

Intravitreal aflibercept in wet AMD (VIEW1 & VIEW2) - 2012



Objective

To compare the efficacy and safety of monthly and every-2-month dosing of intravitreal aflibercept (after 3 months loading) with monthly dosing of ranibizumab in treatment of wet AMD

Methods

Design: Double-masked, multicenter, parallel-group, active-controlled, randomized trials

Sample Size: 2419

Treatment Groups:

- Aflibercept 0.5 mg monthly
- Aflibercept 2.0 mg monthly
- Aflibercept 2.0 mg every-2-months after 3-monthly loading doses
- Ranibizumab 0.5 mg monthly

Outcome Measures:

- Maintain vision at 52 weeks (<15 letters lost ETDRS)
- Reduction in active CNV area
- Additional letters gained

Results

Point 1: All 3 aflibercept treatment regimens were non-inferior to monthly ranibizumab

- All aflibercept groups preserved vision at 52 weeks (VIEW1: 95.1%, 95.9%, and 95.1%) (VIEW2: 95.6%, 96.3%, 95.6%) comparative to monthly ranibizumab (94.4%)

Point 2: All 3 aflibercept regimens resulted in similar secondary outcomes compared to monthly ranibizumab

- Similar reduction of mean active CNV area in aflibercept (-6.0, -4.2, -5.2) vs ranibizumab: -4.2
- Similar gains in EDTRS letters at 52 weeks in aflibercept (10.9, 6.9, 7.9) vs ranibizumab: +8.1
- Similar reduction in central retinal thickness in aflibercept (-116.5, -115.6, -128.5) vs ranibizumab: -116.8

Point 3: Ocular and systemic adverse events were similar across all treatment groups

TLDR: Aflibercept dosed monthly, or every-2-months, resulted in similar visual and anatomic outcomes as monthly ranibizumab, meaning that an extended (q8 weeks) injection regimen worked for wet AMD