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Eylea FDA Approval History

FDA Approved: Yes (First approved November 18, 2011)
Brand name: Eylea
Generic name: aflibercept
Dosage form: Injection
Company: Regeneron Pharmaceuticals, Inc.
Treatment for: Macular Degeneration, Macular Edema, Diabetic Retinopathy

Eylea (aflibercept) is a VEGF inhibitor indicated for the treatment of patients with neovascular (wet) age-related macular degeneration, macular edema following retinal vein occlusion, diabetic macular edema, and diabetic retinopathy.

Development Timeline for Eylea

Date	Article	
Aug 13, 2019	Approval	**FDA Approves Eylea (aflibercept) Injection Prefilled Syringe**
May 13, 2019	Approval	**FDA Approves Eylea (aflibercept) Injection for Diabetic Retinopathy**
Aug 17, 2018	Approval	**FDA Approves New Eylea (aflibercept) Injection Dosing Schedule in Wet Age-Related Macular Degeneration**
Mar 25, 2015	Approval	**FDA Approves Eylea (aflibercept) for Diabetic Retinopathy in Patients with Diabetic Macular Edema**
Jul 29, 2014	Approval	**Eylea (aflibercept) Injection Receives FDA Approval for the Treatment of Diabetic Macular Edema**
Nov 18, 2011	Approval	**FDA Approves Eylea for Wet Age-Related Macular Degeneration**
Aug 17, 2011		Regeneron Announces Review of Biologics License Application for Eylea (aflibercept injection) Extended by Three Months by FDA
Jun 17, 2011		Regeneron Announces Eylea (aflibercept ophthalmic solution) Receives Unanimous Recommendation for Approval for Treatment of Wet AMD from FDA Advisory Committee

Further information

Always consult your healthcare provider to ensure the information displayed on this page applies to your personal circumstances.