

Results of a Phase 1, Open-Label, Dose-Escalation Study of THR-149 for the Treatment of DME

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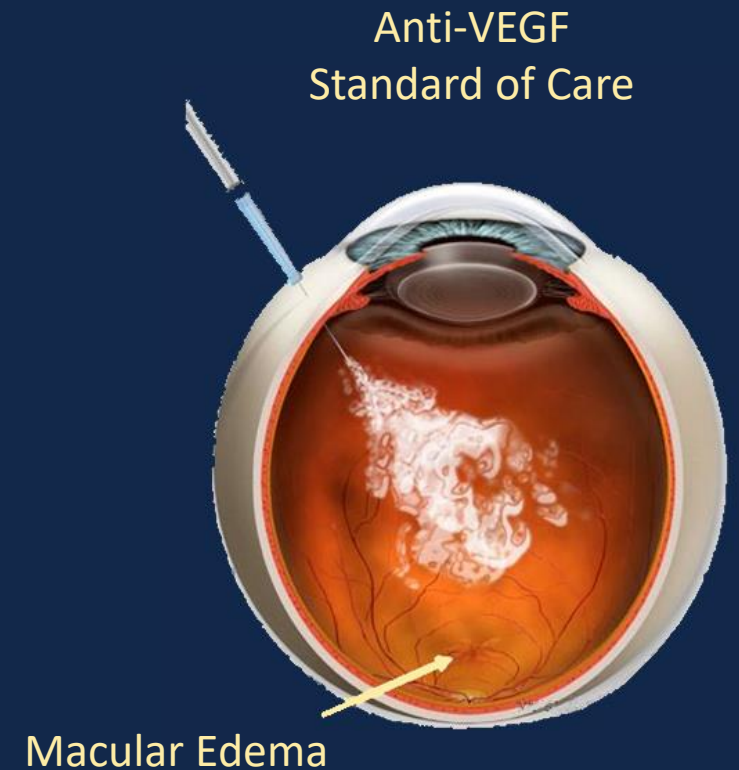
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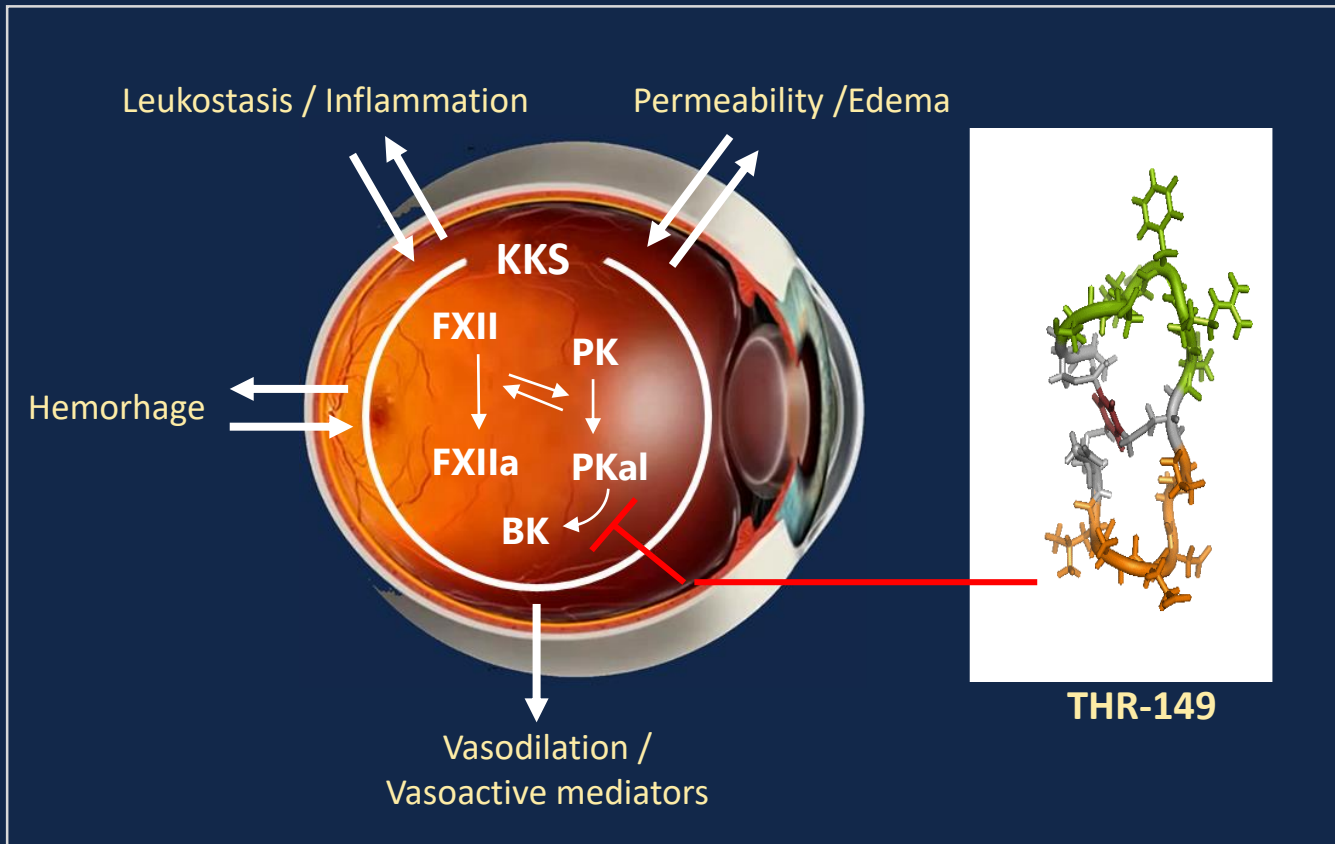
Unmet Medical Need in DME

- Anti-VEGFs are the 1st line treatment for DME
- Up to 40% patients do not adequately respond to anti-VEGF treatment in terms of BCVA and / or CST improvement
⇒ Other pathways involved in development of DME
- Potential side effects after a long-standing VEGF blockade *
 - ✓ VEGF acts as a survival factor for choriocapillaris, retinal neurons, and retinal pigment epithelium
 - ✓ Efficacious inhibition of VEGF can lead to higher incidence of retinal geographic atrophy in the clinic



Targeting Plasma Kallikrein in DME

THR-149 is a Potent Reversible Peptide Inhibitor of Plasma Kallikrein (PKal)

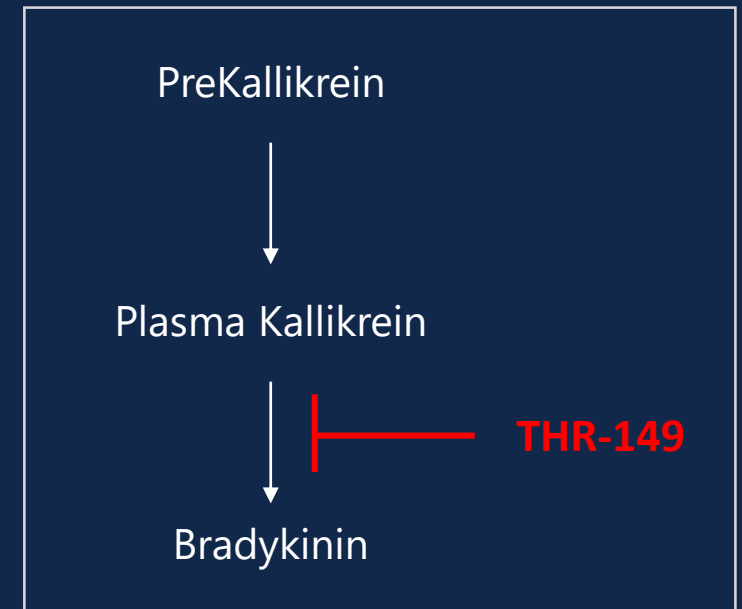
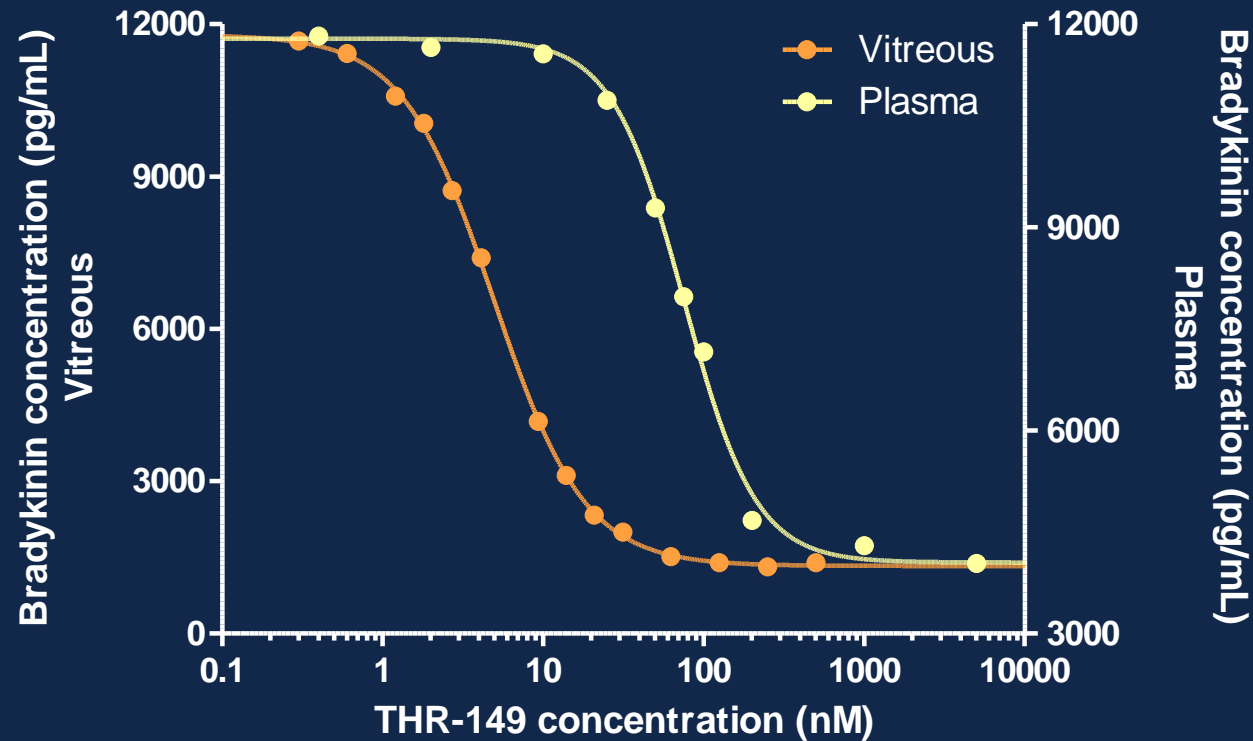


Targeting PKal offers a **VEGF-independent mechanism** for inhibiting DME

- PKal/Kinin System is upregulated under diabetic conditions
- In preclinical models of diabetes, PKal mediates vascular hyperpermeability, leukostasis, inflammation, and micro-hemorrhages
- Evidence for clinical efficacy after PKal inhibition hereditary angioedema and DME

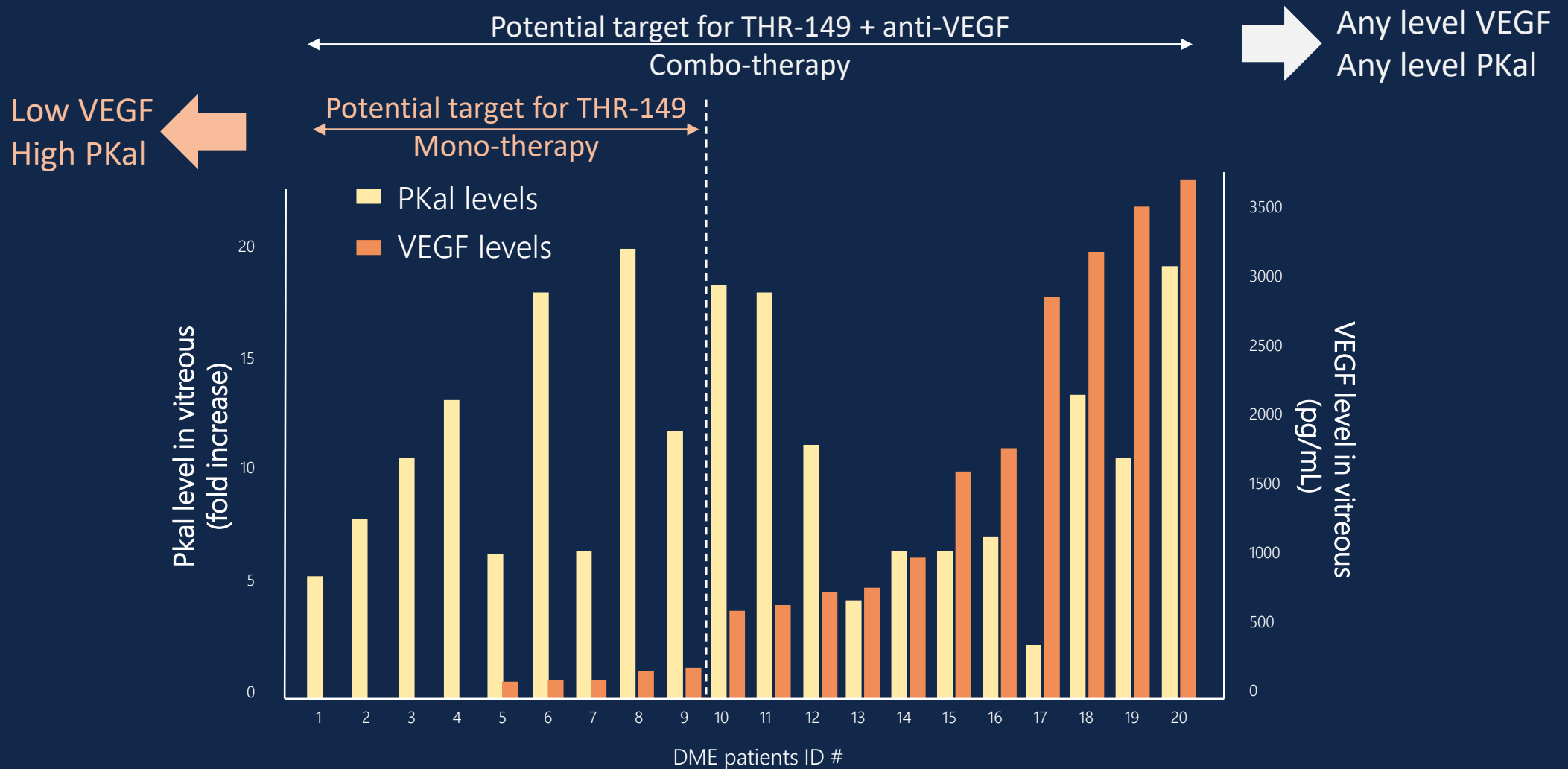
Biochemical Characterization of THR-149

Inhibition of Bradykinin Formation in Kaolin-Activated Human Plasma / Vitreous



THR-149 blocks release of bradykinin in plasma and vitreous

Rationale for Targeting PKal in DME



Start building clinical evidence with THR-149 mono-therapy

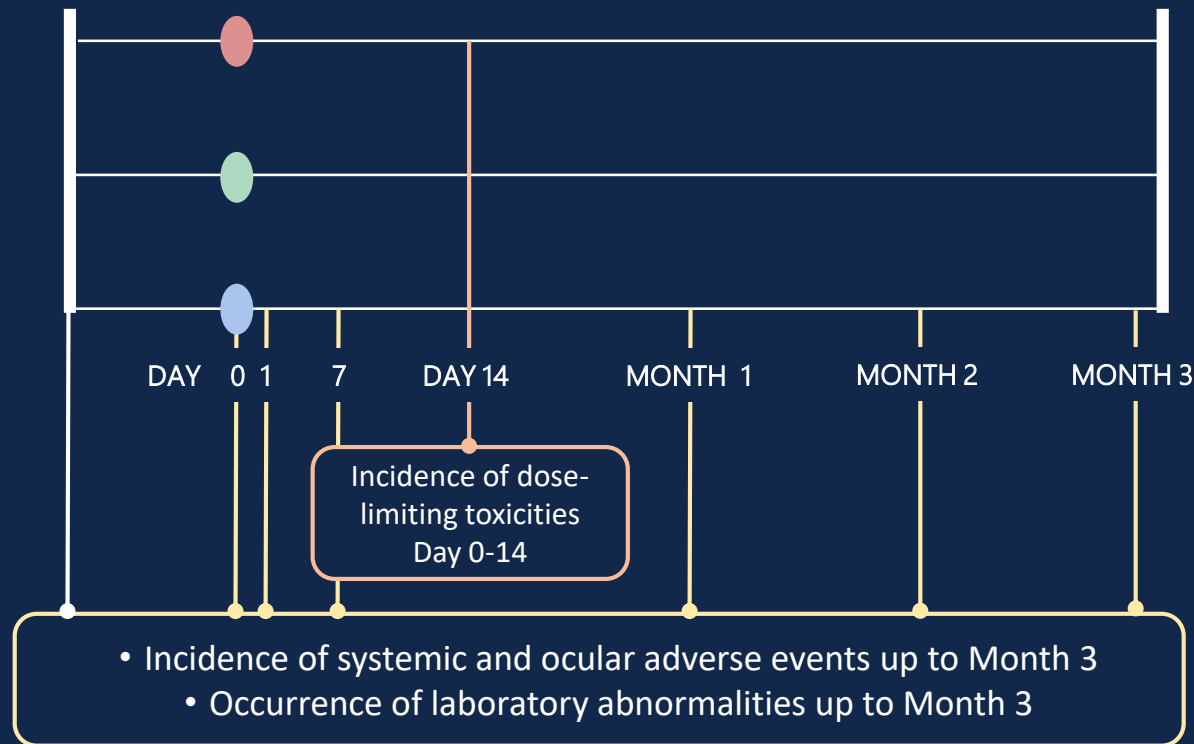
THR-149-001: Study Overview

3+3 Dose-Escalation Study

Study Treatment IVT

Total N = 12 patients

- CI-DME CST >320 μm (Spectralis SD-OCT)
- BCVA ≤ 62 and ≥ 23 letters
- History of response to prior anti-VEGF / corticosteroid treatment



- 0.005mg THR-149 (low dose)
- 0.022mg THR-149 (middle dose)
- 0.13mg THR-149 (high dose)

- Screening
- Primary outcome measure
- Secondary outcome measures

THR-149-001: Baseline Ocular Characteristics in the Study Eye

All Treated Subjects

Characteristic	Low Dose N=3	Middle Dose N=3	High Dose N=6
BCVA (ETDRS letters)			
Mean (SD)	46.0 (9.17)	46.7 (8.62)	43.0 (12.59)
Median	44.0	45.0	43.5
Min, Max	38, 56	39, 56	25, 58
CST (μm)			
Mean (SD)	497.7 (70.04)	539.3 (35.95)	529.5 (120.60)
Median	533.0	551.0	585.0
Min, Max	417, 543	499, 568	373, 626

No relevant imbalances between treatment arms for Baseline BCVA and CST

THR-149-001: Safety Overview

All Treated Subjects

Category	Low Dose N=3	Middle Dose N=3	High Dose N=6
	Number of events	Number of events	Number of events
Death	0	0	0
SAE	0	3	0
DLT	0	0	0
AE leading to withdrawal from study	0	0	0
Treatment-related (drug and / or procedure) AE	0	1	0

- 3 SAEs (non-ocular, nontreatment-related) in 1 subject
- No DLTs
- 1 treatment-related ocular AE

THR-149-001: Adverse Events in the Study Eye

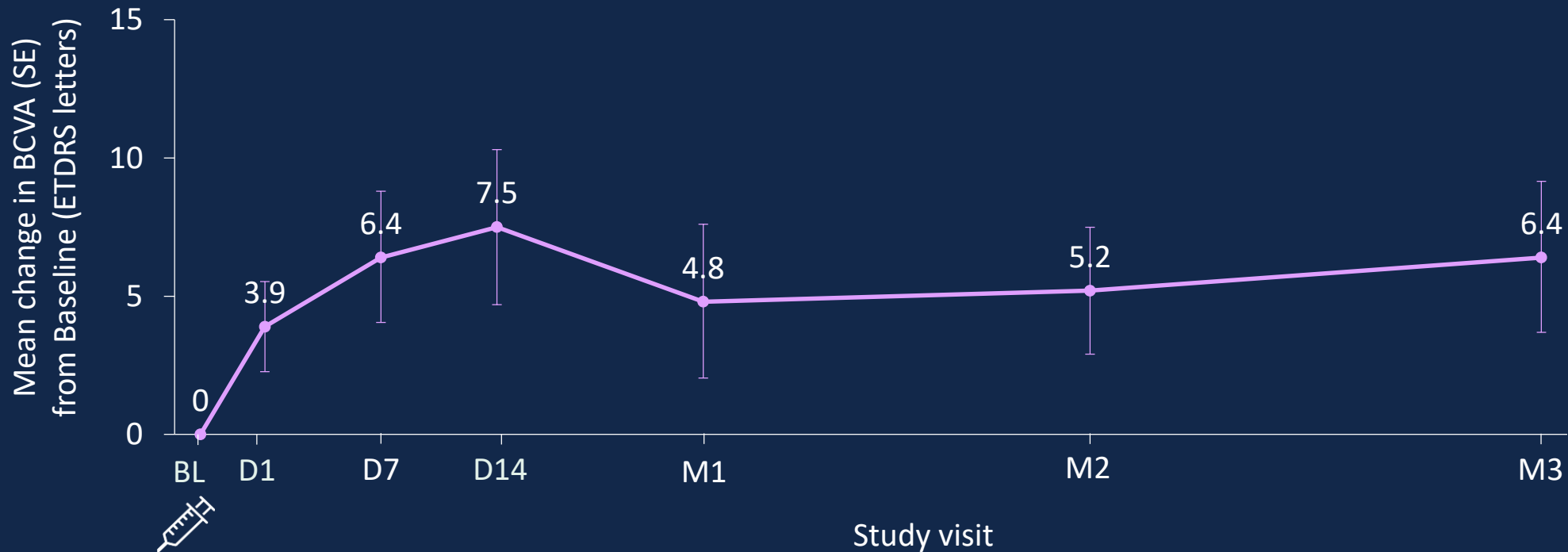
All Treated Subjects

Adverse event	Low Dose N=3	Middle Dose N=3	High Dose N=6
	Number of events	Number of events	Number of events
Anterior chamber inflammation	0	1 ^a	0
Conjunctival hemorrhage	0	1	0
Corneal disorder	0	1	0
Diabetic retinal edema	1	1	1
Eye pain	0	0	1
Macular fibrosis ^b	0	2	0
Vitreous floaters	1	0	0

- 1 ocular AE related to study treatment (likely injection procedure) in the middle dose
- All ocular AEs were likely due to the injection procedure, underlying disease progression, or concomitant diseases

THR-149-001: Mean Change in BCVA From Baseline (Accounted for Rescue)^a

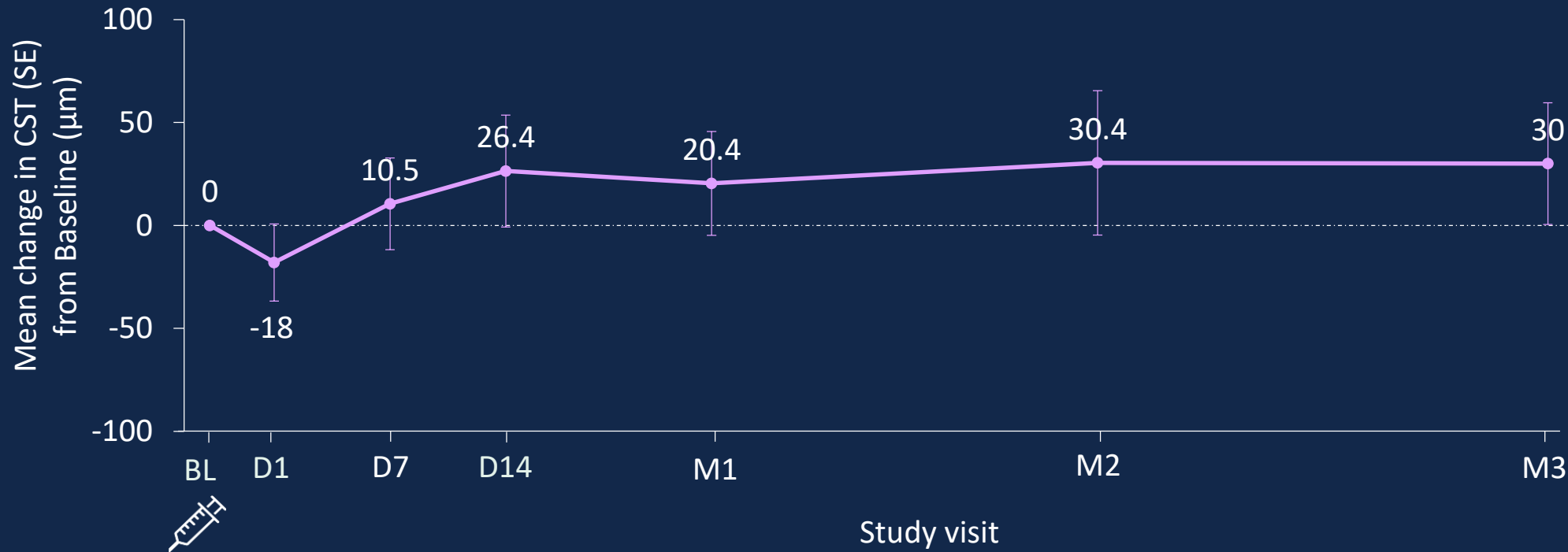
All Treated Subjects, Overall



Mean BCVA gain was fast and maintained until end of study

THR-149-001: Mean Change in CST From Baseline (Accounted for Rescue)^a

All Treated Subjects, Overall



- Marginal impact on mean CST at Day 1 followed by increase until study end
- Mean CST change was minimal and within the variability of measurement

Discussion: Understanding the disconnect between BCVA and CST in study THR-149-001

Existing public data

Known disconnect between BCVA and CST in DME patients:

- DRCR.net, Ophthalmology 2007
- Prior clinical data with PKaI inhibition in DME

Ongoing *quantitative* assessment of THR-149-001 data

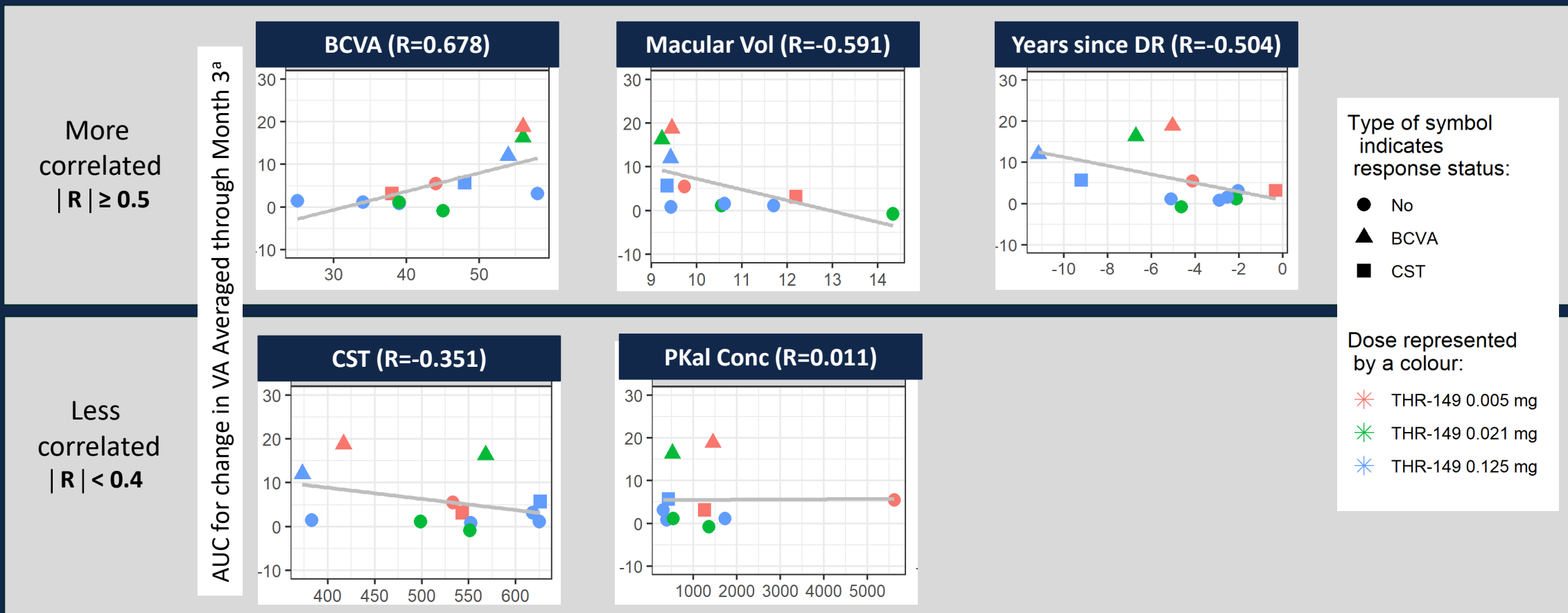
- ✓ Link between **Baseline** characteristics (DR duration, macular volume, PKaI levels in aqueous humor) and BCVA / CST response
- Link **over time** between anatomic parameters (other than CST) with BCVA response
- Impact of VEGF: VEGF levels determination in aqueous humor samples
- Evaluation aqueous humor samples for retinal stress markers

Ongoing *qualitative* assessment of THR-149-001 data

- Image interpretation by experts

Link between Baseline Characteristics and BCVA Response

Study THR-149-001: Exploratory Statistical Analysis

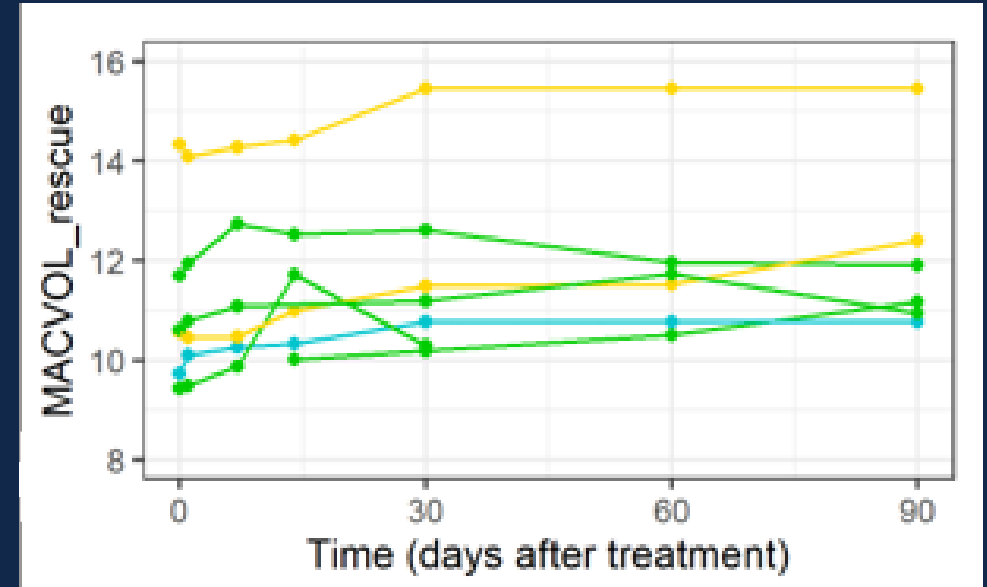
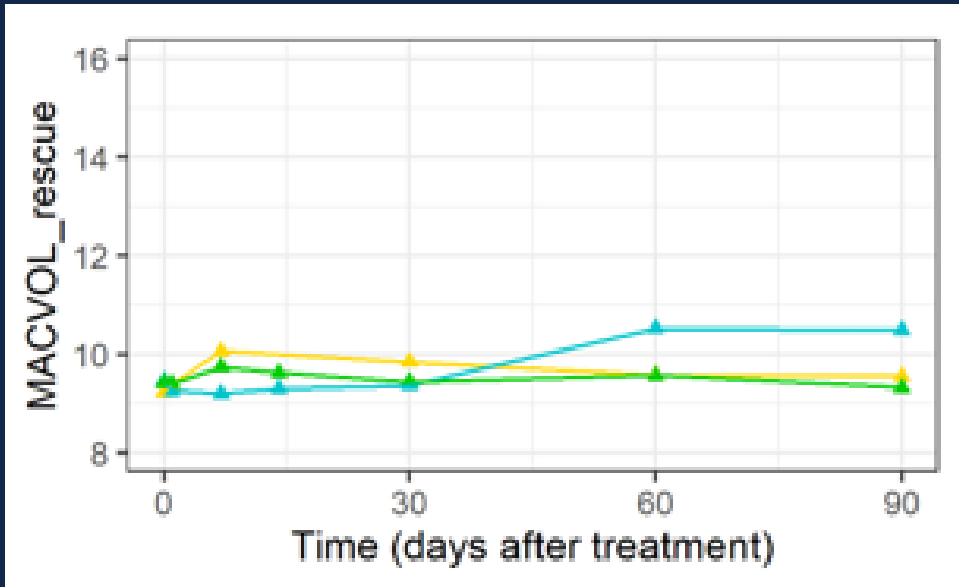


- BL BCVA, macular volume and years since DR are more predictive than BL CST or PKal levels for BCVA response
- Better BL BCVA, lower macular volume and longer duration of DR indicative for BCVA response

THR-149-001: Macular Volume over Time by Subject – by BCVA Response

Subjects defined as BCVA Responders *

Subjects defined as Non-Responders



*

In the BCVA responders, macular volume was maintained over time

Discussion: Understanding Disconnect between Reduction of Retinal Vascular Leakage in Pre-Clinical Model *versus* CST Data Clinical Study

Pre-Clinical Experiments



- Revisit the thickness of the individual retinal layers and visual performance (optomotor) in diabetic rat model
- Detailed study of retinal fluid homeostasis versus vascular permeability / leakage

THR-149-001: Key Take Away Messages

- THR-149 is **safe** and **well tolerated**:
 - No DLTs
 - No ocular SAEs
 - 1 treatment-related ocular AE - considered related to the injection procedure
- Mean BCVA gain was **fast** and **maintained** until end of study:
 - Day 1: 3.9 letters
 - Max at Day 14: 7.5 letters
 - Month 3: 6.4 letters
- **Macular volume** at BL seems to be indicative for BCVA response, and amongst the BCVA responders macular volume was maintained over time

Overall gains noted in BCVA, and improvement in CST in some subjects are encouraging and warrant further clinical research with multiple injections of THR-149