Investor Presentation

July 2025

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The Leader in Sustained Release Drug Delivery for Retinal Disease

DURAVYU™: vorolanib - best-in-class, patent protected TKI in next-generation Durasert E™ sustained delivery technology

LUGANO and LUCIA Phase 3 wet AMD trials fully enrolled >800 patients randomized

Compelling efficacy and safety data in DME from VERONA Phase 2 trial; positive EOP2 meeting with FDA

Favorable safety profile across four clinical trials in 190+ patients

$318M at 3/31/25 – runway into 2027 post Phase 3 pivotal data in 2026

TKI tyrosine kinase inhibitor; wet AMD, wet age-related macular degeneration; DME, diabetic macular edema

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Pipeline Leveraging Bioerodible Durasert E™ Drug Delivery Technology Pursuing Large Market Opportunities

Durasert E™ Programs	Indication	Discovery	Pre-Clin	Phase 1	Phase 2	Phase 3	Anticipated Next Milestone
DURAVYU™ – (vorolanib intravitreal insert) **(f/k/a EYP-1901)**	**Wet AMD**	**LUGANO and LUCIA fully enrolled**					LUGANO topline data expected in mid 2026; LUCIA to closely follow
	DME	**Positive EOP2 meeting with FDA**					Pivotal program in 2026
EYP-2301 – razuprotafib **(TIE-2 agonist)**	**serious retinal diseases**						Tox and PK data

Evaluating additional pipeline opportunities

non-clinical *trial underway*

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wet AMD, wet age-related macular degeneration; DME, diabetic macular edema; EOP2, End-of-Phase 2; PK, pharmacokinetic; GA, geographic atrophy

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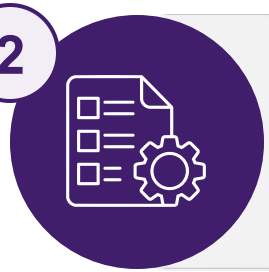
Despite current therapies, patients with wet AMD still lose vision in the long term

1

Many patients with wet AMD are chronically undertreated

>80% of Retina Specialists say undertreatment due to patient noncompliance, scheduling limitations or provider preference for less frequent dosing[1]

2



Chronic disease treated with short acting anti-VEGFs places significant burden on physicians and patients

77% of Retina Specialists say improved durability is most important when choosing a treatment[2].

3

A delay in care/missed visit can result in vision loss

A delay in treatment of only 5.34 weeks resulted in vision loss[3]

4

An increasing lifespan means significantly more injections in a patient's lifetime

Current anti-VEGF treatments are dosed on average every two months in the United States[4]

1. 2022 PAT Survey; 2. 2024 PAT Survey; 3. American Academy of Ophthalmology, The Effect of Delay in Care Among Patients Requiring Intravitreal Injections, Welin Song, BS et al; 4. NIH Current and Upcoming Anti-VEGF Therapies and Dosing Strategies for the treatment of wet AMD: a comparative review, Saira Khanna et al, Dec. 2019

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Diabetic Macular Edema: Large Market Opportunity with Significant Unmet Need for More Durable Treatments

Patients in the US with diabetes by 2030[1]

Develop DME within 10 years[2]

Global branded market by 2030[3]

Delayed/missed treatment visits[4]

Vision loss from missed injection[5]

By 2050, diabetes-related vision loss is expected to cost 500 million US dollars annually[6]

1. William R. Rowley, Clement Bezold, Yasemin Arikan, et al. Diabetes 2030: Insights from Yesterday, Today and Future Trends. PubMed Central. 2017 PMCID: PMC5278808 PMID: 27124621 . 2. Russel Lazarus. Optometrists Network. *Guide to Eye Conditions; Diabetic Macular Edema*. 3. DelveInsights DME Market Report -2030. 4. Monique A. Rose, Meri Vukicevic, Konstandina Koklanis. Adherence of patients with diabetic macular oedema to intravitreal injections: A systematic review. PubMed 2020 PMID: 32829485 DOI: 10.1111/ceo.13845. 5. Lee, R., Wong, T.Y. & Sabanayagam, C. Epidemiology of diabetic retinopathy, diabetic macular edema and related vision loss. Eye and Vis 2, 17 (2015). https://doi.org/10.1186/s40662-015-0026-2. 6. Nagda D, Mitchell W, Zebardast N. The functional burden of diabetic retinopathy in the United States. Graefes Arch Clin Exp Ophthalmol. 2021;259(10):2977–2986. https://doi.org/10.1007/s00417-021-05210-3.

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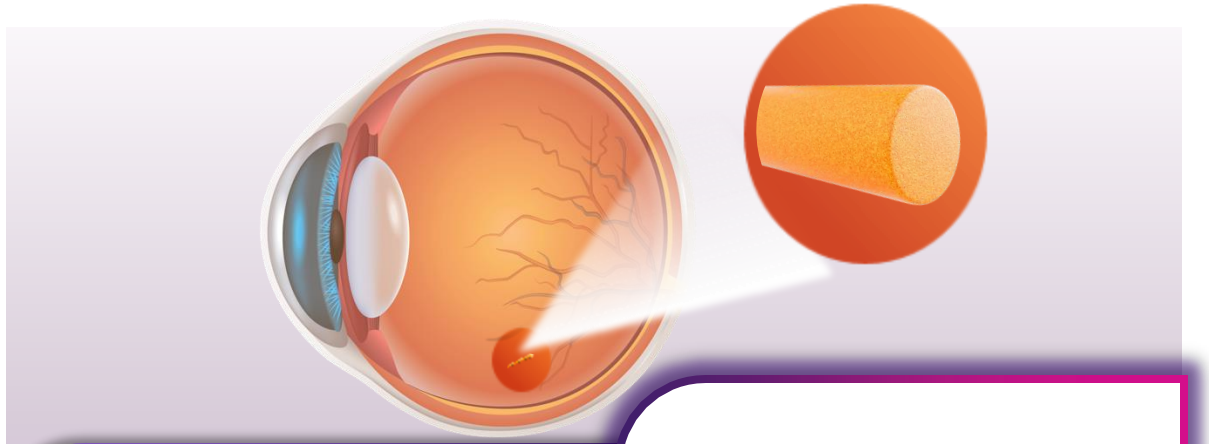
Durasert E™: The Next Generation in Sustained-Release Intravitreal Drug Delivery

Durasert® → **Durasert E™**

Over three decades of innovation in sustained-release drug delivery



Nonerodible

- **Vitrasert®** (ganciclovir IVT)[a]
- **Retisert®** (fluocinolone acetonide)
- **ILUVIEN®** (fluocinolone acetonide)
- **YUTIQ®** (fluocinolone acetonide)

EYP-1901
(vorolanib IVT insert)

Bioerodible
1030 µg payload

DURAVYU (EYP-1901)
(vorolanib IVT insert)

Bioerodible
High payload (1343 µg)

Inserts not drawn to scale, for illustrative purposes only. [a]Vitrasert now discontinued. **IVT**, intravitreal; **PEG**, polyethylene glycol; **PLGA**, poly lactic-co-glycolic acid.

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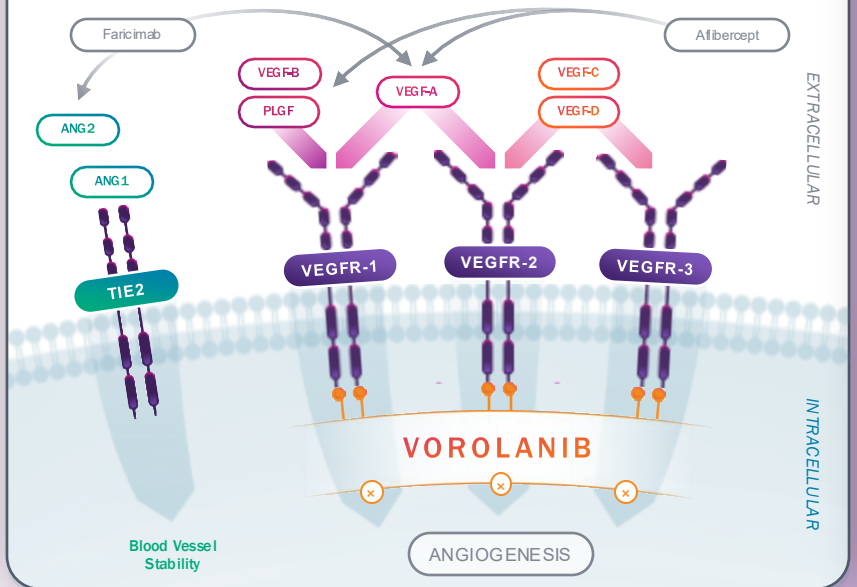
DURAVYU™: Vorolanib in Bioerodible Durasert E™

DURAVYU

Vorolanib: most studied TKI in retinal disease

Novel MOA: Selective and potent TKI that works intracellularly to inhibit all VEGFRs

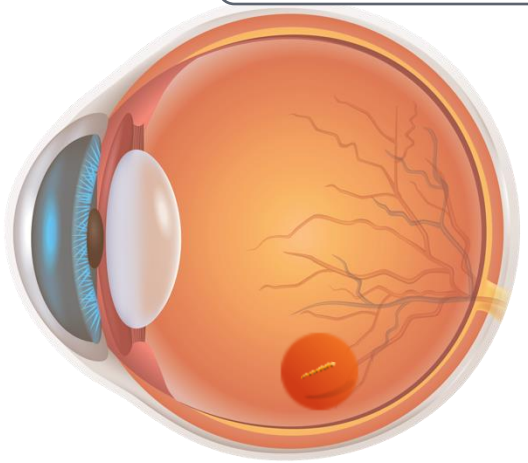


Durasert E

Bioerodible, sustained IVT drug delivery

Insert is 94% drug and 1/5000th of vitreous volume

Contains no PEG or PLGA



Immediate*
No delay - reaches target tissues at therapeutic levels within hours

Consistent*
Delivers a continuous daily therapeutic dose with no fluctuations

Sustained
VEGF receptor inhibition for at least six months

Controlled
Full elution of vorolanib before matrix bioerodes; no free-floating drug particles

Convenient
shipped and stored at ambient temperature; administered in a pre-loaded syringe injector

*Data from preclinical studies; **1.** Bakri SJ, et al. *PLoS One*. 2024;19(6):e0304782. **2.** Patel S, et al. *Ophthalmol Sci*. Sep-Oct 2024;4(5):100527. **3.** Wykoff CC et al. *J Vitreoretin Dis*. Sep-Oct 2024;8(5):577-586. **ANG,** angiopoietin; **IVT,** intravitreal; **TIE2,** tyrosine-protein kinase receptor TIE-2; **TKI,** tyrosine kinase inhibitor; **VEGF(R),** vascular endothelial growth factor (receptor).

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DURAVYU™ Demonstrated Robust Efficacy Outcomes and an Excellent Safety Profile Across Multiple Indications

DURAVYU Has Been Evaluated in >190 Patients to Date Across Four Clinical Trials

17 patients received DURAVYU	**DAVIO** **Phase 1 Wet AMD**	**Favorable safety and tolerability profile** **No safety signals or ocular SAEs related to DURAVYU**	• Stable BCVA and OCT • 74% reduction in treatment burden
102 patients received DURAVYU	**DAVIO 2** **Phase 2 Wet AMD**		• Statistically non-inferior BCVA compared to aflibercept • >80% reduction in treatment burden • Stable OCT
51 patients received DURAVYU	**PAVIA** **Phase 2 NPDR**		• Prevent worsening of disease severity
21 patients received DURAVYU	**VERONA** **Phase 2 DME**		• Met primary and secondary endpoints • Rapid and sustained improvement in vision and anatomical control with fewer injections

BCVA, best-corrected visual acuity; OCT, optical coherence tomography; wet AMD, wet age-related macular degeneration; NPDR, non-proliferative diabetic retinopathy; DME, diabetic macular edema

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Fully Enrolled

LUGANO/LUCIA Phase 3 Pivotal Trials in wet AMD

NON-INFERIORITY VERSUS ON-LABEL AFLIBERCEPT CONTROL

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DURAVYU De-Risked Clinical Approach in Wet AMD

Established FDA approval Path

- Last **four FDA approvals** followed the **noninferiority** pathway
- **On label control arm** – patients randomized on Day 1

Completed robust Phase 2 trial

- **Derisked** and informed **Phase 3**
- **Statistically non-inferior** to on-label aflibercept in tough-to-treat population

Proactive FDA interaction

- Formal **EOP2 and Type C** meetings
- **Written alignment** on NI trial design and -4.5 NI margin

Patient-Centric approach

- **All patients receive treatment** with the goal of maintaining or improving vision
- Trial designed to inform **real-world treatment practices**

nAMD, neovascular age-related macular degeneration; EOP2, End of Phase 2; NI, noninferiority

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Positive Phase 2 DAVIO 2 Results in Wet AMD Support Ongoing DURAVYU Phase 3 Pivotal Program

Achieved All Primary and Secondary Endpoints

DAVIO2 Endpoint	2mg	3mg
☑ **Primary**: Non-inferior change in BCVA vs. aflibercept	- 0.3 letters	- 0.4 letters
☑ **Secondary:** Favorable safety profile	No DURAVYU-related SAEs	
☑ **Secondary:** Reduction in treatment burden vs. 6 mos. prior	89%	85%
☑ **Secondary:** Reduction in treatment burden vs. aflibercept	82%	76%
☑ **Secondary:** Supplement-free up to 6 months	63% 88% of eyes had 0 or 1 supplemental injections	63% 83% of eyes had 0 or 1 supplemental injections
☑ **Secondary:** Anatomical control vs. aflibercept	+12.4um	+5.2um

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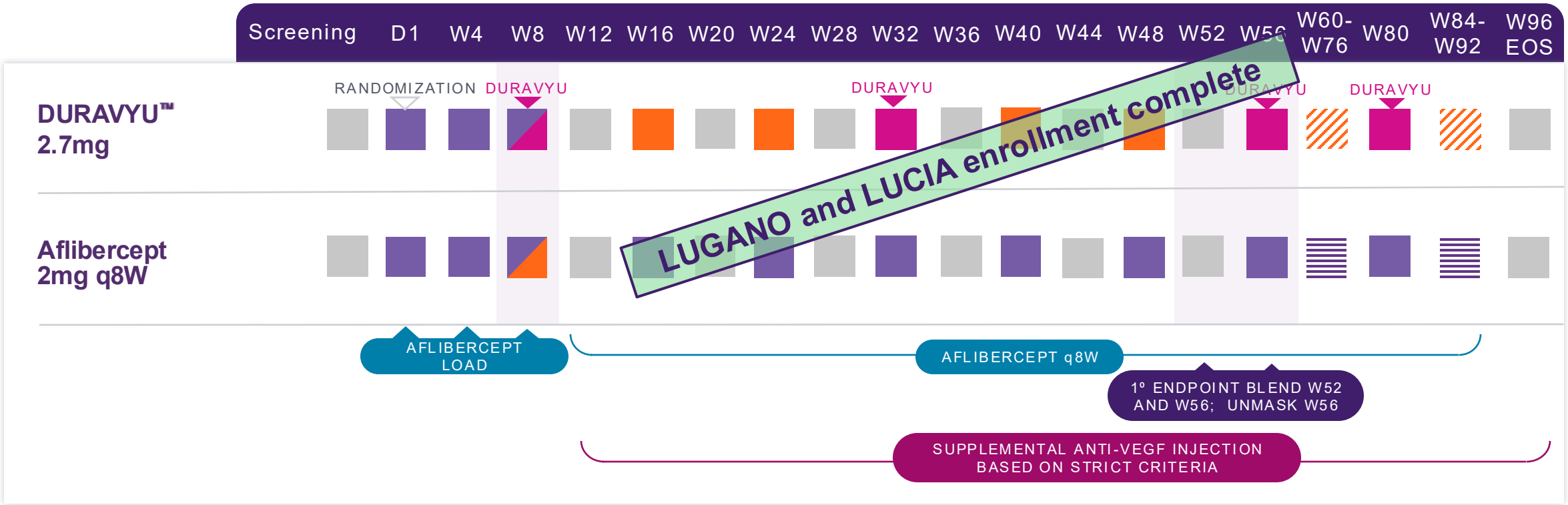
Phase 3 Pivotal Trials - Designed to Evaluate Non-Inferiority of DURAVYU vs On-Label Aflibercept Control



REQUIRED AFLIBERCEPT INJECTION VISIT · VISIT SCHEDULED · DURAVYU™ DOSE · SHAM INJECTION FOR MASKING · AFLIBERCEPT Q8W · SHAM INJECTION FOR MASKING

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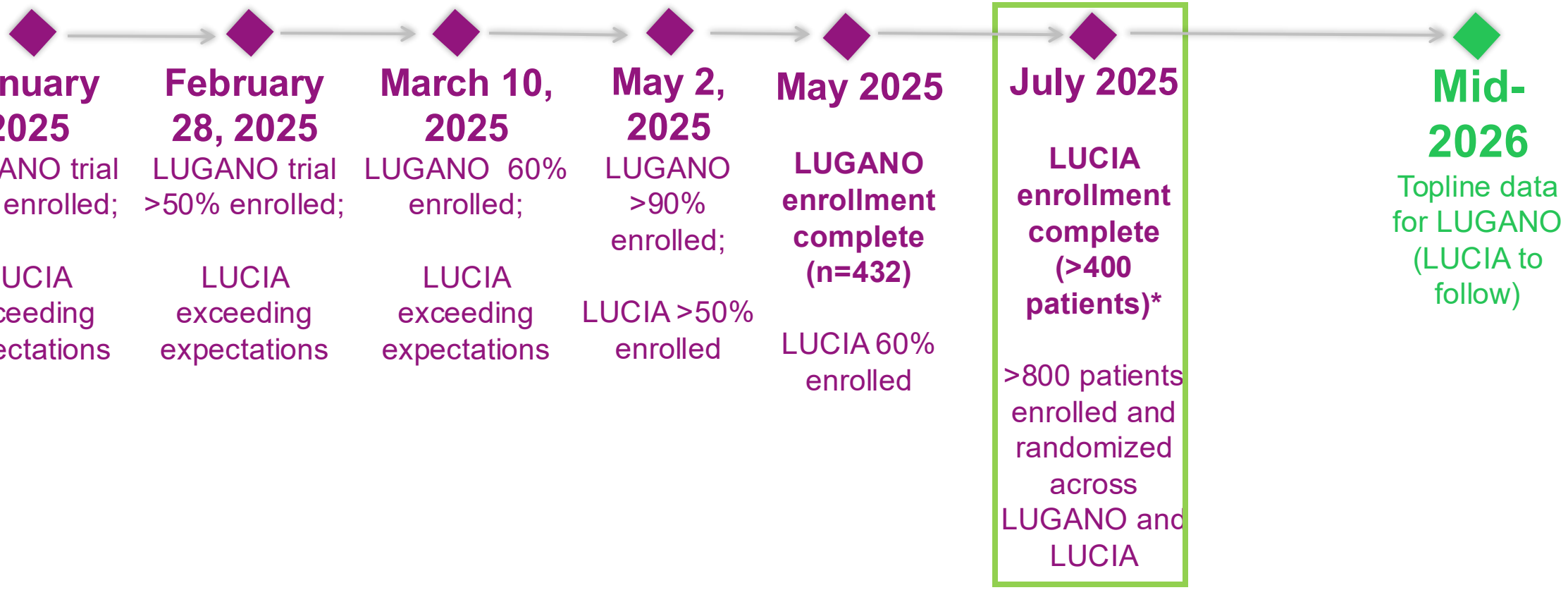
Exceptional Enrollment in Pivotal Phase 3 Wet AMD Program Driven by Strong Enthusiasm for the DURAVYU Pivotal Program



*Screening is closed for the LUGANO pivotal Phase 3 trial. Over 400 patients are enrolled and randomized. Any remaining patients in screening at the time of screening closure may also be randomized into the trial over the next several weeks.

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Phase 2 VERONA Clinical Trial in DME

24 WEEK RESULTS

PRIMARY AND SECONDARY ENDPOINTS ACHIEVED

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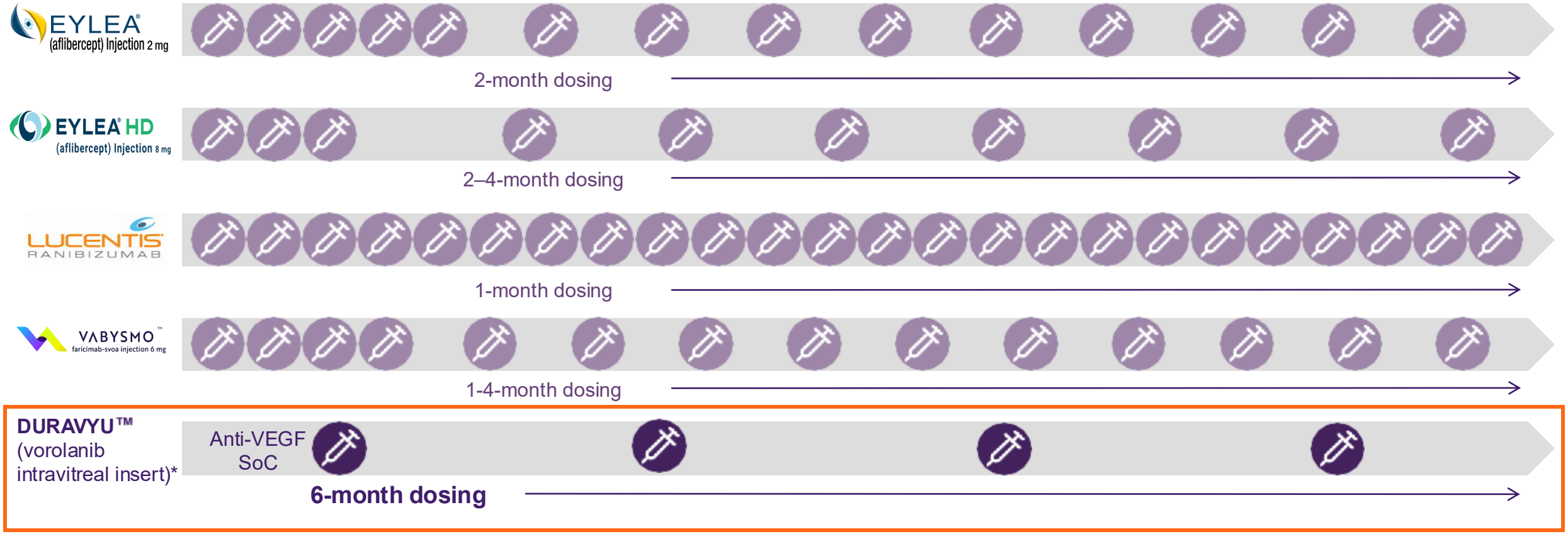
DME, diabetic macular edema

DURAVYU is Uniquely Positioned to Provide an Improved DME Treatment Paradigm Compared to Current Anti-VEGF Therapies

24-Month Treatment Regimens for DME:



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*Potential positioning for DURAVYU, if approved, based on Phase 3 protocol for wet AMD.
SoC, standard of care

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VERONA: DURAVYU demonstrated **stability in vision and anatomical control** with **fewer injections** in patients with DME

DURAVYU 2.7mg EFFICACY:

- **Primary endpoint met** - extended time to first supplemental injection vs. aflibercept control
- **Key secondary endpoints met:**
  - Rapid and sustained **BCVA improvement = +10.1*** letters
  - Rapid and sustained **CST improvement = -76** microns

DURAVYU SAFETY RESULTS:

- **No safety signals or ocular SAEs related to DURAVYU**
- **No cases of:**
  - Impaired vision
  - Endophthalmitis
  - Retinal vasculitis (occlusive or non-occlusive)
  - Intraocular inflammation (IOI)
  - Insert migration

*Data excludes one outlier patient. Outlier patient was removed from analysis because the patient missed multiple visits including the Week 20 visit resulting in vision loss of >20 letters at the Week 24 visit
DME, diabetic macular edema; BCVA, best-corrected visual acuity; CST, central subfield thickness; SAEs, serious adverse events.

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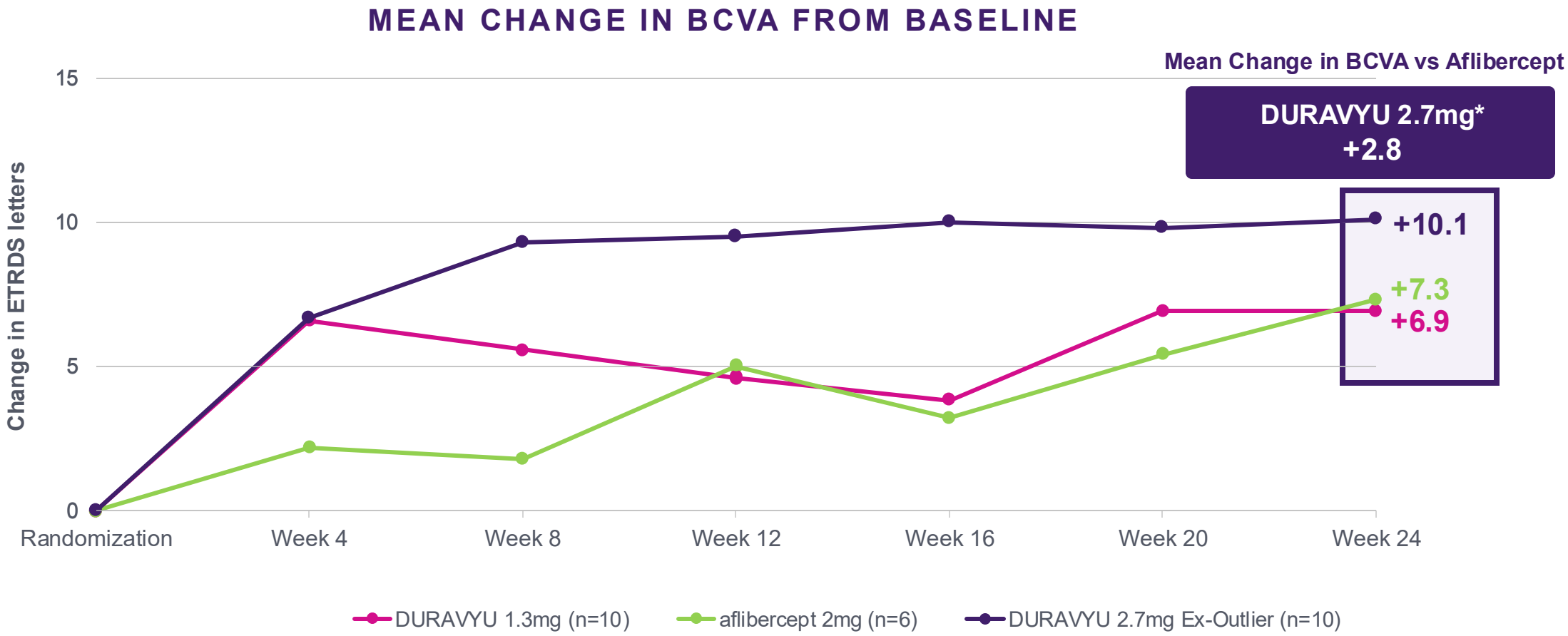
DURAVYU 2.7mg* Improved +10 Letters vs. Baseline; Nearly +3 Letters Better than Control at 24 Weeks



*Data excludes one outlier patient. Outlier patient was removed from analysis because the patient missed multiple visits including the Week 20 visit resulting in vision loss of >20 letters at the Week 24 visit. BCVA, best-corrected visual acuity; VA, visual acuity

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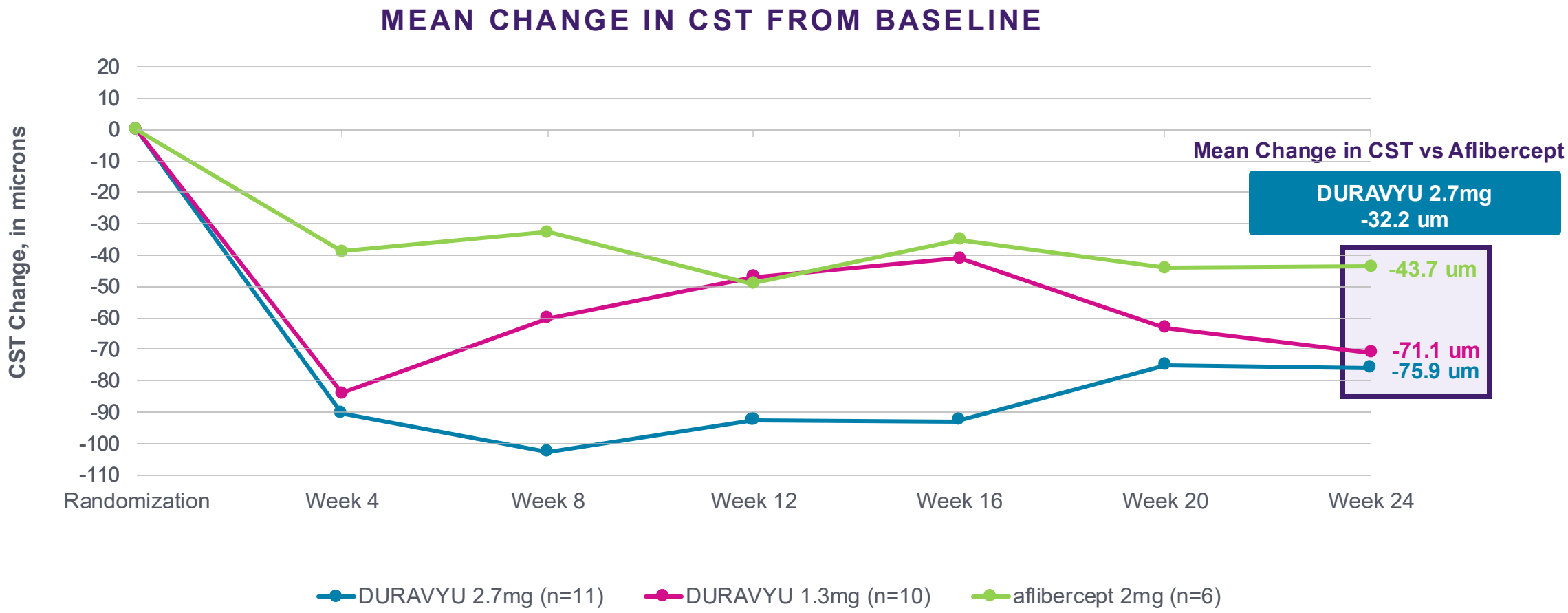
DURAVYU 2.7mg Demonstrated Improved Anatomical Control with 74% More Drying than Aflibercept Control



CST: central subfield thickness
Source: Independent reading center results available as of the full 24-week data cut.

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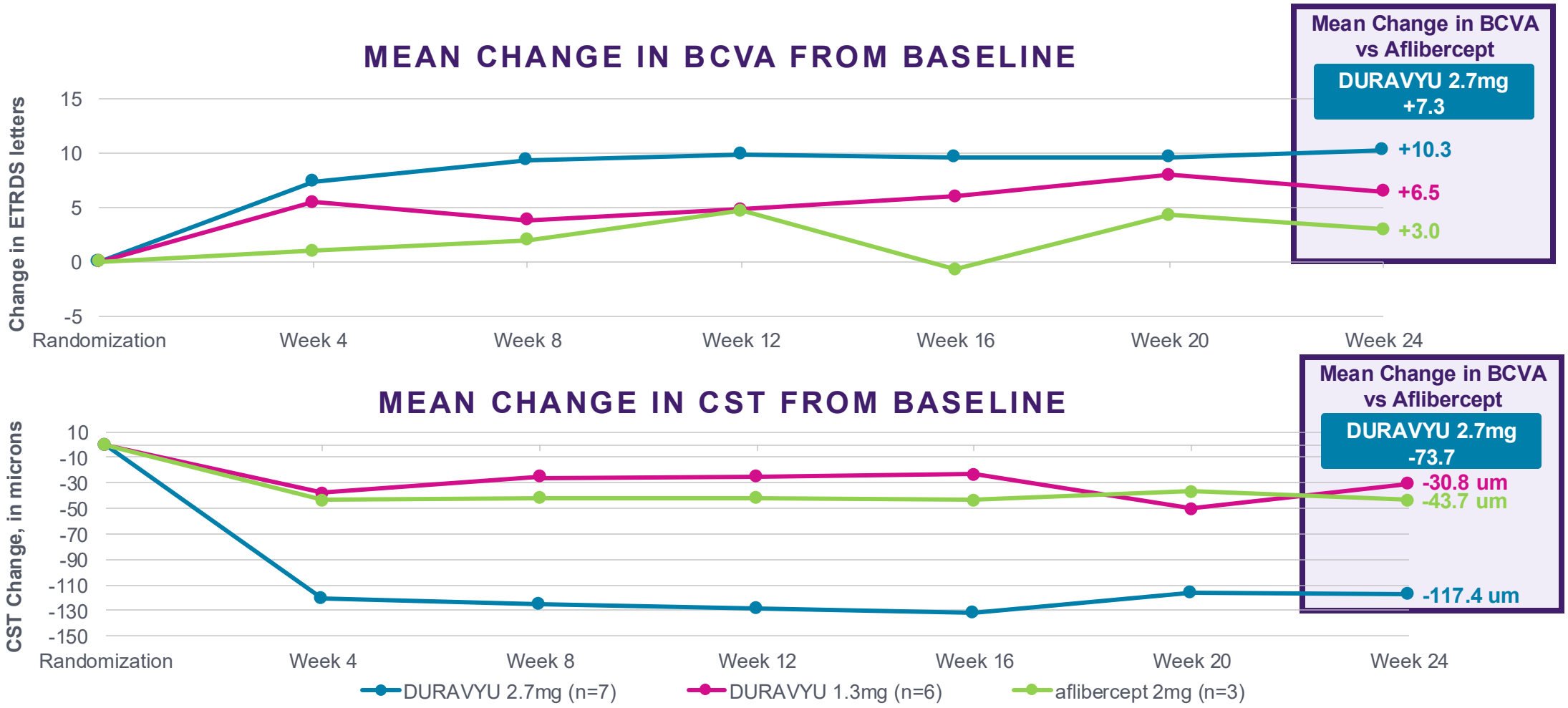
Sub-Group of Supplement-Free Patients Demonstrated Eyes Treated with DURAVYU 2.7mg had Better Visual and Anatomical Outcomes



Supplement-free is defined as patients who did not receive a supplement at any point during the study.

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BCVA, best-corrected visual acuity; VA, visual acuity

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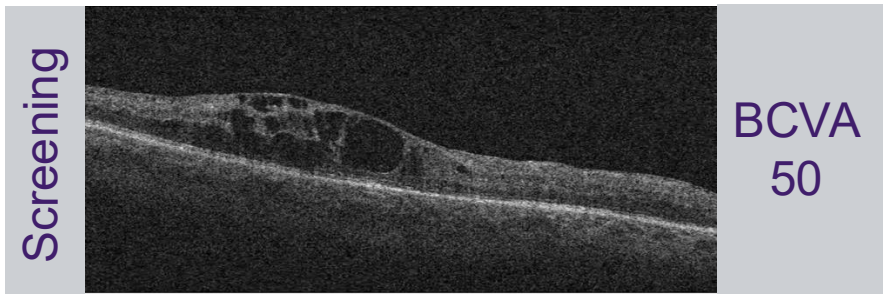
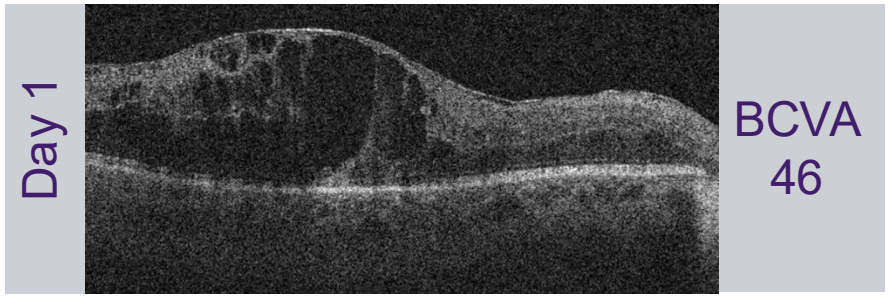
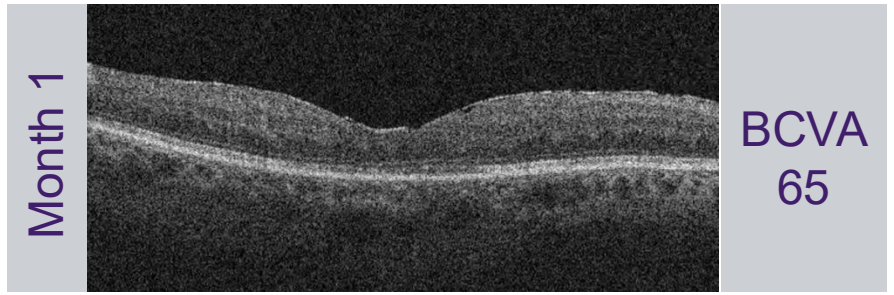
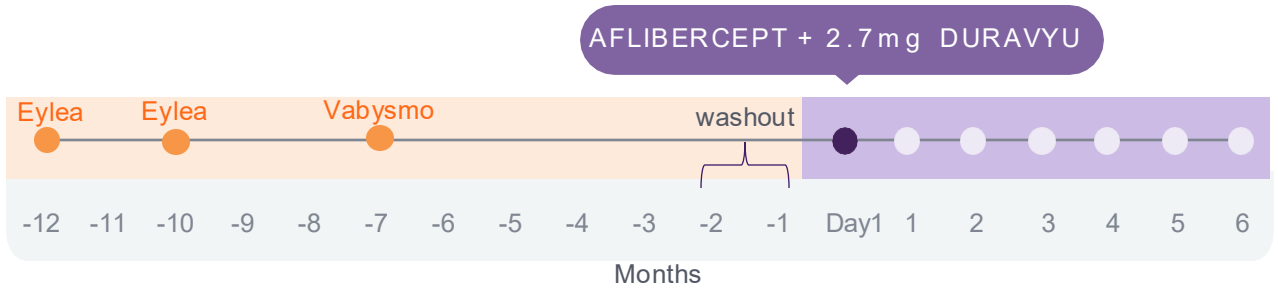
VERONA Case Study 1 - Rapid Drying with Improved Vision After Single DURAVYU 2.7mg Treatment and No Supplementation



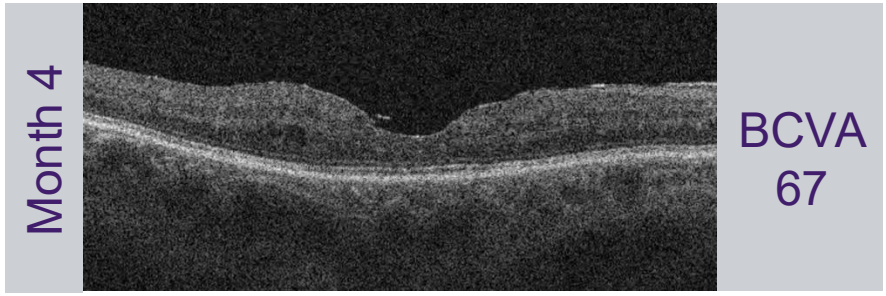
Day 1 (2.7mg DURAVYU insert)

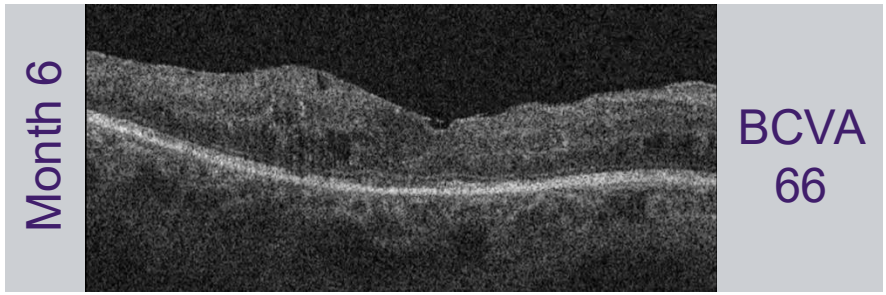






Case 1: SOC Anti-VEGF Injections Before and After Treatment





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BCVA, best-corrected visual acuity

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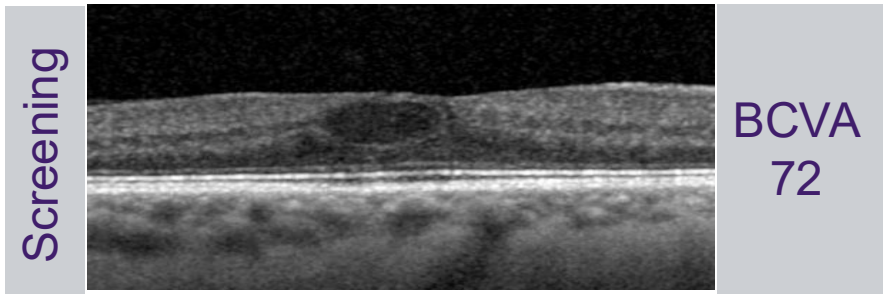
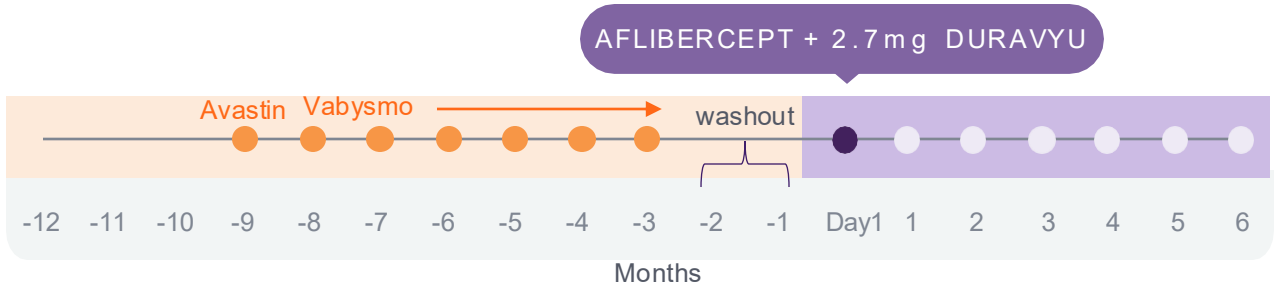
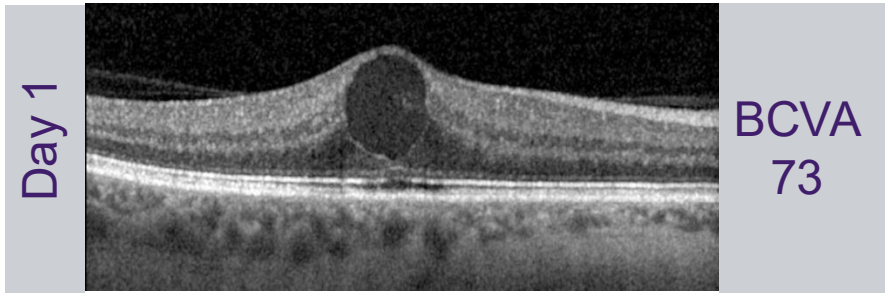
VERONA Case Study 2 – Better VA and Drying Than Vabysmo After Single DURAVYU 2.7mg Treatment and No Supplementation

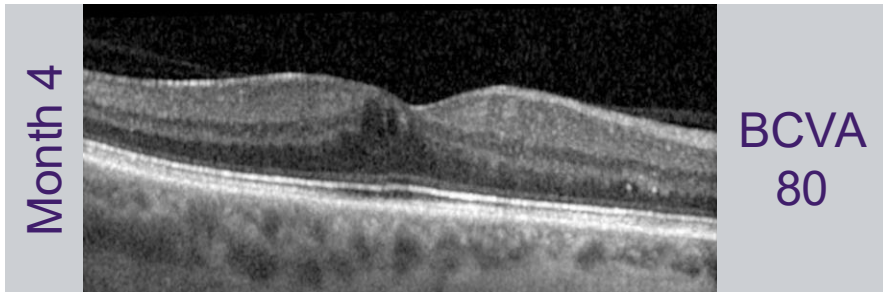


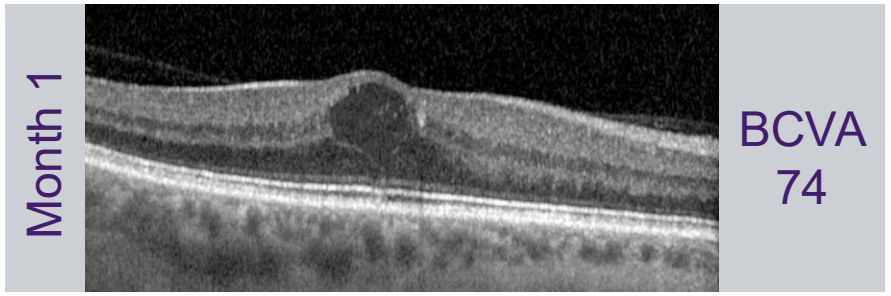


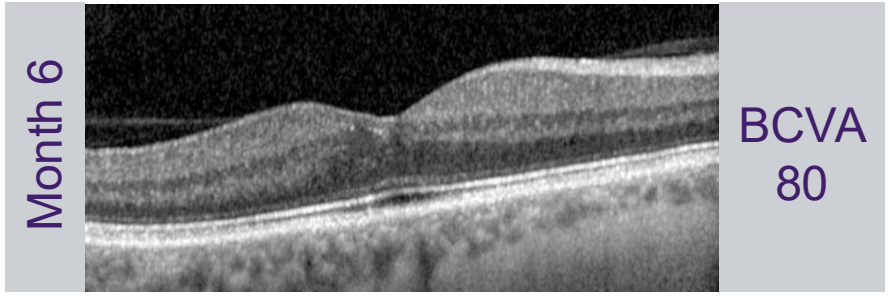
Case 2: SOC Anti-VEGF Injections Before and After Treatment

Day 1 (2.7mg DURAVYU insert)









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VA, visual acuity; BCVA, best-corrected visual acuity

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VERONA: Phase 2 Clinical Trial Summary

- ✓ **Primary endpoint met** for both DURAVYU doses
- ✓ Demonstrated **rapid** and **clinically meaningful results without a full load** of aflibercept
- ✓ Significant improvement in vision and anatomic results paired with a **meaningful reduction in treatment burden**
- ✓ Sub-group analysis of supplement-free eyes demonstrated **results are driven by treatment with DURAVYU** and not by supplemental injections
- ✓ Continued **favorable safety** and tolerability
- ✓ **Positive End-of-Phase 2 meeting with FDA**; pivotal program in 2026

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State-Of-The-Art cGMP Manufacturing Facility to Support DURAVYU through Potential NDA Filing and Global Commercial Production

- 41,000sf commercial manufacturing
- Located in Northbridge, MA
- Built to EYPT specifications by landlord preserving upfront cash investment
- Built to US FDA and EU EMA standards
- DURAVYU registration batches underway to support future NDA filing

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cGMP, current good manufacturing practices; FDA, Federal Drug Administration; EMA, European Medicines Agency; cGMP, current good manufacturing practice

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On Track for Continued Execution And Well-Funded Through Key Anticipated DURAVYU Milestones

DURAVYU™		
✓	Positive interim VERONA data	October 2024
✓	First patient dosed – LUGANO	October 2024
✓	First patient dosed – LUCIA	December 2024
✓	VERONA Phase 2 DME full 24-week data	February 2025
✓	VERONA Phase 2 subgroup analyses	March 2025
✓	Completed enrollment of LUGANO Phase 3 trial in wet AMD	May 2025
✓	FDA EOP2 meeting for DME	July 2025
✓	Full enrollment of LUCIA Phase 3 trial in wet AMD	3Q 2025
☐	**DME pivotal plans**	**2H 2025**
☐	**Topline data for LUGANO/LUCIA**	**2026**

Corporate		
✓	Fred Hassan appointed to Board of Directors	September 2024
✓	Northbridge manufacturing facility grand opening	October 2024
✓	Reginald Sanders, MD appointed to Board of Directors	January 2025
✓	Initiated DURAVYU registration batches in Northbridge	June 2025

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DME, diabetic macular edema; EOP2, End of Phase 2; wet AMD, wet age-related macular degeneration; SAB, Scientific Advisory Board

EYEPOINT

Investor Presentation

July 2025

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