

#### INTRAVITREAL AFLIBERCEPT IN REFRACTORY AGE-RELATED MACULAR DEGENERATION

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## Introduction

Age-related macular degeneration, or AMD, is a leading cause of blindness in developed countries.

Intravitreal Aflibercept is approved in the treatment of **neovascular age-related macular degeneration** (AMD). The duration of its effect is higher than Ranibizumab and Bevacizumab but the effectiveness in patients who do not respond to these treatments is unknown.

**Design:** Prospective observational study in patients diagnosed with AMD treated with Aflibercept, who didn't respond to Ranibizumab and/or Bevacizumab previously.

Methods

Variables: BCVA (ETDRS optotypes) and CRT obtained on optical

# Purpose

Evaluate the efficacy of intravitreal Aflibercept on visual acuity (BCVA) and central retinal thickness (CRT) in refractory AMD.

Results

A total of **27 eyes from 24 patients** were included, 20 of them were females. The median age was 76.98  $\pm$  9,9 years old. The 12,5% of the patients received bilateral treatment.

The 22.2% of the eyes were previously treated with

coherence tomography (OTC 3-D, Topcon Corporation).

**Statistics:** Wilcoxon test for paired data and Student t test for paired data (Stata/IC 12, StataCorp LP, Texax, USA), considering significant if p value <0.05.

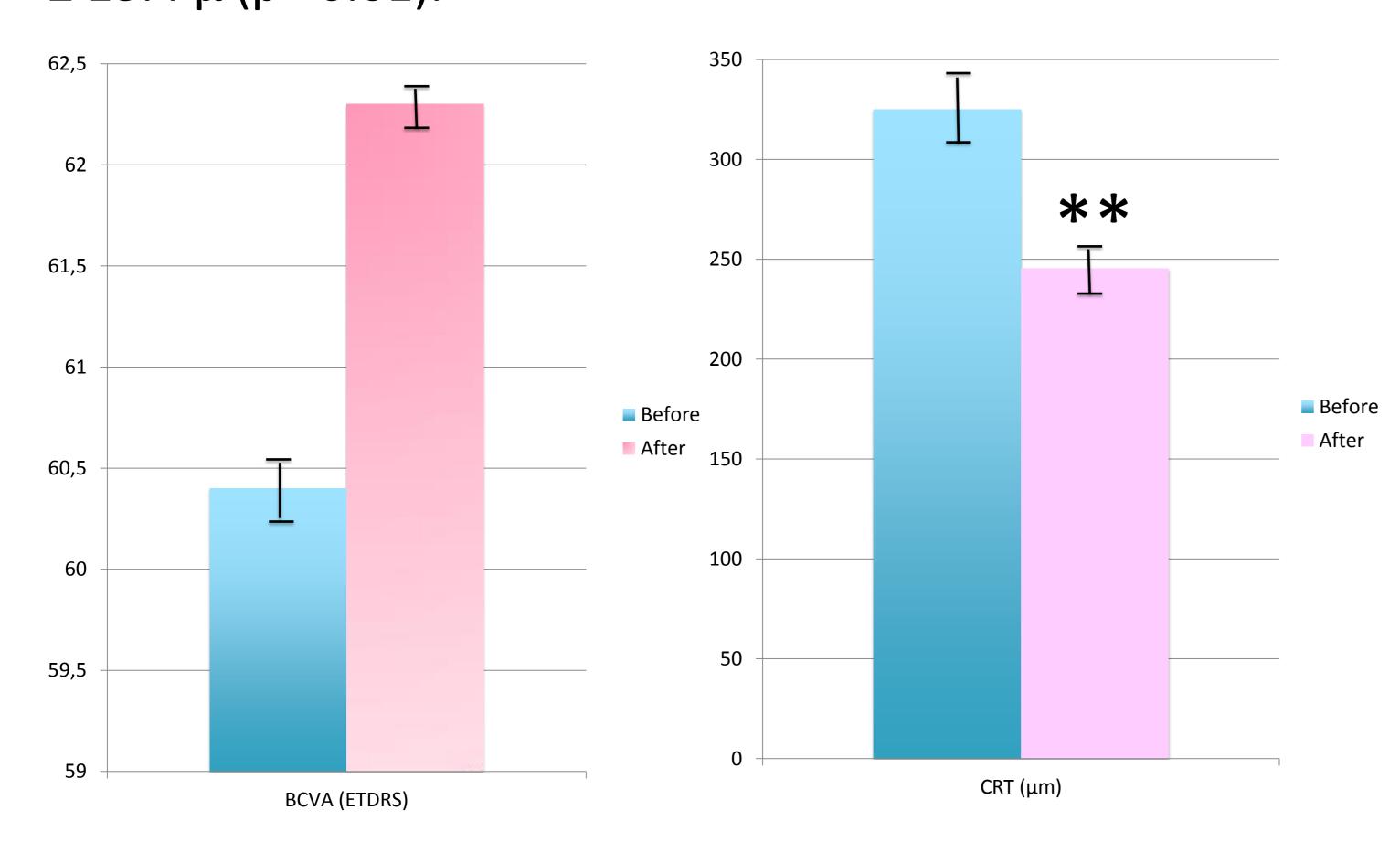
# Discussion

Aflibercept is a new VEGF inhibitor that differs from Ranibizumab and Bevacizumab by a **higher affinity to VEGF**, a **longer half-life** and its action blocking the placental growth factor (**PIGF**).

In phase III trials Aflibercept was shown to be **non-inferior** to Ranibizumab treatments and elicited the same outcome as a monthly Ranibizumab injection regimen when administered **bimonthly** following 3 monthly injections over a 52-week period.

photodynamic therapy (PDT), 70.4% with Ranibizumab (average 7.5 injections/eye, range 2-24) and 77.8% with Bevacizumab (average 8 injections/eye, range 1-33). The 52% of the eyes received both Ranibizumab and Bevacizumab.

The average of Aflibercept injections per eye was 3.3 (1-10). In the 63% of the eyes there was an increase of BCVA, the 7.5% of the eyes maintained previous BVCA and the 29.6% lost vision. ETDRS before and after treatment was 60.4  $\pm$  2.5 vs 62.3  $\pm$  3.2 letters (p = 0.0504). The 89% of eyes experiment a decrease in the CRT measured by OCT, 324.9  $\pm$  22.2 vs 245.5  $\pm$  13.4  $\mu$  (p <0.01).



This study indicates that an **improved anatomical** outcome can be achieved in patients with persistent macular exudation despite prior treatment with either Ranibizumab or Bevacizumab . Even though Aflibercept was able to resolve persistent macular exudate in our patient sample it remains controversial if a dry macula after anti-VEGF treatment allows for regained visual acuity.

## Conclusions

- **Aflibercept is effective in refractive AMD.**
- We obtain better anatomical response than visual.
- Due to the duration of the study we expect a greater effect with repeated administrations

**Figure 1.** Mean differences in best corrected visual acuity (BCVA) before and after the treatment with intravitreal aflibercept. **Figure 2.** Mean differences in central retinal thickness (CRT) before and after the treatment with intravitreal aflibercept.

#### References

- 1. Stewart MW. Aflibercept (VEGF Trap-Eye) for the treatment of exudative agerelated macular degeneration. Expert Rev Clin Pharmacol. 2013;6(2):103-13.
- 2. Frampton JE. Aflibercept for intravitreal injection: in neovascular age-related macular degeneration. Drugs Aging. 2012;29(10):839-46.