



Changes in Macular Perfusion after ILUVIEN implant for Diabetic Macular Edema: an OCTA study

Prof. Francesco Bandello, MD, FEBOphth

Reparto di Oculistica - IRCCS Ospedale San Raffaele Università Vita-Salute San Raffaele (Milano)

Financial disclosure

ALLERGAN, BAYER, BOEHRINGER-INGELHEIM, FIDIA SOOFT, HOFMANN

LA ROCHE, NOVARTIS, NTC PHARMA, SIFI, THROMBOGENICS, ZEISS

Summary slide

- This OCTA study evaluated changes in macular perfusion in patients affected by diabetic macular edema (DME) and treated with ILUVIEN® (fluocinolone acetonide intravitreal implant) 0.19 mg
- The qualitative and quantitative assessment demonstrated that this treatment is not associated with worsening in retinal perfusion
- Areas of reperfusion may be related to reversible retinal vessel closure secondary to leukostasis

Background

- Recent evidences suggest that macular perfusion doesn't modify after 12 months of intravitreal aflibercept therapy
- Because nonperfusion is expected to progress in diabetic retinopathy, this finding may represent a beneficial association between anti–VEGF therapy and macular vascular density

Association of Intravitreal Aflibercept With Optical Coherence Tomography Angiography Vessel Density in Patients With Proliferative Diabetic Retinopathy A Secondary Analysis of a Randomized Clinical Trial

Ahmed Roshdy Alagorie, MD^{1,2,3}; Muneeswar Gupta Nittala, MPhil^{1,2}; Swetha Velaga, MPhil^{1,2}; Brenda Zhou, MD⁴; Alexander M. Rusakevich, MD⁴; Charles C. Wykoff, MD, PhD⁴; SriniVas R. Sadda, MD^{1,2}

> Author Affiliations

JAMA Ophthalmol. Published online June 25, 2020. doi:10.1001/jamaophthalmol.2020.2130

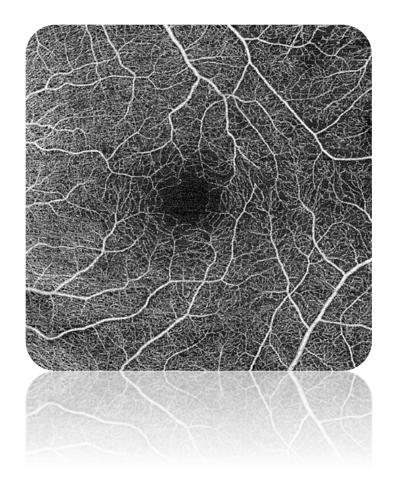
Purpose

To investigate changes in macular
 perfusion in patients affected by
 diabetic macular edema (DME) and
 treated with ILUVIEN® (fluocinolone
 acetonide intravitreal implant) 0.19 mg



Methods

- Ten patients (10 eyes) older than 18 years of age and with type 2 non-proliferative DR and DME at baseline were included
- Nice patients (9 eyes) without disease were included for comparison
- All patients were treated with the ILUVIEN® implant
- In order to investigate macular perfusion changes,
 patients had <u>two OCTA scans</u>: (i) baseline, and (ii) 4-month FU OCTA images



Results – Qualitative grading

Loss of perfusion Reperfusion **Baseline** 4-month FU

The qualitative grading demonstrated that treatment was associated with both areas of loss of perfusion and regions of reperfusion

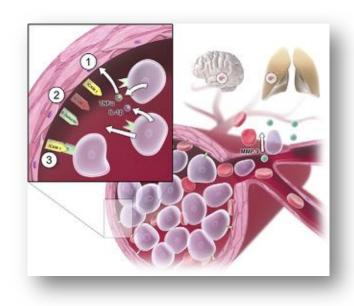
Results – Quantitative analysis

		Baseline	4-month FU	P value
DR eyes treated with Iluvien	Parafoveal perfusion density	64.1±1.8%	66.1±2.9%	0.013
	Perifoveal perfusion density	64.4±2.1%	65.2±2.6%	0.024
DR eyes without treatment	Parafoveal perfusion density	63.7±2.3%	63.1±4.4%	0.875
	Perifoveal perfusion density	64.0±4.1%	64.2±3.7%	1.0

The quantitative analysis proved a slight improvement in macular perfusion at the 4-month FU visit

Conclusions

- This study confirms recent OCTA evidences that intravitreal treatments are not associated with worsening in retinal perfusion
- Reversible retinal vessel closure (areas of reperfusion) may be related to leukostasis
- Inflammation suppression with Iluvien is associated with areas of reperfusion and an overall no worsening of macular perfusion







Thanks for your attention