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Infographic: Efficacy and Safety of High-Dose Intravitreal Aflibercept 8 mg in Patients with Treatment-Naïve Neovascular Age-Related Macular Degeneration: Week 96 Results from the Phase 3 PULSAR Trial

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Fig. 1 In the phase 3 PULSAR trial, aflibercept 8 mg demonstrated similar visual and anatomic improvements with extended dosing intervals and a similar safety profile to aflibercept 2 mg in patients with neovascular age-related macular degeneration.

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All authors equally contributed to the concept and creation of the visual abstract.

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COMPETING INTERESTS

J-FK is a consultant for 4DMT, AbbVie, Adverum, Apellis, Bayer, Boehringer Ingelheim, Carl Zeiss Meditec, Eyepoint Pharma, Ocular Therapeutix, Roche, SeaBeLife, and Thea; and is a member of the Data Safety Monitoring Board at Complement Therapeutics and Novo Nordisk. SL is an employee of and holds stock/stock options in Bayer Consumer Care AG. PL is a consultant for 4DMT, AbbVie, Adverum, Annexon, Apellis, Bausch + Lomb, Bayer, Biogen, Boehringer Ingelheim, Celltrion, EyePoint Pharmaceuticals, Genentech, I-Care, MSD, Novartis, Ocular Therapeutix, Outlook Therapeutics, and Roche.

ADDITIONAL INFORMATION

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