

**Aflibercept (Eylea®)**

**Place of Service**

**Office Administration  
Outpatient Facility Infusion  
Administration  
Infusion Center Administration**

**HCPC:** J0178 per 1mg

**NDC:** 61755-005-02

**Condition(s) listed in policy (see criteria for details)**

- Diabetic Macular Edema (DME), including diabetic retinopathy in patients with DME
- Neovascular (WET) age-related macular degeneration (AMD)
- Macular edema (ME) following central or branch retinal vein occlusion

**AHFS therapeutic class:** Selective vascular endothelial growth factor antagonist

**Mechanism of action:** Aflibercept is a recombinant fusion protein that acts as a soluble decoy receptor that binds VEGF-A and PGF, thereby inhibiting the binding and activation of these cognate VEGF receptors

**(1) Special Instructions and Pertinent Information**

**Covered under the Medical Benefit,** please submit clinical information for prior authorization review via fax.

**(2) Prior Authorization/Medical Review is required for the following condition(s)**

**All requests for Eylea® (aflibercept) for conditions NOT listed in section 3 must be sent for clinical review and receive authorization prior to drug administration or claim payment.**

**(3) The following condition(s) DO NOT require Prior Authorization/Preservice**

**All requests for Eylea® (aflibercept) for conditions NOT listed in section 3 must be sent for clinical review and receive authorization prior to drug administration or claim payment.**

**Diabetic macular edema (DME), including diabetic retinopathy in patients with DME**

**Covered Doses**

2mg (0.05ml) intravitreal injection every month

**Coverage Period**

Yearly

**ICD-9:**

362.0X

**ICD-10:**

**E08.311, 321X, 331X, 341X, 351X,**

**E09.311, 321X, 331X, 341X, 351X,**

**E10.311, 321X, 331X, 341X, 351X,**

**E11.311, 321X, 331X, 341X, 351X**

**E13.311, 321X, 331X, 341X, 351X**

**Macular edema (ME) following central or branch retinal vein occlusion (RVO)**

**Covered Doses**

2mg (0.05ml) intravitreal injection every month

**Coverage Period**

Yearly

**ICD-9:**

362.35, 362.36

**ICD-10:**

H34.8110-8112, H34.8120-8122 H34.8130- 8132, H34.8190-8192,  
H34.8310-8312, H34.8320-8322, H34.8330-8332, H34.8390-8392

**Neovascular (WET) age-related macular degeneration (AMD)****Covered Doses**

2mg (0.05ml) intravitreal injection every month

**Coverage Period**

Yearly

**ICD-9:**

362.52 Exudative senile macular degeneration

**ICD-10:**

H35.3210-3213  
H35.3220-3223  
H35.3230-3233  
H35.3290-3293

**(4) This Medication is NOT COVERED for the following condition(s)**

The following conditions and other indications not listed in this policy do not meet the coverage criteria established by the Blue Shield of CA P&T Committee and are NOT-COVERED. Please refer to the user guide for more information.

**(5) Additional Information**

How supplied:

Each carton contains:

- One single use 3mL glass vial designed to deliver 0.05ml of 40mg/ml of Eylea
- One 19-gauge x 1 ½ -inch, 5 micron, filter needle for withdrawal of the vial contents
- One 30-gauge x ½ -inch needle for intravitreal injection
- One 1-mL syringe for administration

**(6) References**

- Eylea® Package Insert, Regeneron Pharmaceuticals, Inc. 2017
- AHFS®. Available by subscription at <http://www.lexi.com>
- DrugDex®. Available by subscription at <http://www.micromedexsolutions.com/home/dispatch>

**(7) Policy Update**

Dates of last review: 3Q2017

Date of next review: 3Q2018

Changes from previous policy version:

- No change to policy following routine review.

