

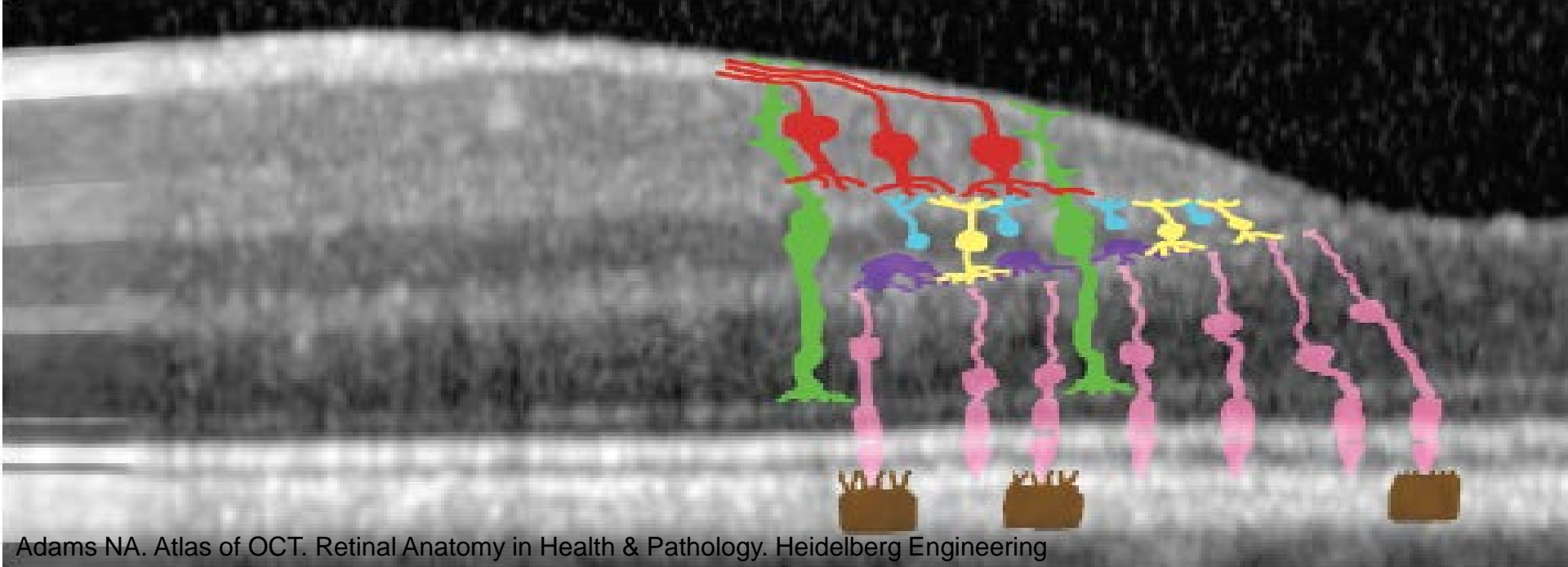
## **Afternoon Session**

**Stephanie Klemencic, OD**

**Yannek Leiderman, MD**

**Janice McMahon, OD**

**Ahmed Aref, MD**

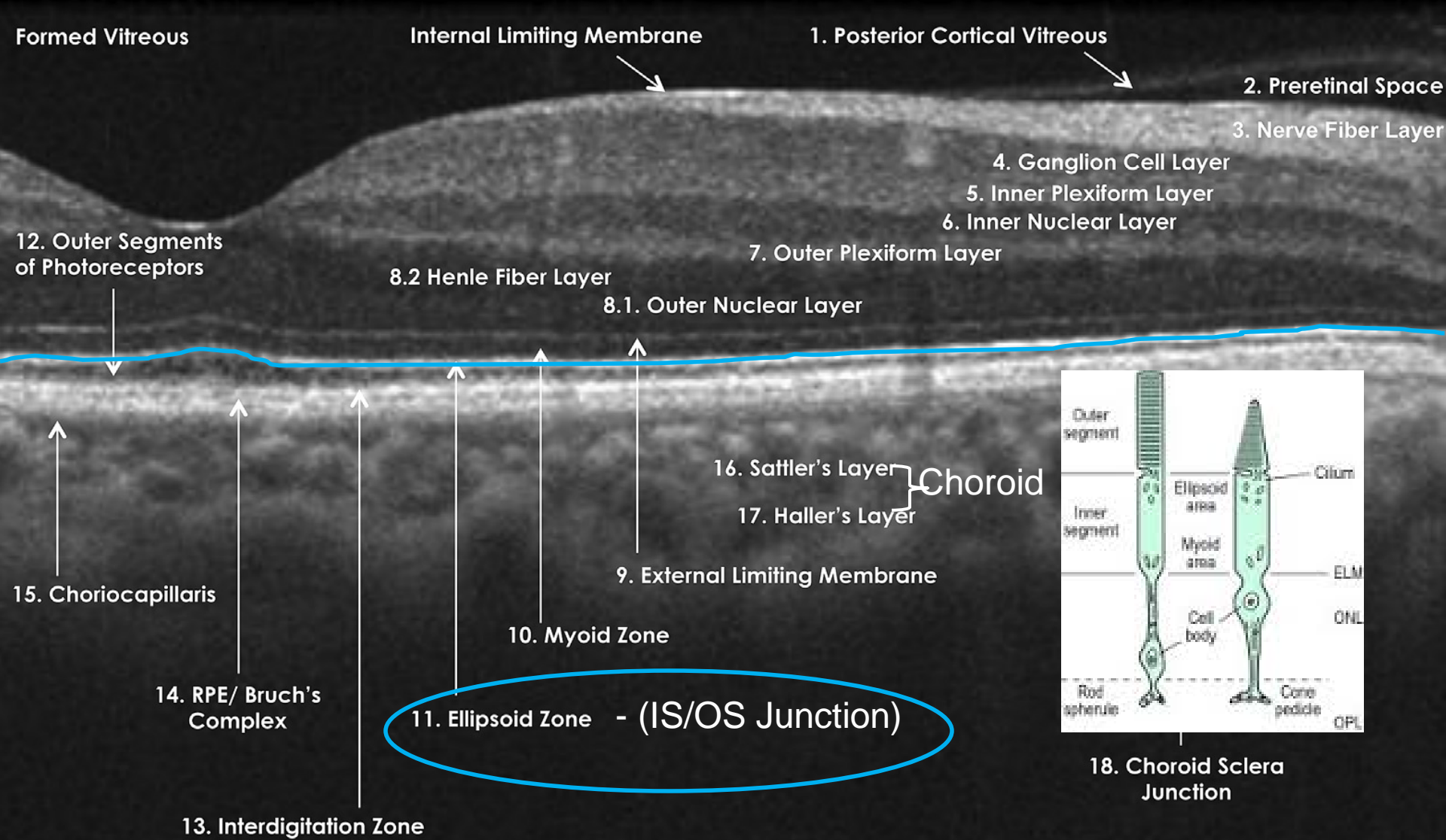


# RETINA UPDATES

Yannek Leiderman, MD, PhD  
University of Illinois Chicago  
Illinois Eye and Ear Infirmary

Stephanie Klemencic, OD  
Illinois College of Optometry  
Illinois Eye Institute

# International Nomenclature for OCT Meeting Consensus Normal OCT Terminology

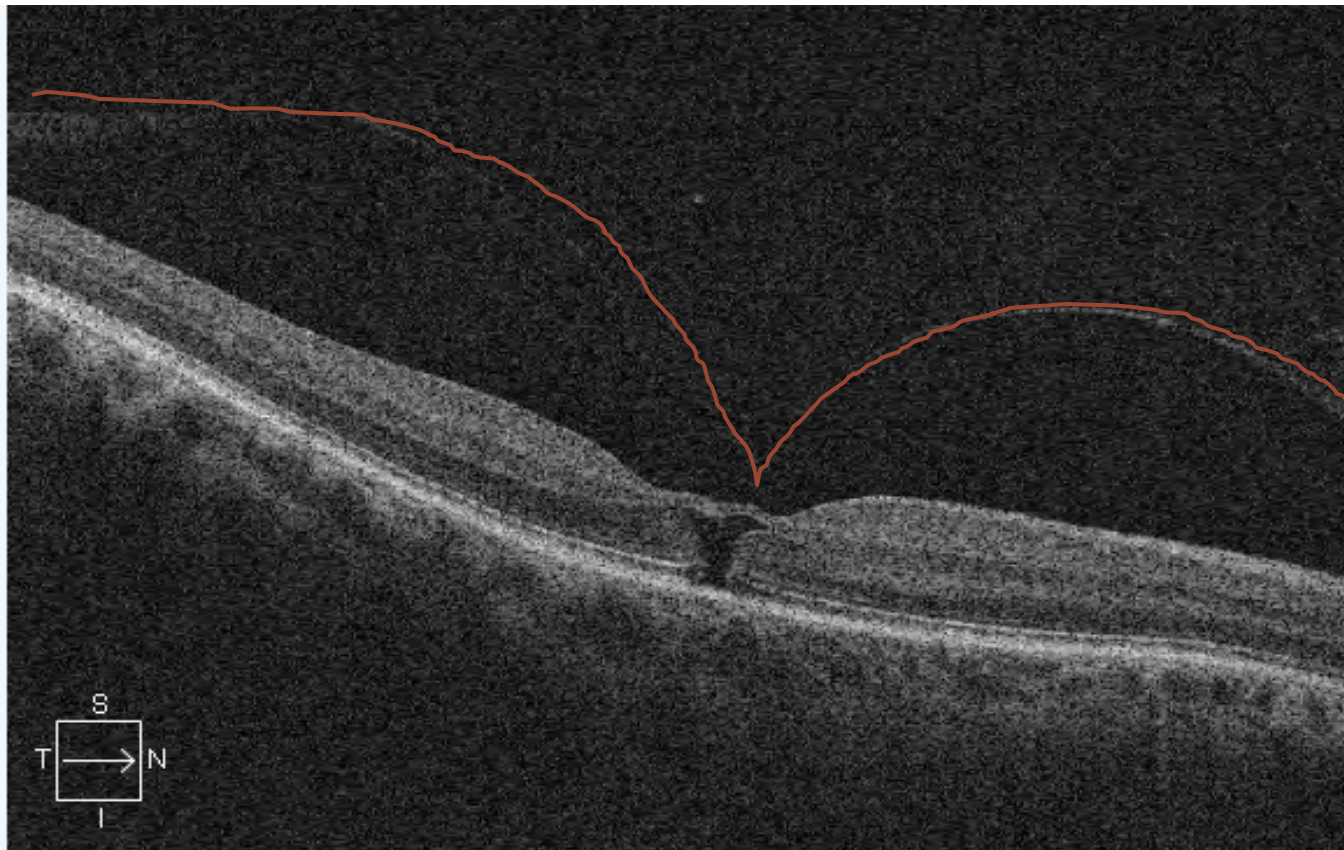


# Case #1: 46 y.o. AAF

BCVA: OD 20/30

OS 20/20

“Severe” visual distortion OD x 2 weeks





Pseudohole

Full thickness

Impending hole

Partial thickness

VMA

VMT

# Vitreo-Macular Interface

Stage 1

Medium

ERM

Small

Stage 0

Lamellar Hole

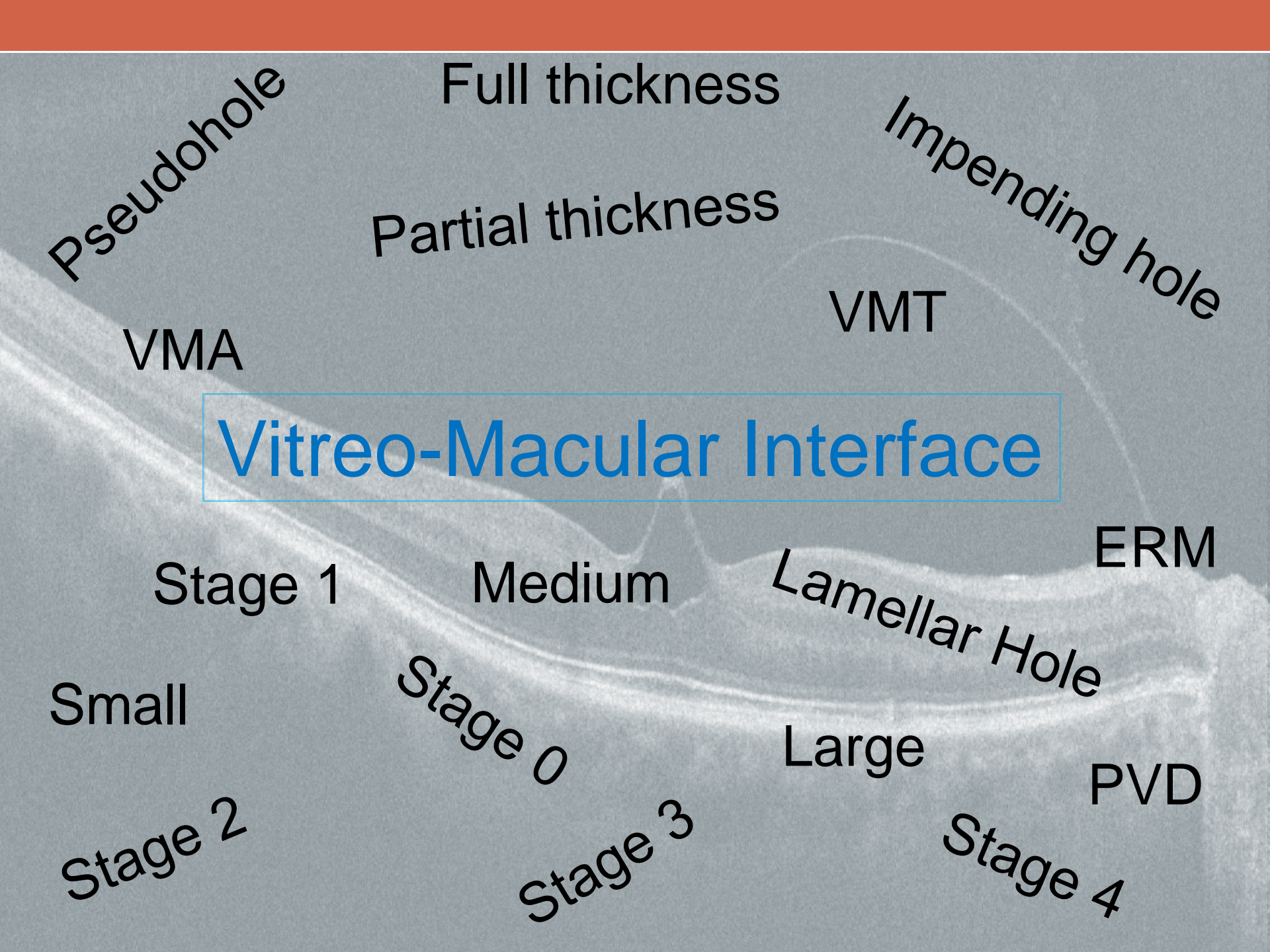
Stage 2

Stage 3

Large

PVD

Stage 4



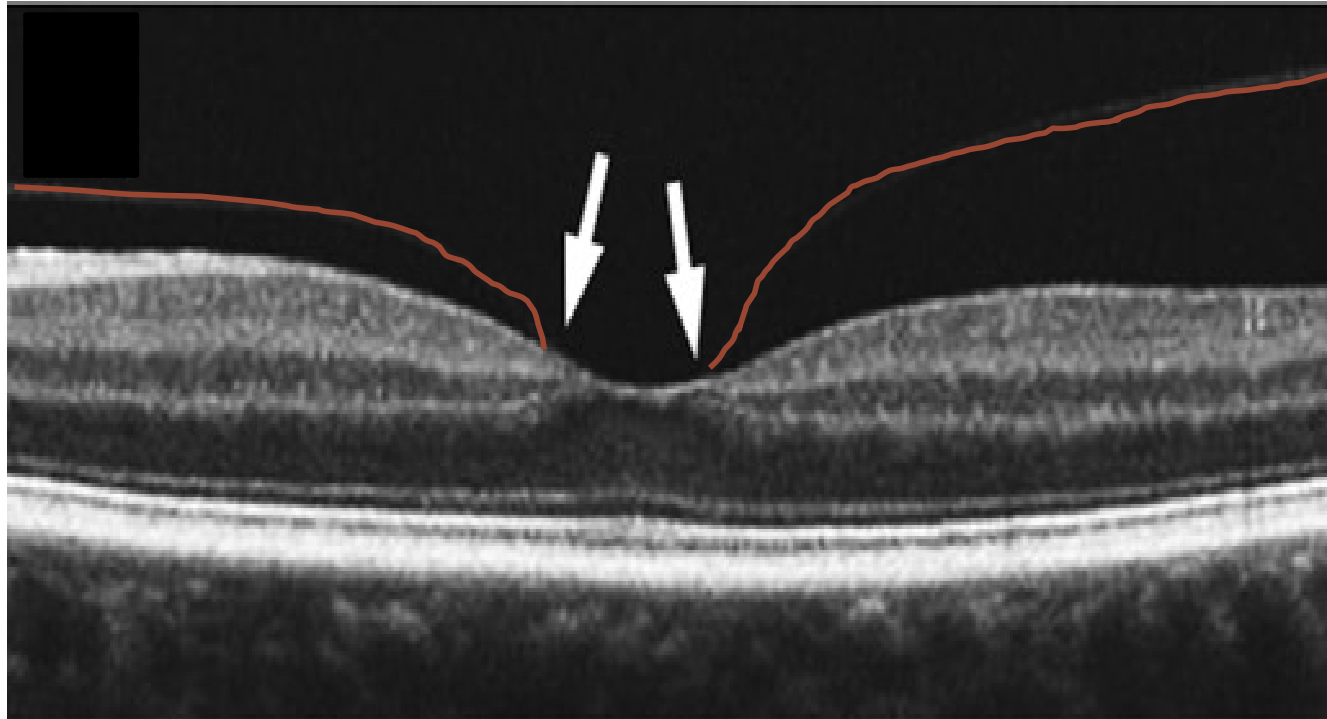
# OCT Classification of Macular Holes

Full Thickness Macular Hole (FTMH) Stages	IVTS Classification
Stage 0	VMA
Stage 1: Impending hole	VMT
Stage 2: Small hole	Small or medium FTMH with VMT
Stage 3: Medium hole	Medium or large FTMH with VMT
Stage 4: FTMH with PVD	Small, medium or large FTMH without VMT

“Smaller” hole = < 400um

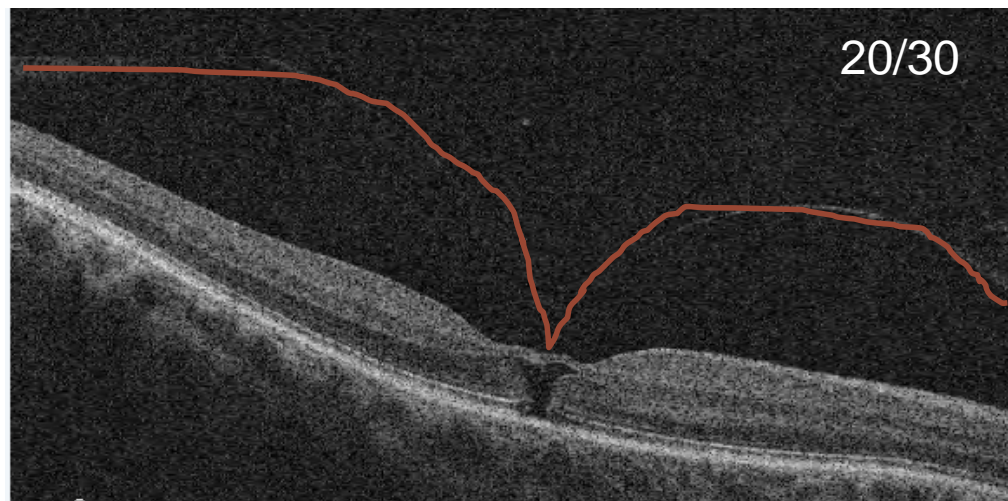
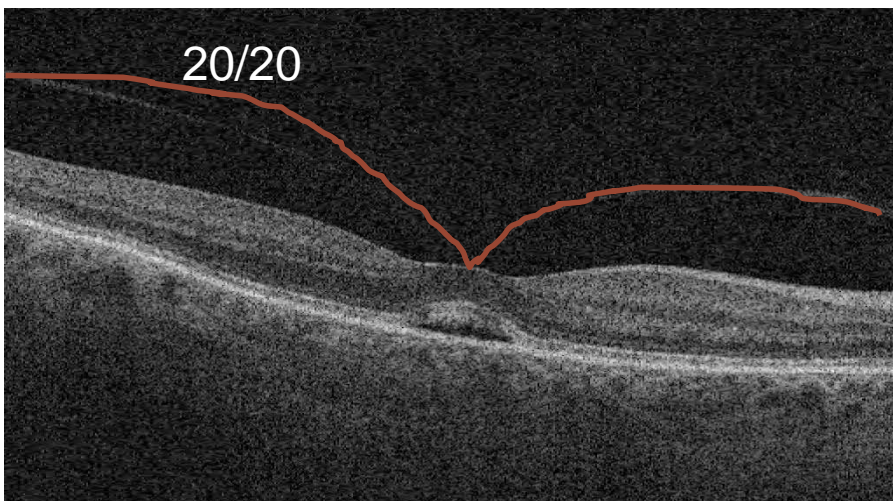
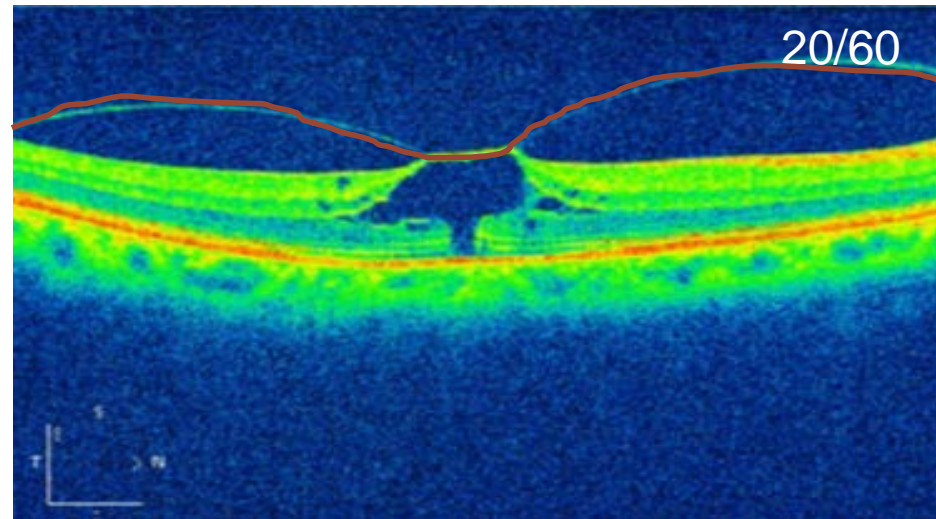
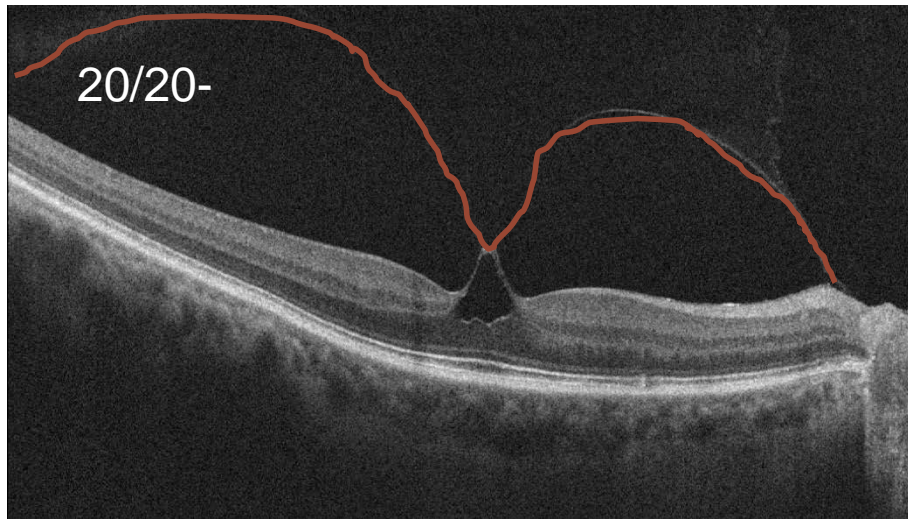
# Vitreo-Macular Adhesion (VMA)

Normal foveal contour and contents/thickness



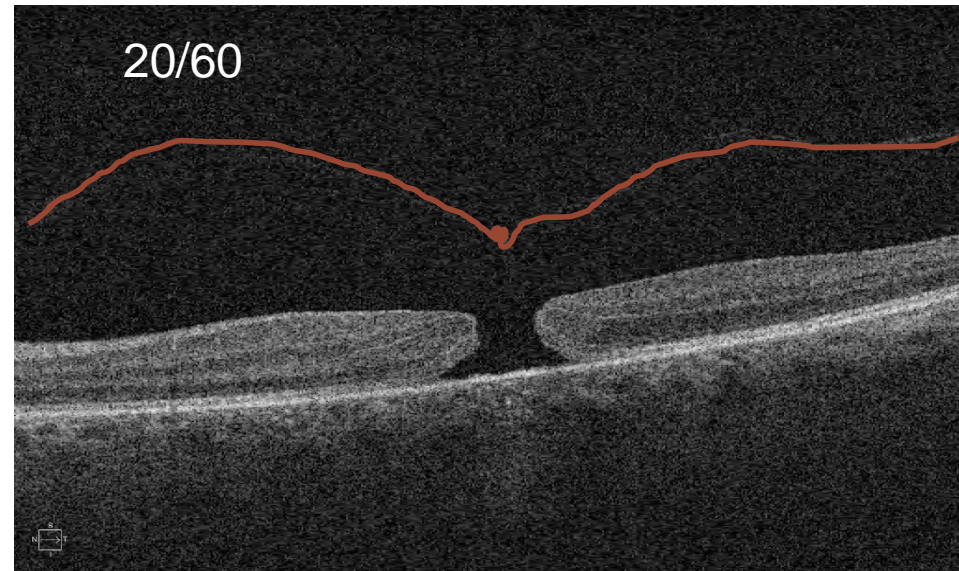
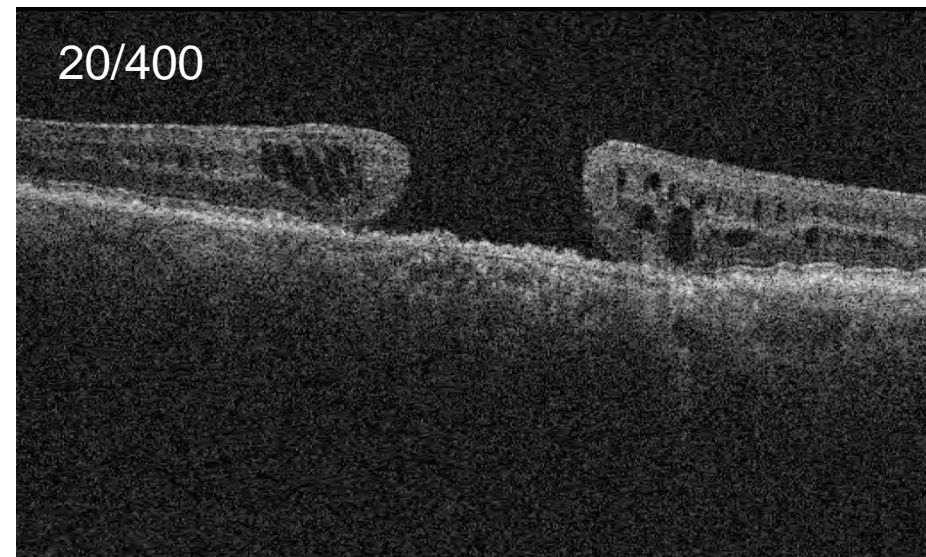
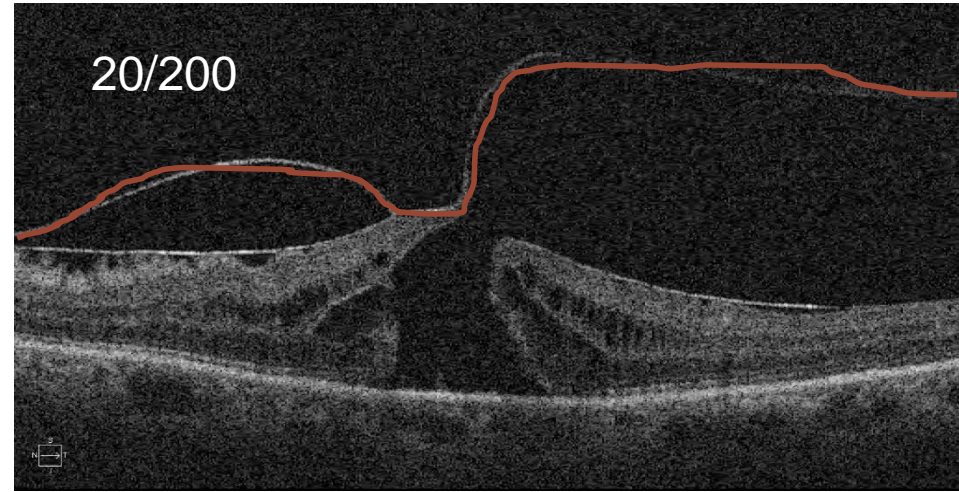
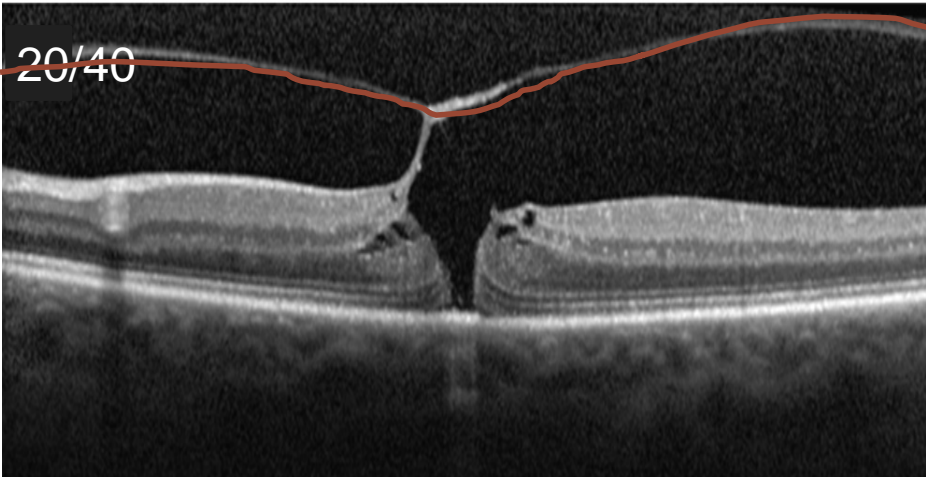


# Stage 1/VMT: 50% spontaneously resolve

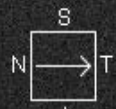
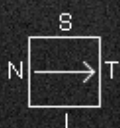
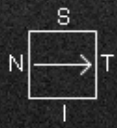
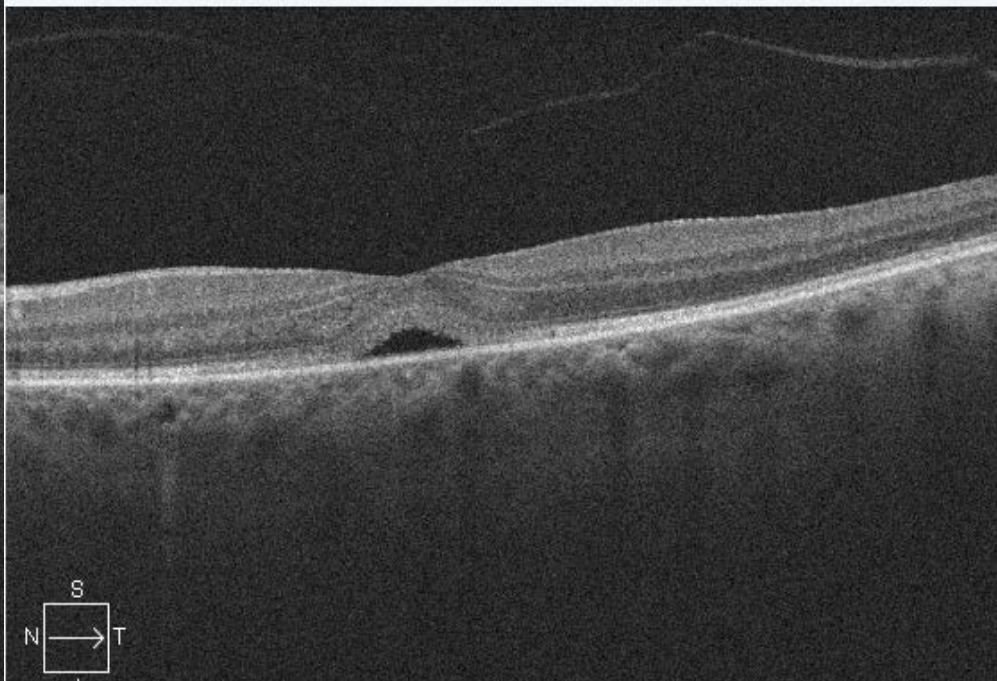
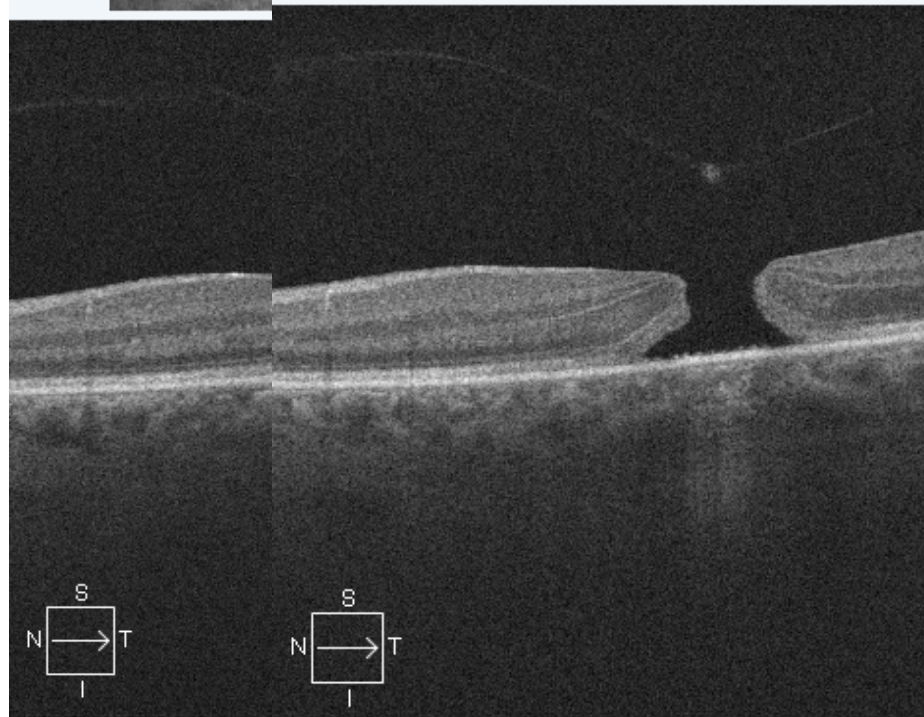
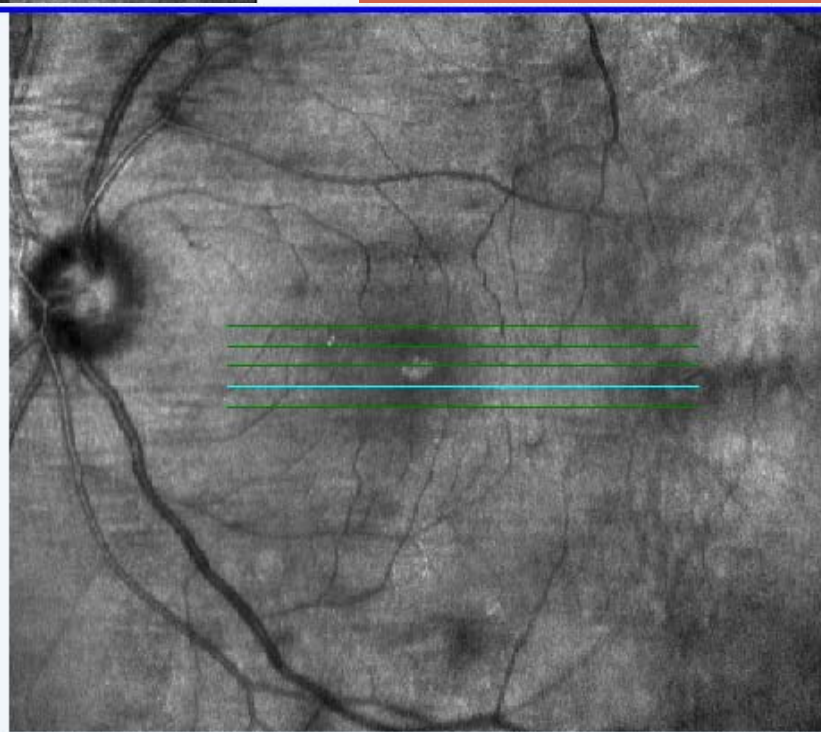
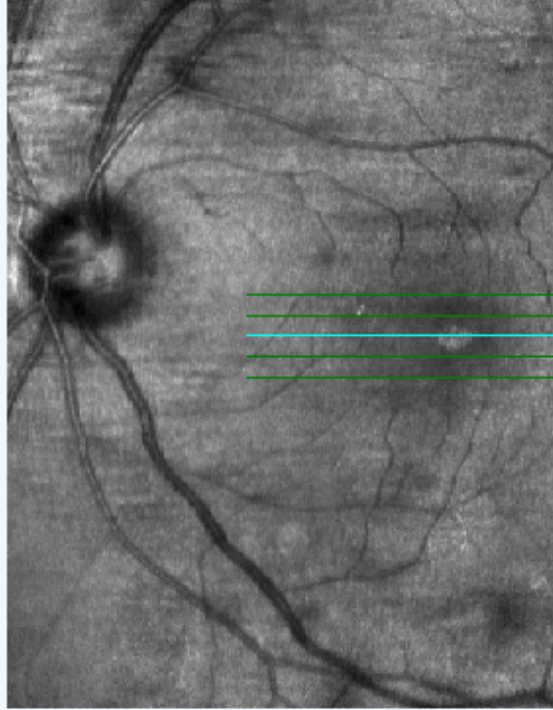




# Stages 2, 3, 4/small, med, large +/- VMT: Full Thickness





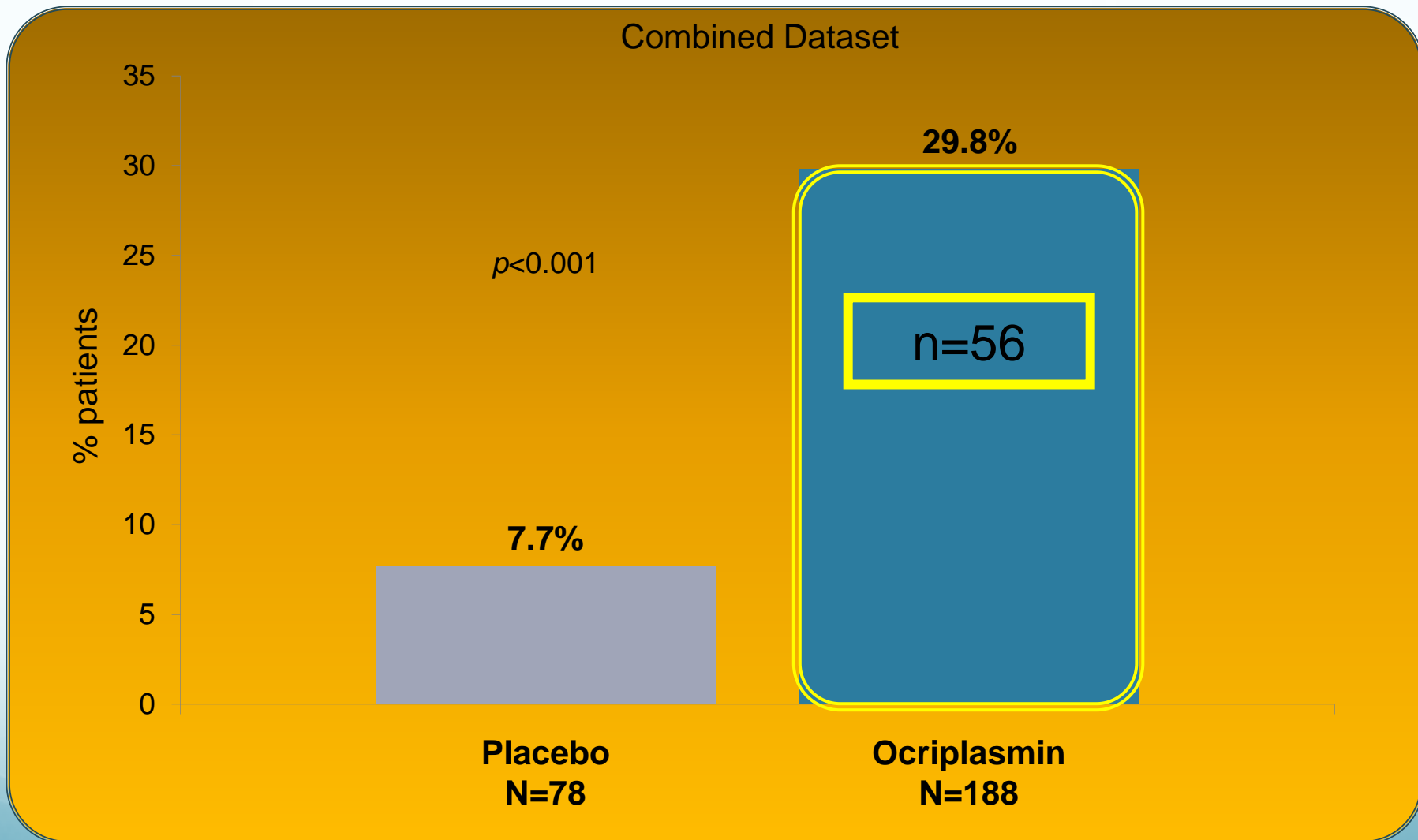


# OCT Classification of Macular Holes

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Stage 4: FTMH with PVD	Small, medium or large FTMH without VMT

“Smaller” hole = < 400um

# Proportion of VMT Patients with VMA Resolution at Day 28

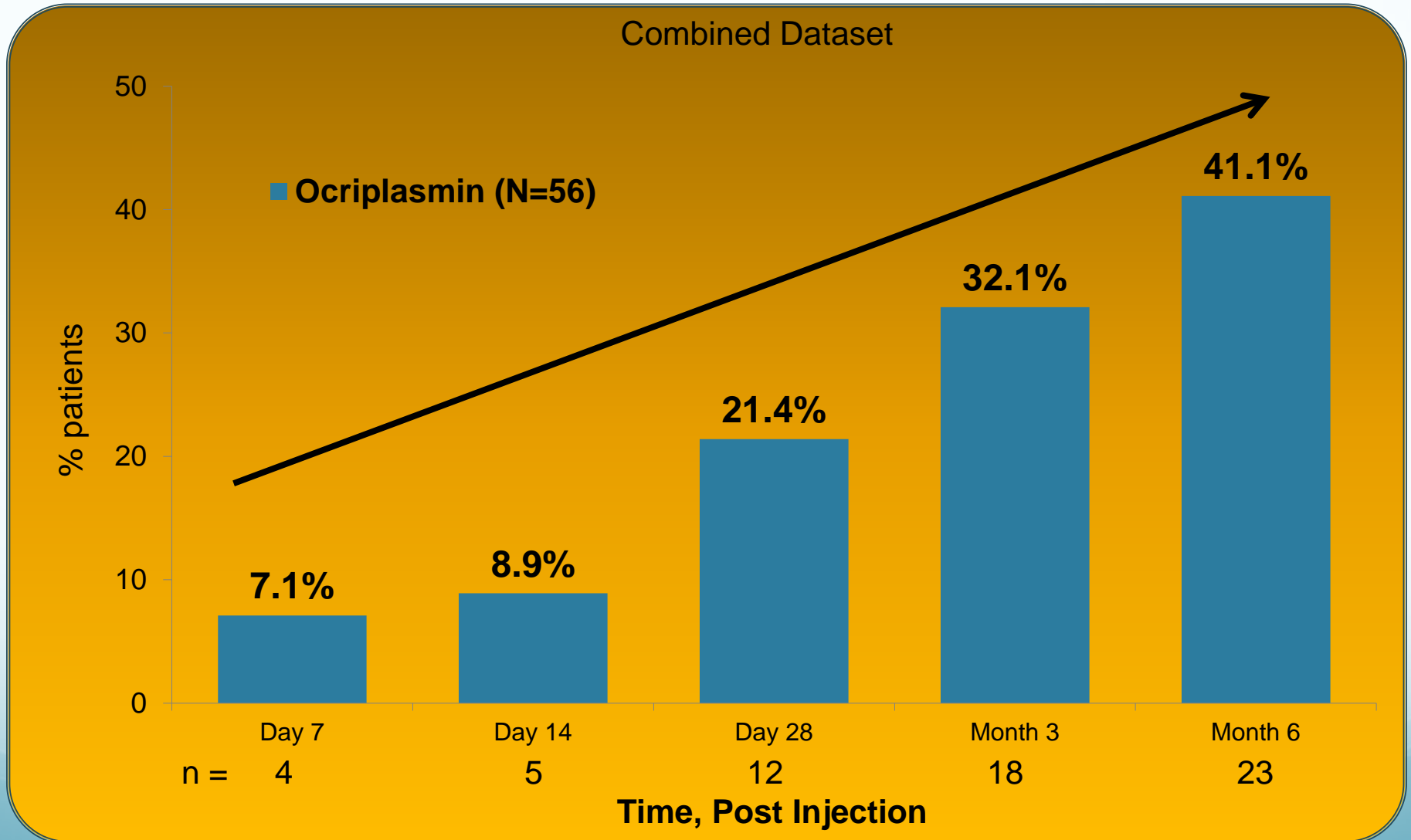


VMT: vitreomacular traction; VMA: vitreomacular adhesion.

Data on file, ThromboGenics.

# Proportion of Patients Gaining $\geq 2$ Lines VA

(Ocriclasmin-Treated VMT Patients with VMA Resolution at Day 28)

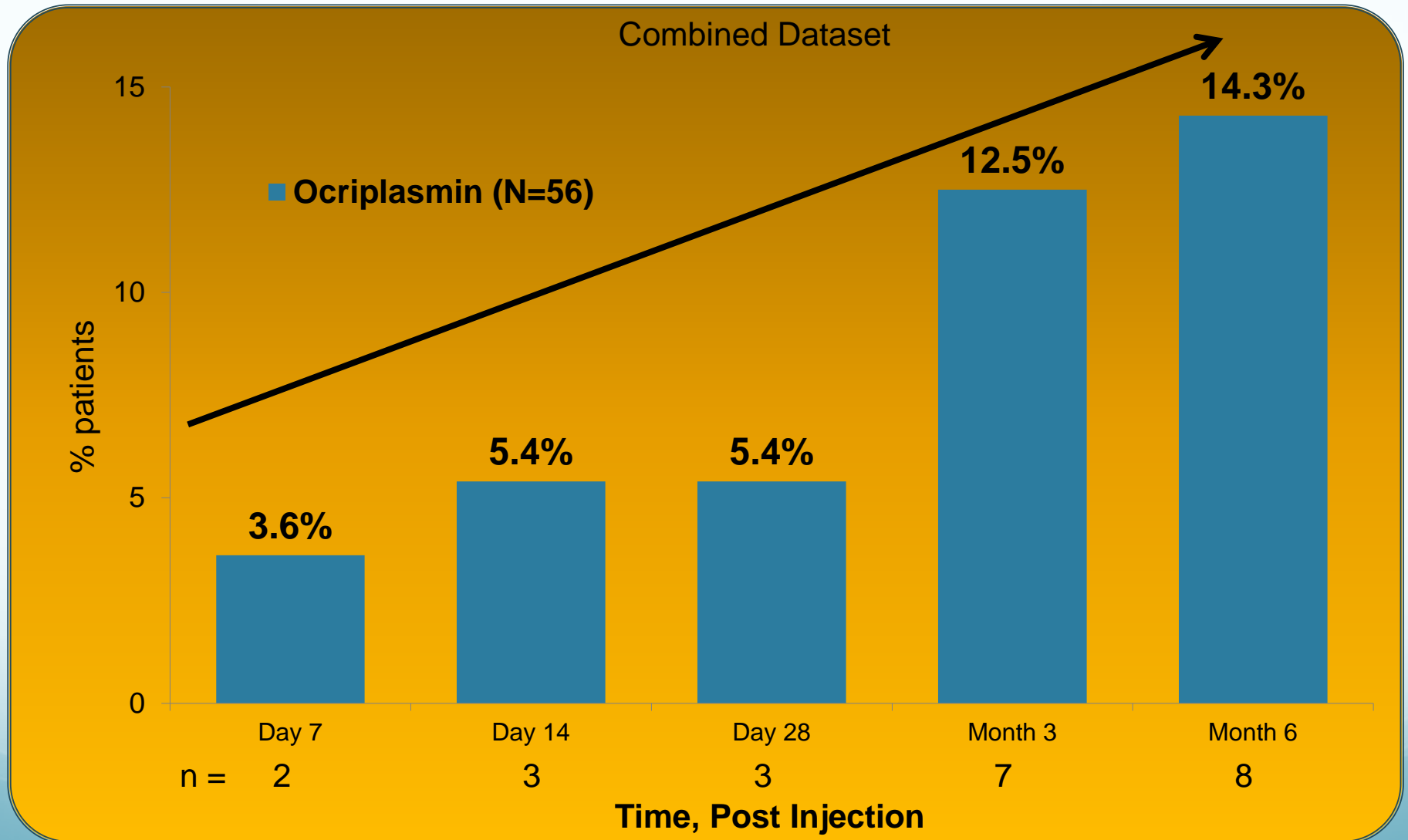


VA: visual acuity; VMT: vitreomacular traction; VMA: vitreomacular adhesion.

Data on file, ThromboGenics.

# Proportion of Patients Gaining $\geq 3$ Lines VA

(Ocriclasmin-Treated VMT Patients with VMA Resolution at Day 28)



VA: visual acuity; VMT: vitreomacular traction; VMA: vitreomacular adhesion.

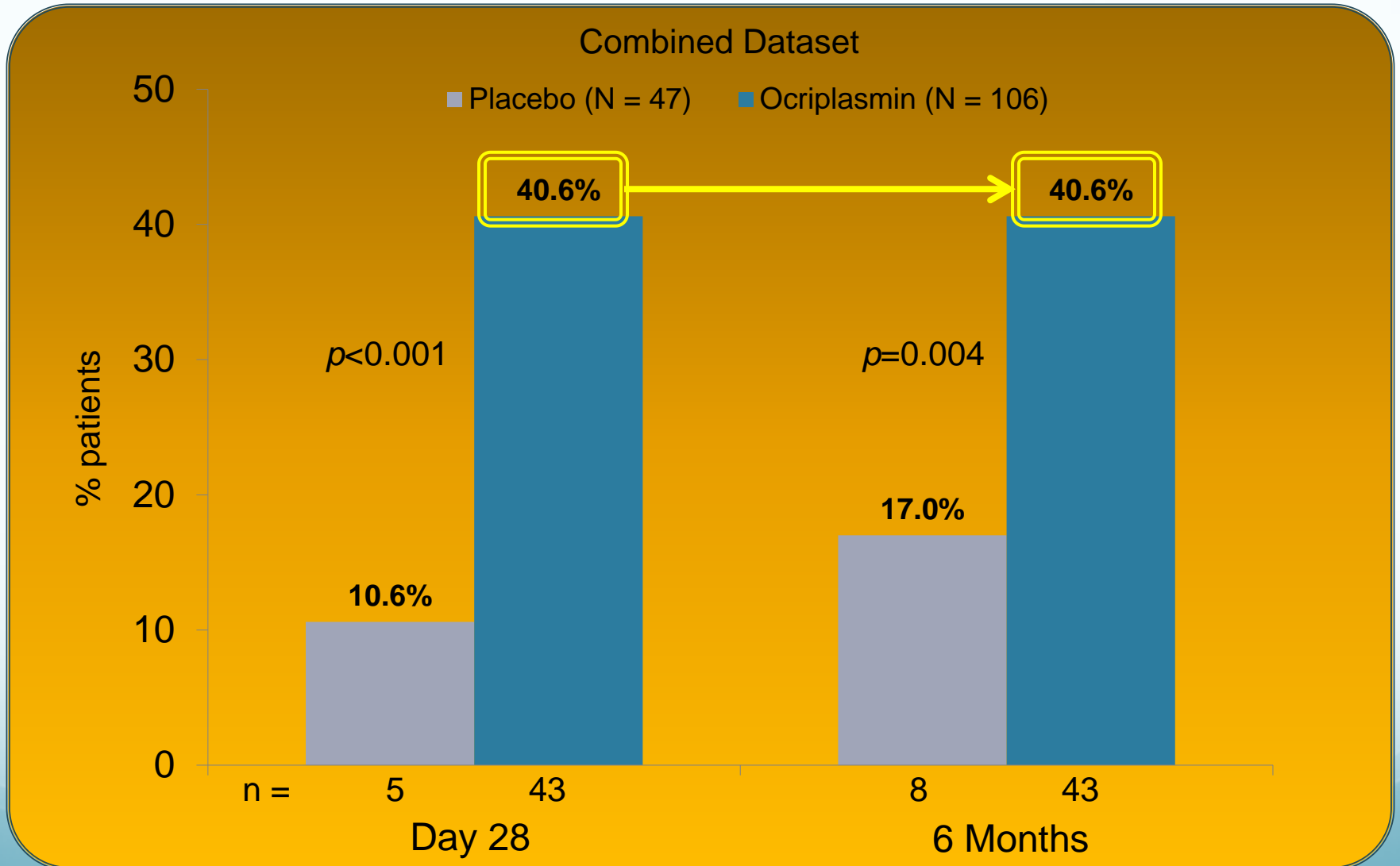
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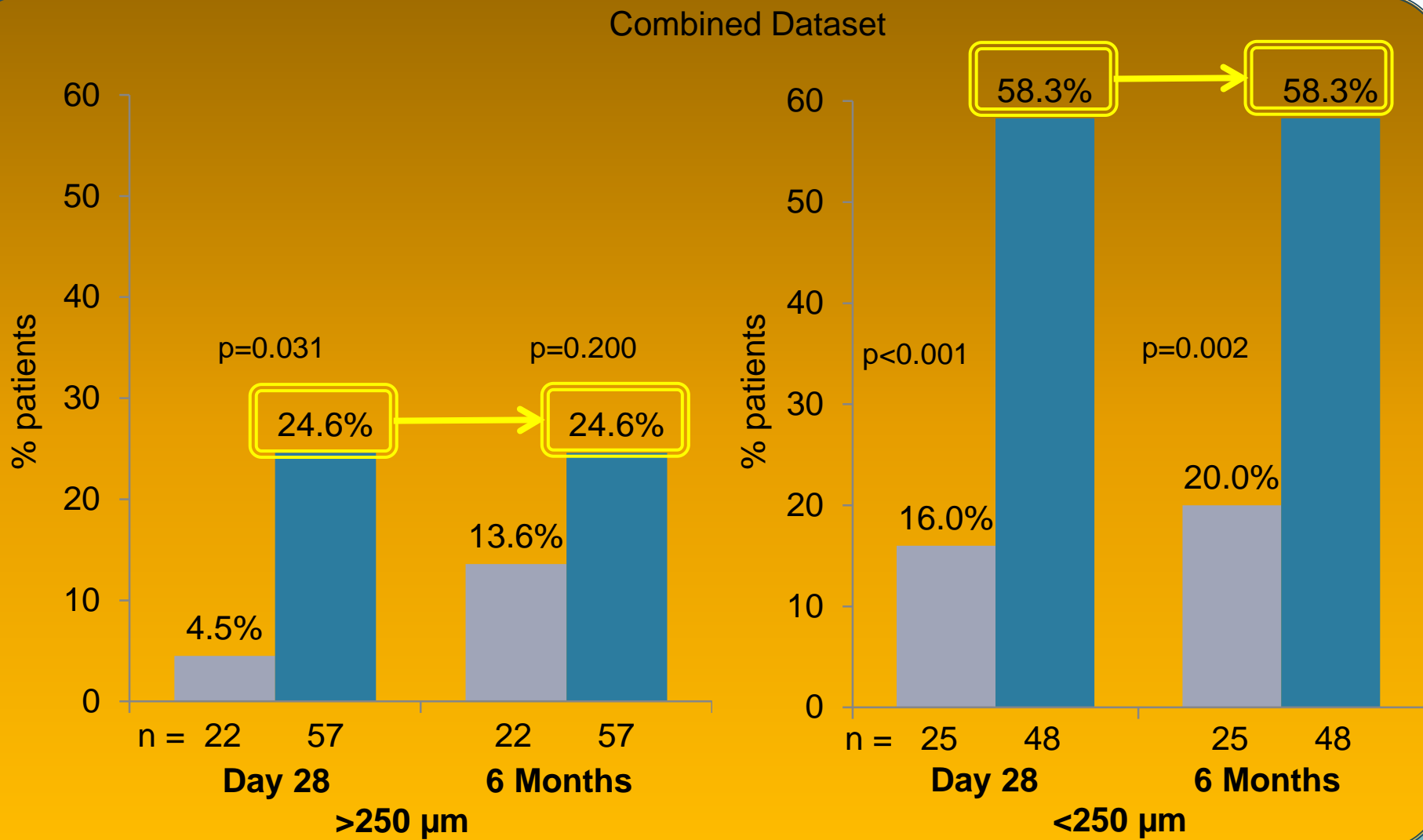
# Macular Hole Subgroup\*

## Responder Analysis

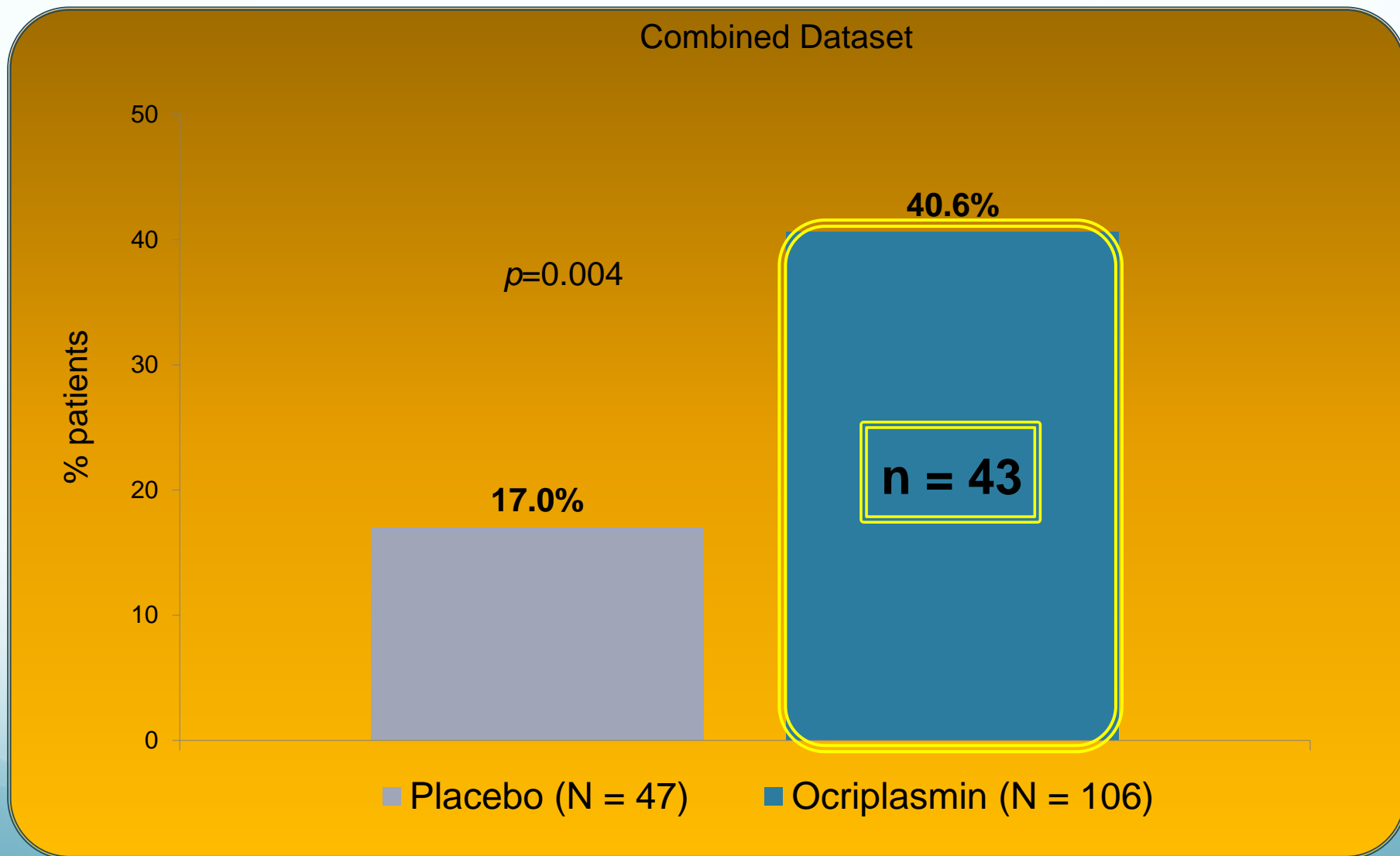
# Proportion of Patients with FTMH Closure (without vitrectomy)



# Proportion of Patients with FTMH Closure by Size at Baseline (without vitrectomy)

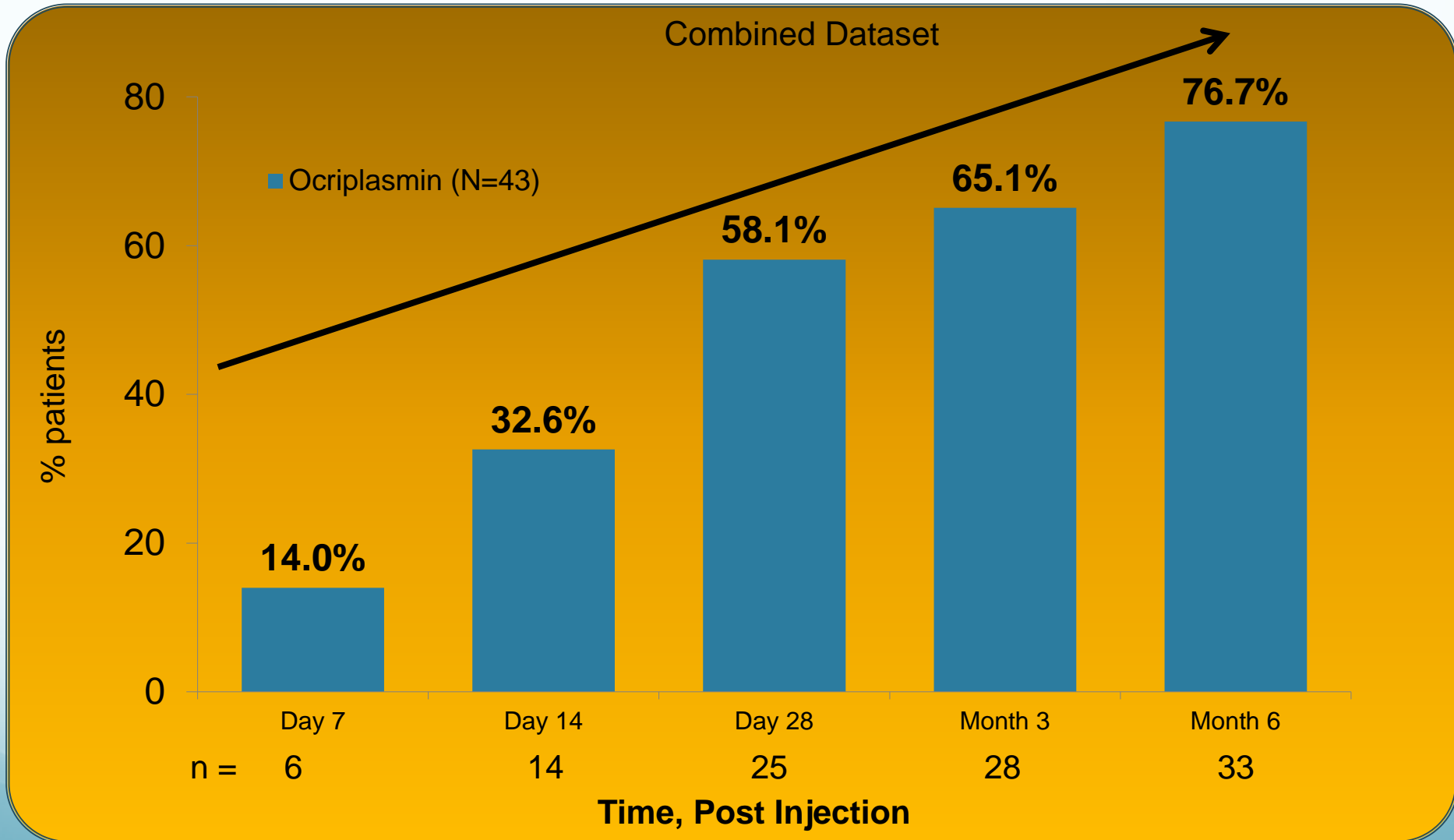


# Proportion of Patients with FTMH Closure at Month 6 (without vitrectomy)



# Proportion of Patients Gaining $\geq 2$ Lines VA

(Ocristasmin-Treated Patients with FTMH Closure at Month 6 without Vitrectomy)



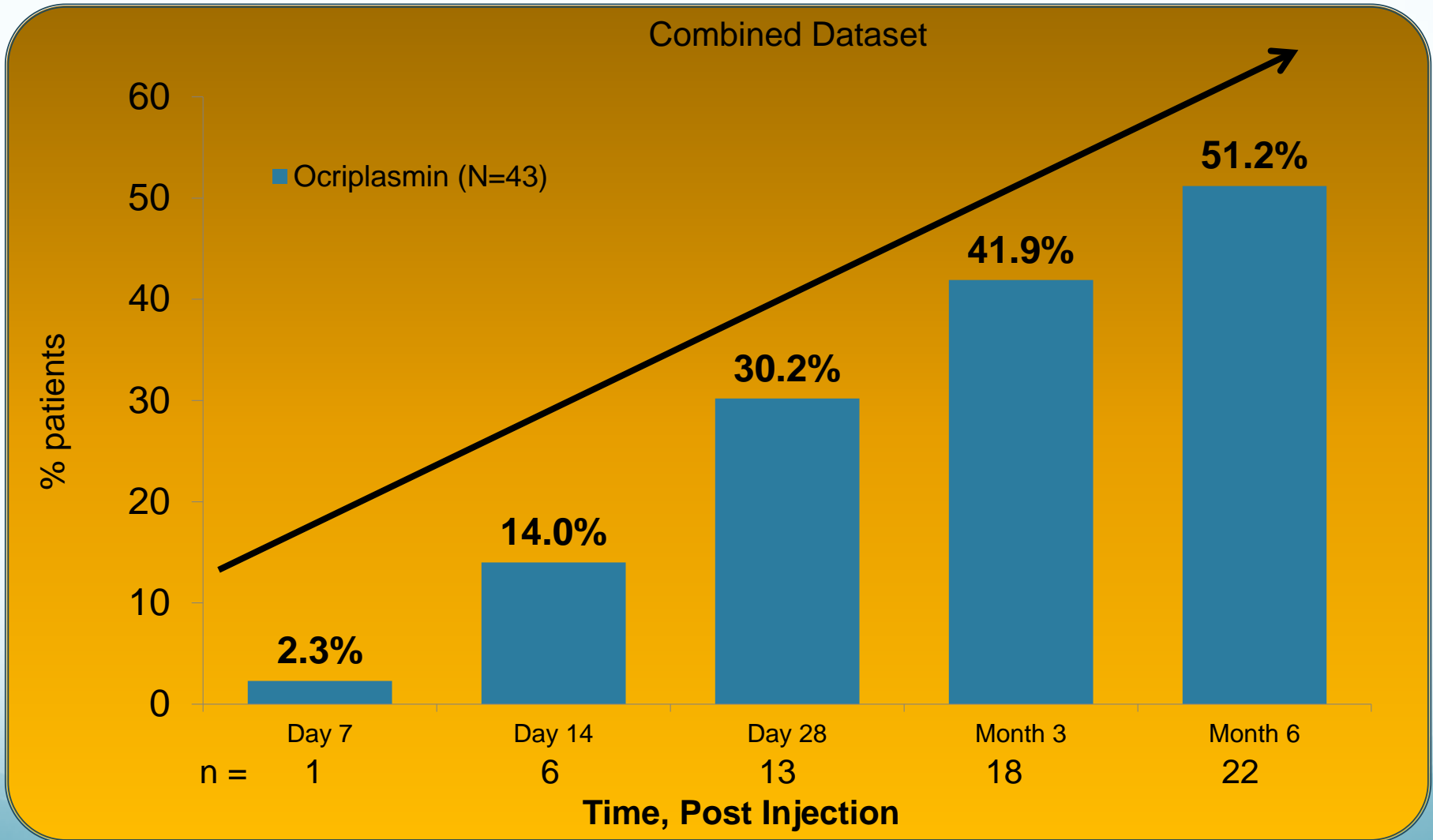
VA: visual acuity, FTMH: full thickness macular hole

Data on file, ThromboGenics



# Proportion of Patients Gaining $\geq 3$ Lines VA

(Ocriclasmin-Treated Patients with FTMH Closure at Month 6 without Vitrectomy)



VA: visual acuity; FTMH: full thickness macular hole

Data on file, ThromboGenics

# FTMH

## Vitrectomy vs. Ocriplasmin

- **Pros**

- High (>85%) success rate
- Known long-term side effects

- **Cons**

- Surgical risks
- Face down positioning(+/-)
- Cataract formation

- **Pros**

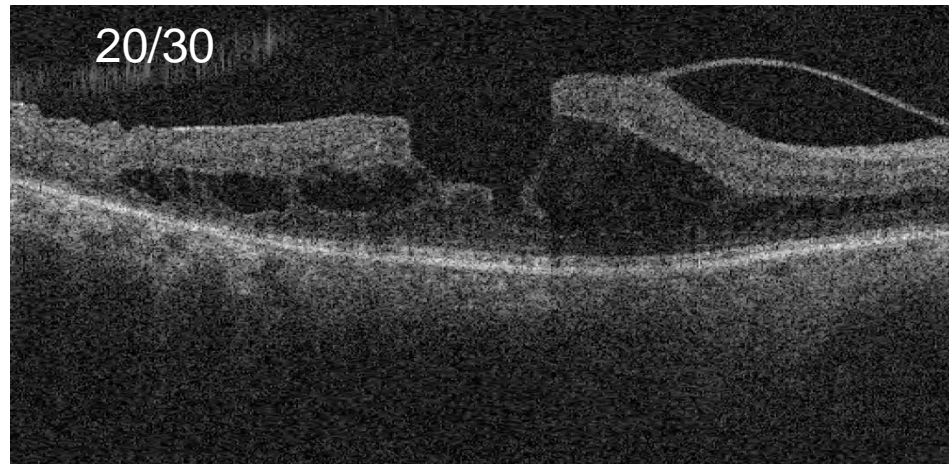
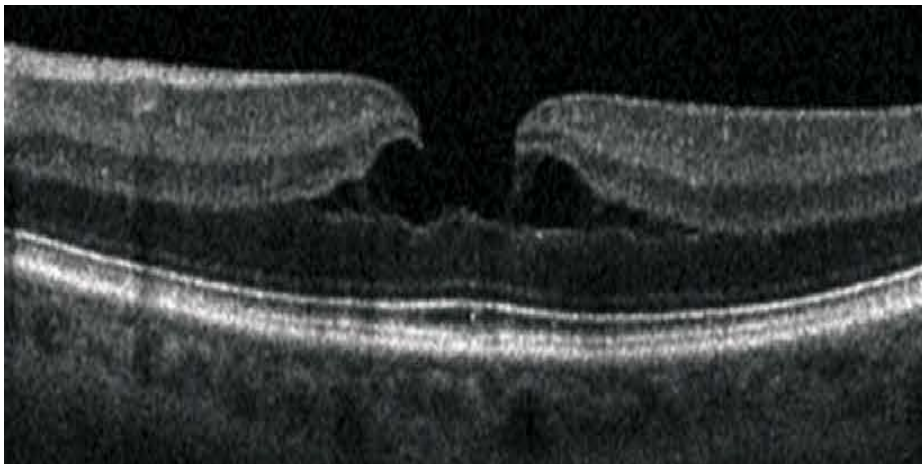
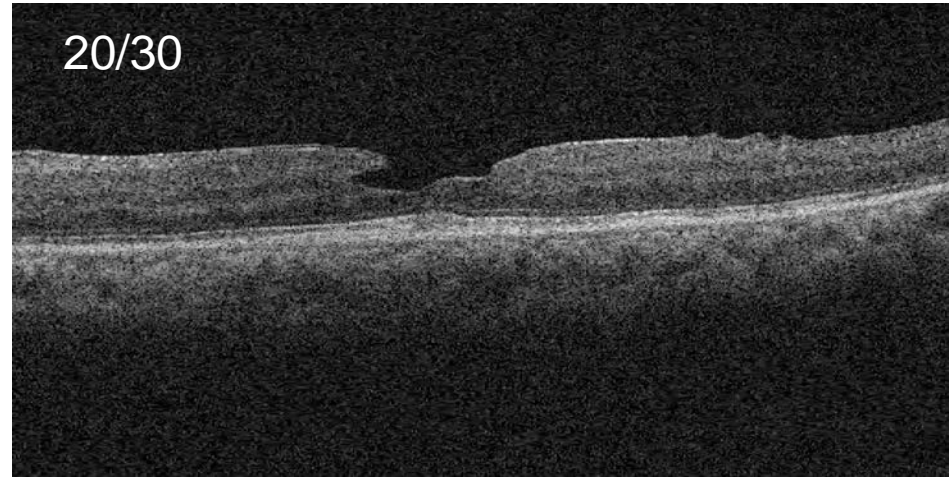
- In-office procedure
- Less \$ if successful
- Avoid surgery/earlier cataract formation if successful
- No face down positioning

- **Cons**

- Lower (~40%) success rate
- Surgery needed if fails
- Long term side effects not known
- Risks: tears, dyschromatopsias
  - Avoid in high risk RD patients

# Lamellar Holes

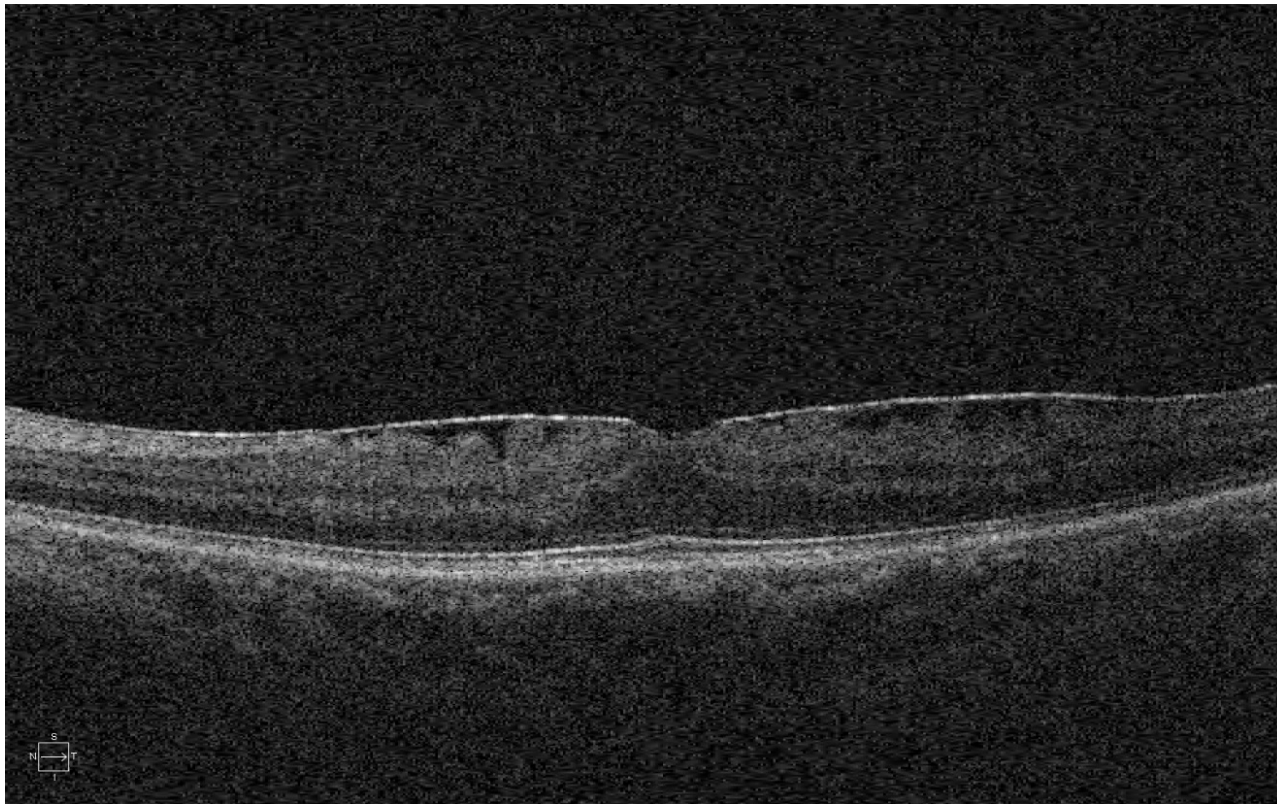
- Aborted macular hole
- “Partial thickness”
- Criteria:
  1. Irregular foveal contour
  2. Defect/break inner fovea
  3. Intra-retinal split (schisis) – can be variable
  4. Intact photoreceptors



# Macular Pseudohole

- Criteria

1. ERM
2. No loss of tissue
3. Heaped foveal edges
4. Near normal foveal thickness

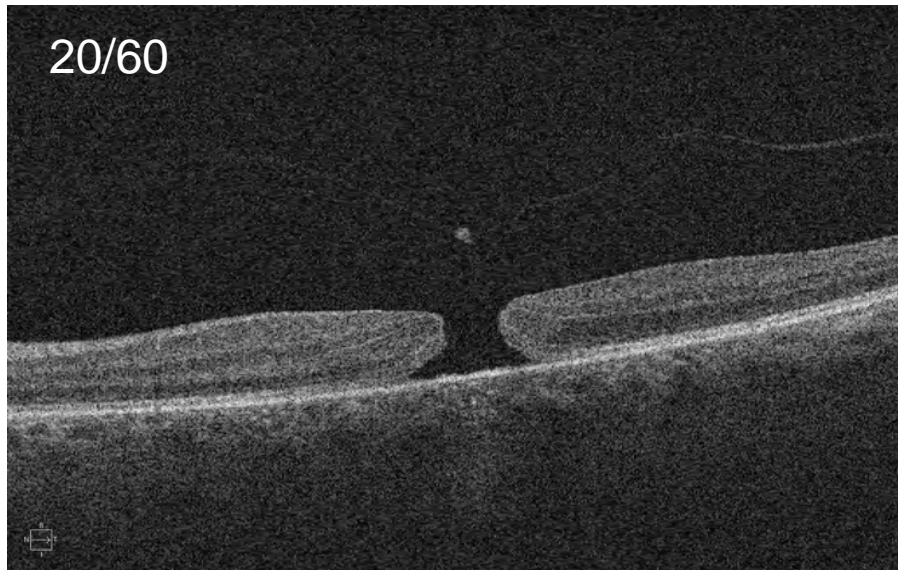




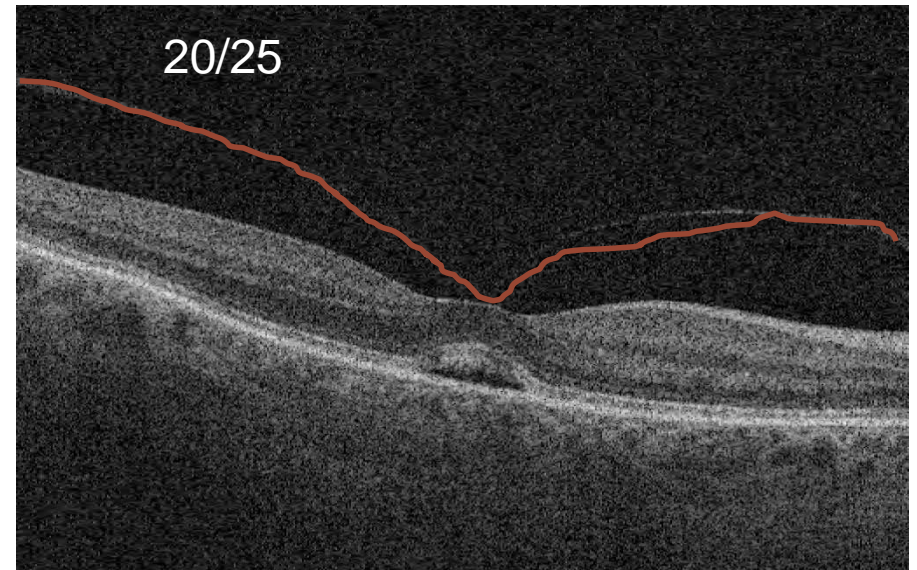
# Stage 0/Impending Macular Hole

- FTMH one eye + VMA/VMT in fellow eye
- Fellow eye at increased risk for FTMH

OD: FTMH

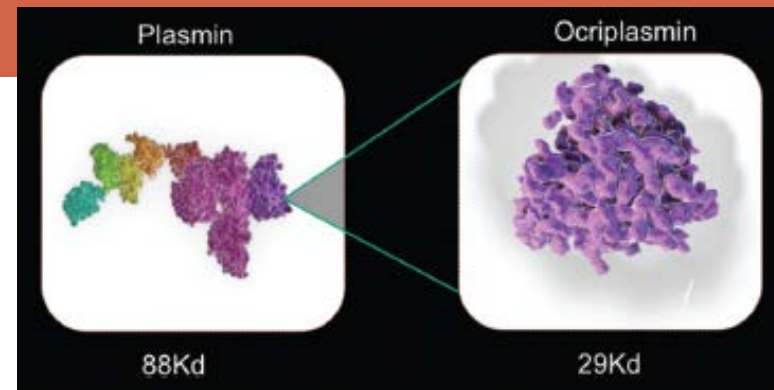


OS: Impending mac hole





# Ocriplasmin (Jetrea)



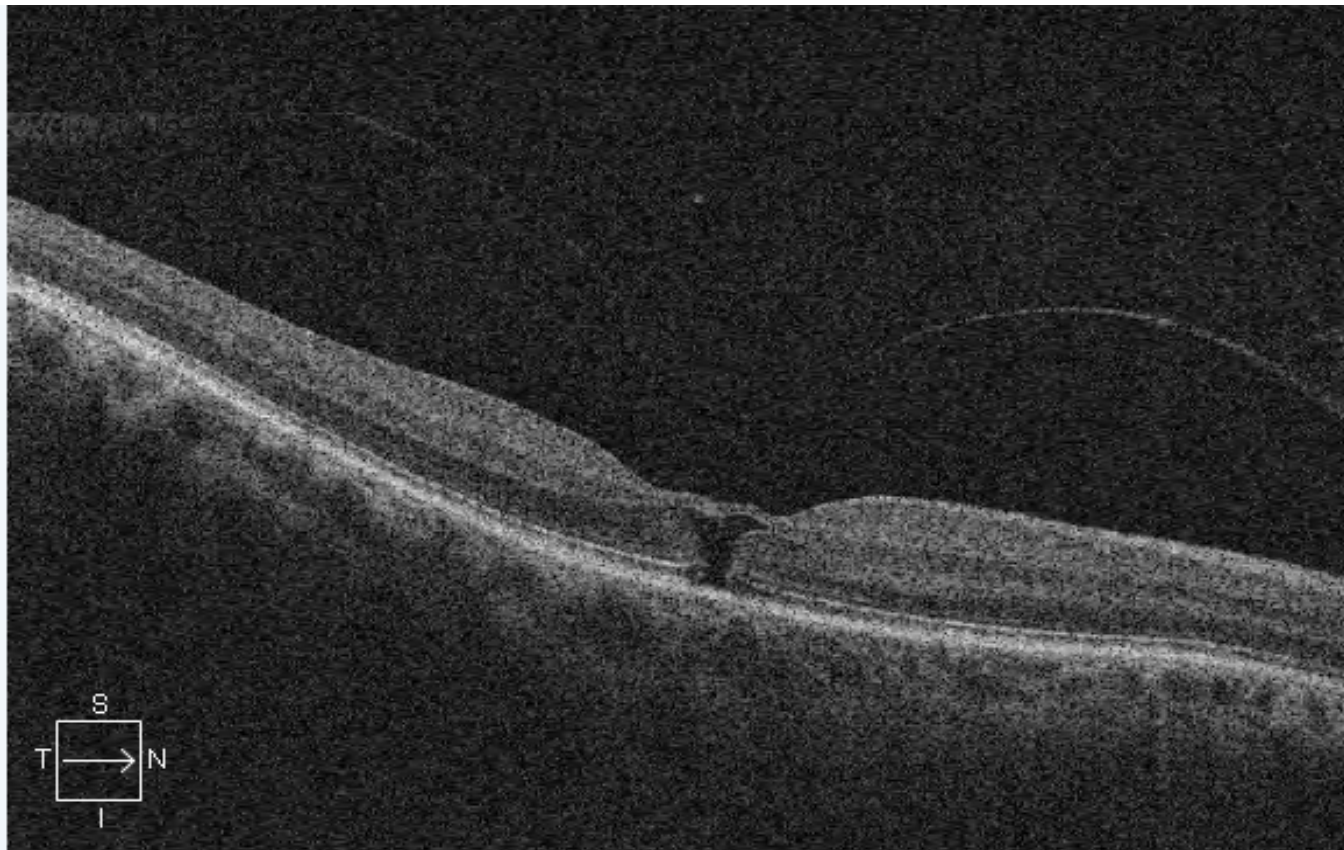
- “Symptomatic Vitreo-macular Adhesion”
- FDA Approved 2012
- MIVI-TRUST Trials
  - FTMH closure (~40%) vs. placebo (~10%)
  - VMT resolution (~29%) vs. placebo (~8%)
  - Smaller holes/more focal adhesion did better
  - Can’t have ERM
- Recommend treatment for:
  - Symptomatic VMT or smaller macular holes (<400 um) with VMT

# Back to our patient: Case #1: 46 y.o. AAF

BCVA: OD 20/30

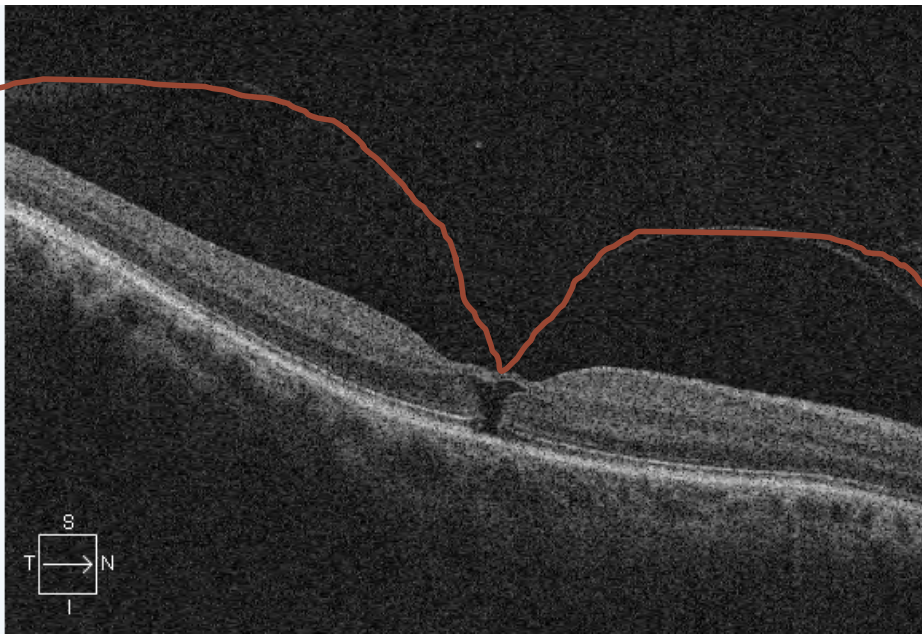
OS 20/20

“Severe” visual distortion OD x 2 weeks

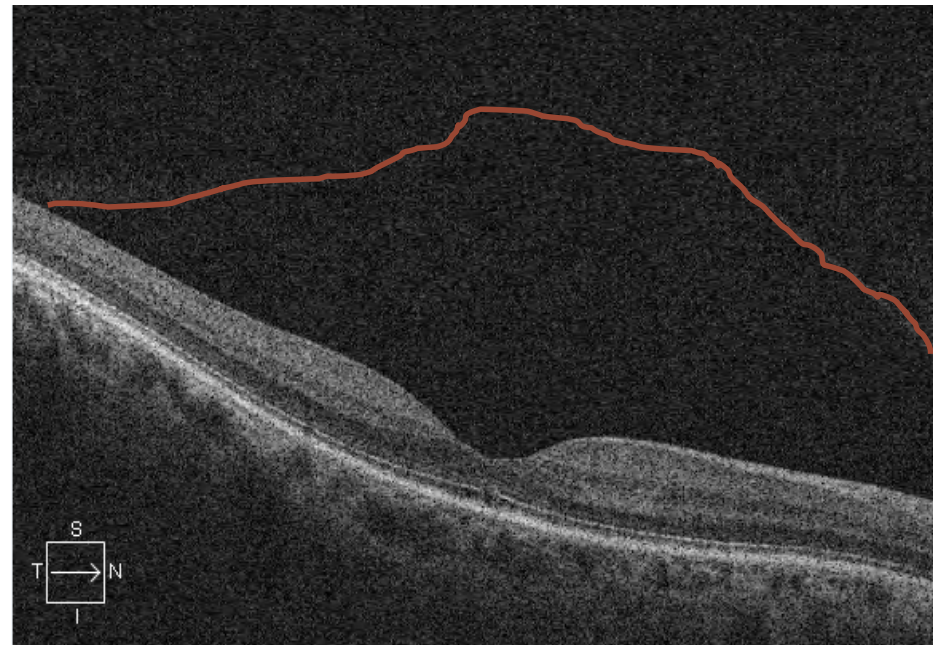


# Back to our patient...

OD: 8/2/2014 20/30



OD: 9/5/2014 20/25

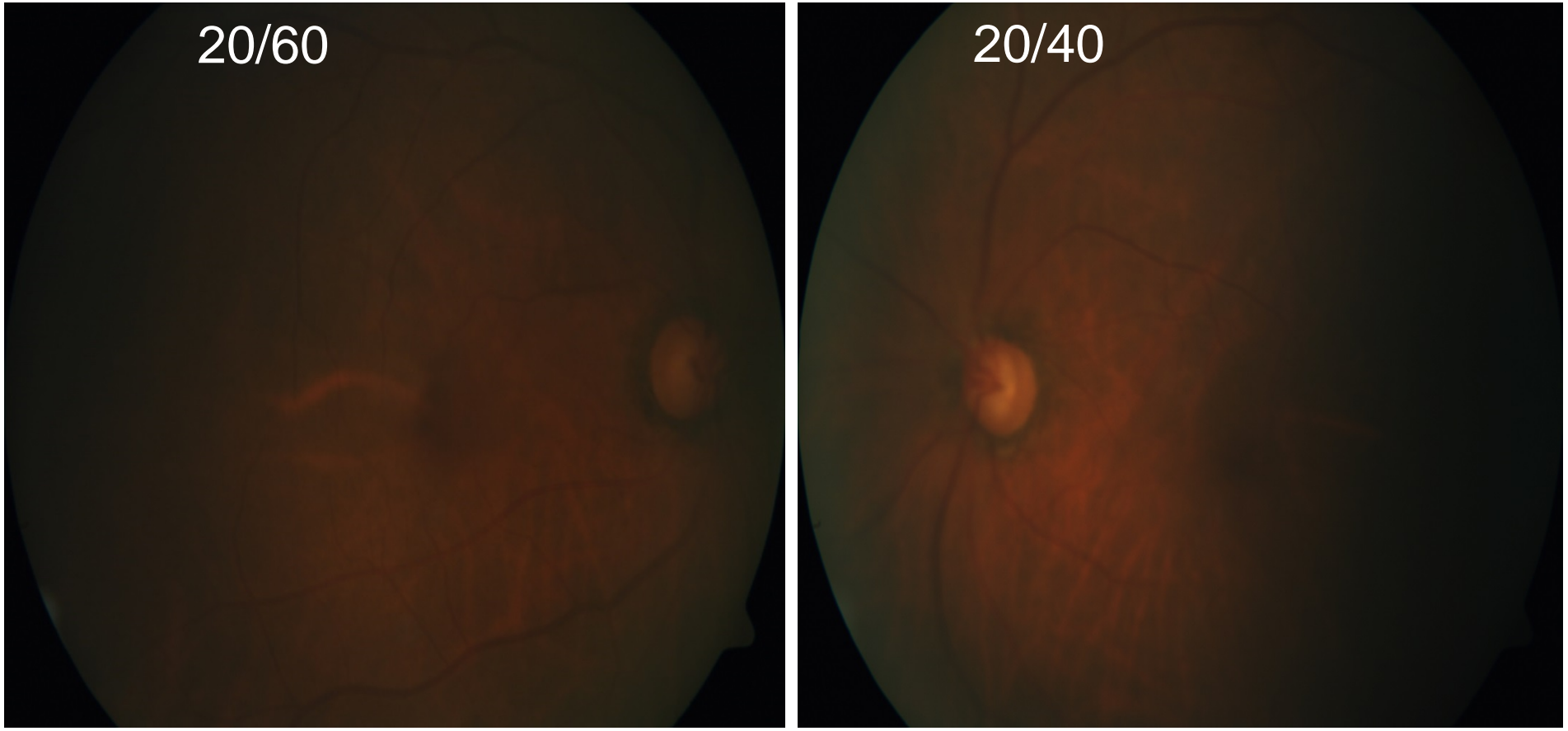


*She chose: MONITOR*

# Case #2: 80 y.o. AAF Requesting Cataract Extraction

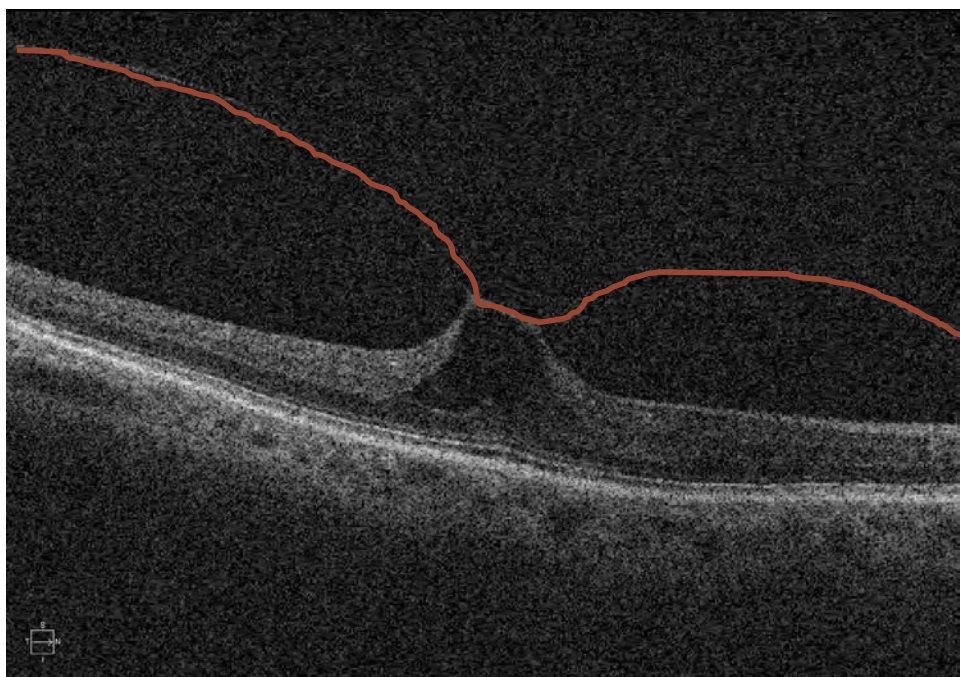
20/60

20/40

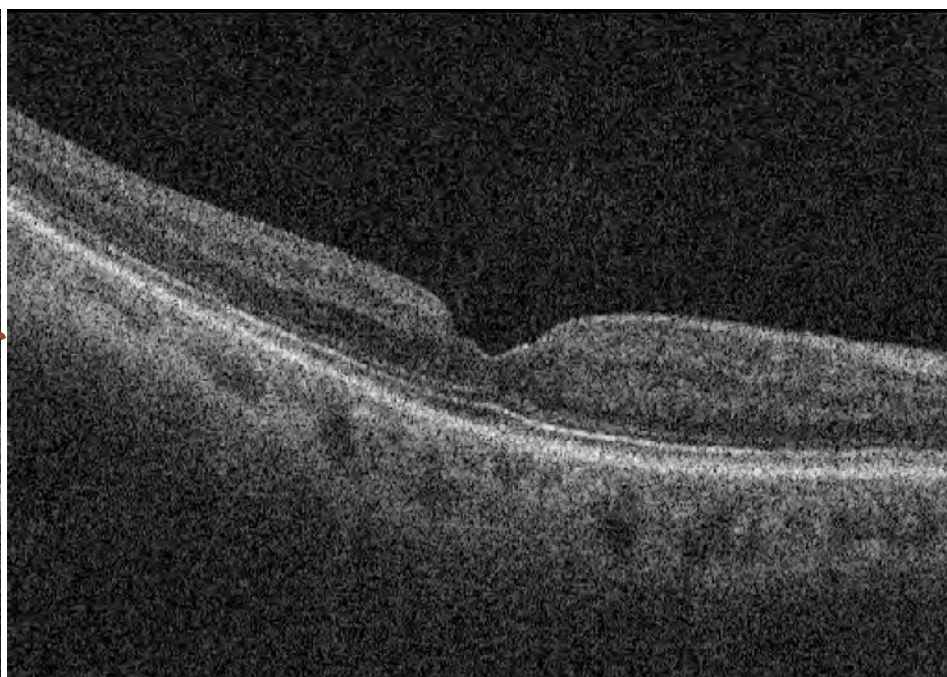




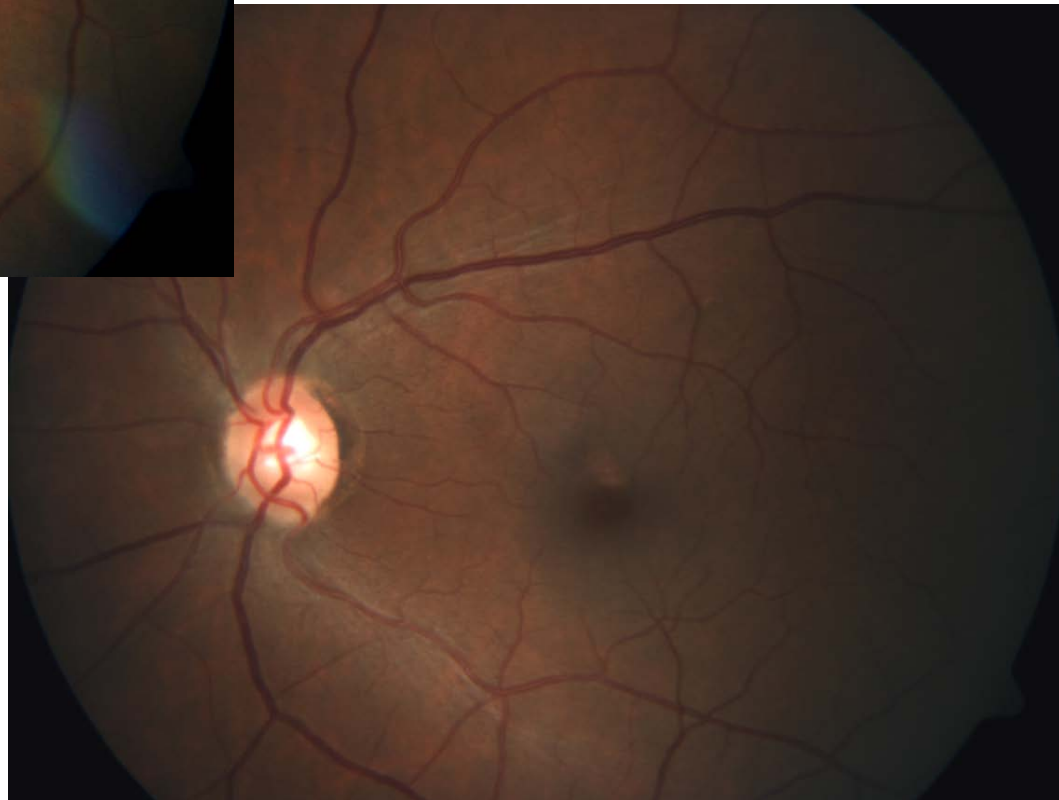
OD



OS

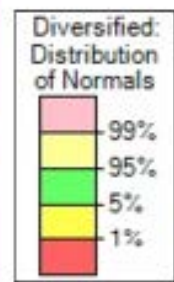
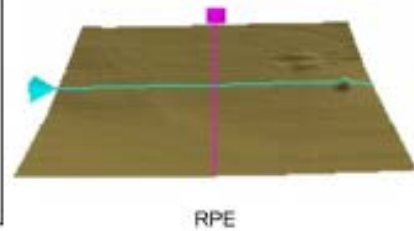
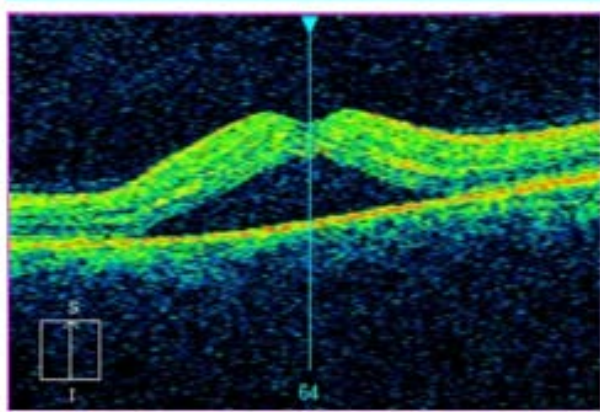
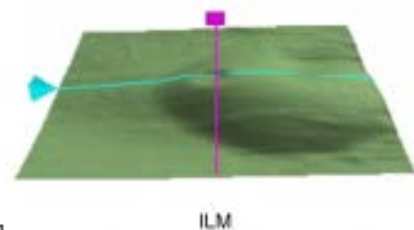
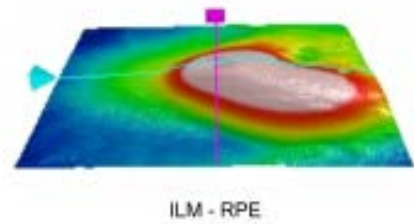
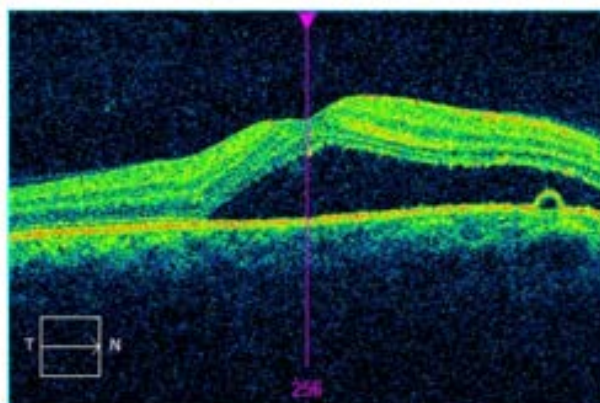
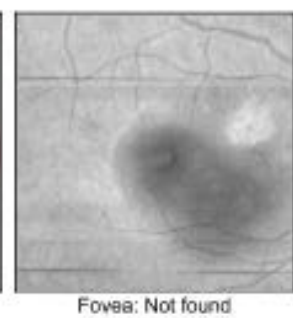
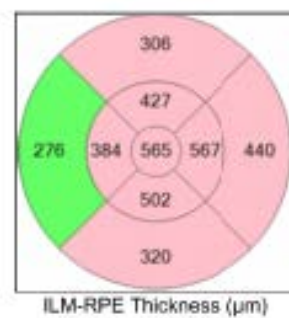
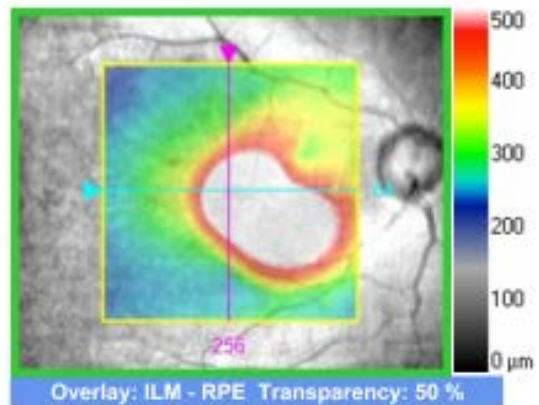



# Case #3



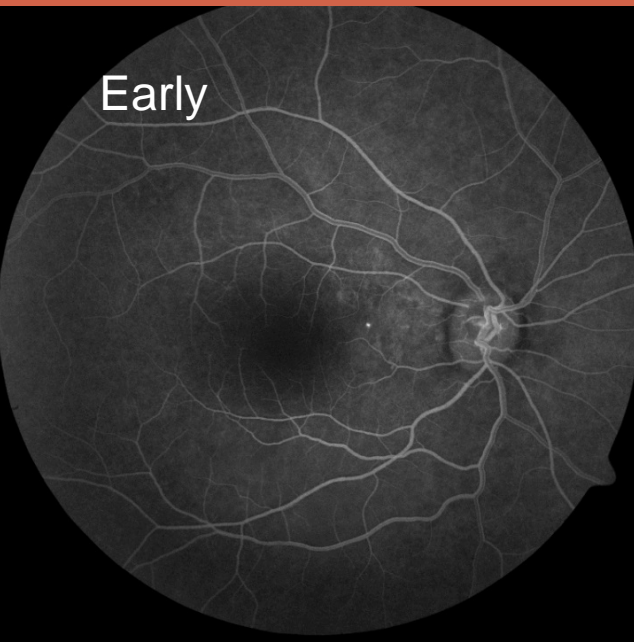
31 y.o. attorney decreased vision  
X 3 weeks

VA 20/50 OD  
20/20 OS



	Central Subfield Thickness ( $\mu\text{m}$ )	Cube Volume ( $\text{mm}^3$ )	Cube Average Thickness ( $\mu\text{m}$ )
ILM - RPE	565	12.7	351







# Question 1

- A 31 y/o attorney complains of decreased vision OD ( $VA_{cc}$  20/50) for 3 weeks. Examination and imaging findings are most consistent with central serous chorioretinopathy. Select the most appropriate course at this time:
  - a) Inquire about steroid-containing medication use and observe
  - b) Perform fluorescein-angiography-guided thermal laser photocoagulation
  - c) Perform fluorescein-angiography-guided photodynamic therapy
  - d) Initiate oral therapy with mifepristone
  - e) Recommend lifestyle modification...consider career change

# Central Serous Chorioretinopathy

- Serous macular detachment
- ‘Idiopathic’
- ♂ >> ♀ 9:1
- Possible hyperopic shift
- Risk factors: stress ( ‘type-A personality’ )
- Exogenous/endogenous steroids

# Central Serous Chorioretinopathy

- May have serous RPE detachment (PED)
- CSCCR characteristic fluorescein angiography: expansile pinpoint leakage
- Smoke stack configuration in 5-10%
- Multifocal patterns
  - More common in medication-induced or systemic disease

# Central Serous Chorioretinopathy

- 90% of patients exhibit spontaneous resorption of subretinal fluid and recover ‘good’ vision without treatment



- Complaints of metamorphopsia and objective defects in contrast sensitivity are common
- ~ 1/3 will experience recurrence
- Laser speeds recovery, does not change visual outcome



## CSCR – Therapeutic Interventions

- Thermal laser photocoagulation
  - Speeds recovery..does not change visual outcome
- Photodynamic therapy with verteporfin (PDT)
- Subthreshold diode micropulse photocoagulation
- Intravitreal bevacizumab (Avastin)
- Mifepristone

## CSCR – Therapeutic Interventions

- Br J Ophthalmol. 2010 Jul 19. Lim JW et al.
- **Comparative study of patients with central serous chorioretinopathy undergoing focal laser photocoagulation or photodynamic therapy**
- Compared with focal laser, half-dose PDT may facilitate earlier resolution of macular detachment and earlier recovery of central retinal function
- However, at 3 months after treatment and thereafter, no difference in anatomical and functional recovery was noted between the two modalities of treatment

# CSCR – Therapeutic Interventions

- Ophthalmology. 2008 Oct;115(10):1756-65.
- Chan WM, Lai TY, Lai RY, Liu DT, Lam DS.
- **Half-dose verteporfin photodynamic therapy for acute central serous chorioretinopathy: one-year results of a randomized controlled trial**
- Thirty-seven (94.9%) eyes in the verteporfin group compared with 11 (57.9%) eyes in the placebo group showed absence of subretinal fluid at the macula at 12 months (P = 0.001)
- **CONCLUSIONS: Photodynamic therapy with half-dose verteporfin is effective in treating acute symptomatic CSC, resulting in a higher proportion of patients with absence of exudative macular detachment and better visual acuity compared with placebo.**

# Question 1

- A 31 y/o attorney complains of decreased vision OD (VA<sub>cc</sub> 20/50) for 3 weeks. Examination and imaging findings are most consistent with central serous chorioretinopathy. Select the most appropriate course at this time:
  - a) **Inquire about steroid-containing medication use and observe**
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# Initial presentation

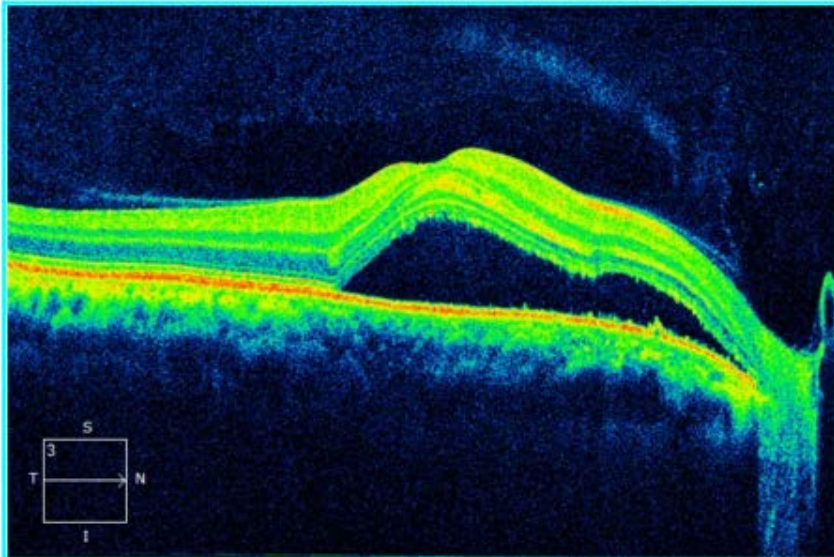
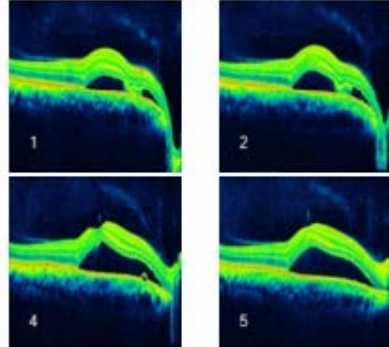
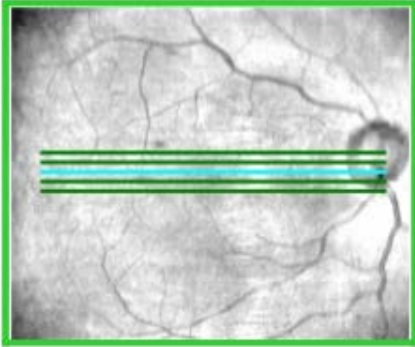
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OD  OS

Scan Angle: 0°

Spacing: 0.25 mm

Length: 9 mm



# 1 month

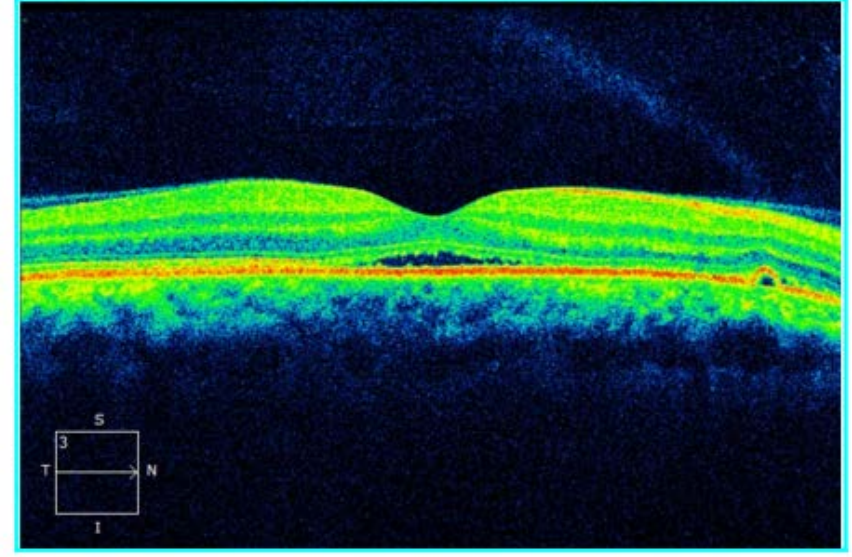
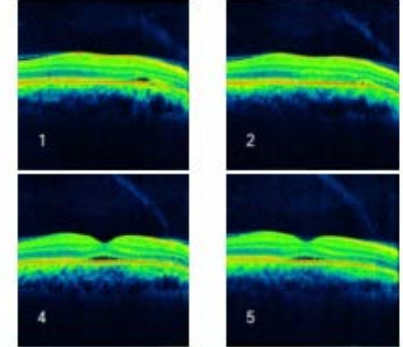
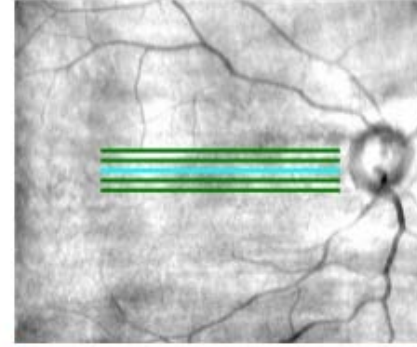
High Definition Images: HD 5 Line Raster

OD  OS

Scan Angle: 0°

Spacing: 0.25 mm

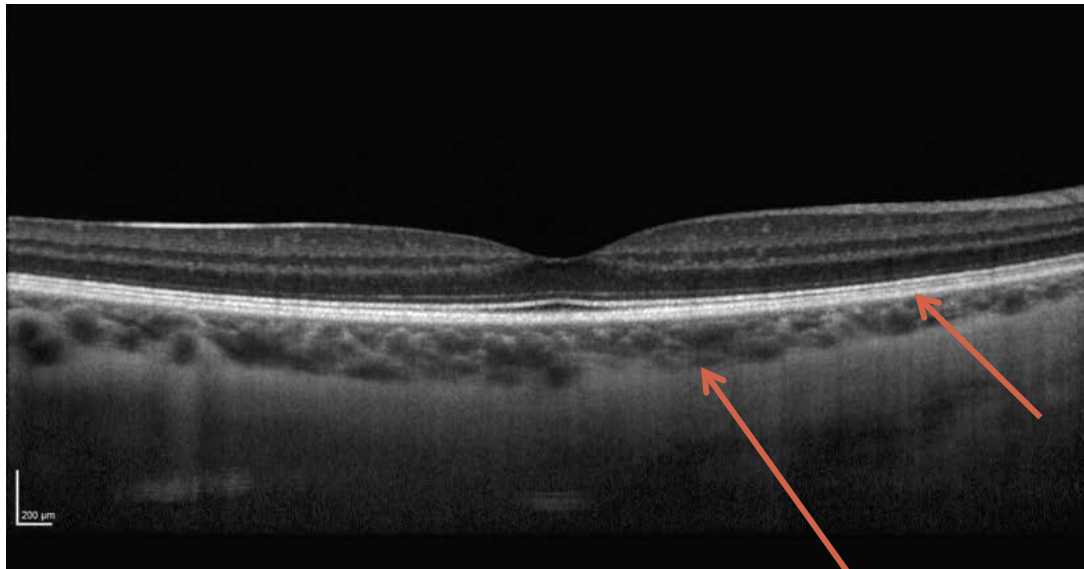
Length: 6 mm



# Enhanced Depth Imaging

- Increased visualization of choroidal anatomy

Normal

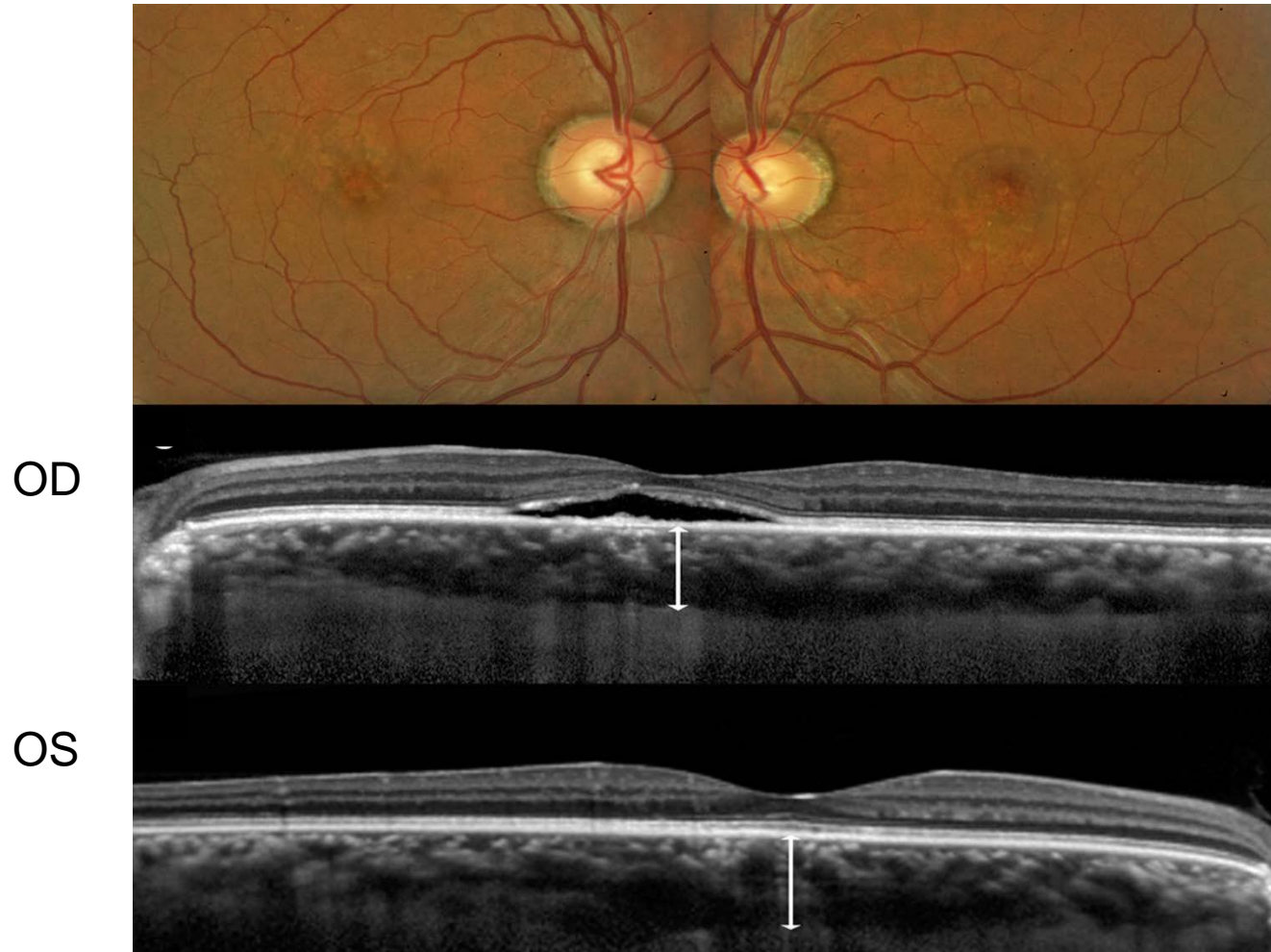


Outer border of RPE/  
Bruch's complex

Choroid/sclera junction

# Enhanced Depth Imaging CSCR

- Choroidal hyperpermeability/thickening in BOTH eyes



# Central Serous Chorio-Retinopathy (CSCR)

- Treatments
  - Observation
  - Laser photocoagulation
  - PDT
- Referral if:
  - 3-4 months duration
  - Recurrent with reduced VA
  - VA other eye reduced from CSCR
  - Vocation/visual needs – willing to take risk
    - permanent scotoma
  - CNV



# Case #4: 73 y.o. AAF

OD: 20/25



OS: 20/25



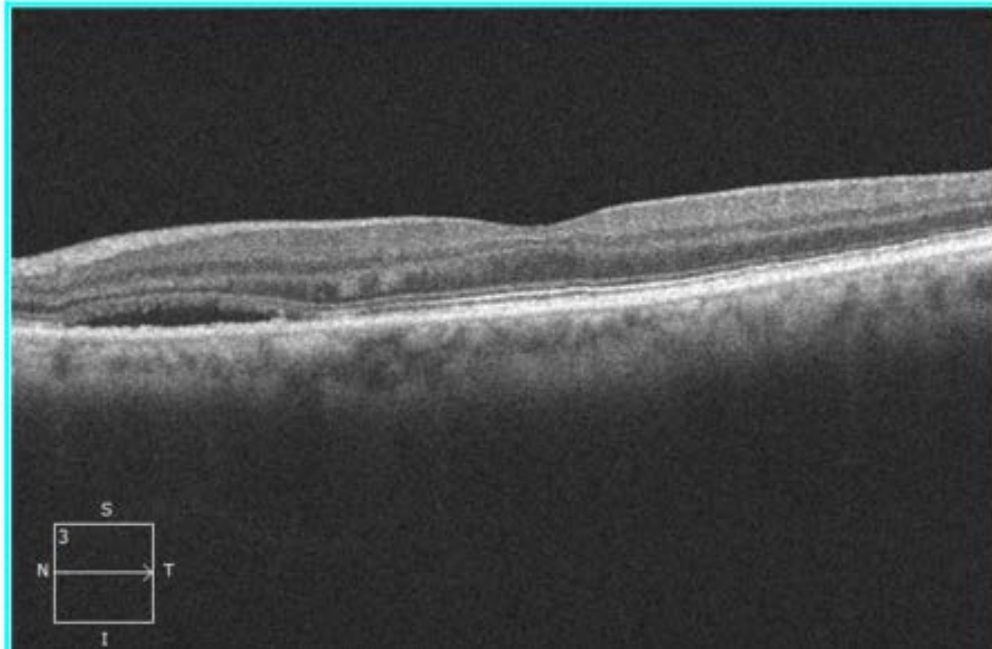
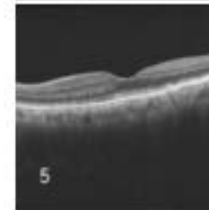
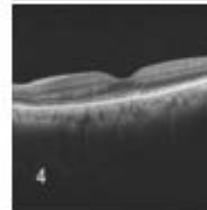
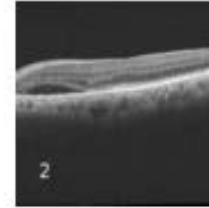
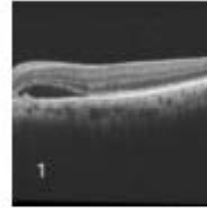
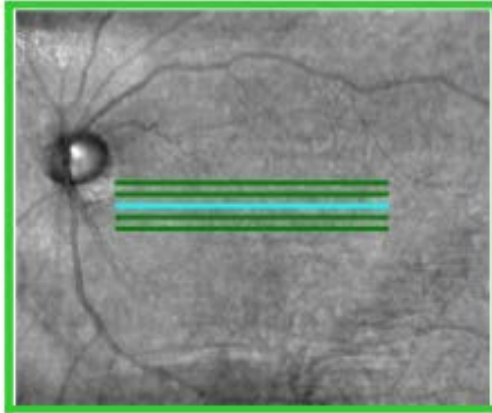
# High Definition Images: HD 5 Line Raster

OD  OS

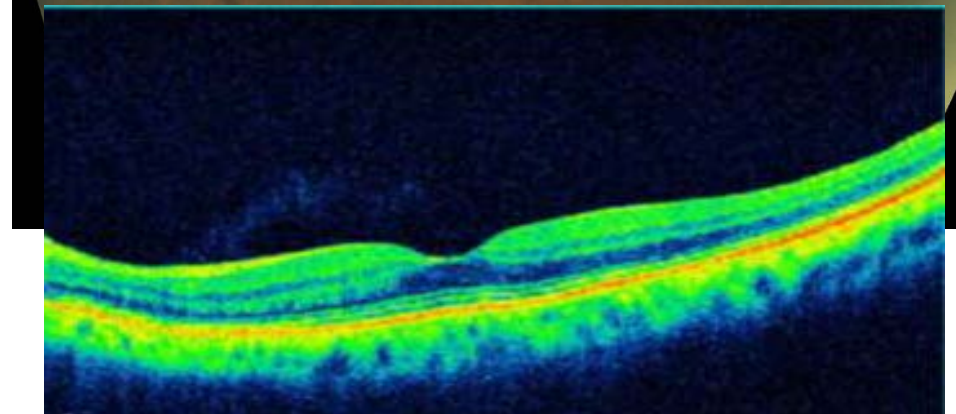
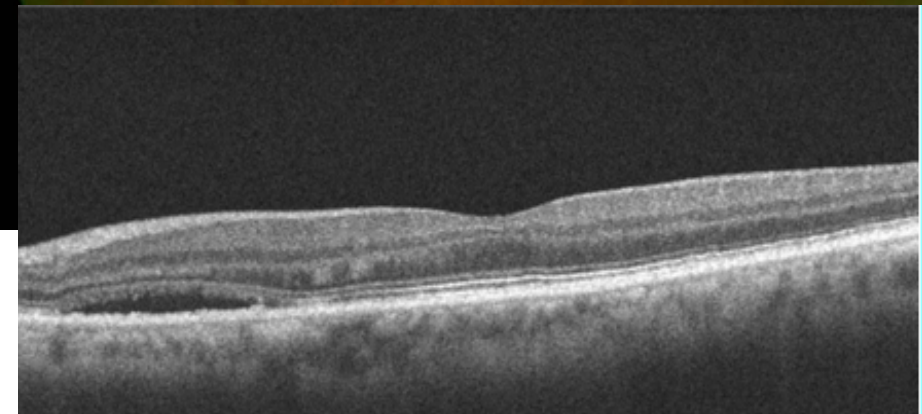
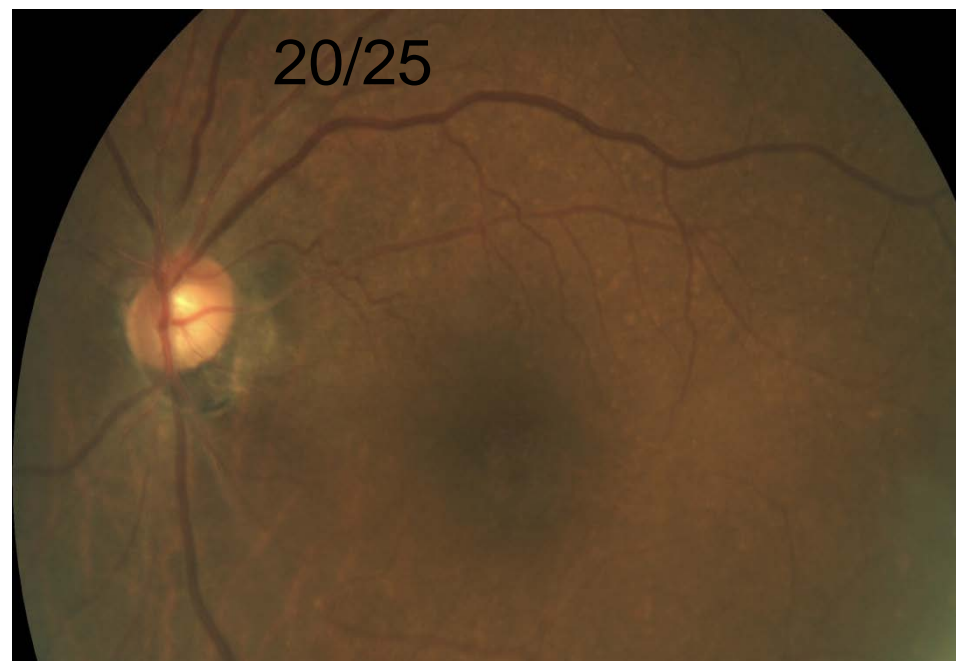
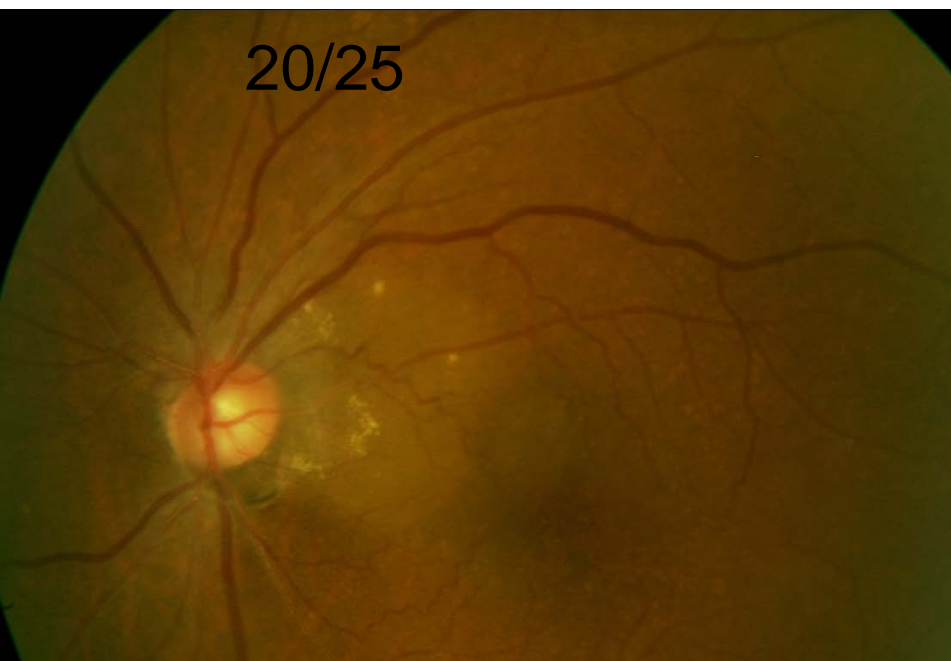
Scan Angle: 0°

Spacing: 0.25 mm

Length: 6 mm

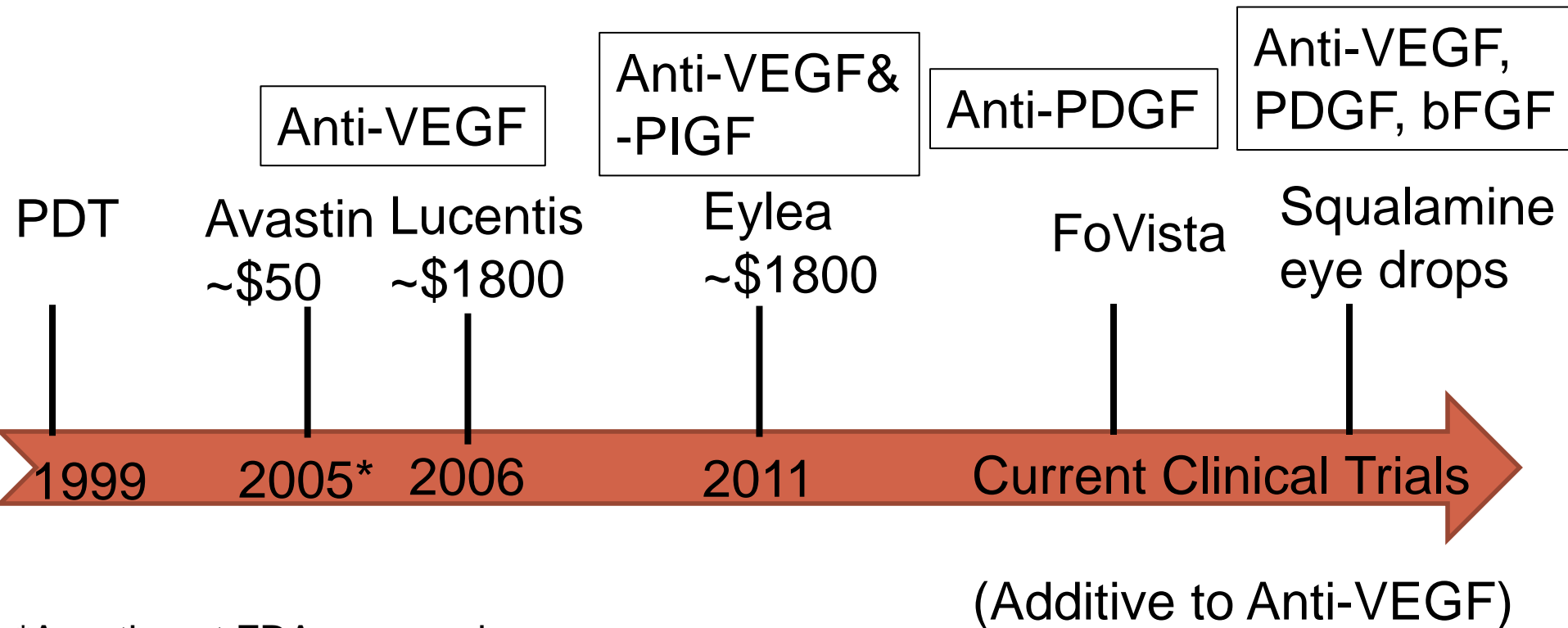


# s/p Avastin injection x 1



# Exudative AMD – polypoidal variant

- Early CNV detection/referral imperative – DON'T DELAY!
  - Typically within within 1 week or less



\*Avastin not FDA approved



THANK YOU!!!!!!

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## Glaucoma Update Fall 2014 Continuing Education

Janice McMahon, OD, FAAO      Ahmad A. Aref, MD  
Associate Professor of      Assistant Professor of  
Optometry      Ophthalmology

October 27, 2014  
Chicago, IL

## Disclosures – Dr. McMahon

No disclosures.

## Disclosures – Dr. Aref

C: New World Medical, Inc.

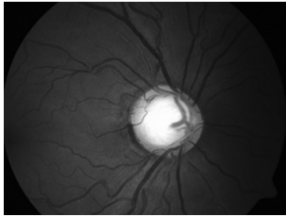
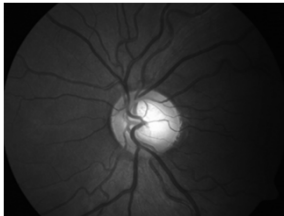
L: Alcon Laboratories, Carl Zeiss Meditec

R: Akorn Pharmaceuticals

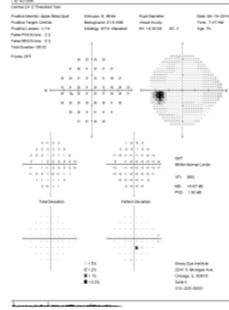
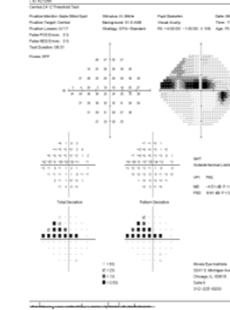
## Case Presentation

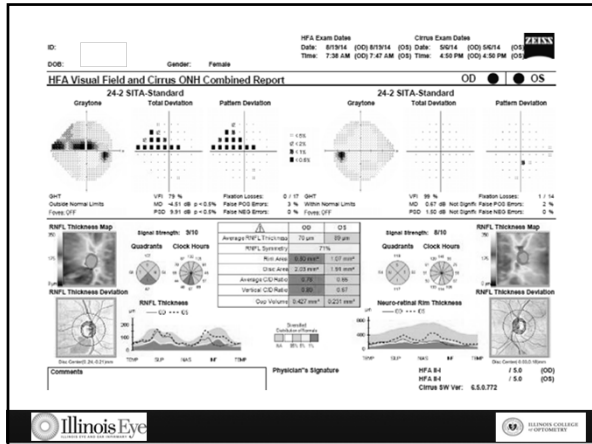
- 67 year old female
- Following for 3 years with early POAG
- IOP pre-treatment mid-20s
- Achieved high teens, OD>OS
- Single medication, prostaglandin qhs OU

## ONH

## Visual fields



## Case Presentation

- VA 20/40 OD, OS
- Mild cataract with glare complaint
- BAT ↓VA

## Management options

- Add second topical medication
- Consider laser
- Consider surgical intervention

## Medical Therapy

<h3 style="text-align: center;">Advantages</h3> <ul style="list-style-type: none"> <li>• Non-invasive</li> <li>• Better perceived by patient</li> <li>• Efficacious</li> </ul>	<h3 style="text-align: center;">Disadvantages</h3> <ul style="list-style-type: none"> <li>• Side effects</li> <li>• Dosing Schedule</li> <li>• Contraindications</li> <li>• Drug interactions</li> </ul>
--	--

## Adjunctive Medical Therapy

- Nearly 40% of patients in Ocular Hypertension Treatment Study required  $\geq 2$  meds to reach target IOP
- Options
  - Beta-blockers
  - Alpha-agonists
  - Carbonic anhydrase inhibitors

### Meta-analysis of the Efficacy and Safety of $\alpha_2$ -Adrenergic Agonists, $\beta$ -Adrenergic Antagonists, and Topical Carbonic Anhydrase Inhibitors With Prostaglandin Analogs

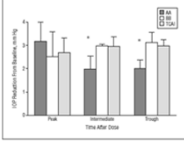
Angelo P. Tanna, MD, Ajay W. Rademaker, PhD, William C. Stewart, MD, Robert M. Feldman, MD

- Results of 10 prospective, randomized, parallel or crossover clinical trials compiled and analyzed
- Primary outcome: Mean IOP reduction from baseline
- Peak, intermediate, and trough IOP lowering efficacy also compared
- Adverse events investigated

### Meta-analysis of the Efficacy and Safety of $\alpha_2$ -Adrenergic Agonists, $\beta$ -Adrenergic Antagonists, and Topical Carbonic Anhydrase Inhibitors With Prostaglandin Analogs

Angelo P. Tanna, MD, Alfred W. Raskemaker, PhD, William C. Stewart, MD, Robert M. Feldman, MD

- IOP-lowering from baseline equivalent among all adjunctive therapies ( $p=0.22$ )
- IOP lowering efficacy at intermediate and trough time points less with alpha agonists
- Greater risk of adverse events with alpha-agonists



The bar chart displays the percentage of patients achieving IOP lowering at four time points: Peak, Intermediate, Trough, and Night. The treatments compared are Prostaglandin Analogs (PA),  $\alpha_2$ -Adrenergic Agonists (A),  $\beta$ -Adrenergic Antagonists (B), and Topical Carbonic Anhydrase Inhibitors (CAI). Error bars represent standard deviation. Significance markers are present above the bars.

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### Comparing Diurnal and Nocturnal Effects of Brinzolamide and Timolol on Intraocular Pressure in Patients Receiving Latanoprost Monotherapy

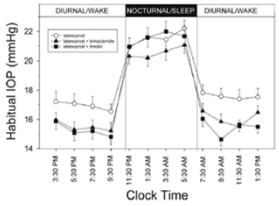
John H. K. Liu, PhD, Felipe A. Medeiros, MD, PhD, J. Rigby Slight, MD, Robert N. Wrensch, MD

- Prospective, open-label, crossover trial of 26 patients with glaucoma or ocular hypertension who were receiving treatment with latanoprost qhs
- IOP measured q2 hrs post-randomization to timolol vs. brinzolamide
- IOPs measured in sitting and supine positions during diurnal time periods and in supine position during nocturnal time period

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### Comparing Diurnal and Nocturnal Effects of Brinzolamide and Timolol on Intraocular Pressure in Patients Receiving Latanoprost Monotherapy

John H. K. Liu, PhD, Felipe A. Medeiros, MD, PhD, J. Rigby Slight, MD, Robert N. Wrensch, MD



The line graph plots Habitual IOP (mmHg) on the y-axis (ranging from 14 to 22) against Clock Time on the x-axis (ranging from 3:30 PM to 1:30 PM). Three data series are shown: Brinzolamide + Timolol (open circles), Brinzolamide + Latanoprost (filled circles), and Brinzolamide + Timolol + Latanoprost (open squares). The graph is divided into three phases: DIURNAL/WAKE, NOCTURNAL/SLEEP, and DIURNAL/WAKE. Error bars are included for each data point.

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### Effects of Systemic $\beta$ -blocker Therapy on the Efficacy and Safety of Topical Brimonidine and Timolol

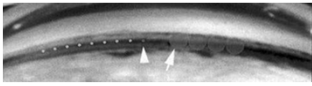
Joel S. Schuman, MD for the Brimonidine Study Groups 1 and 2

- Post-hoc evaluation of data from prospective clinical trials comparing brimonidine and timolol
- Timolol treated subjects concurrently taking systemic B-blockers experienced less IOP-lowering effect ( $P \leq .041$ )
  - $4.41 \pm 0.51$  vs.  $6.23 \pm .18$  mm Hg at peak

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### Laser Therapy

- Argon laser trabeculoplasty described by Wise and Witter in 1979
- Both argon & selective laser trabeculoplasty act to lower IOP by enhancing aqueous outflow facility



(Photo Courtesy of Mark Latina, MD)

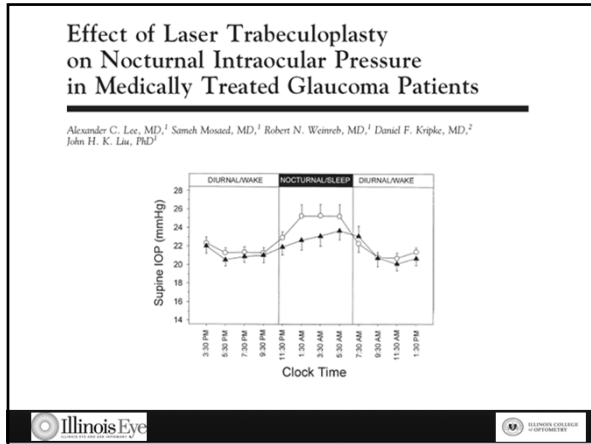
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### Effect of Topical Prostaglandin Analog Use on Outcome Following Selective Laser Trabeculoplasty

WARREN J. SCHERER

- Retrospective review of 113 eyes with POAG treated with SLT
- IOP outcomes slightly better in prostaglandin users compared to non-users ( $P < 0.02$ )

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### Which is better ALT or SLT?

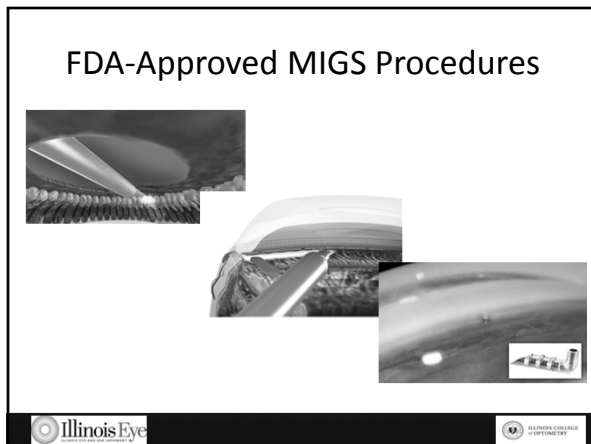
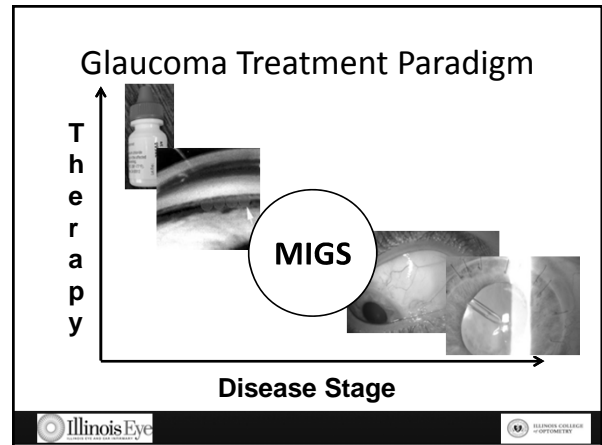
- Similar efficacy<sup>1</sup>
- Similar safety profile<sup>1</sup>
- SLT repeatable<sup>2</sup>, but ALT is not<sup>3</sup>

ALT

SLT

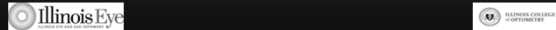
<sup>1</sup>Bovell AM, et al. Can J Ophthalmol. 2011;46:408-413.  
<sup>2</sup>Riansuwan L, et al. Poster 77. AGS, 2007.  
<sup>3</sup>Richter CU, et al. Ophthalmology. 1987;94(9):1085-9.

- ### Microinvasive Glaucoma Surgery
- Ab interno microincisional approach
  - Minimally traumatic to target tissue
  - At least modest IOP-lowering efficacy
  - High safety profile
  - Rapid recovery with minimal impact on quality of life
- Saheb H, Ahmed II. Curr Opin Ophthalmol 2012; 23:96-104.



- ### Trabectome
- Ablation of trabecular meshwork and inner wall of Schlemm's canal
  - FDA-approved in 2004
  - Indications
    - All stages of glaucoma
    - Requires adequate visualization of trabecular meshwork
-

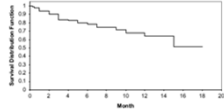

## Trabectome Video



### Combined cataract extraction and trabeculotomy by the internal approach for coexisting cataract and open-angle glaucoma: Initial results

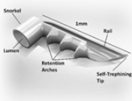
Brian A. Francis, MD, MS, Don Minckler, MD, MS, Laurie Dustin, Shahem Kawji, MD, Jason Yeh, MD, Arthur Sit, MD, Samir Mousad, MD, Murray Johnstone, MD, and the Trabectome Study Group

- Mean IOP reduced from  $20 \pm 6.3$  mm Hg to  $15.5 \pm 2.9$  mm Hg at year with mean medicine reduction of  $1.44 \pm 1.29$  meds

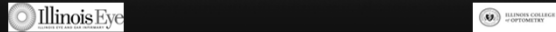



## iStent

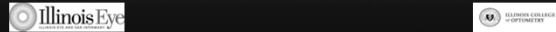
- Titanium L-shaped stent implanted ab-interno
- Bypasses trabecular meshwork by connecting anterior chamber with Schlemm's canal to enhance aqueous outflow



- FDA approved in 2012
  - Indicated for treatment of mild to moderate glaucoma in conjunction with cataract surgery



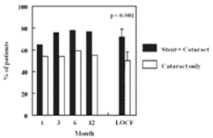
## iStent video



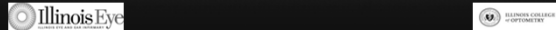
### Randomized Evaluation of the Trabecular Micro-Bypass Stent with Phacoemulsification in Patients with Glaucoma and Cataract

Thomas W. Samuelson, MD,<sup>1</sup> L. Jay Katz, MD,<sup>2</sup> Jeffrey M. Webb, PharmD,<sup>3</sup> Yi-Jing Dai, PhD,<sup>4</sup> Jane Ellen Giamporcaro, BS,<sup>5</sup> for the US iStent Study Group\*

- 1-year results:
  - 72% in treatment group vs. 50% in phaco alone group with unmedicated IOP  $\leq 21$  mm Hg ( $P < 0.001$ )




Ophthalmology 2011;118:459-467.



## Postoperative Care after MIGS

- Perform gonioscopy to visualize surgical result
- Fluctuations in IOP managed accordingly
- Hyphema possible during week 1
- VA stabilizes after 1-2 weeks
- Suture remains in place





### Postoperative Trabectome

- Topical antibiotic x1 week
- Topical corticosteroid x4 weeks
- Maintain glaucoma meds x4-6 weeks
- Pilocarpine qid x6-8 weeks
  
- Follow up at 1 day, 1 week, 3 weeks, 2 months



### Postoperative iStent

- Topical antibiotic x1 week
- Topical corticosteroid x4 weeks
- Maintain glaucoma meds x4-6 weeks
  
- Follow up at 1 day, 1 week, 3 weeks, 2 months



### Goal

- Achieve lower IOP with minimal risk and less significant impact to the patient .



**Thank you**

