

Real-World Treatment Outcomes With Anti-VEGF Therapy in Patients With Retinal Vein Occlusion in the UK

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Disclosures

Financial Disclosures

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- ▶ PA: Employee: Roche Products Ltd.
- ▶ IS: Employee: F. Hoffmann-La Roche Ltd.
- ▶ LPP, GCC: Employee: Genentech, Inc.

Study and Product Disclosures

- ▶ This study was a noninterventional, retrospective, secondary data use study, leveraging data from Medisoft electronic health records. The study was considered exempt from institutional review board review as the research involved only the collection of existing data, which had been de-identified and are unable to be traced
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Treatment Patterns May Contribute to Outcomes in Patients With RVO



Suboptimal vision outcomes in clinical practice with standard-of-care anti-VEGF injections in RVO may, in part, **reflect fewer injections** received in clinical practice¹⁻³



There are also **limited data** on **long-term outcomes** up to 5 years of treatment of RVO

What are the anti-VEGF treatment patterns and long-term visual acuity outcomes in patients with macular edema secondary to RVO?

Real-world outcomes in patients being treated in routine clinical practice in the UK



Study Design: Retrospective Observational Study

- **Medisoft electronic health record** data from **2013 to 2023**
- Contributed by **16 UK National Health Service trusts**



Study Eyes: Inclusion Criteria

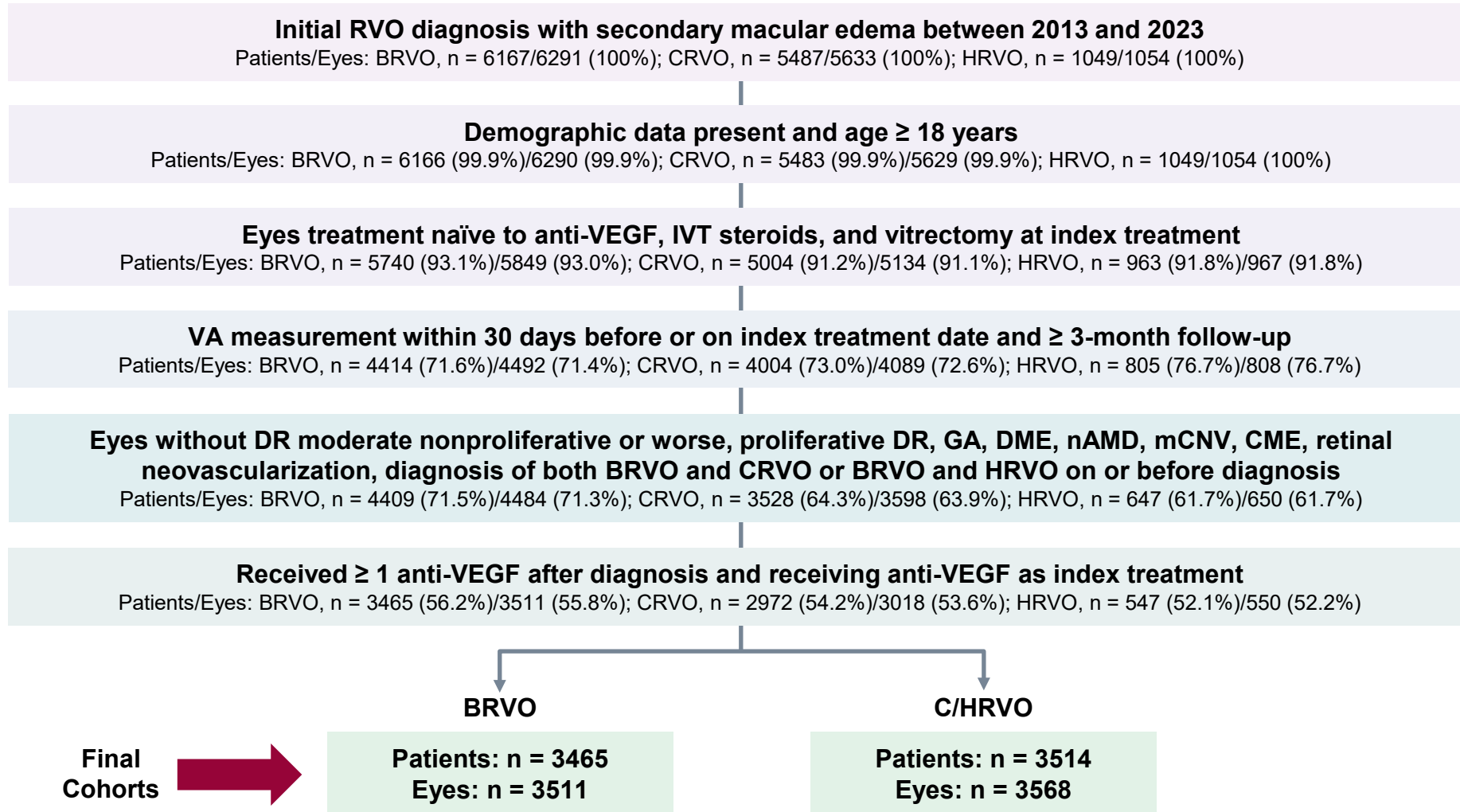
- ✓ **Macular edema secondary to BRVO, CRVO, or HRVO**
- ✓ **Treatment naïve to anti-VEGFs, intravitreal steroids, vitrectomy, laser therapy**
- ✓ **≥ 1 intravitreal anti-VEGF injection** on or after diagnosis
- ✓ **VA measured at index injection**
- ✓ **≥ 3 months follow-up after index injection**



Data Analyses

- **Observation period: Up to 60 months** after index injection
- **Treatment patterns** and **VA outcomes** were **evaluated descriptively**
- **CRVO and HRVO eyes** were **analyzed together**

BRVO and C/HRVO Final Cohorts

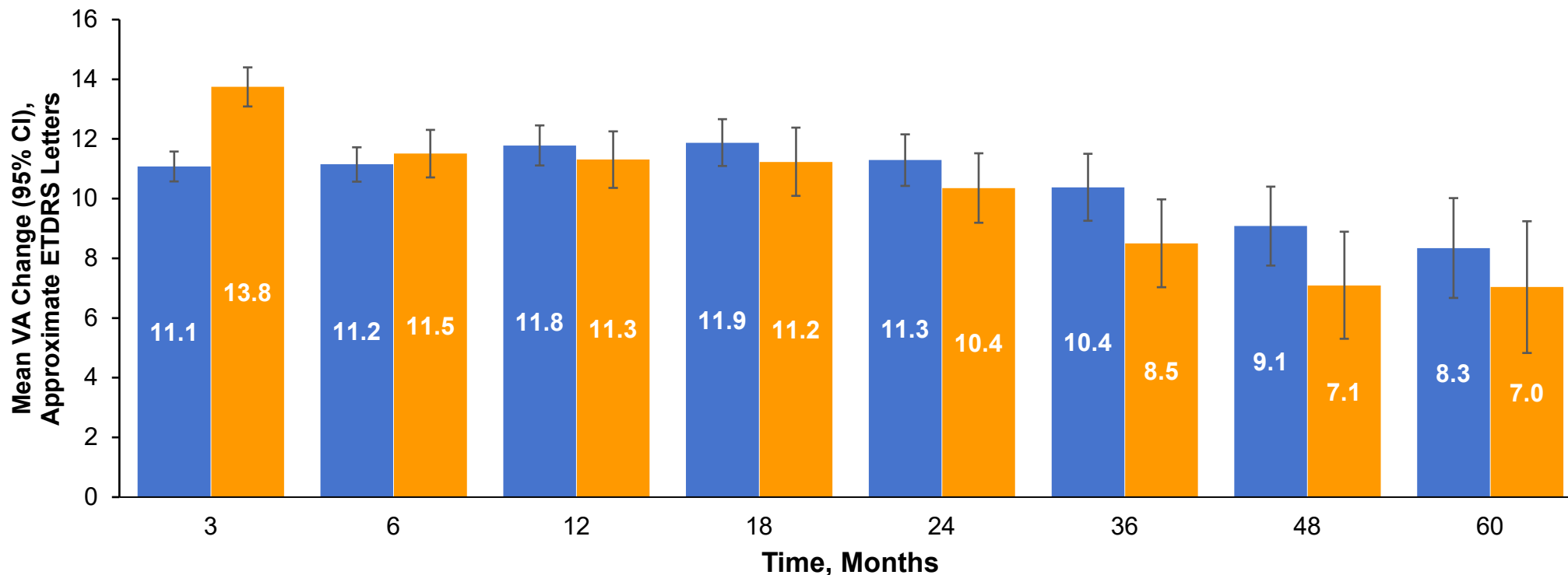


Baseline Demographics and Clinical Characteristics

| Baseline Characteristics (Patient-Level) | BRVO, n = 3465 | C/HRVO, n = 3514 |
|--|----------------|------------------|
| Age at index date, mean years (SD) | 71.5 (12.2) | 73.6 (12.4) |
| Sex, n (%) | | |
| Female | 1852 (53.5) | 1671 (47.6) |
| Race, n (%) | | |
| White | 2846 (82.1) | 2979 (84.8) |
| Asian | 89 (2.6) | 75 (2.1) |
| Black | 20 (0.6) | 27 (0.8) |
| Other/Mixed | 27 (0.8) | 26 (0.7) |
| Not stated | 483 (13.9) | 407 (11.6) |
| IMD decile, n (%) | | |
| 1–4 | 1080 (31.2) | 1184 (33.7) |
| 5–10 | 2367 (68.3) | 2306 (65.6) |
| Not stated | 18 (0.5) | 24 (0.7) |
| Eye Characteristics (Eye-Level) | BRVO, n = 3511 | C/HRVO, n = 3568 |
| Phakic lens status, n (%) | 2619 (74.6) | 2752 (77.1) |
| Duration since diagnosis at index treatment, days (SD) | 27.3 (110.8) | 24.9 (116.8) |
| VA at index treatment (analysis groups), n (%) | | |
| ≤ 35 ETDRS letter score | 564 (16.1) | 1494 (41.9) |
| ≥ 70 ETDRS letter score | 908 (25.9) | 412 (11.6) |
| ≥ 85 ETDRS letter score | 61 (1.7) | 22 (0.6) |

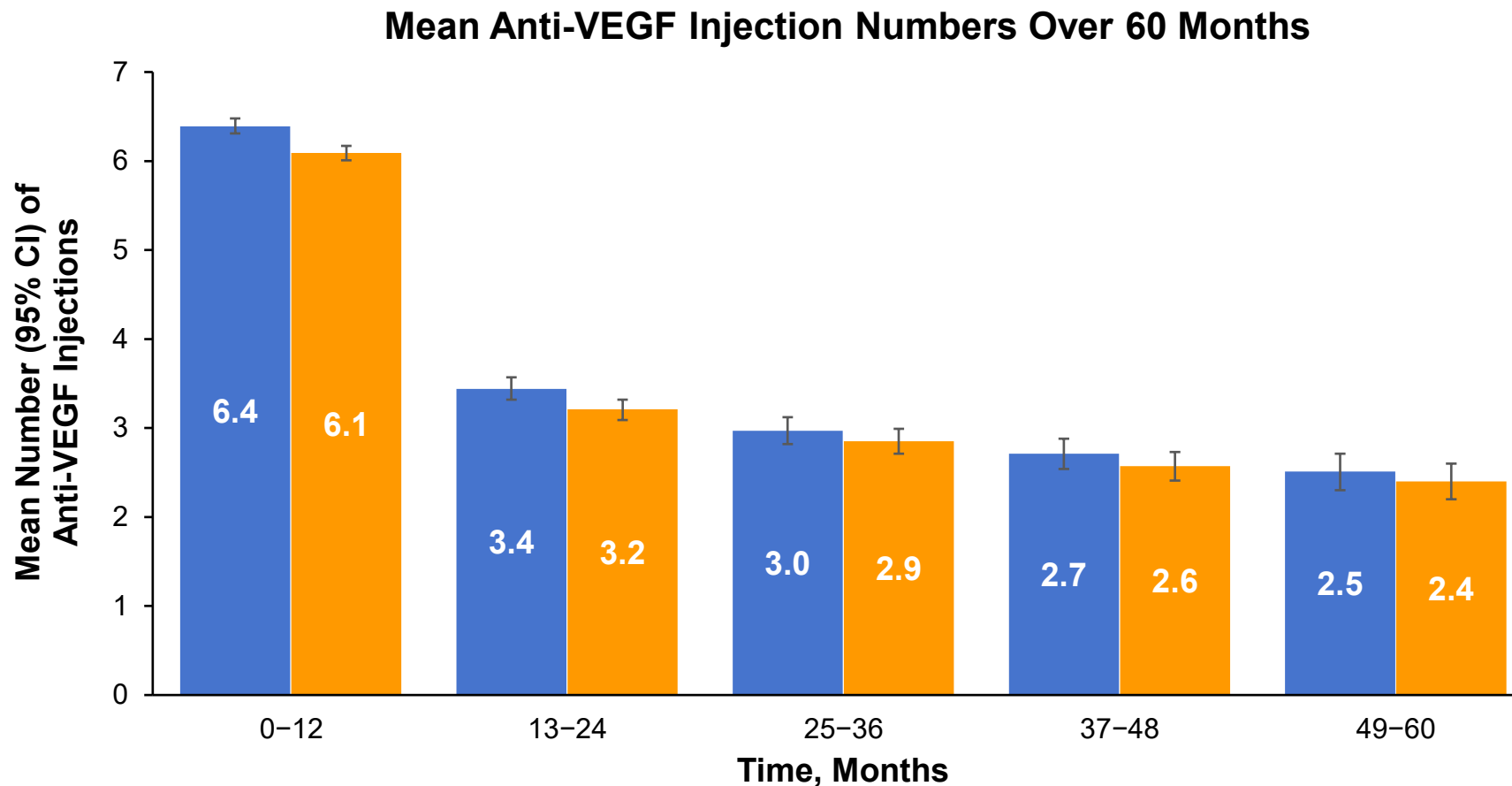
On Average, VA Gains Were Observed After Index Anti-VEGF Injection in BRVO and C/HRVO Eyes; VA Gains Were Lower With Longer Follow-Up

Mean VA Change From Index Anti-VEGF Injection Over 60 Months



| | | | | | | | | | |
|--------|-------------|------|------|------|------|------|------|------|------|
| BRVO | n | 3052 | 2642 | 2137 | 1723 | 1654 | 1058 | 769 | 497 |
| | Baseline VA | 55.4 | 55.8 | 56.1 | 56.4 | 56.4 | 56.9 | 58.0 | 58.0 |
| C/HRVO | n | 3127 | 2587 | 2091 | 1642 | 1651 | 1065 | 734 | 498 |
| | Baseline VA | 39.7 | 40.2 | 40.9 | 41.5 | 41.5 | 42.6 | 43.7 | 43.8 |

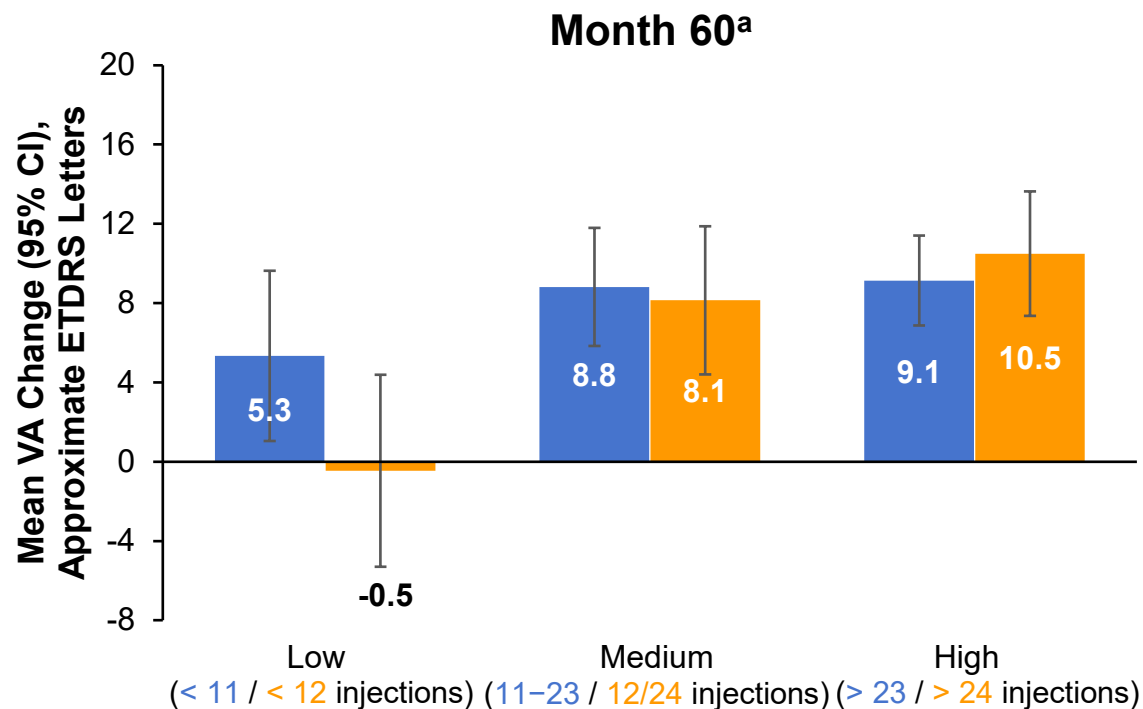
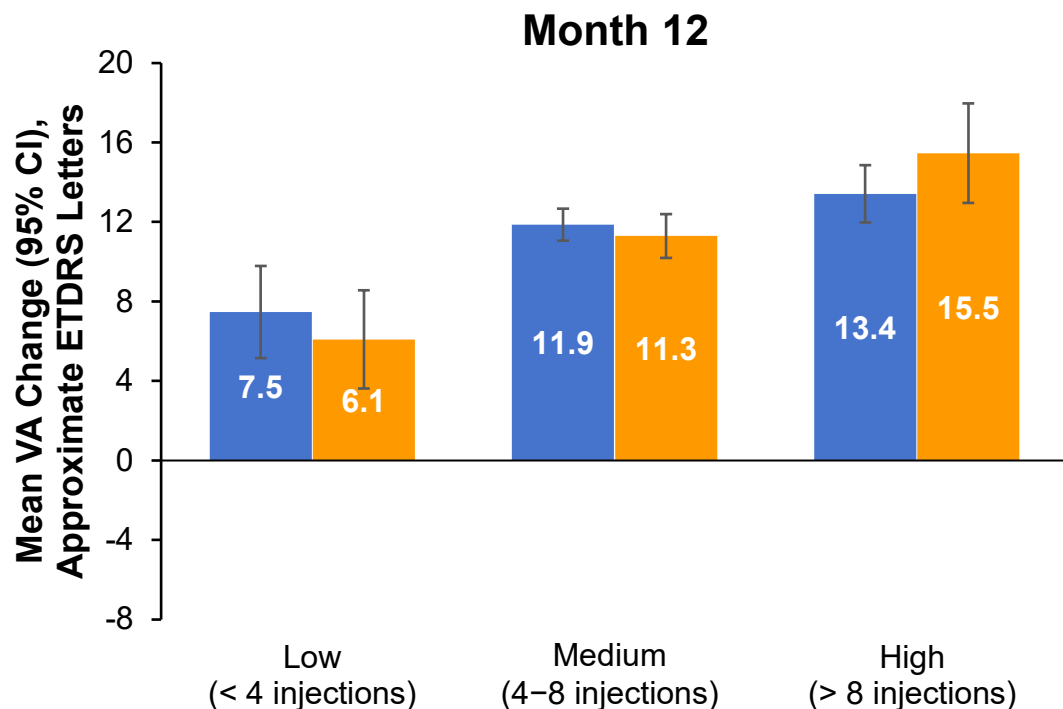
Number of Anti-VEGF Injections Was Highest in the First Year and Lower in Years 2–5 in BRVO and C/HRVO Eyes



| | | | | | | |
|--------|---|------|------|------|------|-----|
| BRVO | n | 2856 | 2007 | 1369 | 1012 | 673 |
| C/HRVO | n | 2879 | 2055 | 1420 | 993 | 679 |

VA Change From Index Injection to Months 12 and 60 Was Lower in BRVO and C/HRVO Eyes With Low vs High Injection Numbers

Mean VA Change From Index Anti-VEGF Injection Over 60 Months



| BRVO | n | 202 | 1474 | 461 |
|--------|---|-----|------|-----|
| C/HRVO | n | 237 | 1553 | 301 |

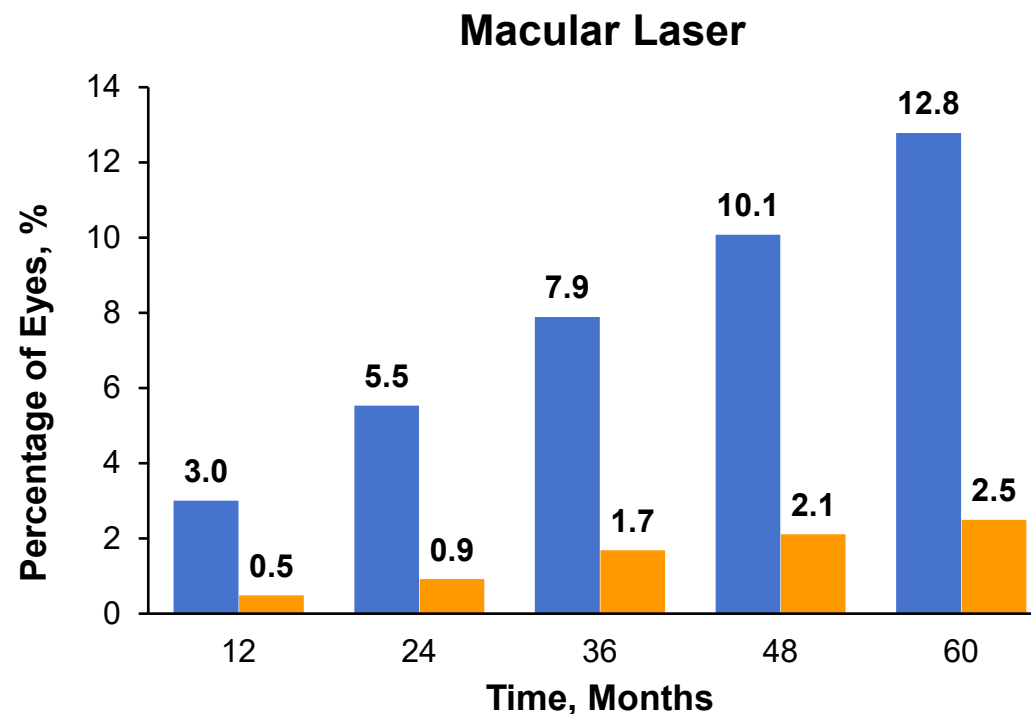
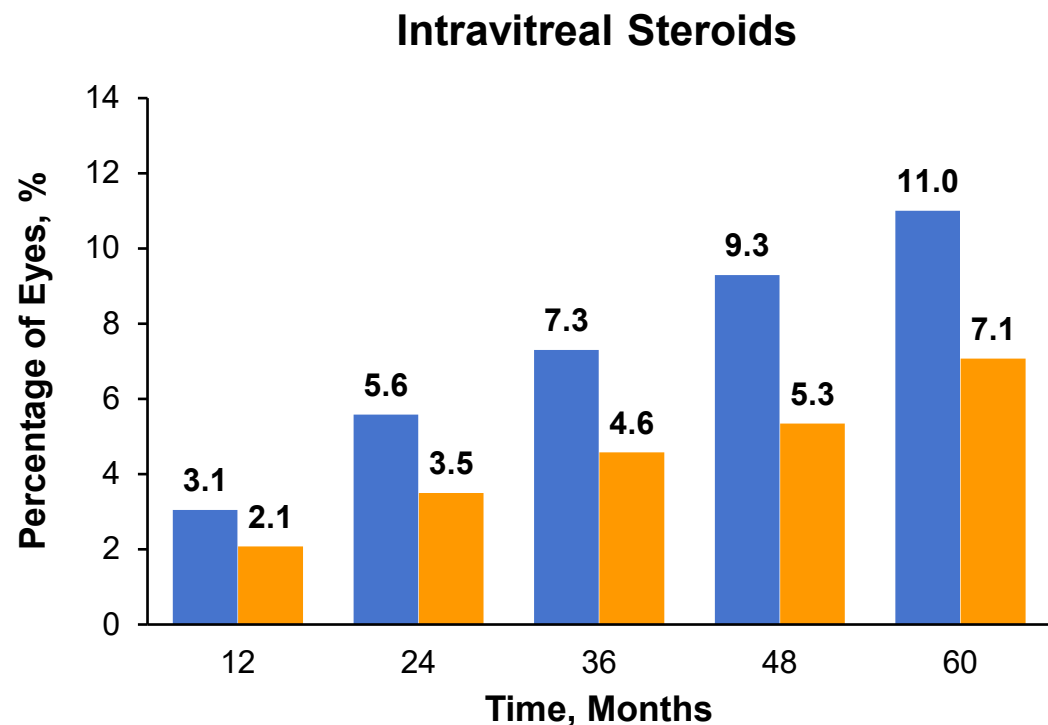
| BRVO | n | 90 | 167 | 240 |
|--------|---|-----|-----|-----|
| C/HRVO | n | 117 | 187 | 194 |

^a Injection number cut-offs were different between BRVO (low: < 11; median: 11-13; high: > 23) and C/HRVO (low: < 12; median: 12-24; high: > 24).

VA measurements were collected within ± 30 (up to 18 months) or ± 60 (from 24 to 60 months) days from date of interest to increase likelihood of collecting a measurement for each eye. The reading closest to the date of interest was used in eyes with 2 readings within the buffer window. Categories for injection counts over 60 months are based on tertiles. BRVO, branch retinal vein occlusion; C/HRVO, central/hemiretinal vein occlusion; CI, confidence interval; ETDRS, Early Treatment Diabetic Retinopathy Study; VA, visual acuity; VEGF, vascular endothelial growth factor.

Intravitreal Steroids and Macular Laser Treatments Were More Common Among BRVO and C/HRVO Eyes With Longer Follow-Up

Percentage of Eyes Receiving Treatment Over 60 Months



| | | | | | | |
|--------|---|------|------|------|------|-----|
| BRVO | n | 2856 | 2007 | 1369 | 1012 | 673 |
| C/HRVO | n | 2879 | 2055 | 1420 | 993 | 679 |

| | | | | | | |
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| BRVO | n | 2856 | 2007 | 1369 | 1012 | 673 |
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Conclusions



These data from the United Kingdom over **60 months** show that, on average, **anti-VEGF injections improve vision** in eyes with macular edema secondary to **RVO**, but eyes may require **long-term treatment to maintain baseline or any improvement in vision**



With **longer follow-up**:

- **VA gains** and the average number of **anti-VEGF injections** were **lower**
- **Intravitreal steroid** and **macular laser** use were **higher**



Greater vision improvements were observed among eyes with a **high vs low number of injections**



Study Limitations

- ▶ **Limited generalizability** beyond the United Kingdom
- ▶ **Missing data** (eg, due to loss to follow-up)
- ▶ **Anatomic outcomes** were not available for assessment



Study Strengths

In contrast to highly selected clinical trial participants, this analysis reflects the **long-term visual outcomes of heterogeneous real-world RVO patients** treated with **anti-VEGF therapy**

These data highlight a need for more durable treatments and long-term monitoring to maintain vision in eyes with RVO