbox"/>            |
| Aetna Small Group Affordable Care Act (SG ACA) | <input type="checkbox"/>            |
| Aetna Health Exchange (AHE)                    | <input type="checkbox"/>            |
| Aetna Individual Lives (IVL)                   | <input type="checkbox"/>            |
| Value (VF)                                     | <input checked="" type="checkbox"/> |

| Formulary   | Applies                  |
|---|--------------------------|
| New to Market (NTM)                                       | <input type="checkbox"/> |
| Standard Formulary Chart (SFC)                            | <input type="checkbox"/> |
| Basic Control Chart Preferred Drug Plan Design (BCC PDPD) | <input type="checkbox"/> |
| Advanced Control Specialty Formulary Chart (ACSFC)        | <input type="checkbox"/> |
| Value Formulary Chart (VFC)                               | <input type="checkbox"/> |
| Medical Benefit   | <input type="checkbox"/> |
| Medical Benefit: Advanced Biosimilars First               | <input type="checkbox"/> |
| Medical Benefit: Managed Medicaid (MMMB)                  | <input type="checkbox"/> |
| Medicare Part B   | <input type="checkbox"/> |
| Medicare Part B: Advanced Biosimilars First               | <input type="checkbox"/> |

# Exceptions Criteria

## Retinal Disorders

This document informs prescribers of preferred products and provides an exception process for targeted products through prior authorization.

These criteria were developed to align with the following: Standard Control Formulary (SF), Standard Control Choice Formulary (SCCF), Advanced Control Specialty Formulary (ACSF), Advanced Control Specialty – Choice Formulary (ACSCF) and Value Formulary (VF).

## Plan Design Summary

This program applies to the retinal disorders products specified in this document. Coverage for targeted products is provided based on clinical circumstances that would exclude the use of the preferred product and may be based on previous use of a product. The coverage review process will ascertain situations where a clinical exception can be made. This program applies to all members requesting treatment with Lucentis and Susvimo. This program applies to members who are new to treatment with Beovu, Eylea, Eylea HD, and Vabysmo for the first time.

Each referral is reviewed based on all utilization management (UM) programs implemented for the client.

## Table. Retinal Disorder Products

Medications considered formulary or preferred on your plan may still require a clinical prior authorization review.

|                     |
|---------------------|
| Reference number(s) |
| 6151-D              |

|           | Products  |
|-----------|---|
| Preferred | <ul style="list-style-type: none"> <li>• Byooviz (ranibizumab-nuna)</li> <li>• Cimerli (ranibizumab-eqrn)</li> </ul>  |
| Target    | <ul style="list-style-type: none"> <li>• Beovu (brolucizumab-dbll)</li> <li>• Eylea (aflibercept)</li> <li>• Eylea HD (aflibercept)</li> <li>• Lucentis (ranibizumab)</li> <li>• Susvimo (ranibizumab)</li> <li>• Vabysmo (faricimab-svoa)</li> </ul> |

## Exception Criteria

### Eylea

Coverage for the targeted product is provided when any of the following criteria are met:

- Member is currently receiving treatment with the targeted product (Eylea), excluding when the requested targeted product is obtained as samples or via manufacturer's patient assistance programs.
- Member has a diagnosis of retinopathy of prematurity.
- Member has a documented inadequate response or intolerable adverse event with either of the preferred products, Byooviz or Cimerli.

### Lucentis

Coverage for the targeted product is provided when the member has had a documented intolerable adverse event to both of the preferred products, Byooviz and Cimerli, and the adverse event was not an expected adverse event attributed to the active ingredient as described in the prescribing information (i.e., known adverse reaction for both the reference product and biosimilar products).

### Susvimo

Coverage for the targeted product is provided when the member has had a documented intolerable adverse event to both of the preferred products, Byooviz and Cimerli, and the adverse event was not an expected adverse event attributed to the active ingredient as described in the prescribing information.

### Beovu, Eylea HD, Vabysmo

Coverage for the targeted product is provided when any of the following criteria are met:

| Reference number(s) |
|---------------------|
| 6151-D              |

- Member is currently receiving treatment with the targeted product, excluding when the requested targeted product is obtained as samples or via manufacturer's patient assistance programs.
- Member has a documented inadequate response or intolerable adverse event with either of the preferred products, Byooviz or Cimerli.

## References

1. Beovu [package insert]. East Hanover, NJ: Novartis Pharmaceuticals Corporation; July 2024.
2. Byooviz [package insert]. Cambridge, MA: Biogen, Inc.; June 2023.
3. Cimerli [package insert]. Redwood City, CA: Coherus BioSciences, Inc.; June 2024.
4. Eylea [package insert]. Tarrytown, NY: Regeneron Pharmaceuticals, Inc.; March 2023.
5. Eylea HD [package insert]. Tarrytown, NY: Regeneron Pharmaceuticals; December 2023.
6. Lucentis [package insert]. San Francisco, CA: Genentech, Inc.; February 2024.
7. Susvimo [package insert]. San Francisco, CA: Genentech, Inc.; April 2022.
8. Vabysmo [package insert]. San Francisco, CA: Genentech, Inc.; July 2024.