

Real-World Management and Outcomes in Retinal Vein Occlusion

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BACKGROUND

- Retinal vein occlusion (RVO) is a common vascular retinal disorder resulting from a blockage of venous outflow from the retina and is a leading cause of vision loss worldwide
- Cystoid macular edema (CME) is the accumulation of fluid within the intracellular spaces of the retina and is most common complication of RVO
- Types of RVO: central, hemispheric, or branched retinal vein occlusion (CRVO, HRVO, and BRVO)
- Neovascular age-related macular degeneration (NV-AMD) and diabetic macular edema (DME) studies have shown that real world outcomes do not always match those demonstrated in randomized clinical trials (RCTs)

PURPOSE

- To evaluate the real world management and outcomes of eyes with retinal vein occlusion (RVO) and cystoid macular edema from five academic retina practices

METHODS

- Multi-center retrospective study of acute RVO eyes from 2011-2015 with at least 1 year follow-up
- Visual acuity (VA), central subfield thickness (CST) on OCT, and management were recorded at 3, 6, 12, and 24 month follow-up
- Exclusion Criteria: Other retinal or choroidal conditions other than cataract surgery
- Treatments:
 - Anti-VEGF: Bevacizumab, Ranibizumab, Aflibercept
 - Steroids: Triamcinolone, Dexamethasone
 - Laser Photocoagulation

Table 1

| | All (n = 262) | CRVO/HRVO (n = 163) | BRVO (n = 99) |
|---------------------------|------------------|------------------------|------------------|
| Mean Age (SD) | 69.9 (12.5) | 69.7 (12.5) | 70.2 (12.4) |
| Male (%) | 106 (40.5) | 65 (39.9) | 41 (41.4) |
| Right eye (%) | 81 (49.7) | 70 (51.1) | 44 (44.4) |
| Diabetes (%) | 96 (36.6) | 61 (37.4) | 35 (35.4) |
| Dyslipidemia (%) | 106 (40.5) | 64 (39.3) | 42 (42.4) |
| Hypertension (%) | 164 (62.6) | 98 (60.1) | 66 (66.7) |
| Hypercoagulable State (%) | 29 (11.1) | 20 (12.3) | 9 (9.1) |
| Glaucoma (%) | 63 (24.0) | 44 (27.0) | 19 (19.2) |
| Phakic (%) | 190 (72.5) | 122 (74.9) | 68 (68.7) |

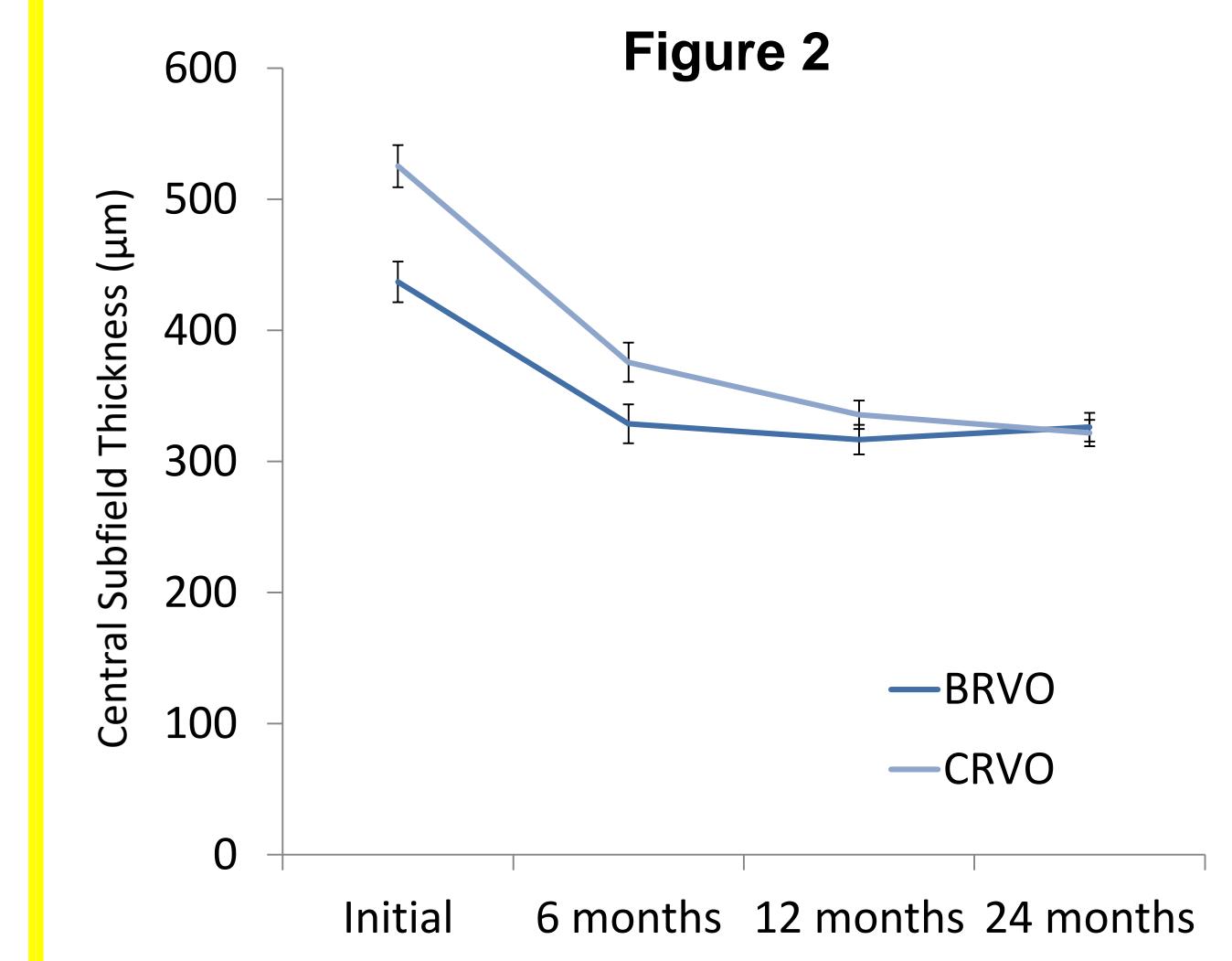
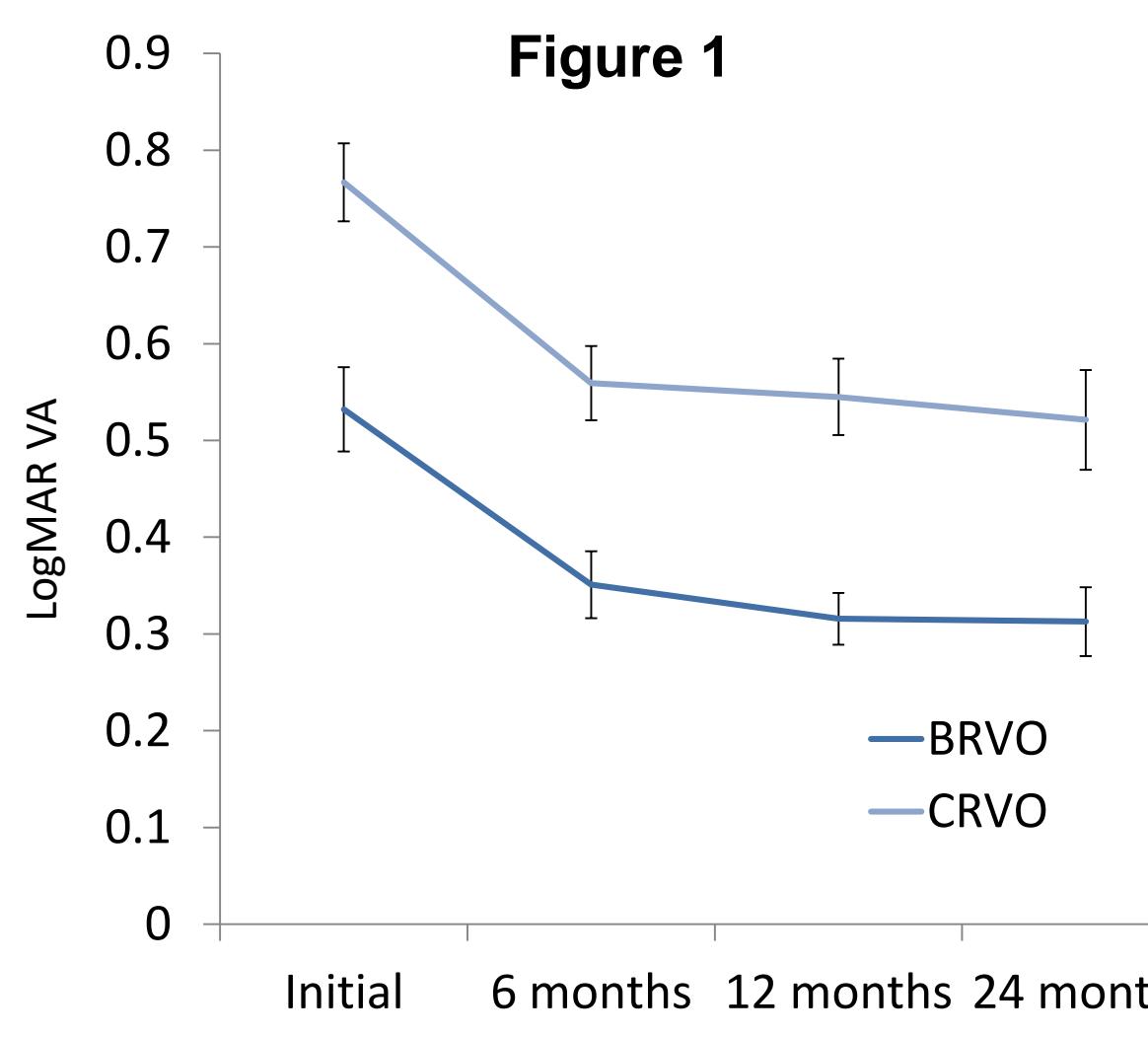
DATA

Table 1

| | IVB only (%) | IVR only (%) | IVA only (%) | Mixed anti-VEGF (%) | IVS only (%) | Mixed IVS and anti-VEGF (%) | Laser only (%) | Mixed laser and Injections (%) |
|-------------------------|--------------|--------------|--------------|---------------------|--------------|-----------------------------|----------------|--------------------------------|
| Initial Visit (n = 262) | 58.8 | 28.6 | 2.7 | 0.0 | 2.3 | 0.0 | 1.1 | 0.0 |
| 1 - 3 mos (n=258) | 53.1 | 30.2 | 4.3 | 3.1 | 1.2 | 0.5 | 0.8 | 0.4 |
| 4 - 6 mos (n = 256) | 37.5 | 29.7 | 8.2 | 2.3 | 1.6 | 0.8 | 4.7 | 4.2 |
| 7 - 12 mos (n = 262) | 36.3 | 27.5 | 11.1 | 2.7 | 3.1 | 1.1 | 6.5 | 3.8 |
| 13 - 24 mos (n = 231) | 32.5 | 22.5 | 16.9 | 5.2 | 6.1 | 0.9 | 5.2 | 3.0 |

Table 2

| Patients with >6 months without treatment | All (n = 94, 36%) | CRVO/HRVO (n = 57, 36%) | BRVO (n = 37, 37.4%) |
|---|----------------------|----------------------------|-------------------------|
| Mean No. of Months to Resolution (SD) | 7.0 (4.9) | 7.0 (4.9) | 7.2 (4.8) |
| Mean No. Anti-VEGF Injections (SD) | 4.4 (3.2) | 4.3 (3.2) | 4.1 (3.0) |
| Mean No. Steroid Injections (SD) | 0.1 (0.4) | 0.1 (0.4) | 0.08 (0.4) |
| Mean No. Lasers (SD) | 0.1 (0.3) | 0.1 (0.3) | 0.3 (0.6) |



RESULTS

- 262 patients met the study criteria
- The demographics and systemic and ocular comorbidities are listed in Table 1
- The management of the patients is summarized in Table 2
- 36% of patients were observed without treatment for >6 months (Table 3)
- There was a significant improvement ($p<0.001$) in VA (Figure 1) and CST (Figure 2) for patients after 1 year
 - BRVO patients:
 - mean logMAR VA improved from 0.52 ± 0.33 to 0.30 ± 0.22
 - CMT improved from 398 ± 111 to 316 ± 115
 - CRVO patients
 - Mean logMAR improved from 0.80 ± 0.51 to 0.53 ± 0.42
 - CMT improved from 503 ± 160 to 321 ± 139

CONCLUSIONS

- RVOs with CME are largely managed using intravitreal anti-VEGF agents
- Real world outcomes of RVOs with CME demonstrate improvement in VA and CMT with treatment, however, these improvements are less than those found in RCTs.

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DISCLOSURES

- Disclosures: AEK: Consultant for Allergan, Alimera Sciences, MYW: None; AVR: Consultant for Allergan; RPS: Consultant for and Grants from Regeneron, Alcon, Genentech, Consultant for Shire, Biogen, Grants from Zeiss; KMW: None; GCY: Consultant for Allergan, Alimera, Zeiss, Southern California Desert Retina, Grants from Alcon and E Matilda Ziegler Foundation, Iridex; K-YL: None; EN: Consultant for Allergan; KCC: None; JPC: Consultant for Allergan, KG: None
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