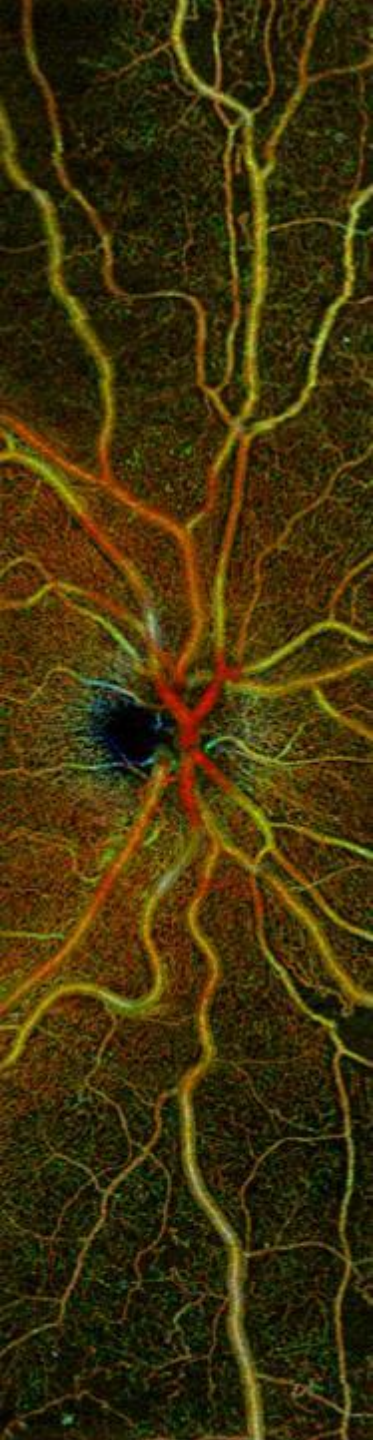




# **OCT ANGIOGRAPHY: IMAGING MOTION**

Rick Trevino, OD, FAAO

Rosenberg School of Optometry



# ONLINE NOTES

- [richardtrevino.net](http://richardtrevino.net)

# CONTACT

- [rctrevin@uiwtx.edu](mailto:rctrevin@uiwtx.edu)

# DISCLOSURES

- No conflicts of interest to disclose.

# RESOURCES

PMID: 33292740

Google:  
OCT Angiography

Greig et al. *Int J Retina Vitreous* (2020) 6:55  
<https://doi.org/10.1186/s12942-020-00262-9>

International Journal  
of Retina and Vitreous

REVIEW

Open Access

## A practical guide to optical coherence tomography angiography interpretation

Eugenia Custo-Greig<sup>1,2</sup>, Jay S. Duker<sup>1</sup> and Nadia K. Waheed<sup>1\*</sup>

### Abstract

**Background:** Optical coherence tomography angiography (OCTA) can image the retinal vasculature in vivo, without the need for contrast dye. This technology has been commercially available since 2014. However, its use has been limited to the research setting. Over time, more clinical publications detail OCTA's use for the study of retinal vascular disease.

**Body:** This review provides an overview of OCT imaging and details the various OCTA techniques and artifacts that arise from practices for acquisition. OCTA has the unique ability to visualize the retinal plexus. Slabs offered in standard for research and clinical use. Lastly, the use of OCTA for the clinical interpretation of age-related macular degeneration, is discussed.

**Conclusion:** OCTA is evolving from a scientific tool to a clinical imaging modality. Image interpretation in a clinical setting.

**Keywords:** Optical coherence tomography angiography, Interpretation

### Background

Optical coherence tomography angiography (OCTA) first became commercially available in 2014 [1]. This technology allowed for the visualization of retinal microvasculature in vivo. Prior to OCTA, fluorescein angiography (FA) and indocyanine green angiography (ICGA) were the mainstay modalities for retinovascular visualization. These imaging modalities generated a 2-dimensional en face view of the retinal vasculature and did not allow for the individual visualization of retinal capillary plexuses. OCTA offered in vivo visualization of the retinal microvasculature in a depth-resolved fashion, without the need for time-consuming dye administration.

Prior to a research of OCTA, age related macular degeneration (AMD) emerged as a leading cause of blindness in the United States. In 1991, the first OCTA study was published, showing the utility in

Functions in 1991 described its utility in retina in

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Box 400, Boston, MA 02111, USA

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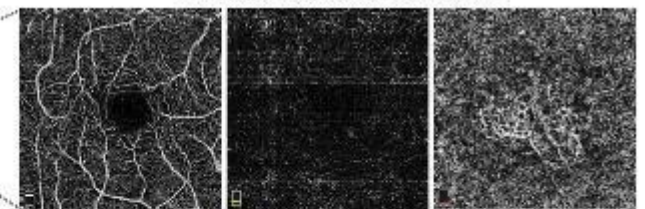


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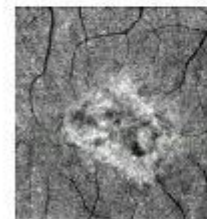
Step 1: Image the right spot



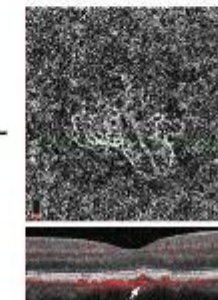
Step 2: Look at the en face OCTA image



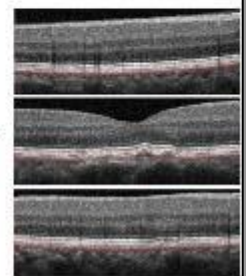
Step 5: Look at the en face intensity image

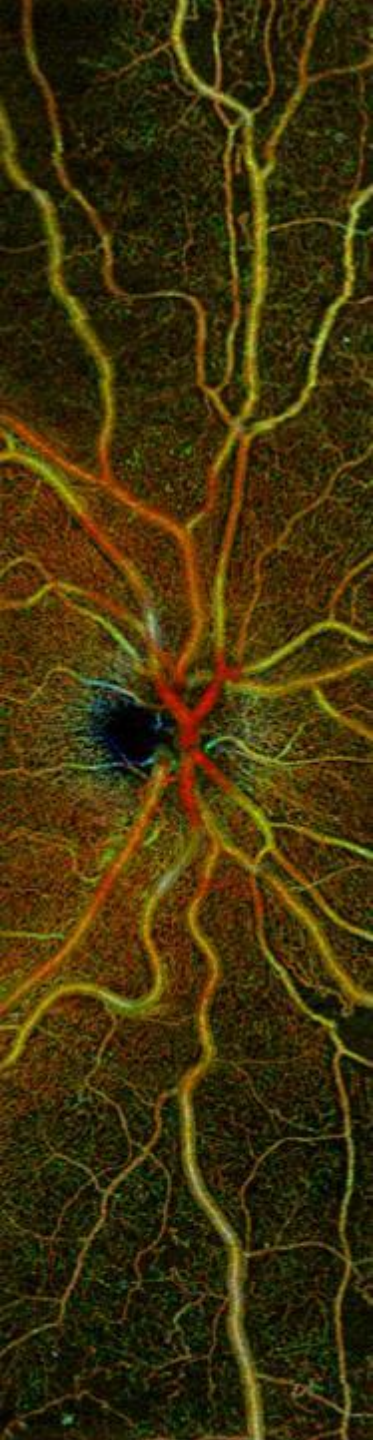


Step 4: Look at the B-scan with flow overlay



Step 3: Look at the segmentation lines

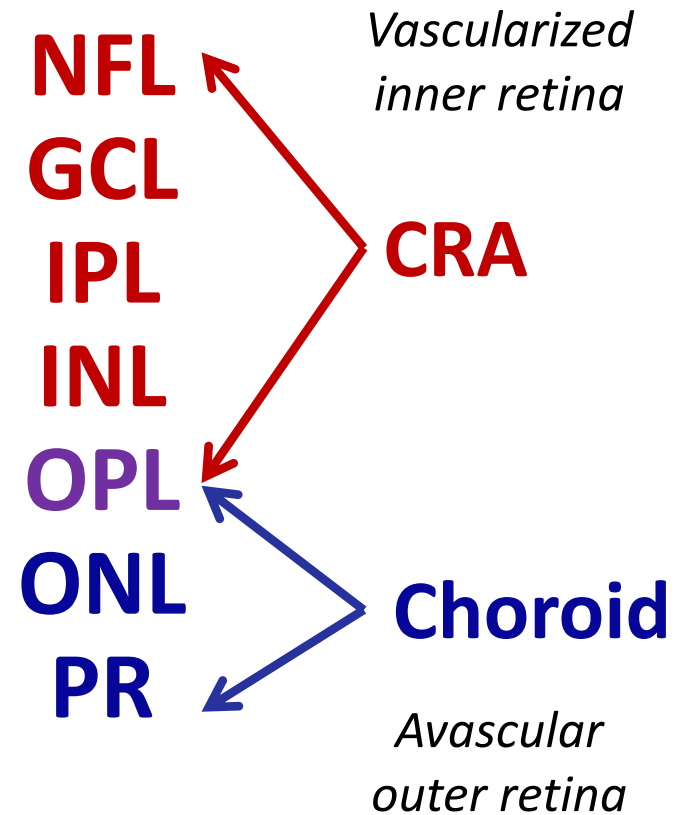
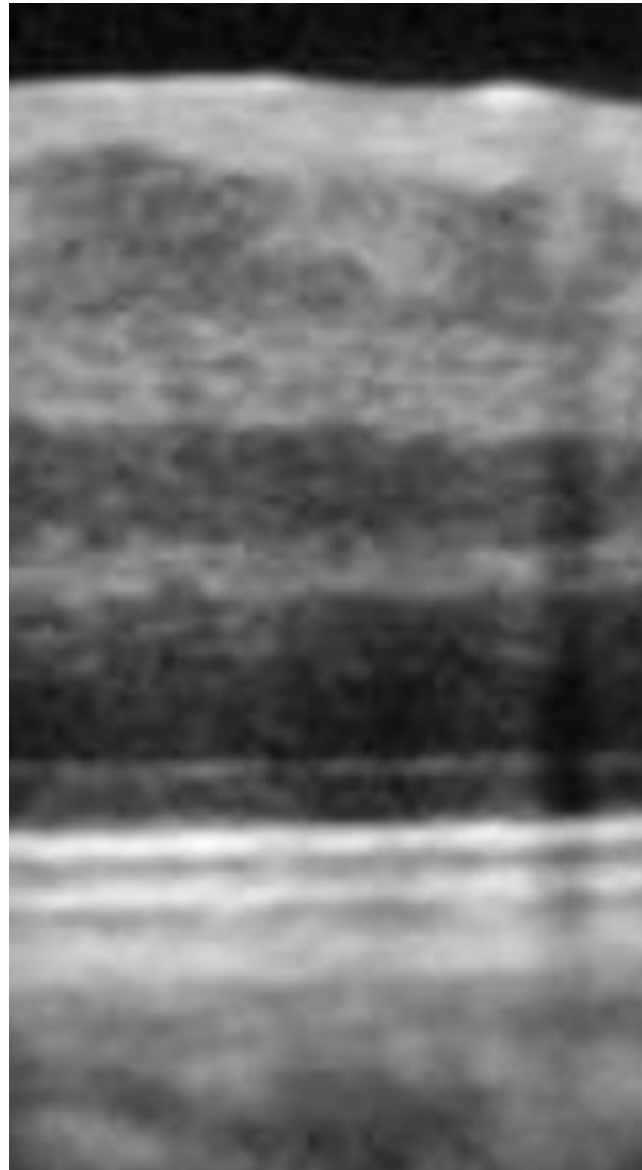
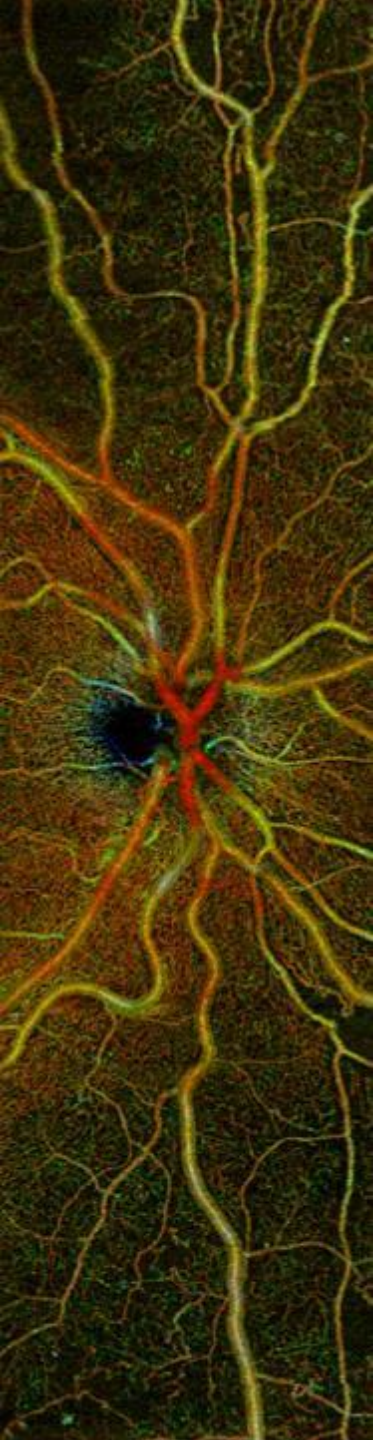




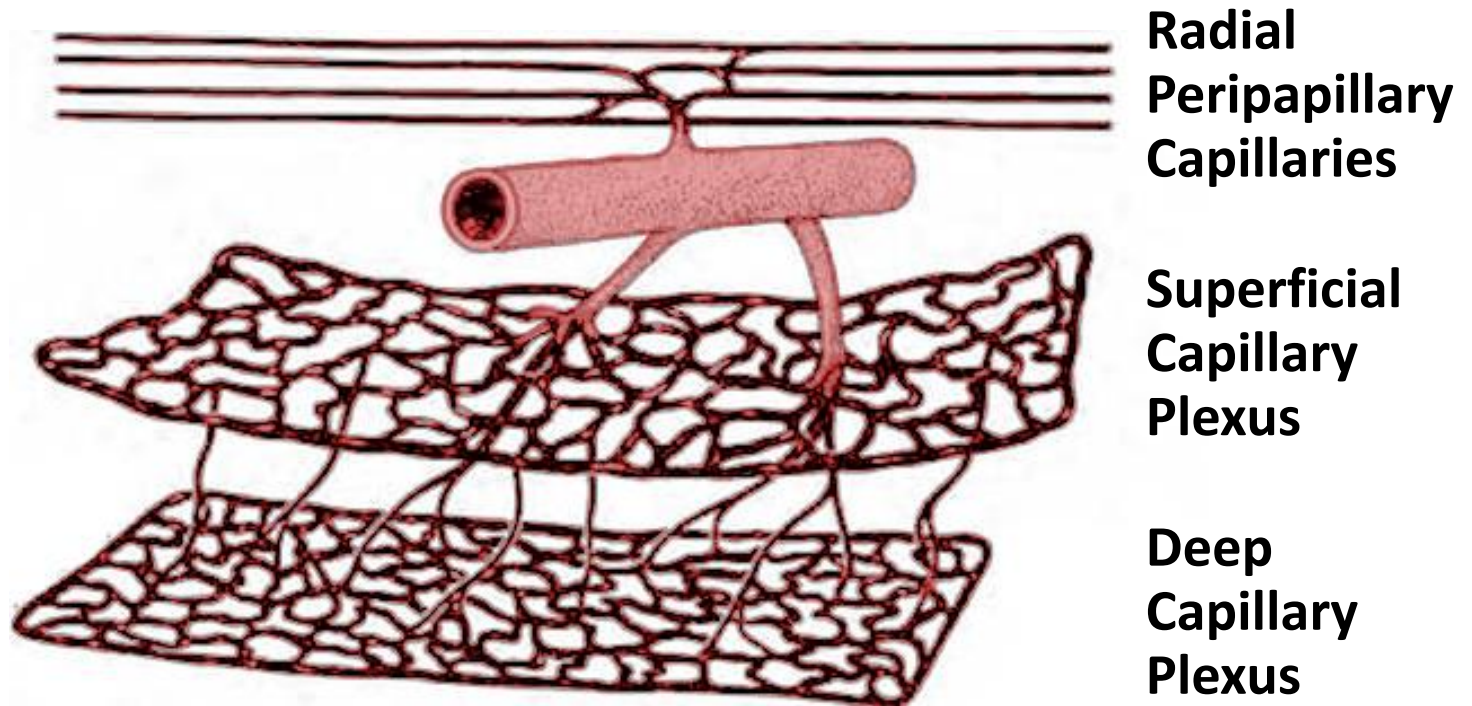
# OVERVIEW

- Retinal blood supply
- Technology (Motion contrast)
- Displays & Interpretation
- OCTA in disease
  - Diabetic retinopathy
  - Venous occlusion
  - CNV (AMD, others)
  - Glaucoma

# RETINAL BLOOD SUPPLY

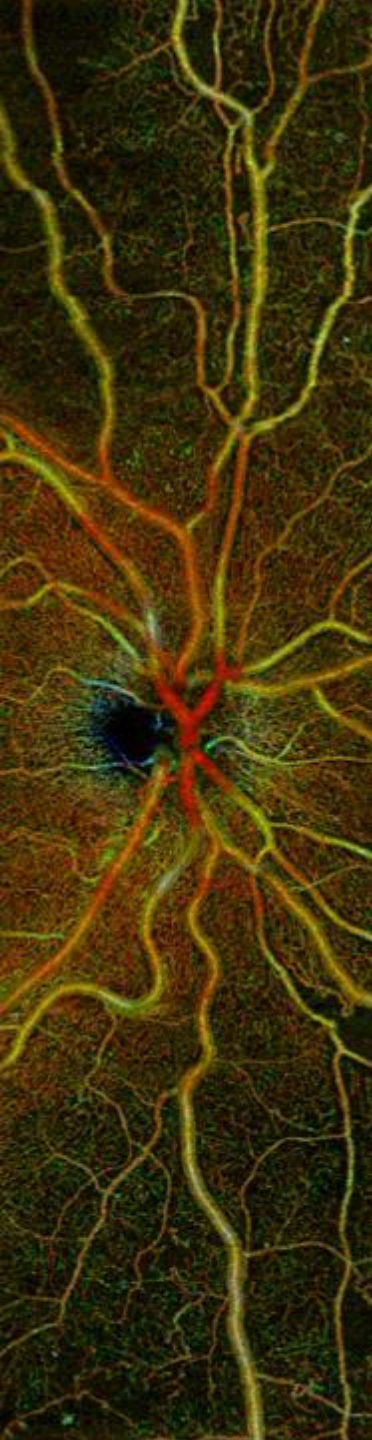


# CENTRAL RETINAL ARTERY



Superficial capillary plexus: Ganglion cells  
Deep capillary plexus: Bipolar cells (INL)

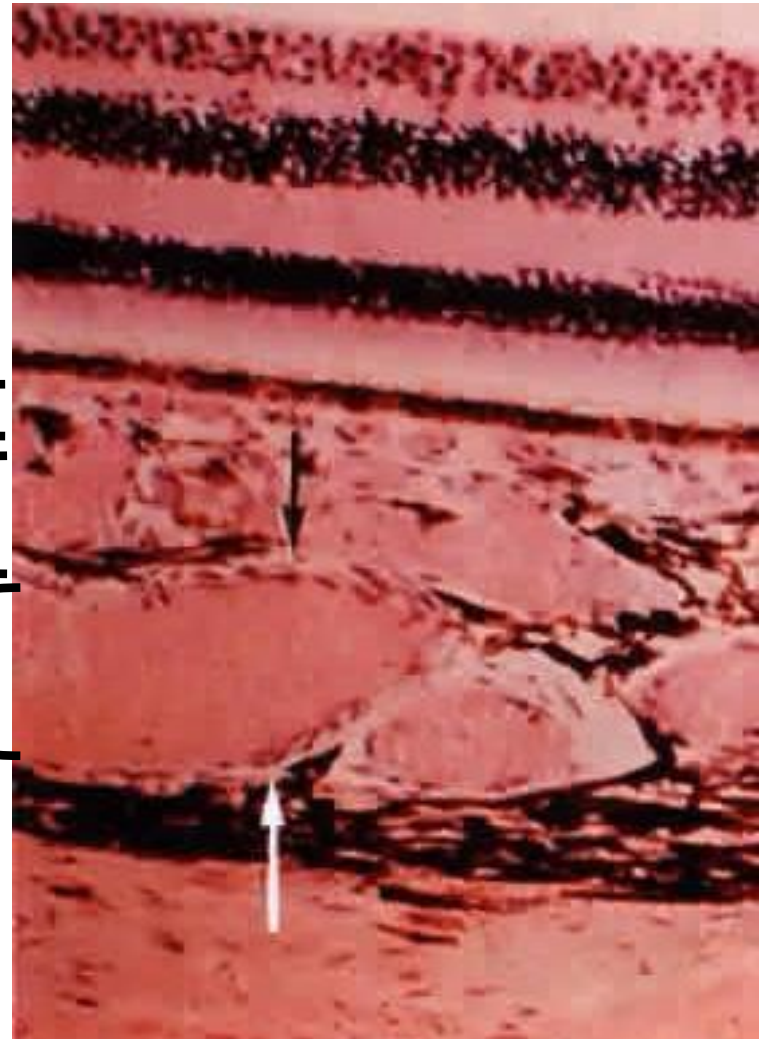
# CHOROID



**Choriocapillaris**

**Sattler's**

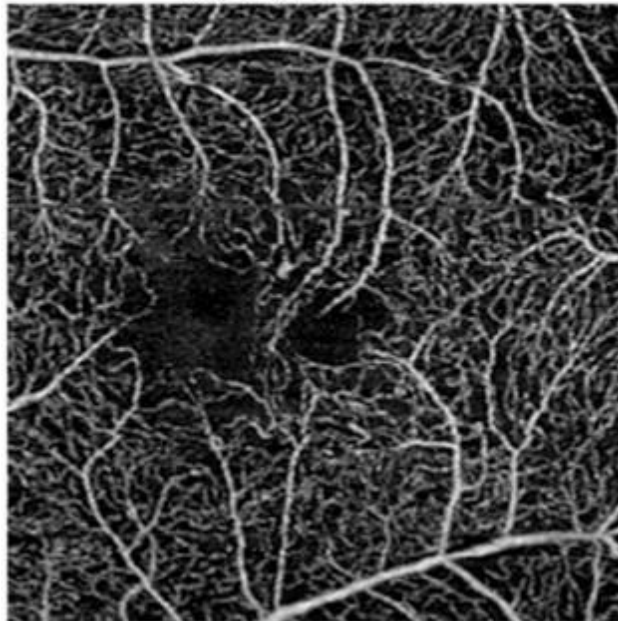
**Haller's**



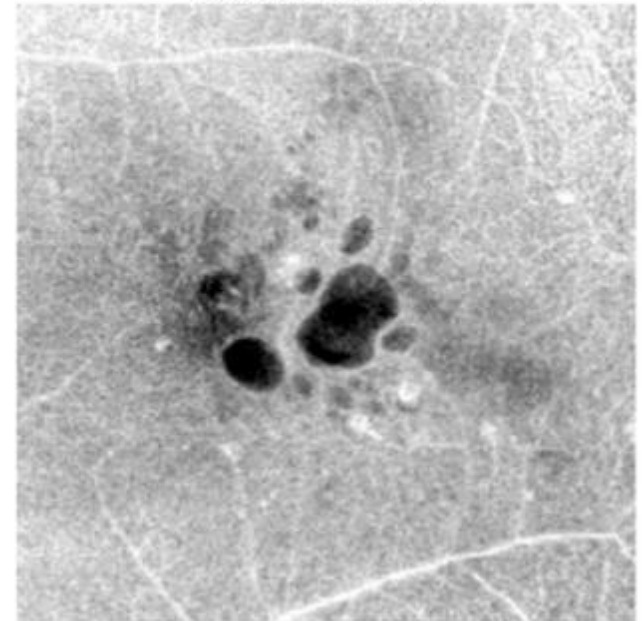
# INTRODUCTION

- Non-invasive “flow” imaging
- 3D volumetric data
- Simultaneous retinal and choroidal imaging
- Structure/ vasculature in tandem

AngioPlex - Superficial

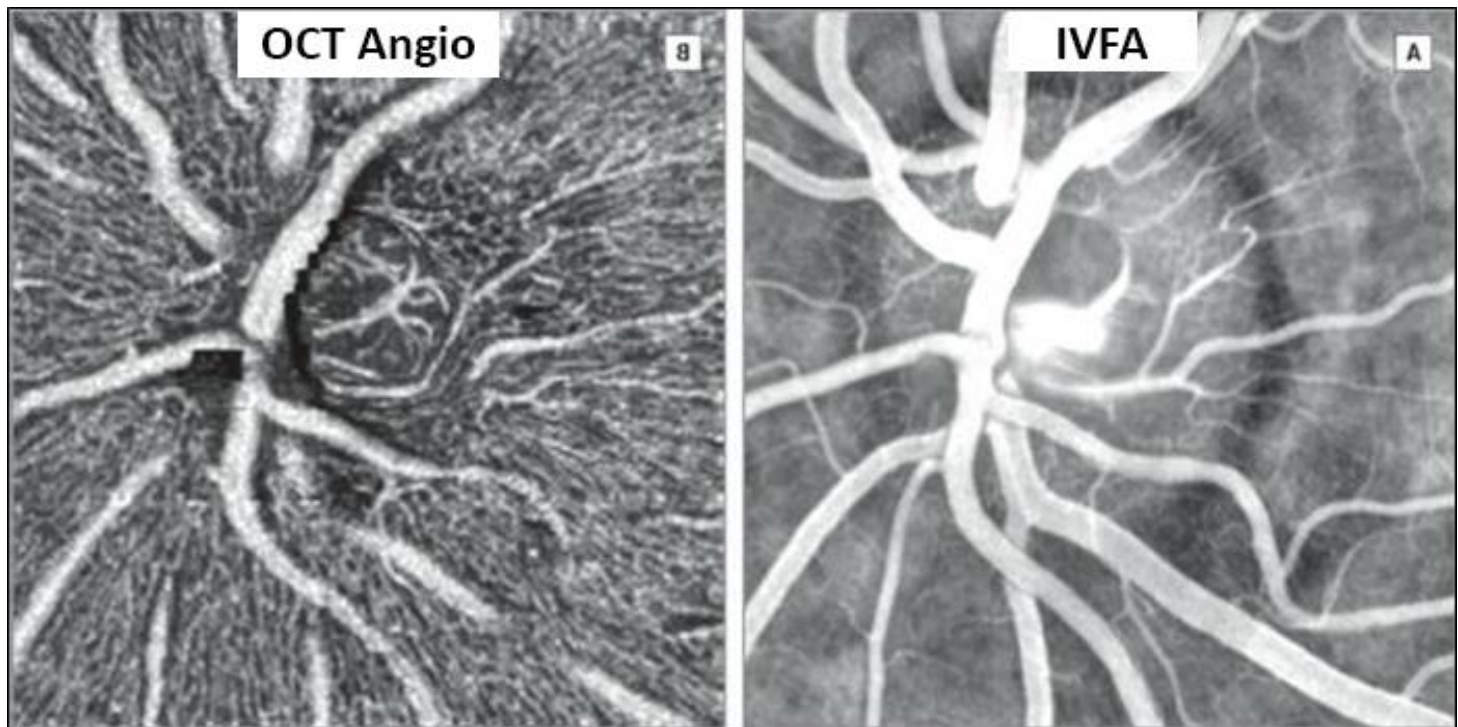


Structure - Superficial



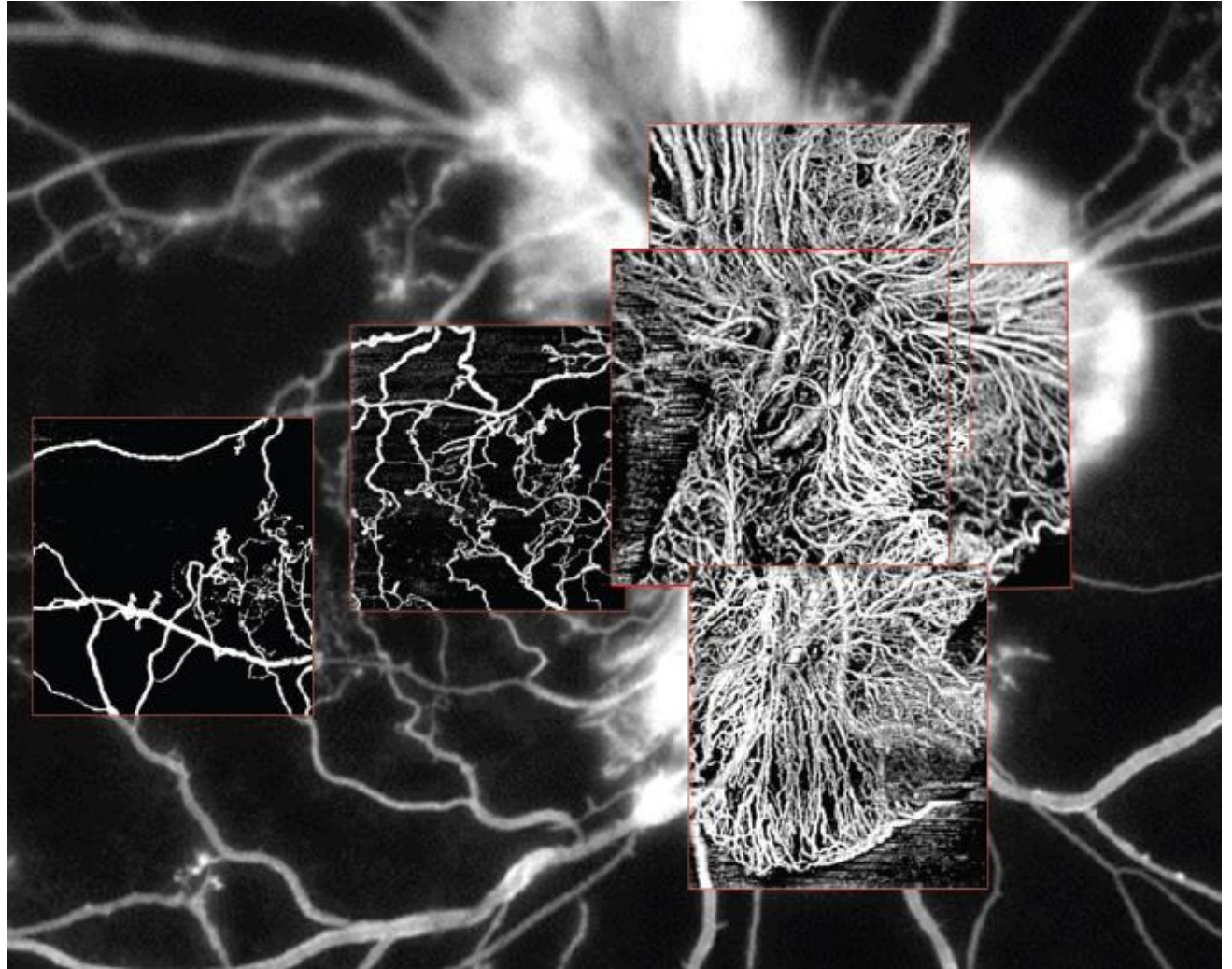
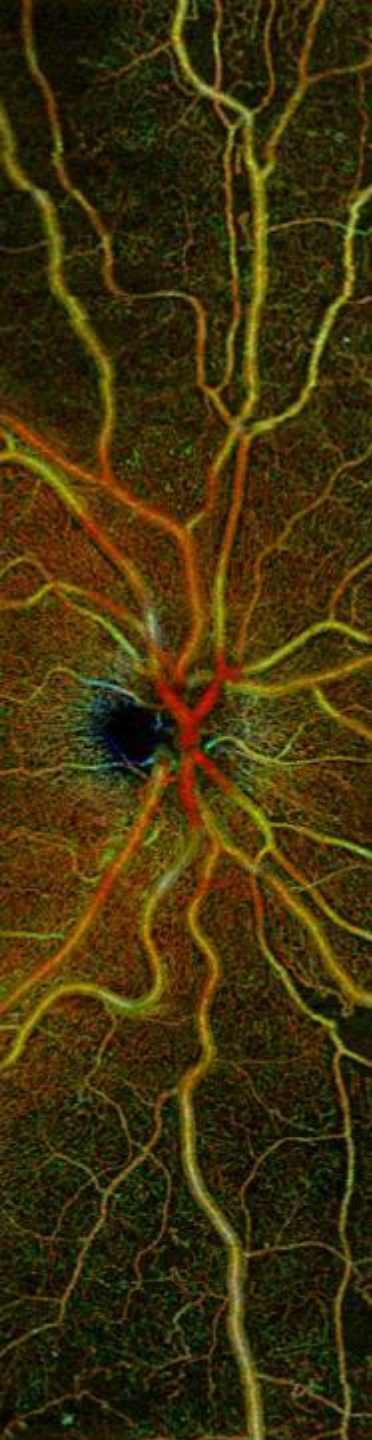
# INTRODUCTION

- Rapid acquisition
- Short term repeatability
- High microvascular resolution



# INTRODUCTION

- Precise delineation/measurement of neo



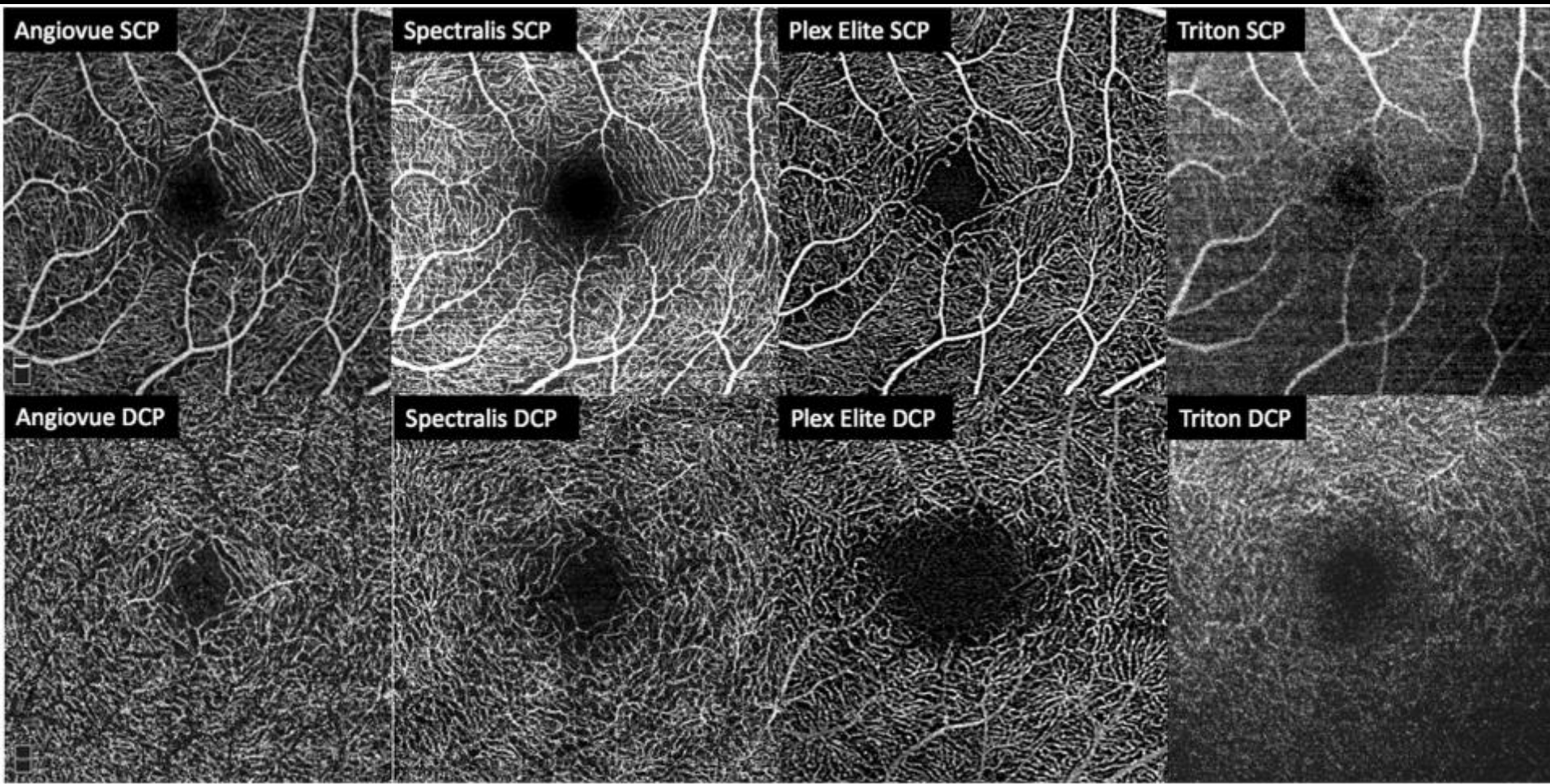
# TECHNOLOGY

## FDA Cleared

- **Zeiss AngioPlex™**
  - Cirrus & Plex Elite 9000
  - Optical micro-angiography
- **Optovue AngioVue™**
  - RTVue XR Avanti Angiovue
  - Split-spectrum amplitude decorrelation angiography
- **Heidelberg Spectralis OCTA Module**
  - Full spectrum amplitude decorrelation angiography
- **Topcon DRI-OCT Triton SS-OCT**
  - OCT angiography ratio analysis

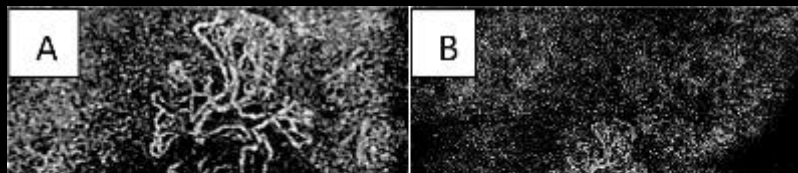


Statistically significant differences exist across devices when measuring the same parameter

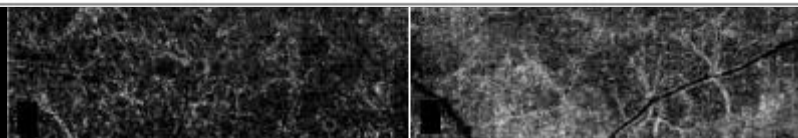
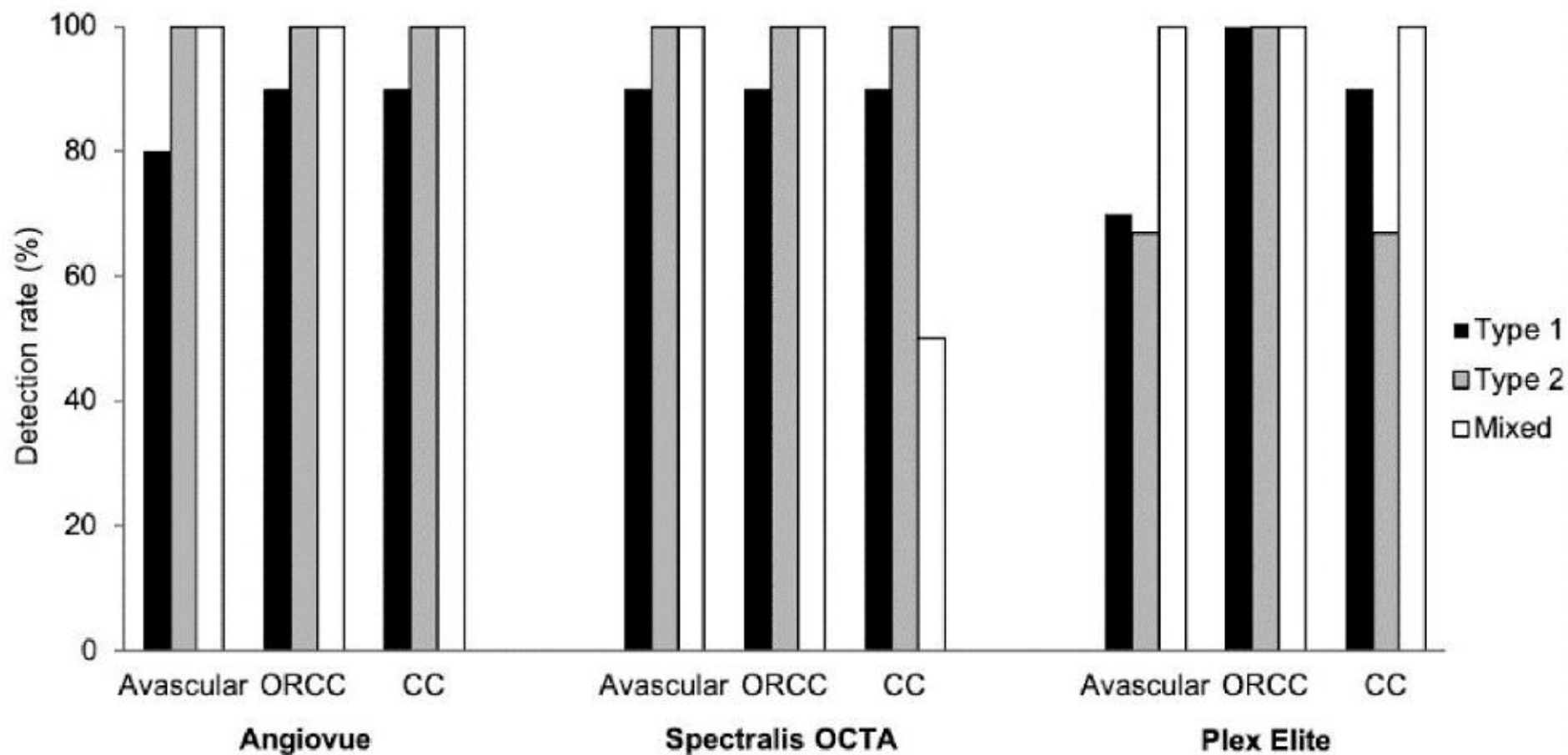


3 x 3

6 x 6

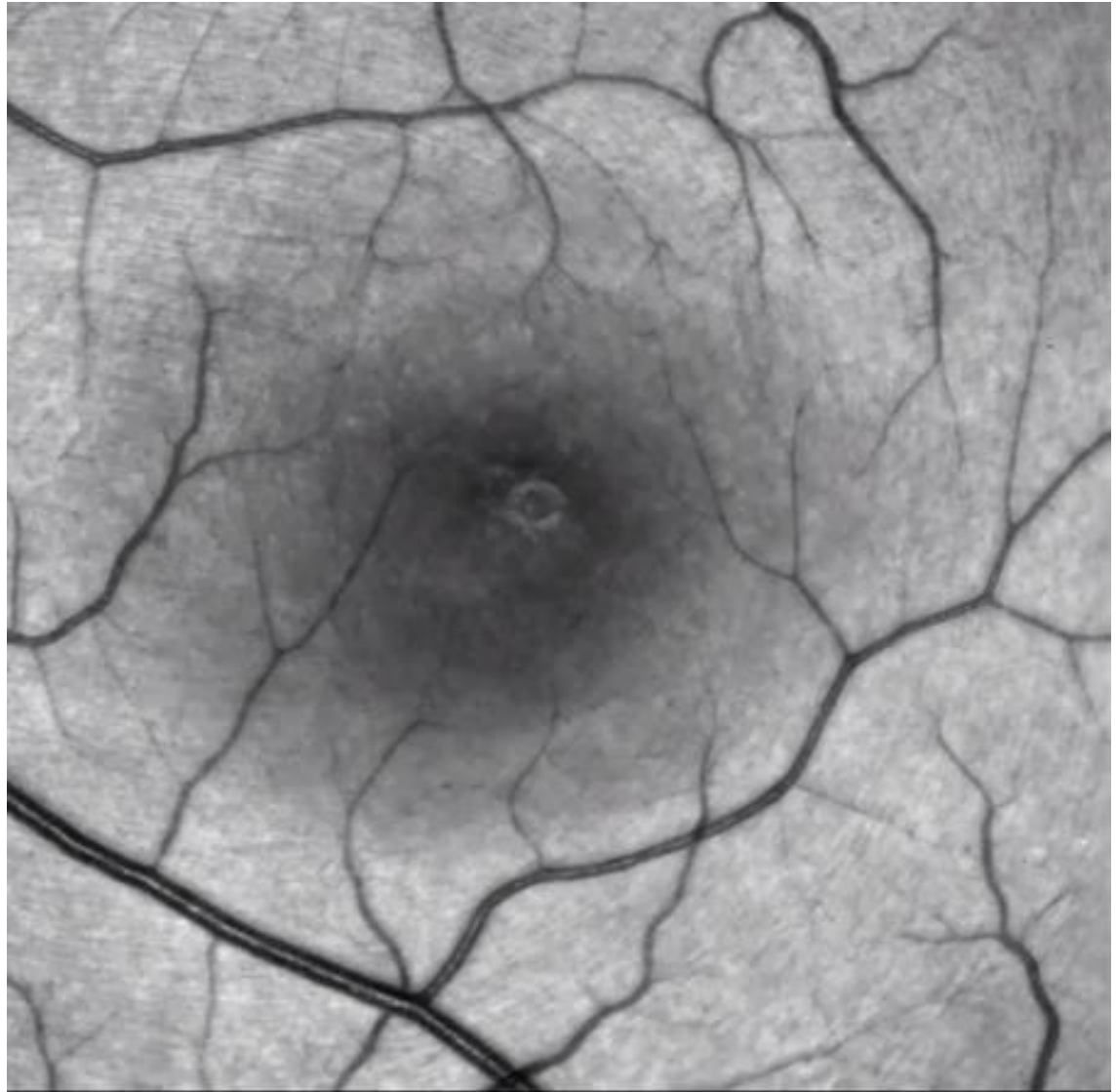
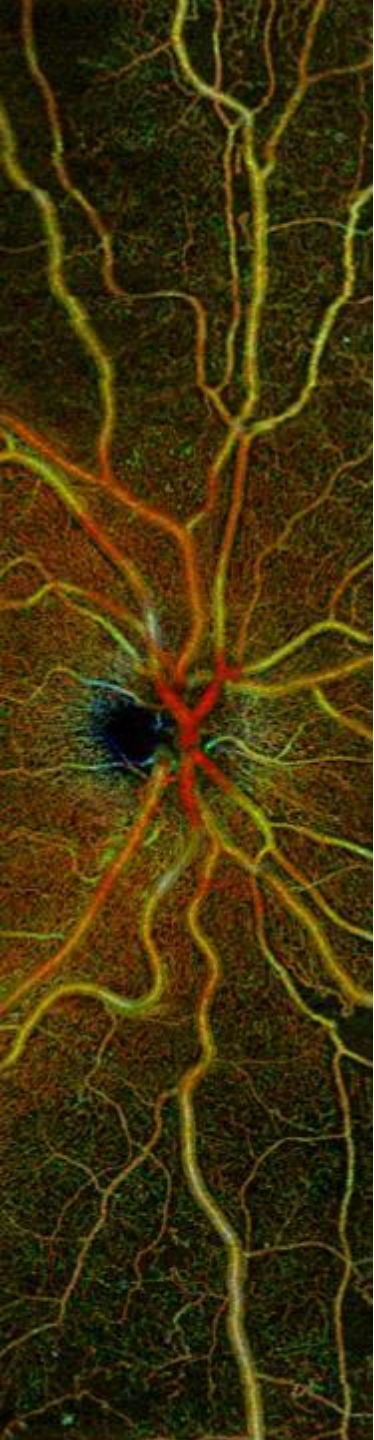


Plex

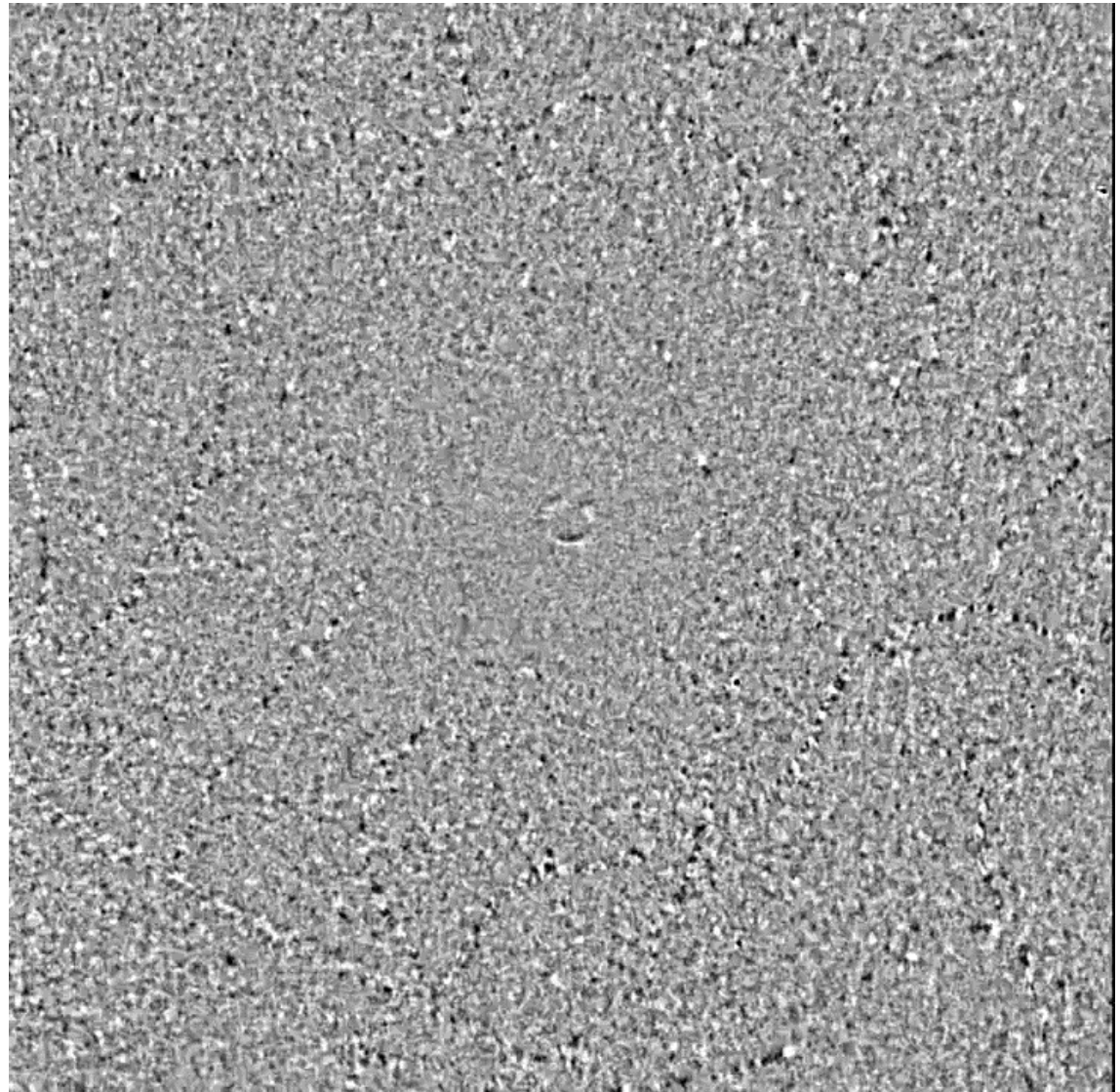
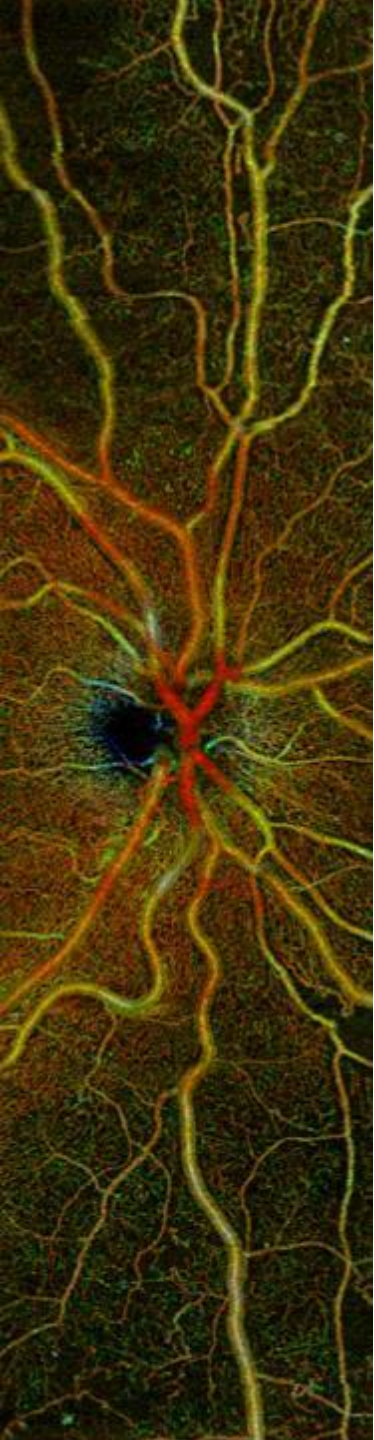


PMID: 33101783

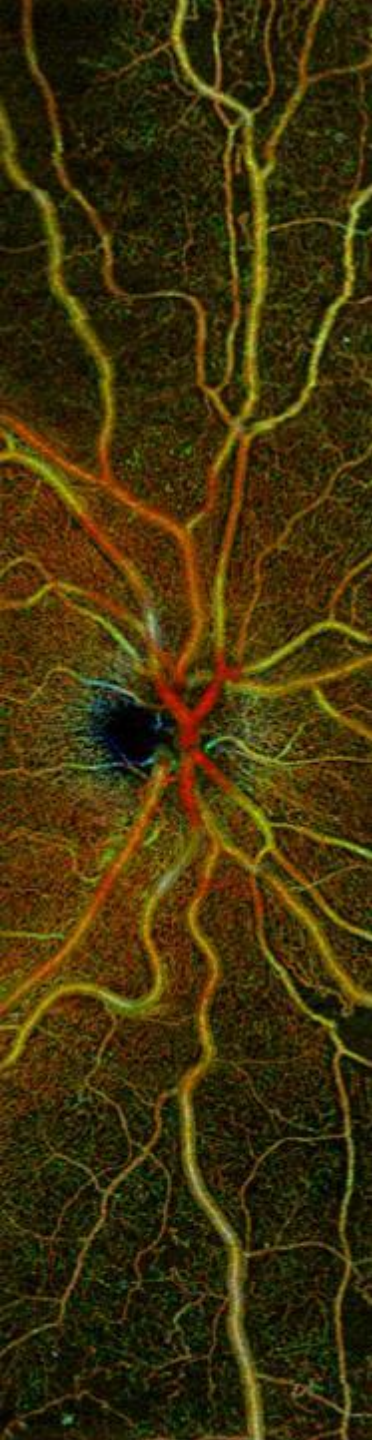
# MOTION CONTRAST



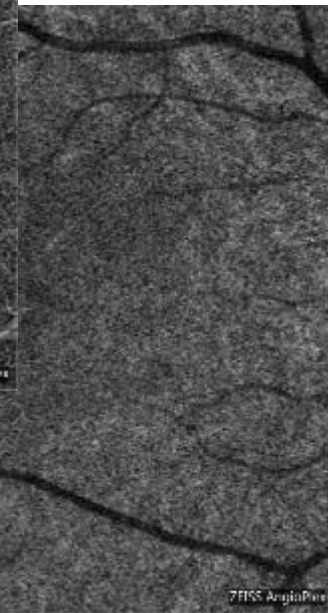
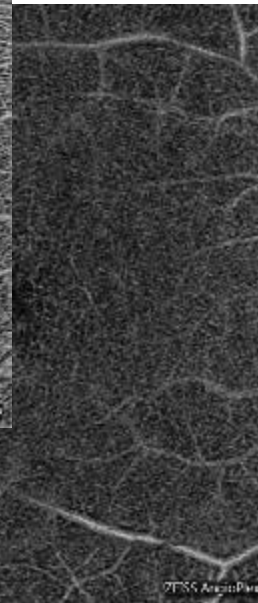
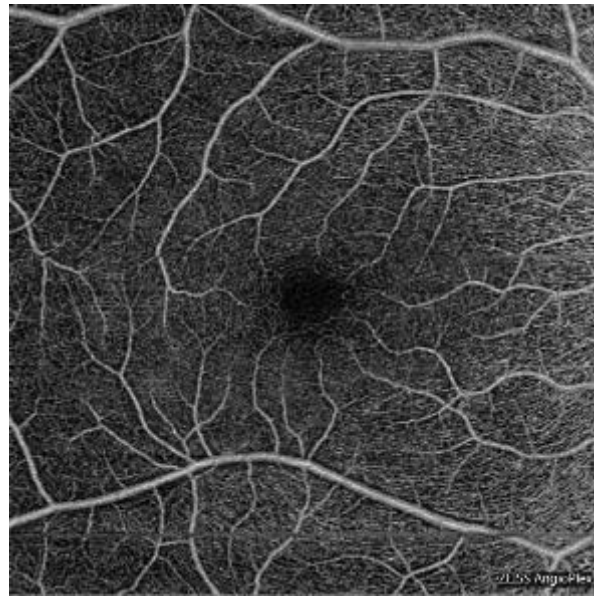
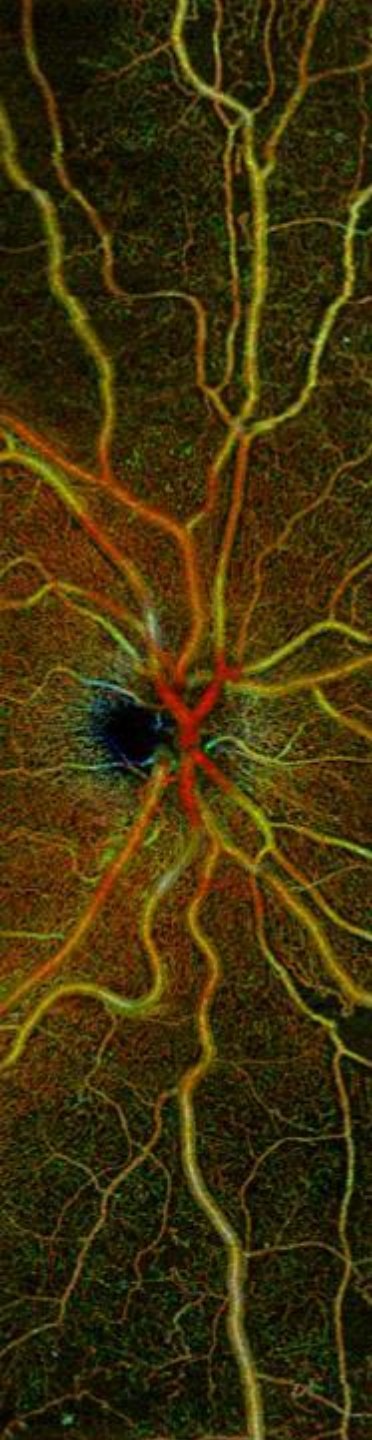
# MOTION CONTRAST



# MOTION CONTRAST



# DISPLAY- En Face

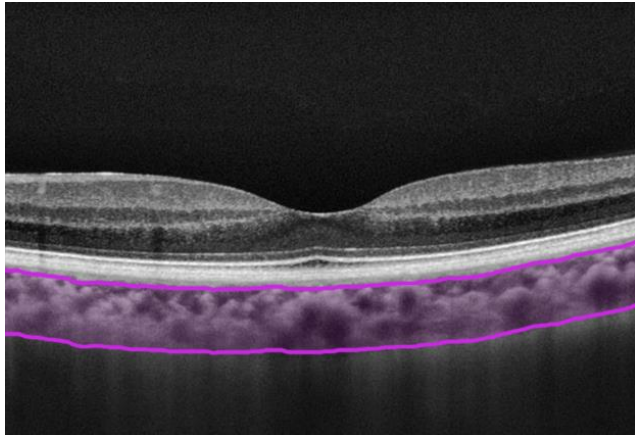


Anterior

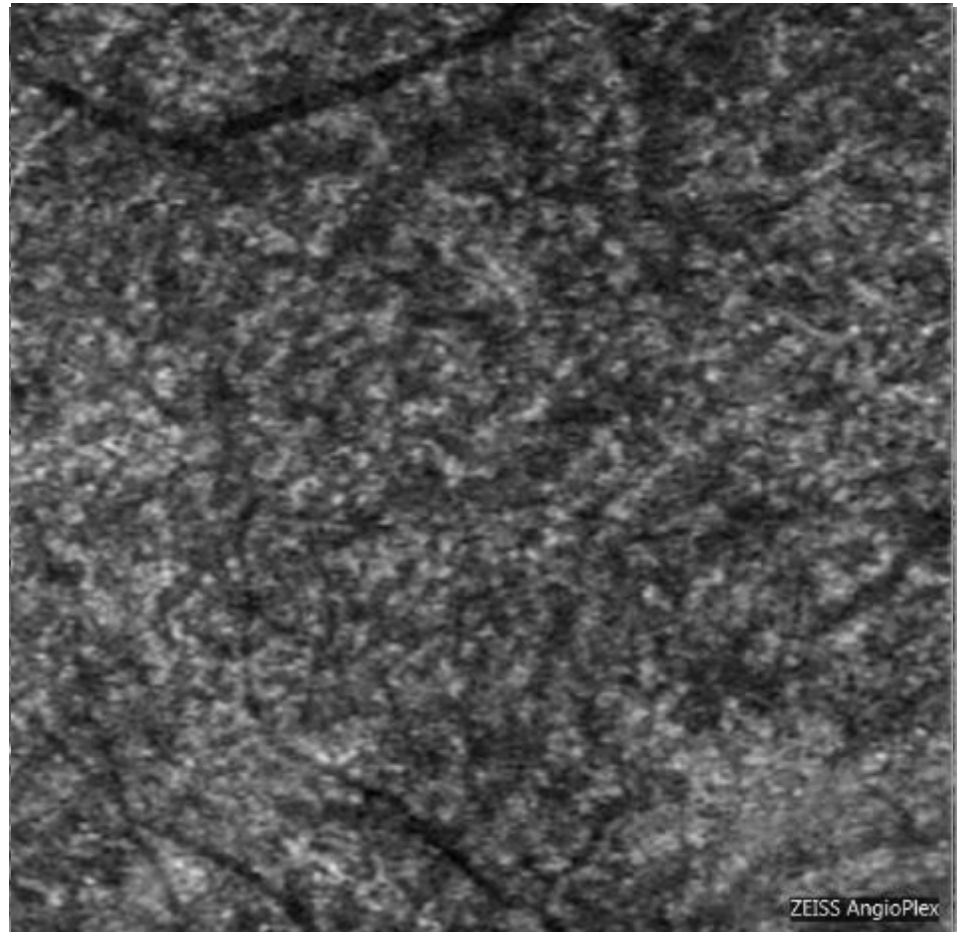
Posterior

# DISPLAY- En Face

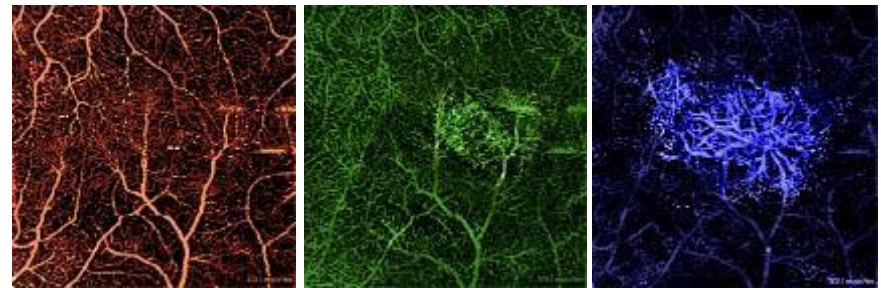
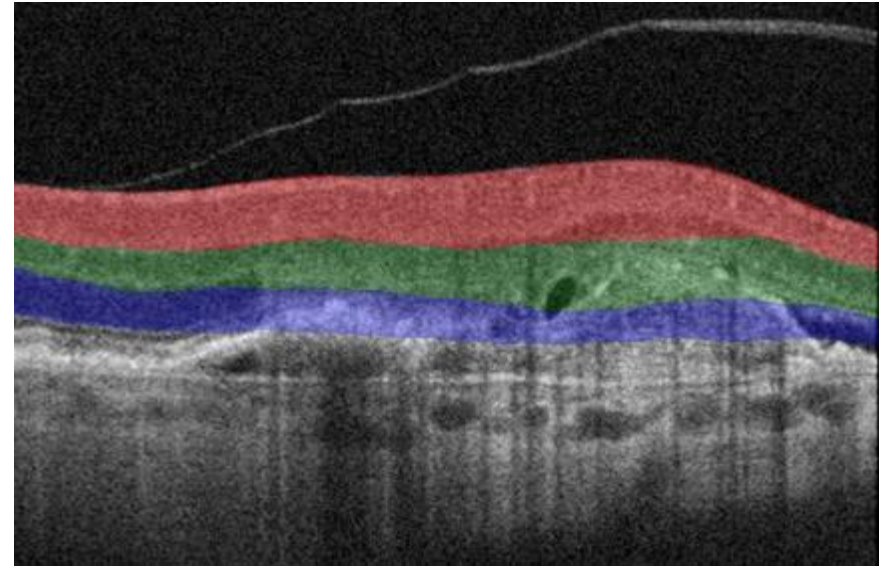
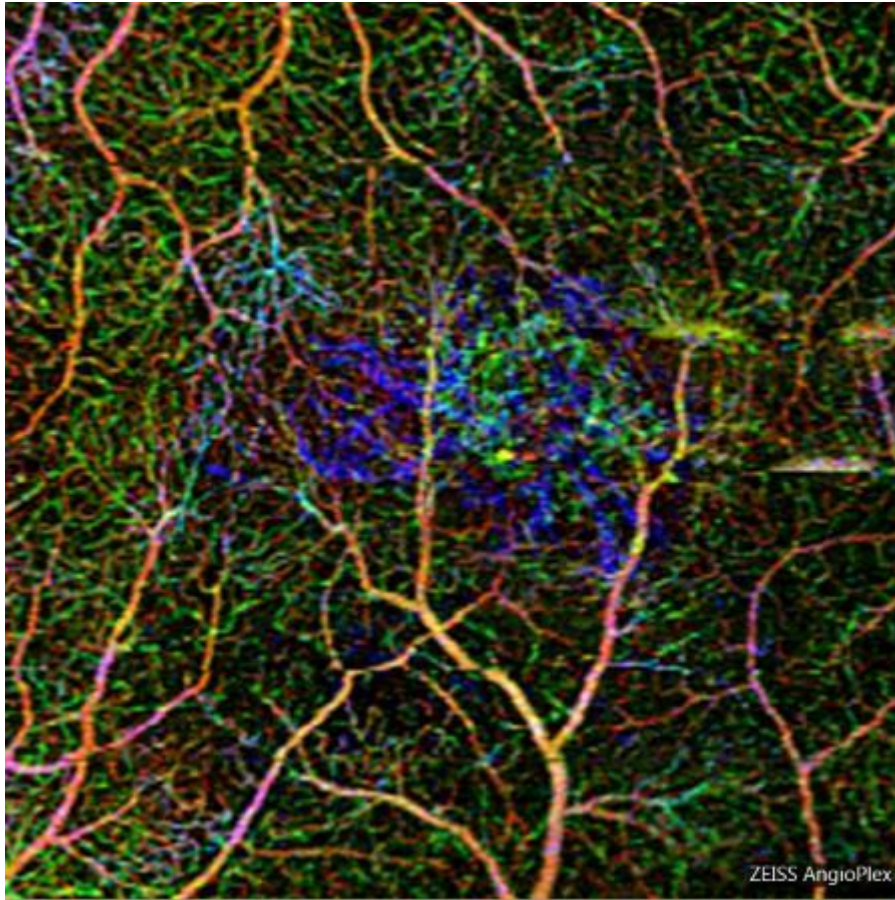
Segmentation lines  
tell you what retinal  
layers are being  
displayed en-face



**Choroid**

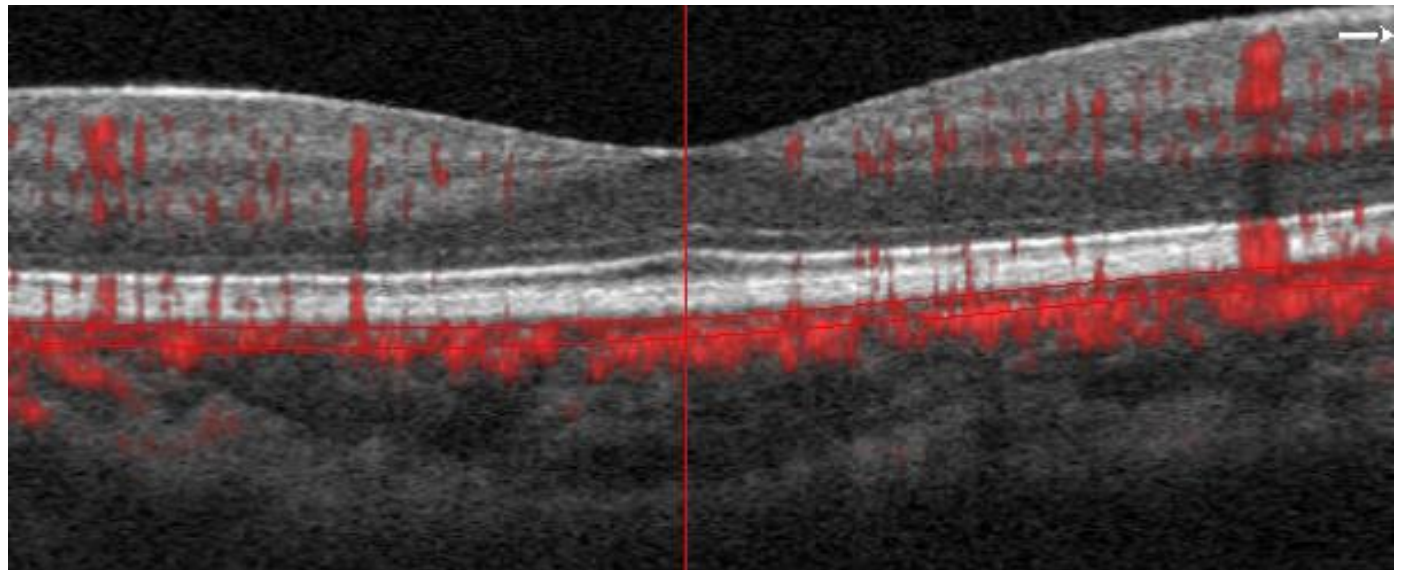
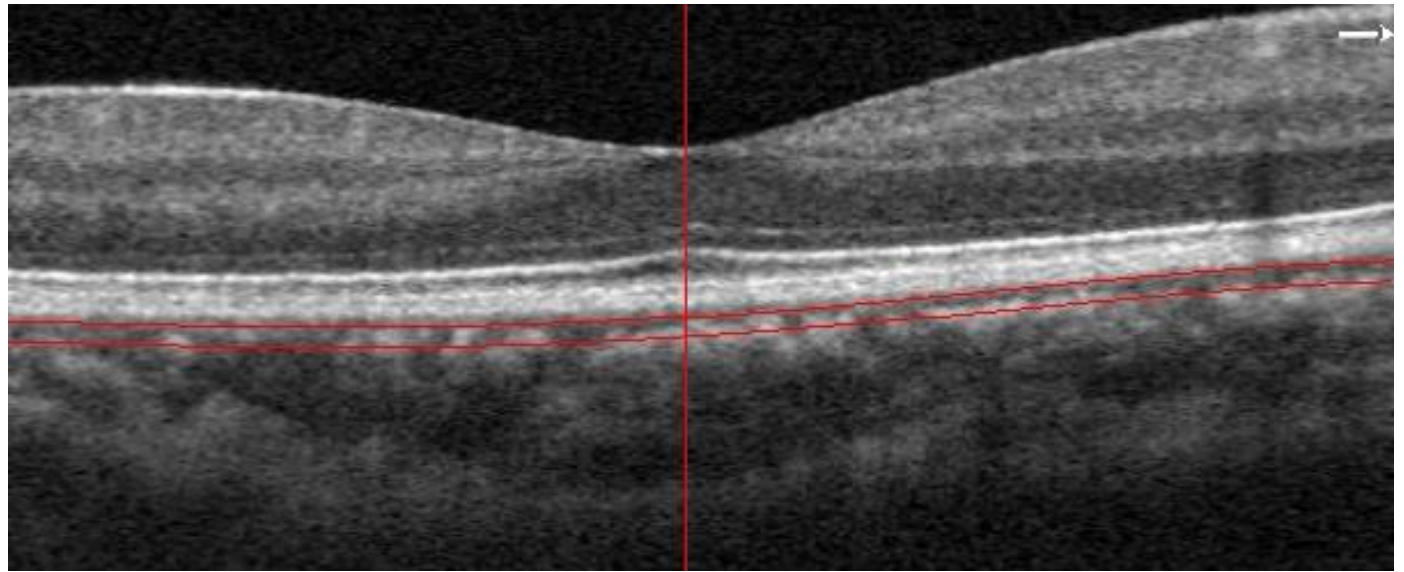
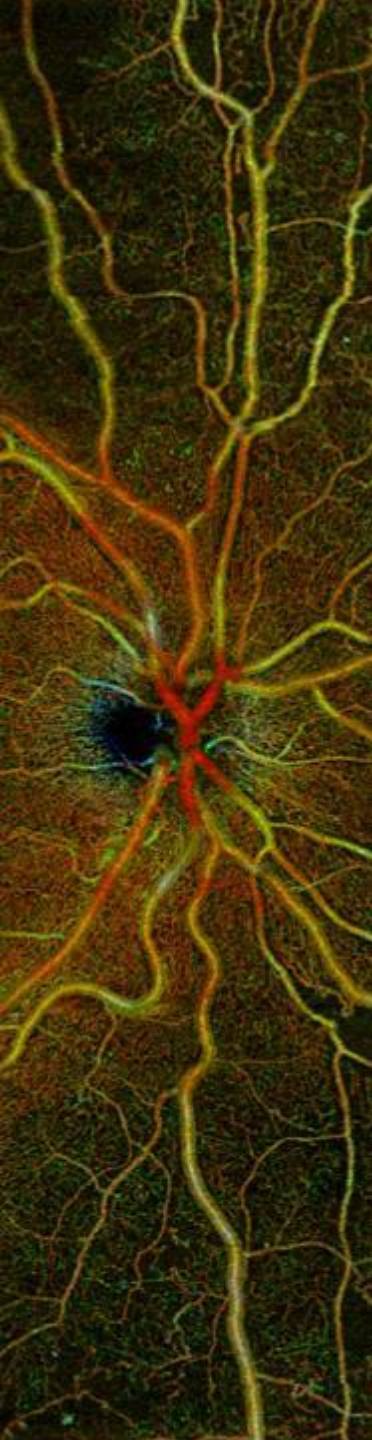


# DISPLAY- Color En Face



S  
D  
A

# DISPLAY- B Scan Overlay



Name:

ID:

DOB:

Gender:

Technician:

Male

Operator, Cirrus

Exam Date:

11/10/2017

Exam Time:

11:22 AM

Serial Number:

4000-8813

Signal Strength:

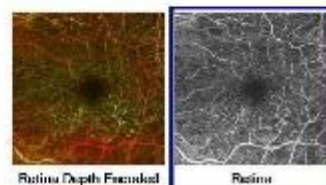
5/10

OS



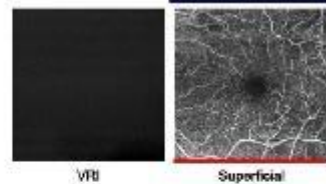
## Angiography Analysis : Angiography 6x6 mm

OD ☐ ☒ OS



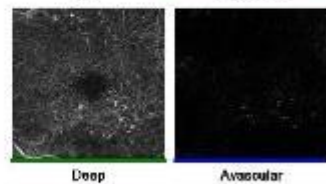
Retina Depth Encoded

Retina



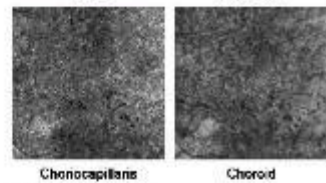
VFI

Superficial



Deep

Avascular



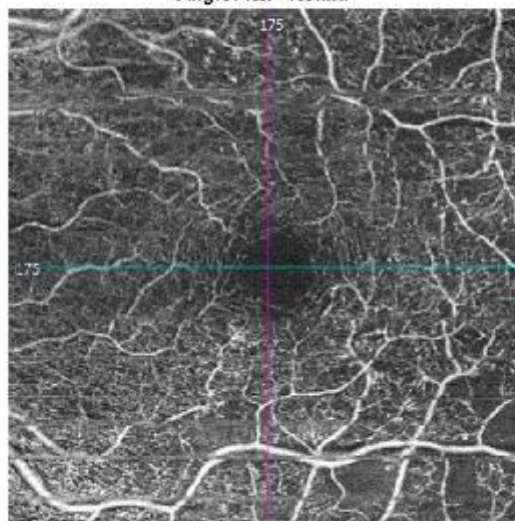
Chonocapillans

Choroid

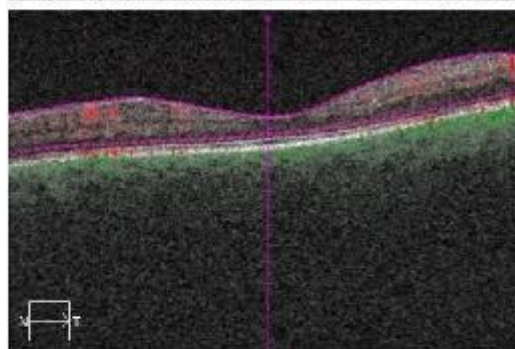
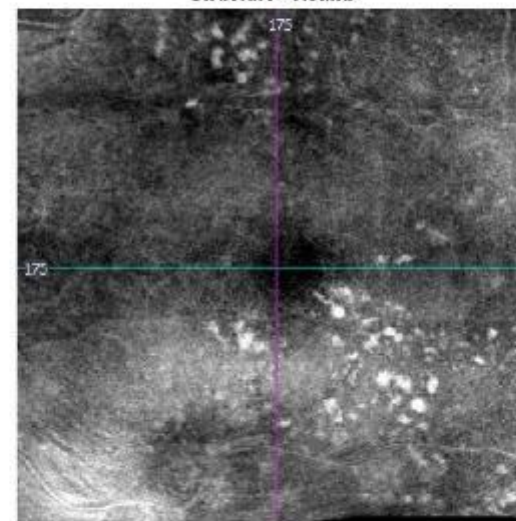


Custom (Global)

AngioPlex - Retina



Structure - Retina



Slice: 175

Current View: Retina

| Reference      | Offset                           |
|----------------|----------------------------------|
| Top: ILM       | <input type="text" value="0"/>   |
| Bottom: RPEfit | <input type="text" value="-70"/> |

Comments

Doctor's Signature

**OCT Angiography Overview**  
SPECTRALIS® Tracking Laser Tomography

**HEIDELBERG  
ENGINEERING**

Patient:  
Patient ID:  
Diagnosis: ---

DOB:  
Exam.: Jul/30/2021  
Comment: ---

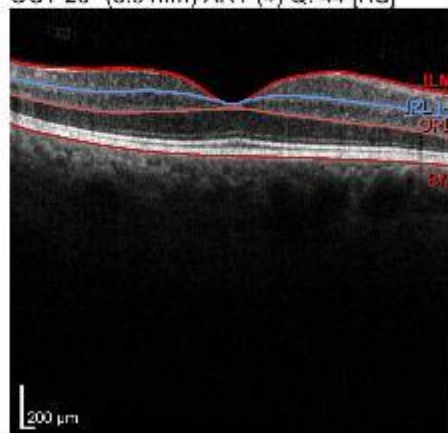
Sex: M

**OD**

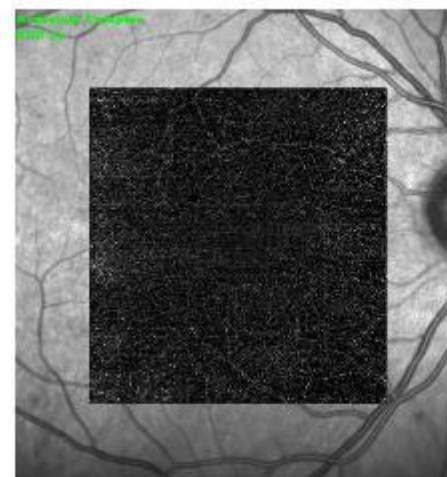
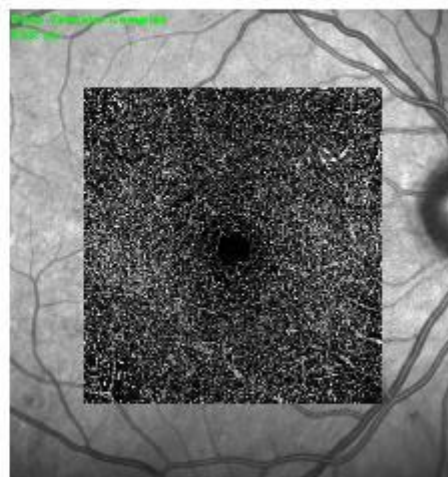
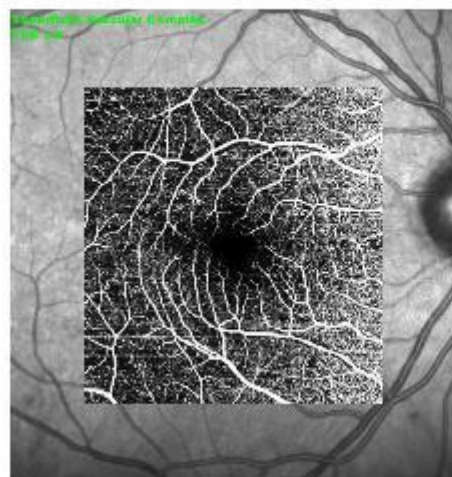
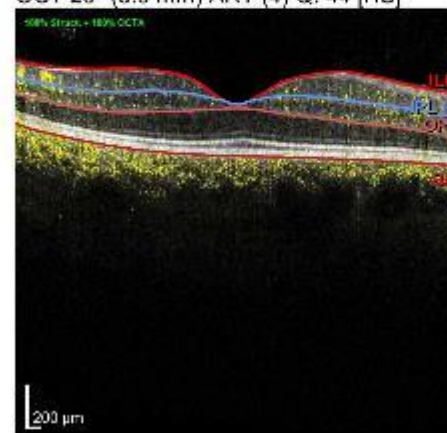
IR 30° ART [HS]



OCT 20° (5.9 mm) ART (4) Q: 44 [HS]



OCT 20° (5.9 mm) ART (4) Q: 44 [HS]



Name:

OS

CZMI



ID:

Exam Date:

11/4/2019

DOB:

Exam Time:

8:27 AM

Gender:

Male

Serial Number:

5000-20078

Technician:

Operator, Cirrus

Signal Strength:

N/A

## Montage Angio Analysis : Montage Angio 6x6 mm

OD ☐ ☒ OS



Superficial  
AngioPlex

**Top**

ILM, Offset: 0

**Bottom**

IPL, Offset: 0

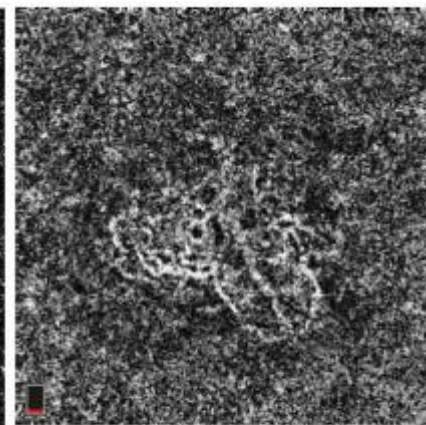
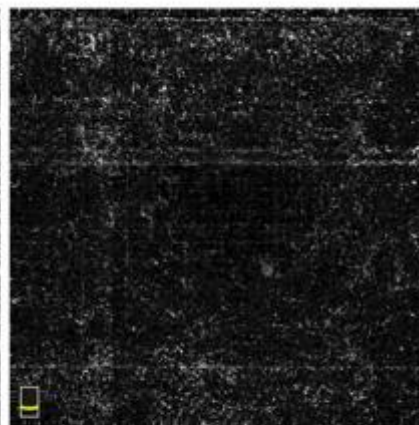
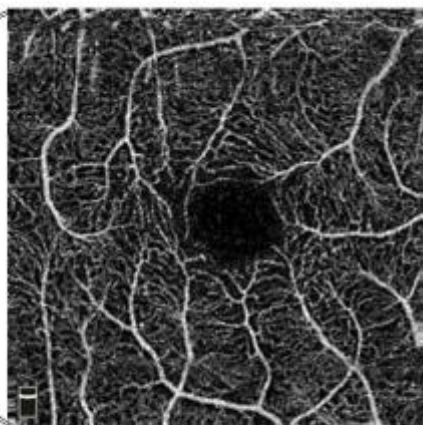
Comments

Doctor's Signature

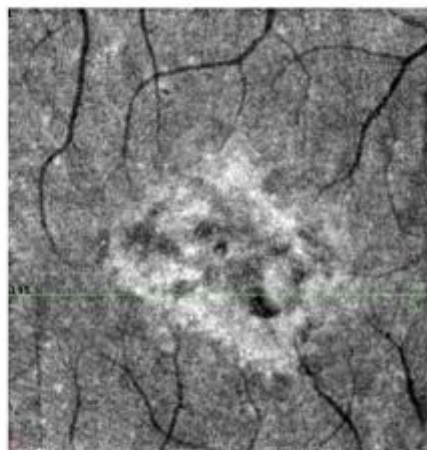
Step 1: Image the right spot



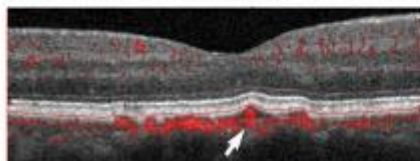
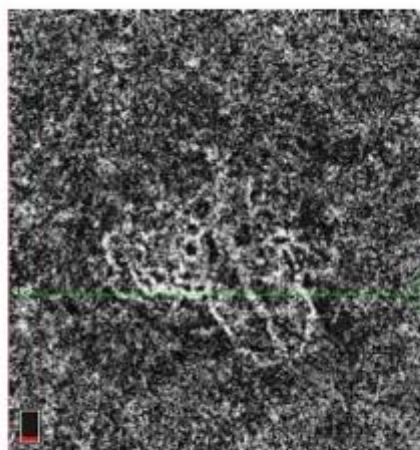
Step 2: Look at the en face OCTA image



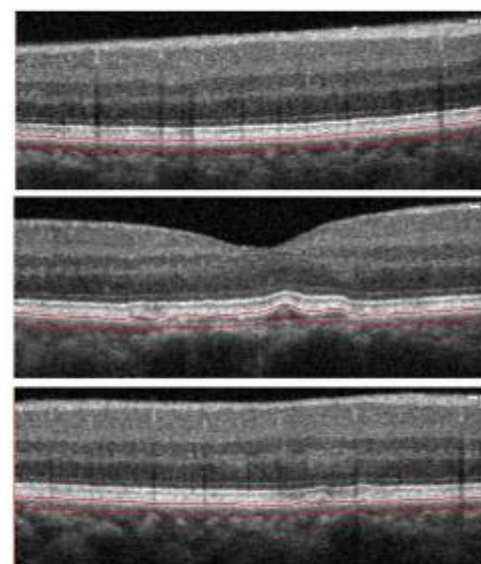
Step 5: Look at the en face intensity image



Step 4: Look at the B-scan with flow overlay

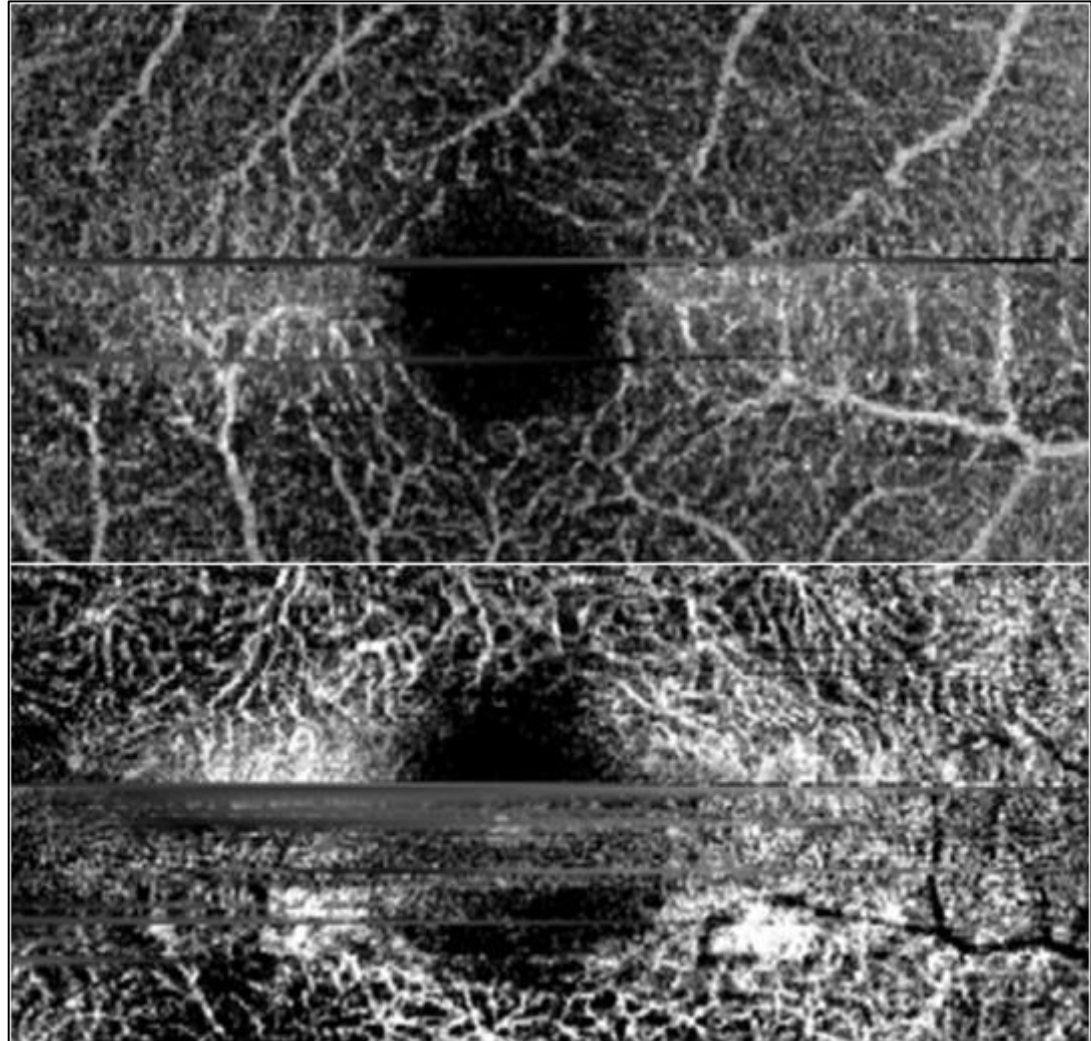
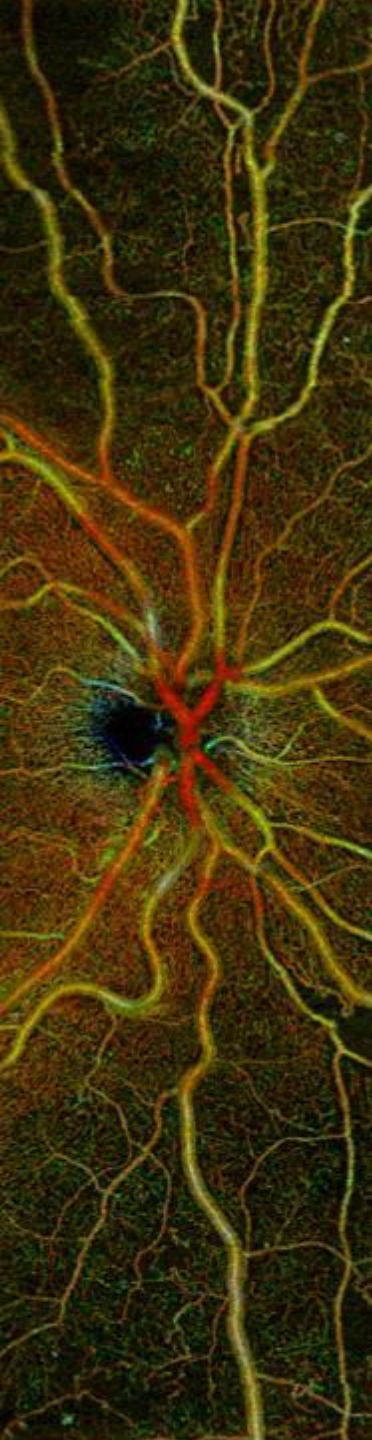


Step 3: Look at the segmentation lines

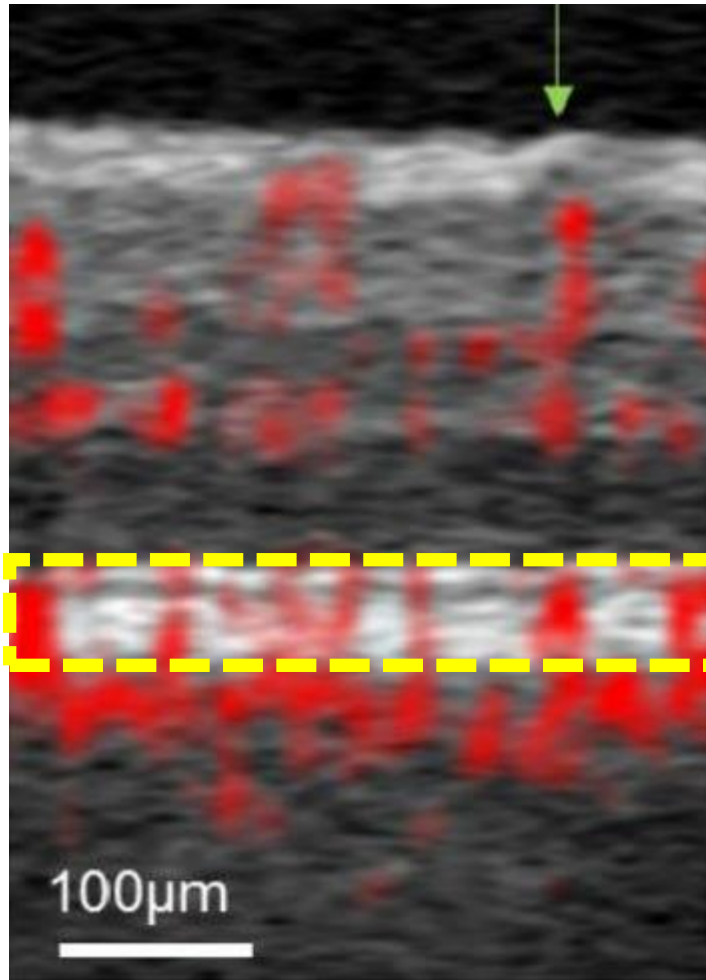
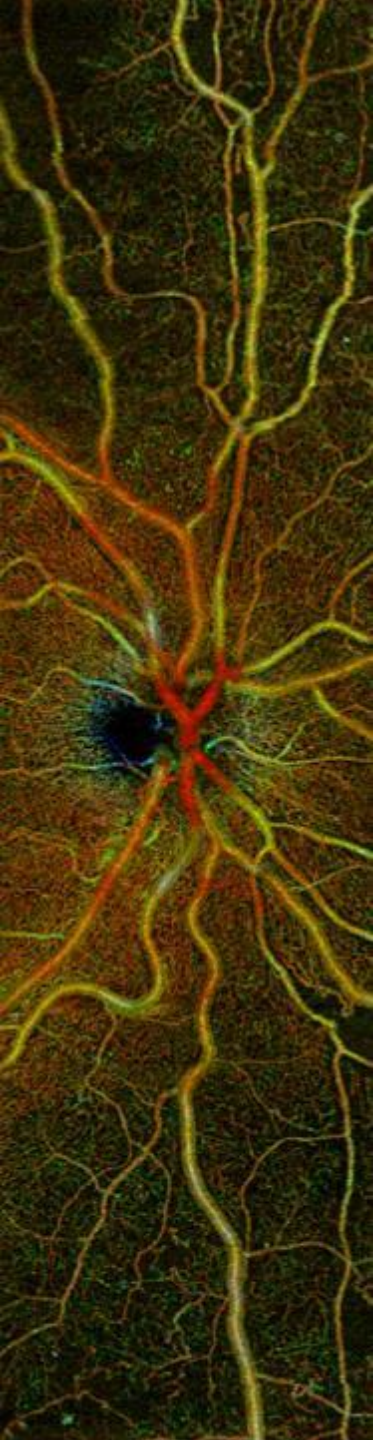


PMID: 33292740

# ARTIFACTS- Motion

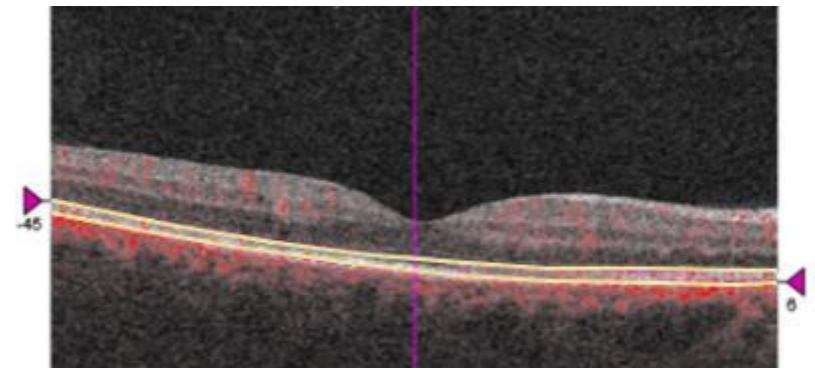
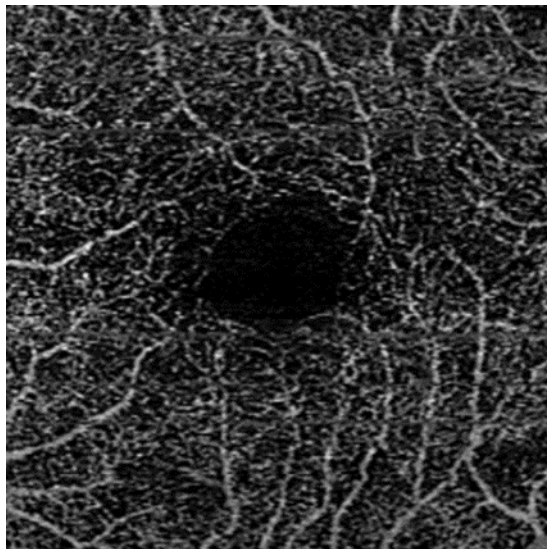
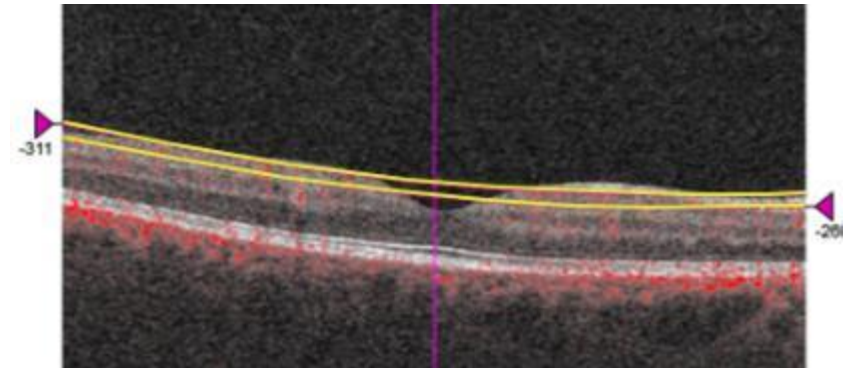
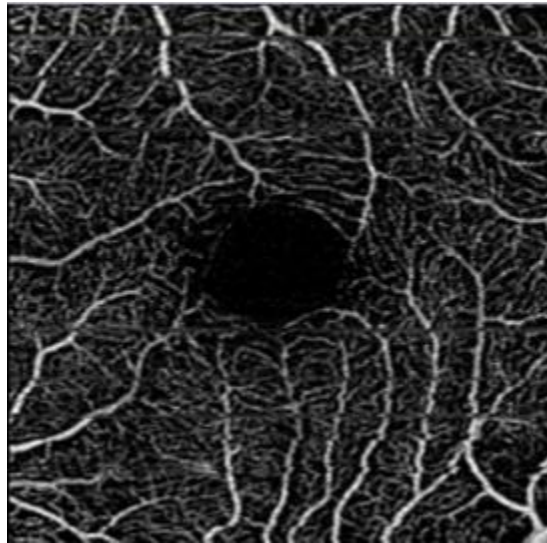
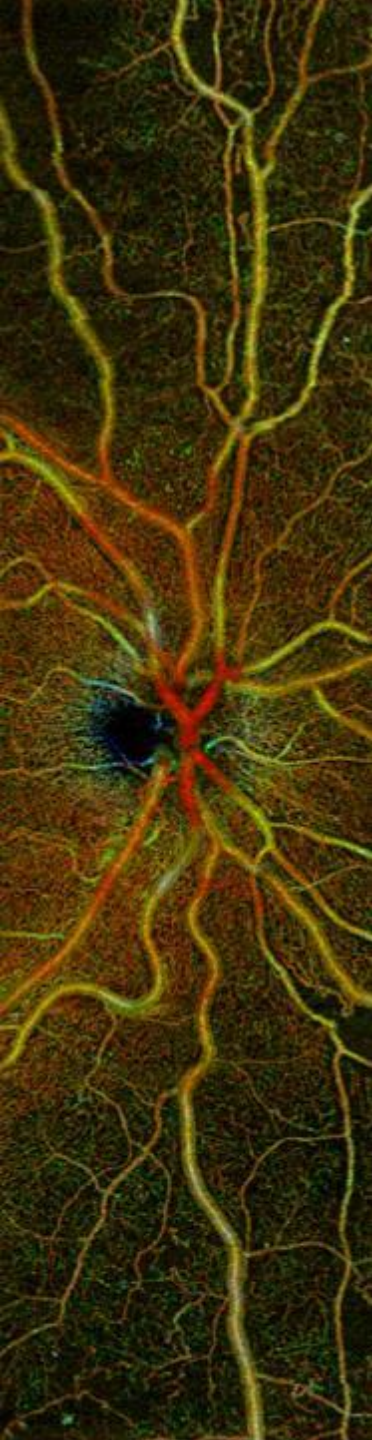


# ARTIFACTS- Projection



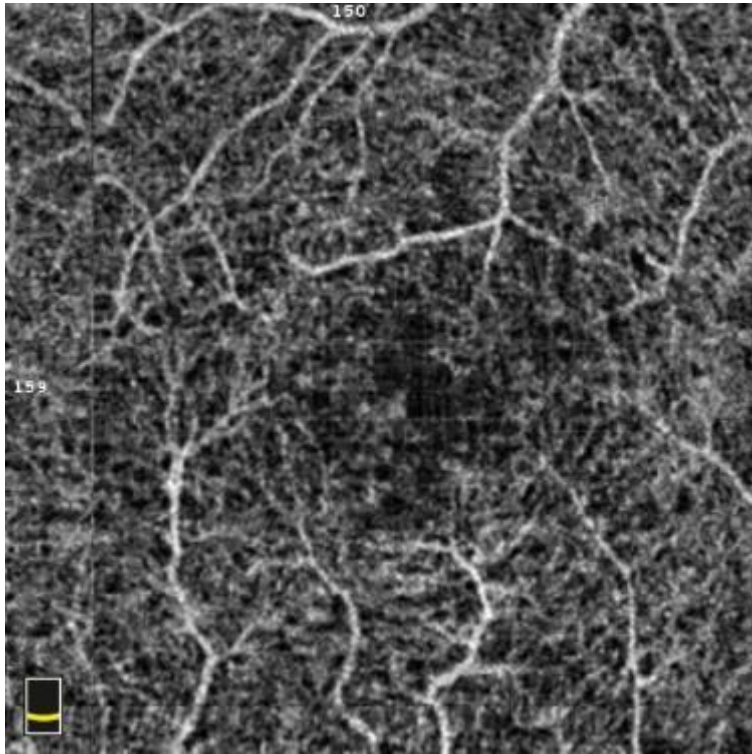
Spaide R, et al. Image Artifacts in OCTA. Retina. 35(11):2163-2180.

# ARTIFACTS- Projection

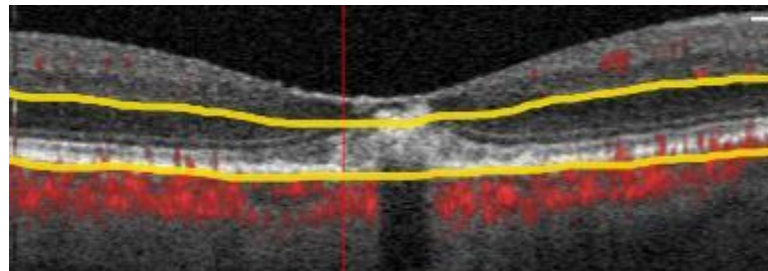
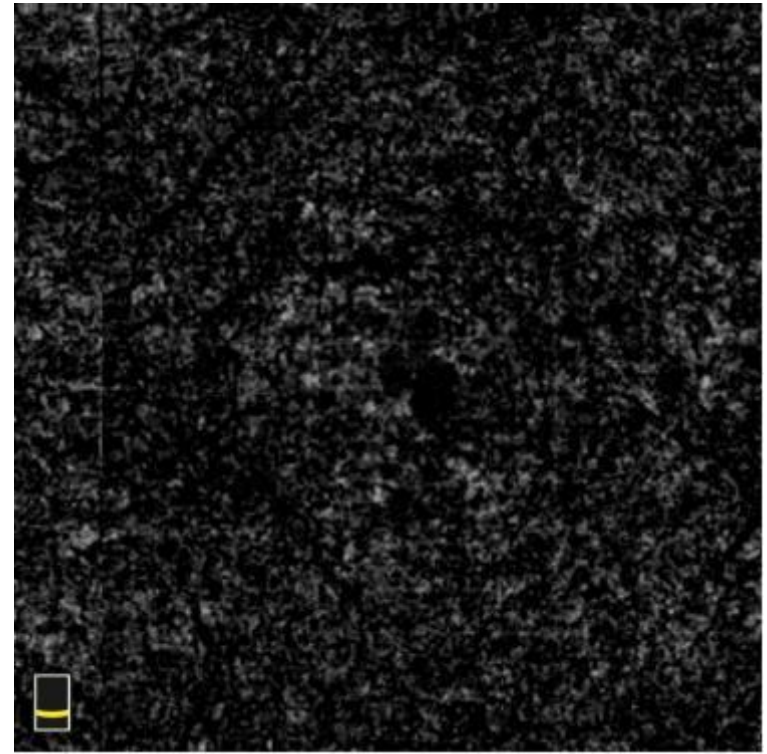


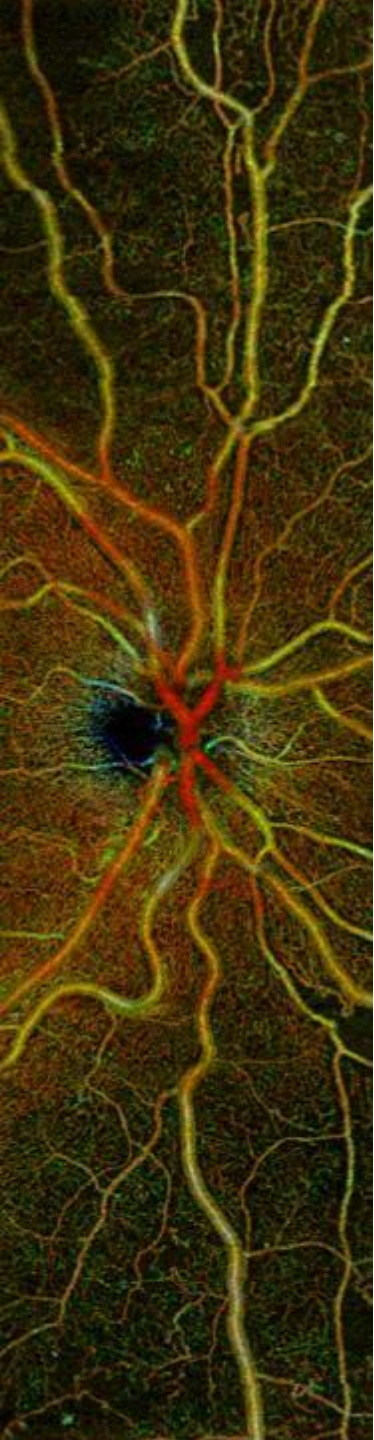
# ARTIFACTS- Projection

Without projection  
artifact removal



With projection  
artifact removal





# QUANTITATIVE ANALYSIS

## FAZ Area

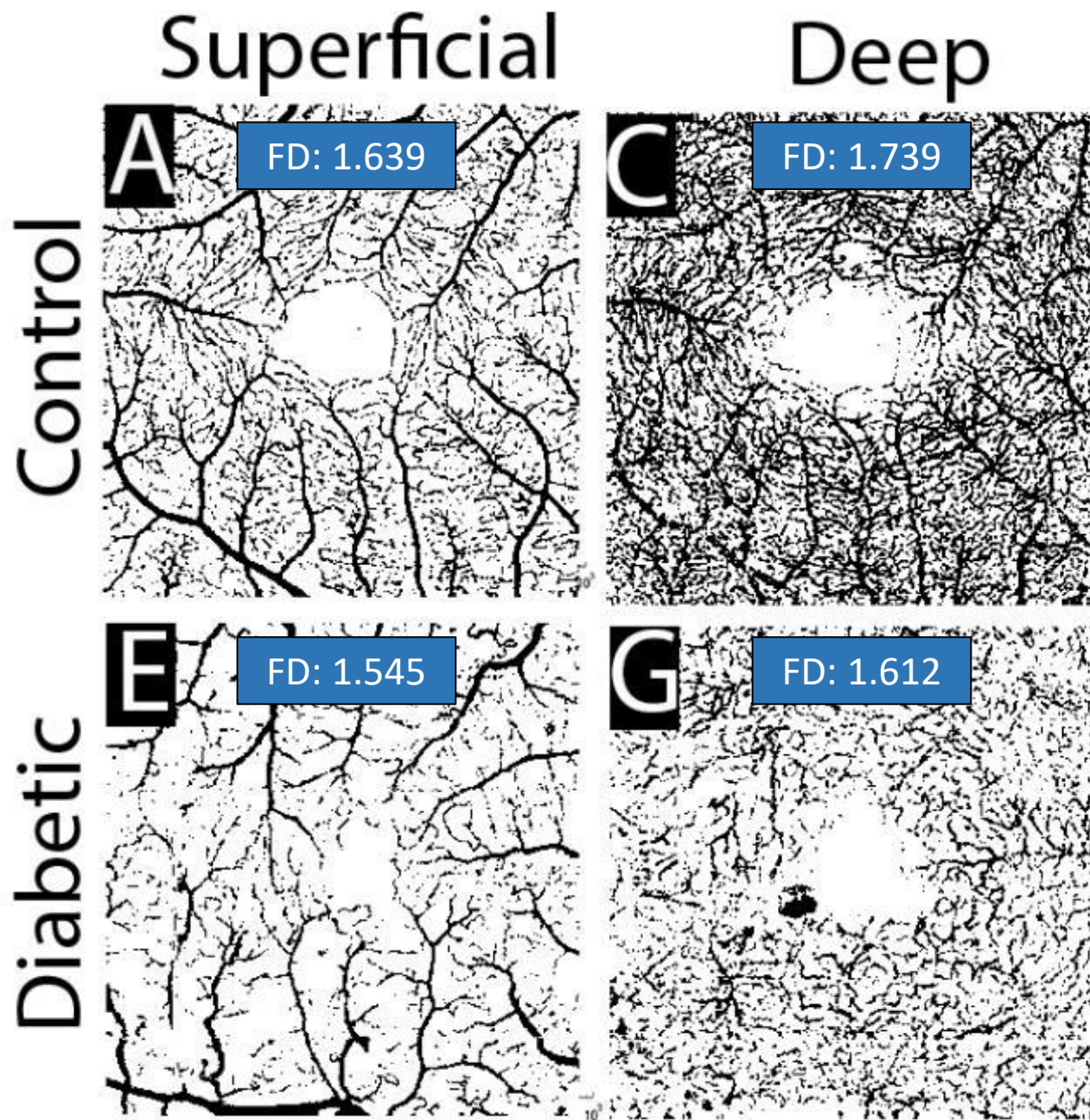
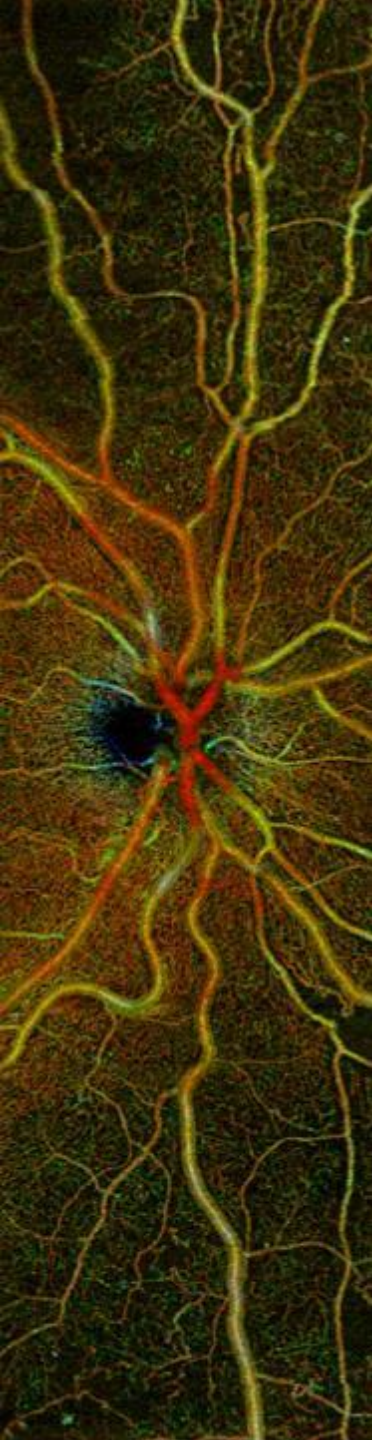
- FAZ area at the level of the superficial capillary plexus

## Vessel Density

- Proportion of image area occupied by blood vessels

## Fractal Dimension

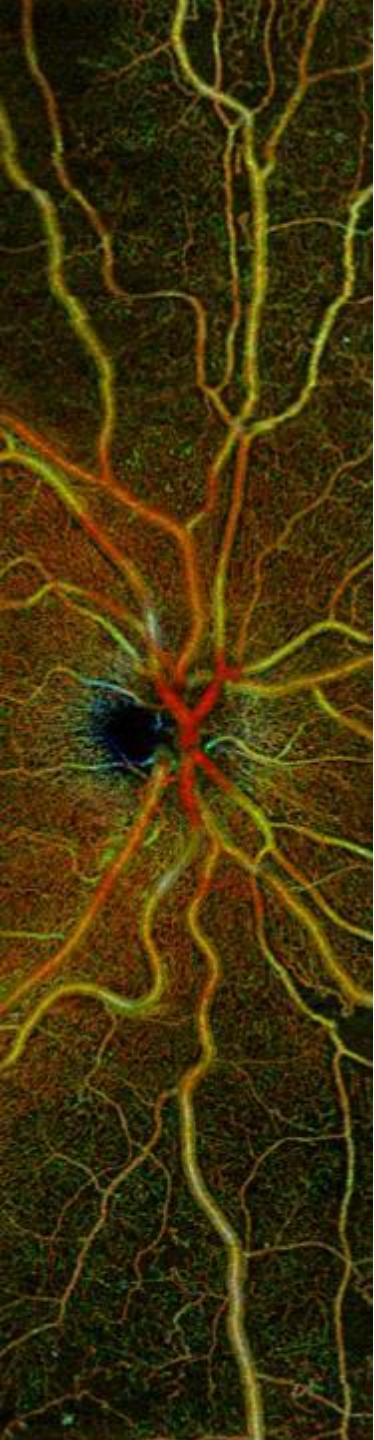
- A mathematical parameter that describes the degree of complexity of the vascular pattern
- 1 = low complexity; 2 = high complexity



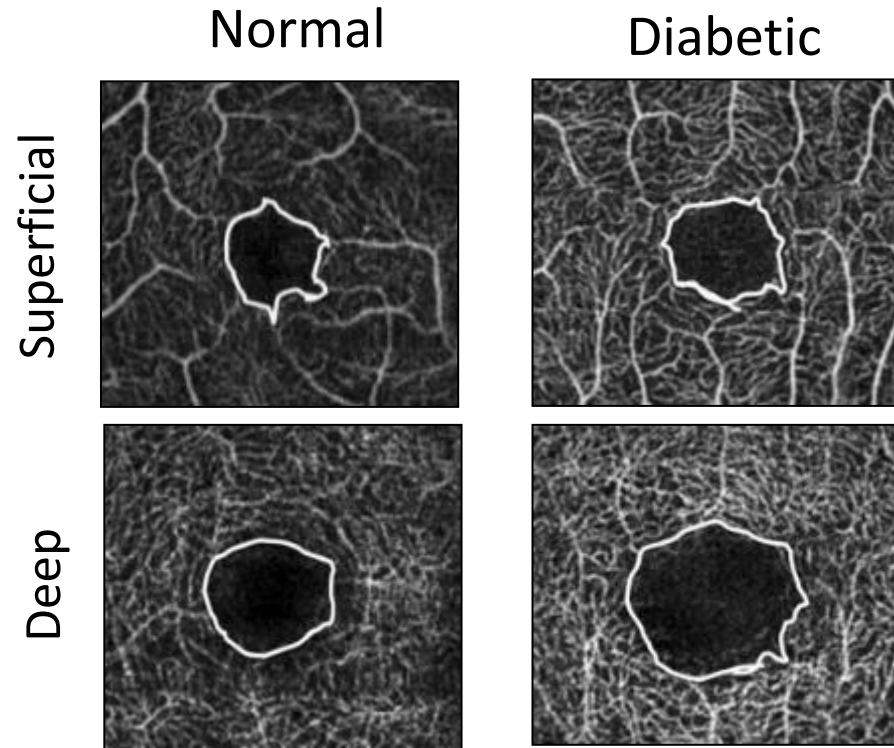
# DIABETIC RETINOPATHY

## OCTA Clinical Applications

- Early diabetic retinopathy detection
- Identify vascular abnormalities (IRMA)
- Identify non-perfusion
- Evaluation of FAZ
- Visualize the deep capillary plexus
- Differentiate IRMA from early NVE
- Localization of microaneurysms
- Follow response to treatment



# DIABETIC RETINOPATHY

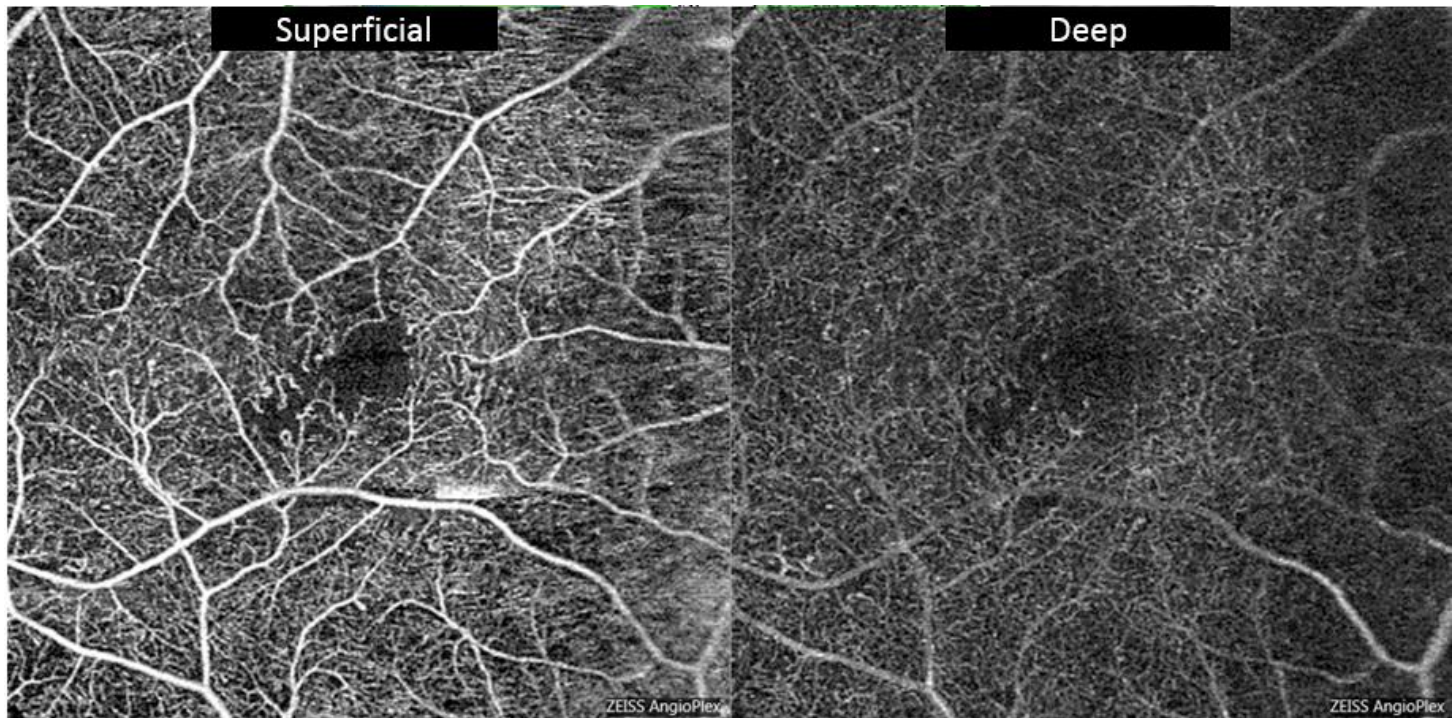
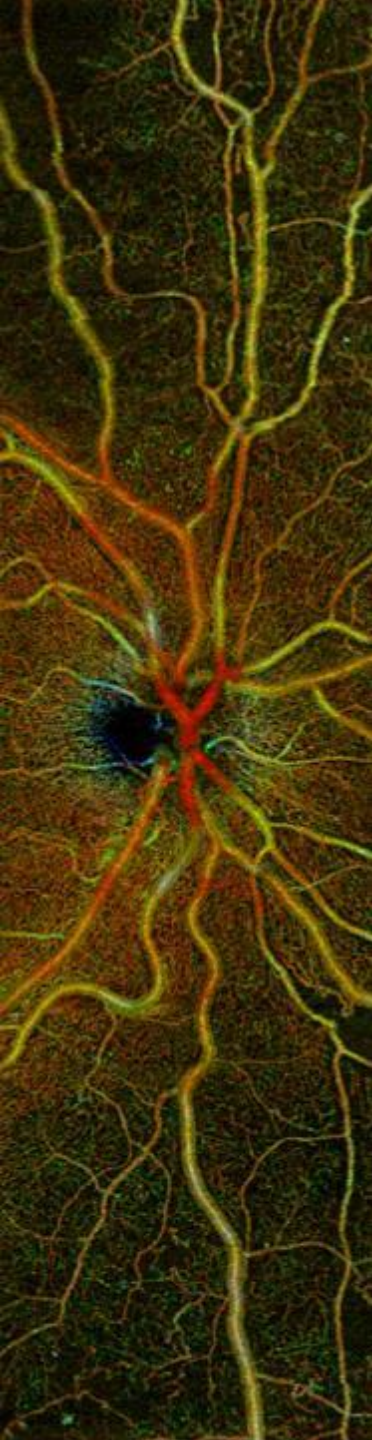


|                                                                 | Control | Diabetics Without DR | Diabetic Retinopathy |
|-----------------------------------------------------------------|---------|----------------------|----------------------|
| <b>Mean FAZ area in the superficial plexus (mm<sup>2</sup>)</b> | 0.25    | 0.37 (p<0.01*)       | 0.38 (p<0.01*)       |
| <b>Mean FAZ area in the deep plexus (mm<sup>2</sup>)</b>        | 0.38    | 0.54 (p<0.01*)       | 0.56 (p<0.01*)       |

\*compared to controls (no sig diff between diabetic grps)

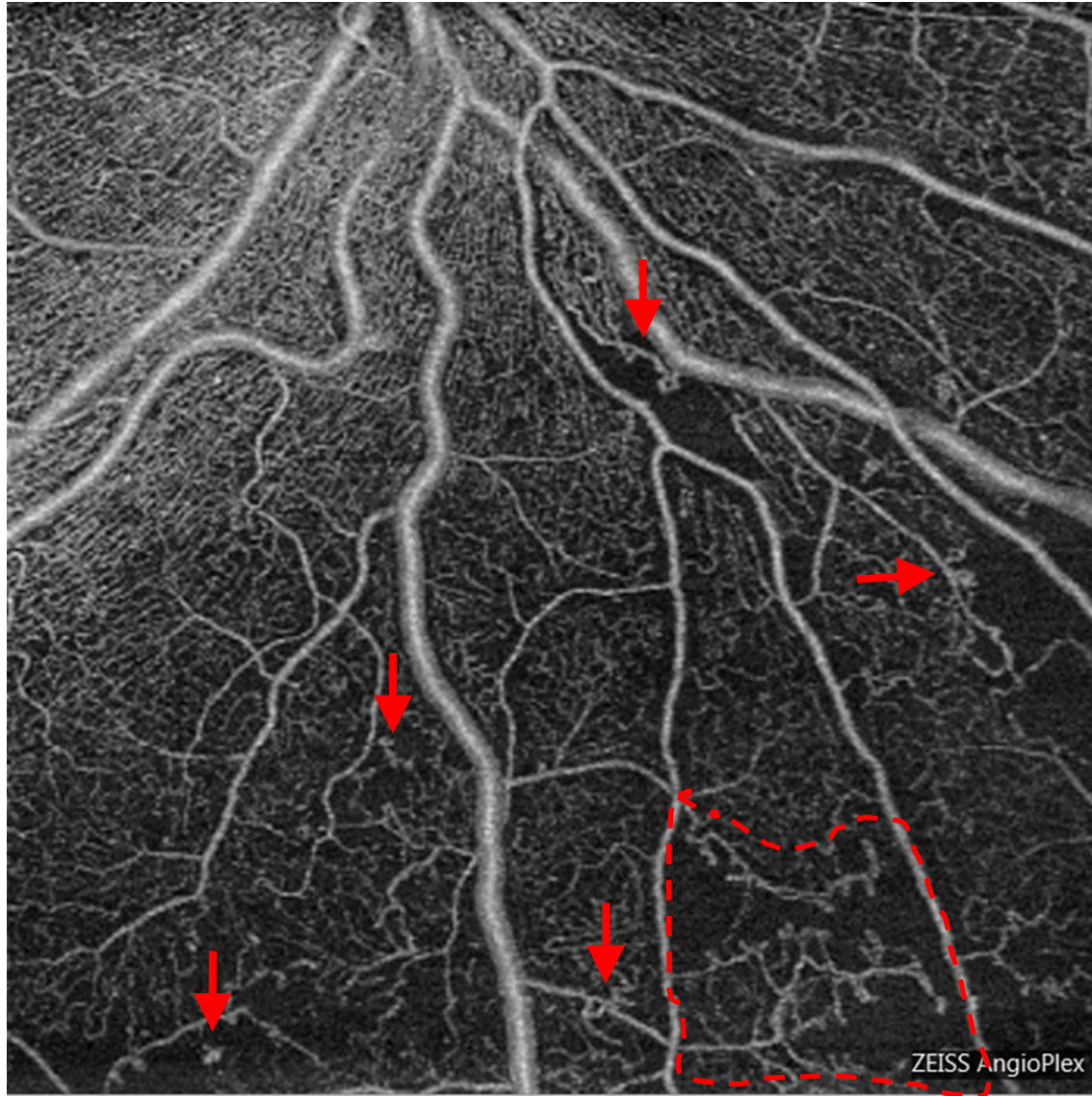
# DIABETIC RETINOPATHY

Mild NPDR with reduction in BCVA



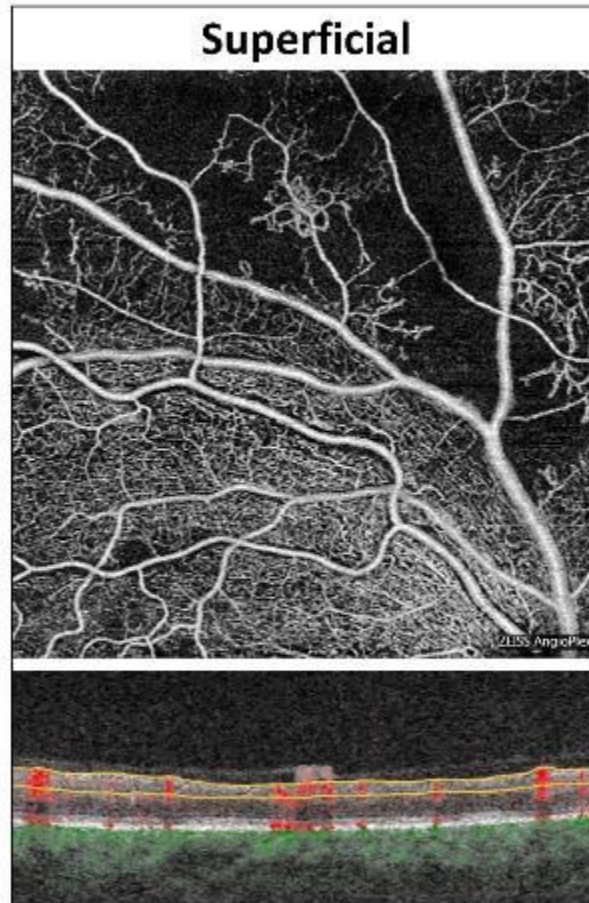
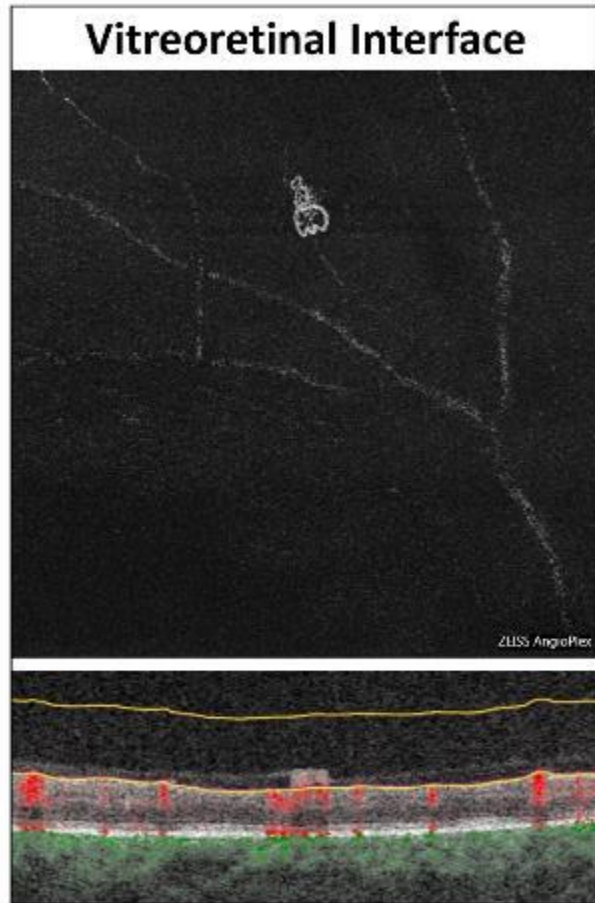
# DIABETIC RETINOPATHY

## Intraretinal microvascular abnormalities (IRMA)



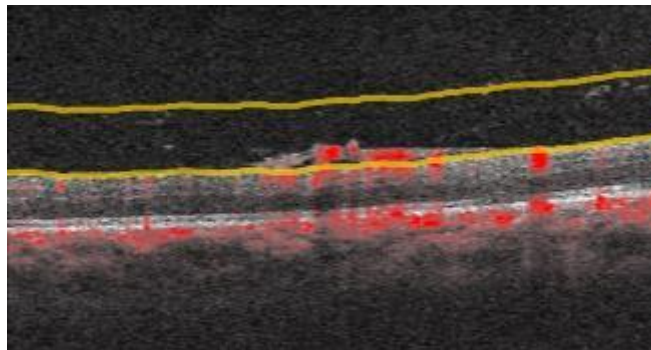
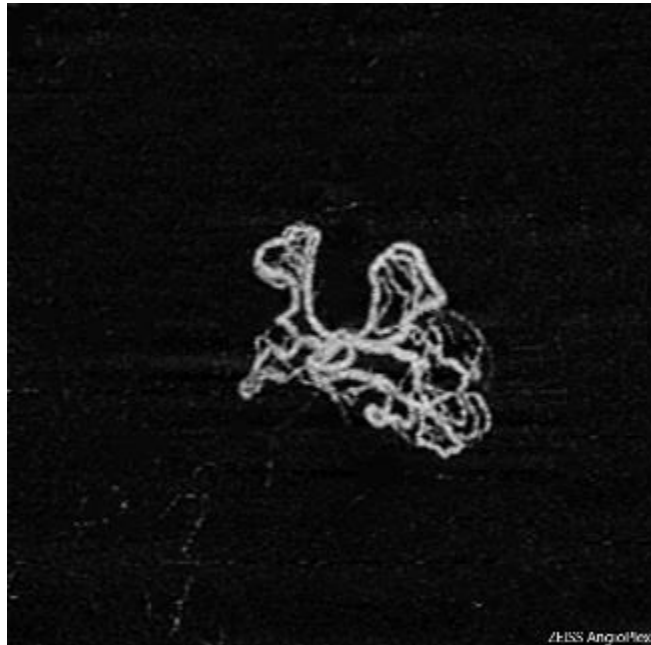
# DIABETIC RETINOPATHY

IRMA or early NVE???

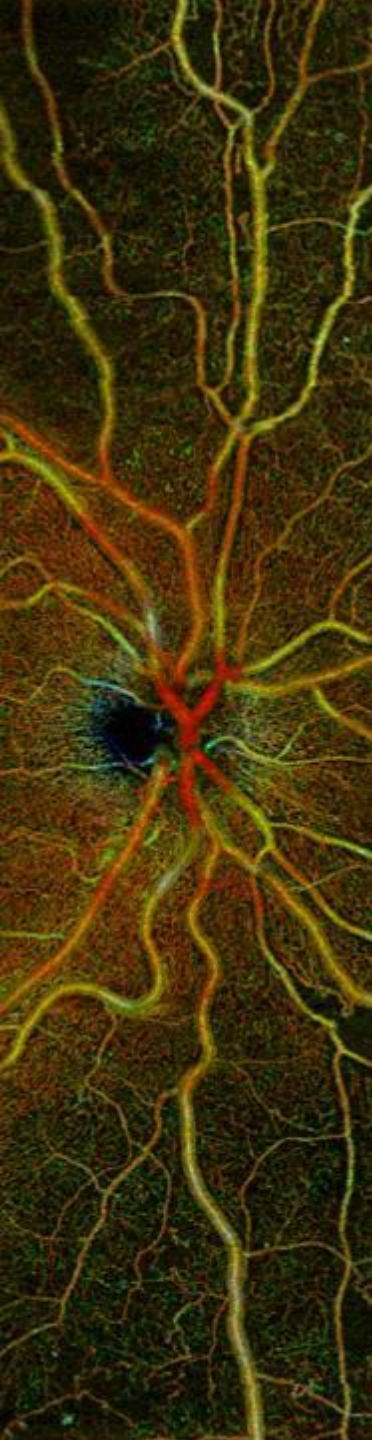
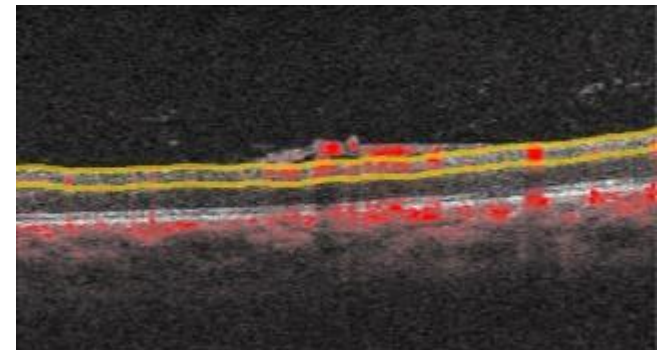
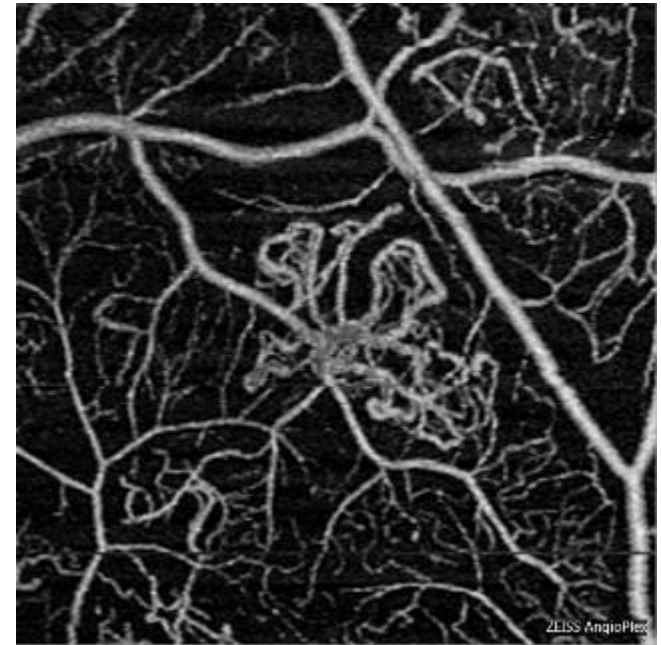


# DIABETIC RETINOPATHY

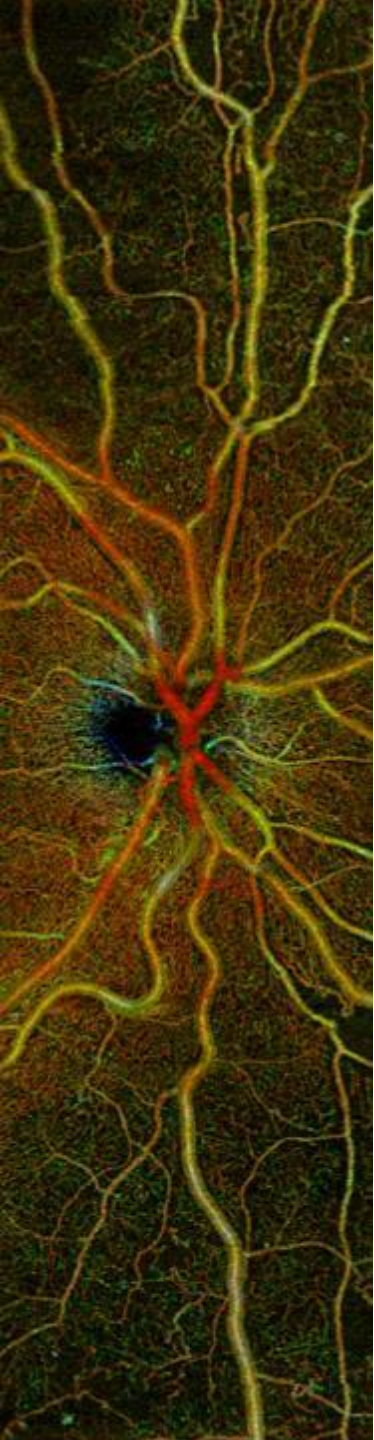
VRI



Superficial



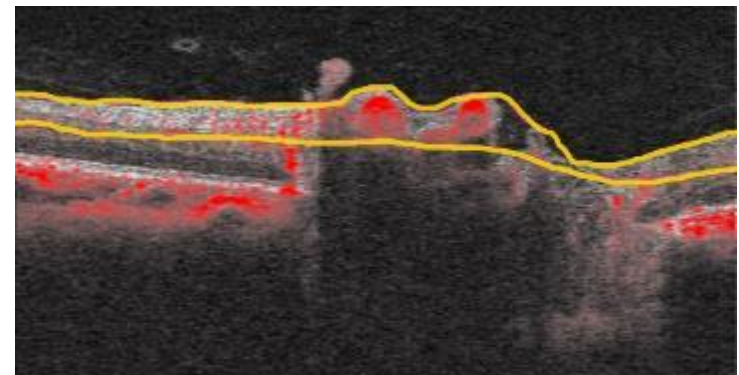
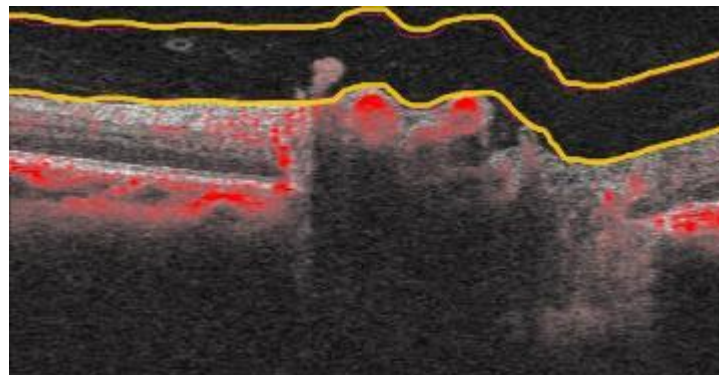
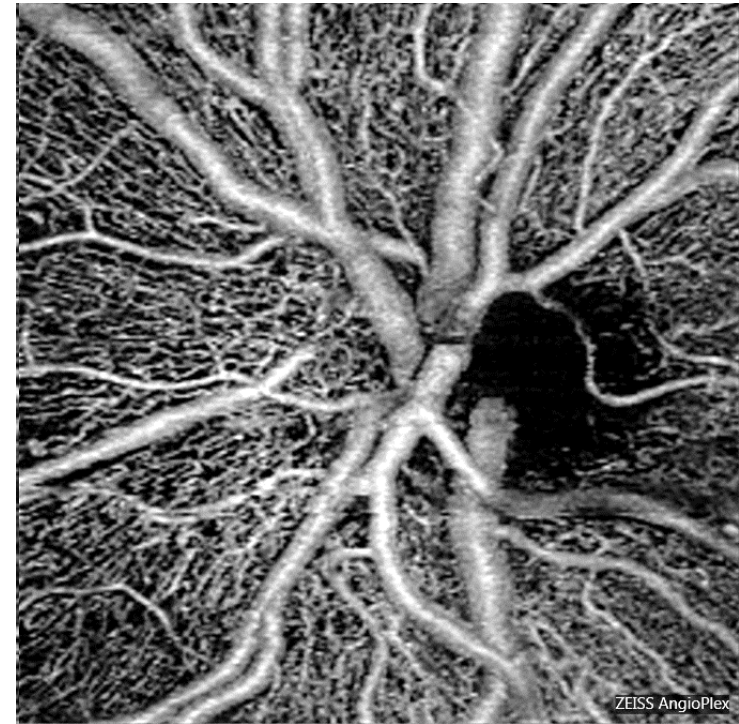
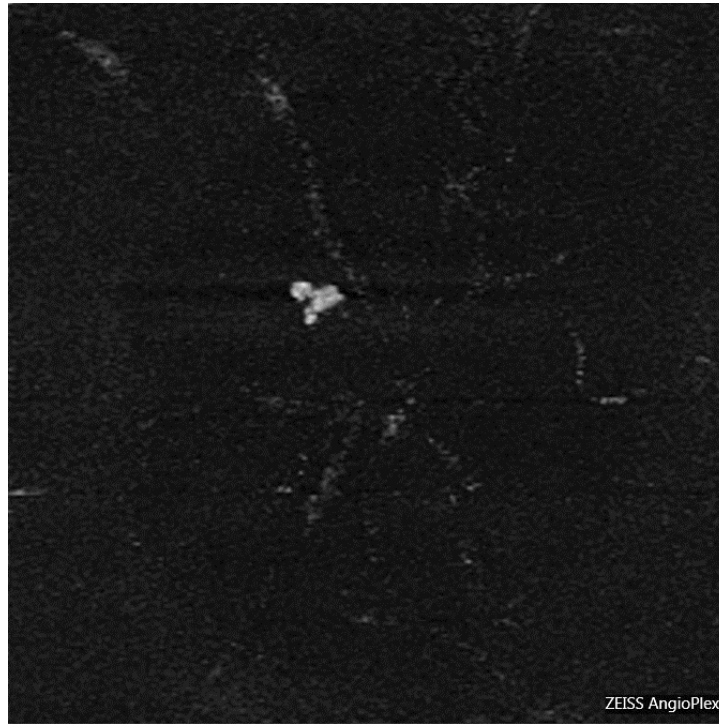
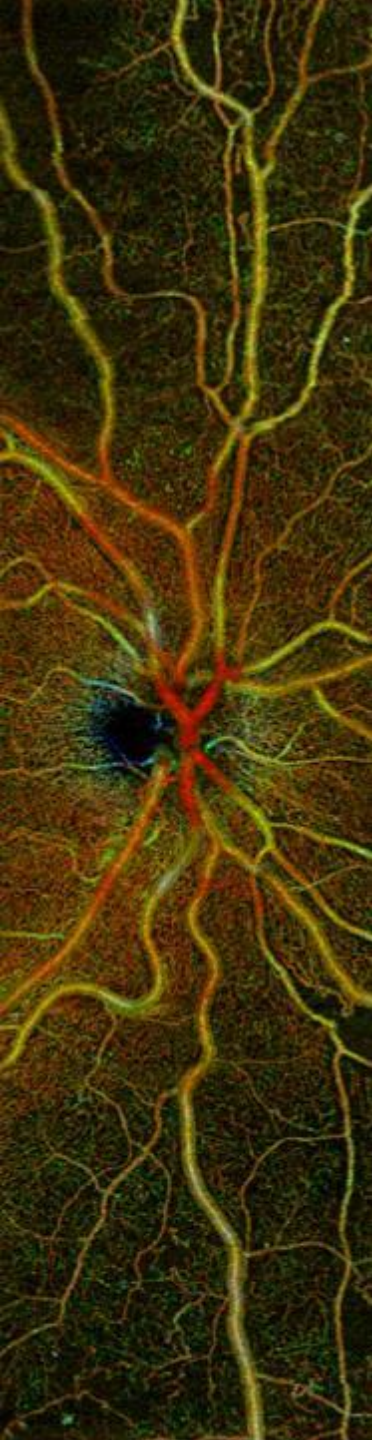
# DIABETIC RETINOPATHY



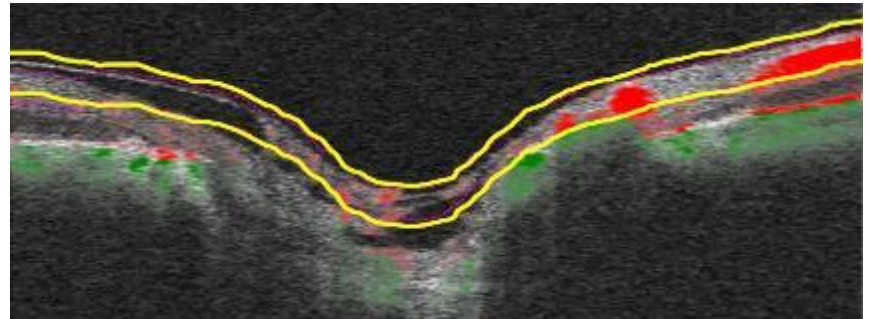
# DIABETIC RETINOPATHY

VRI

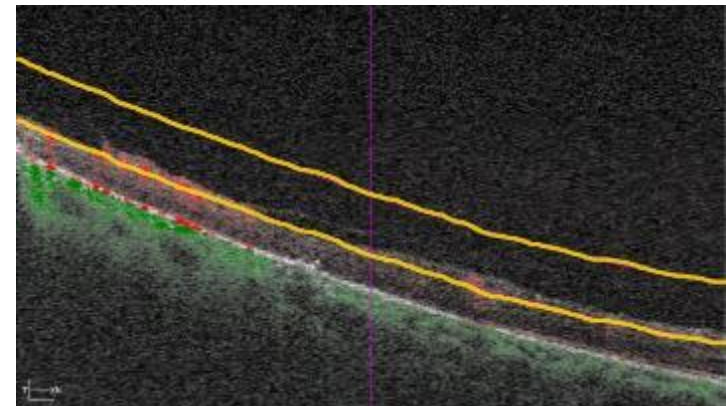
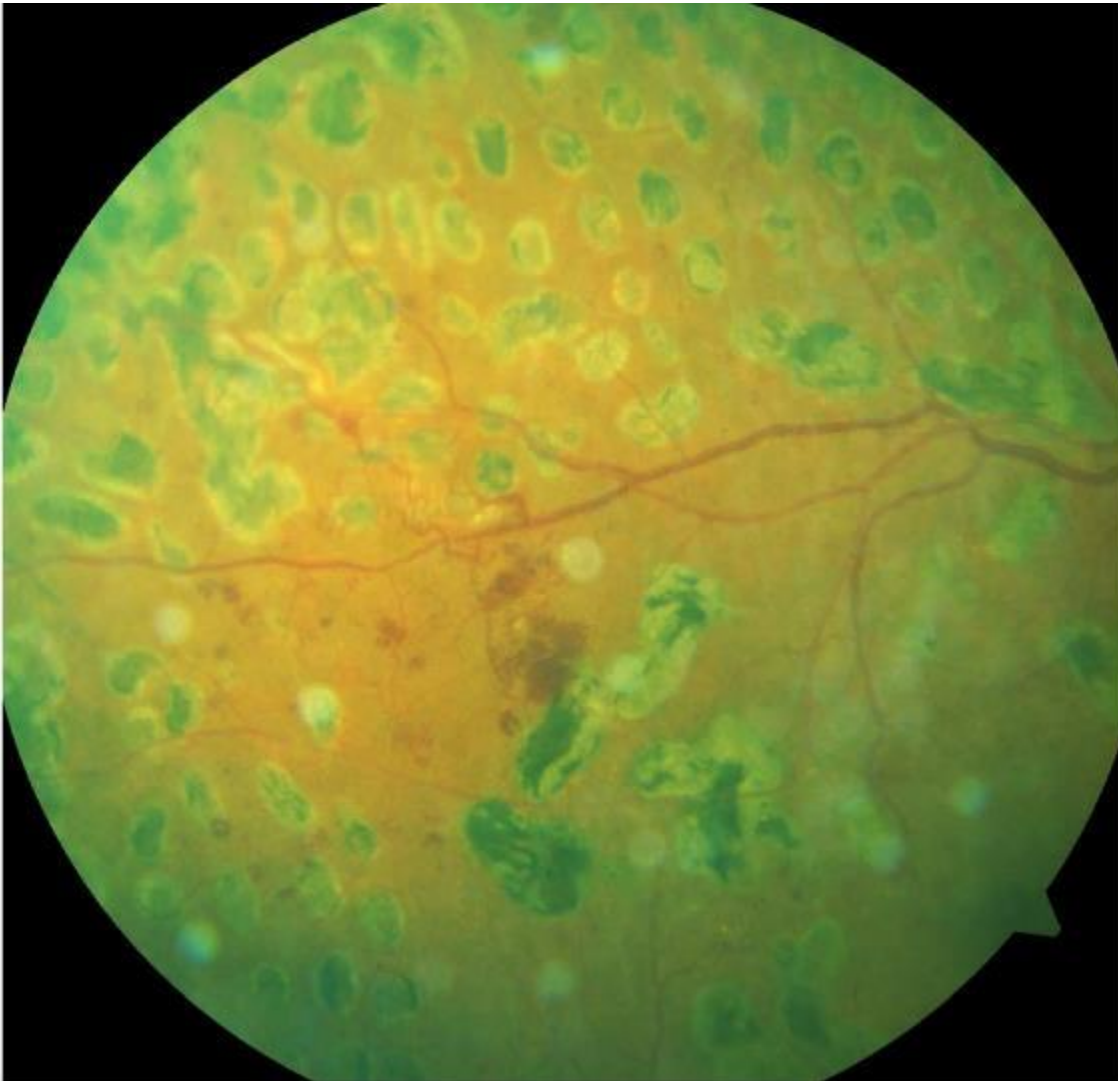
Superficial



# DIABETIC RETINOPATHY



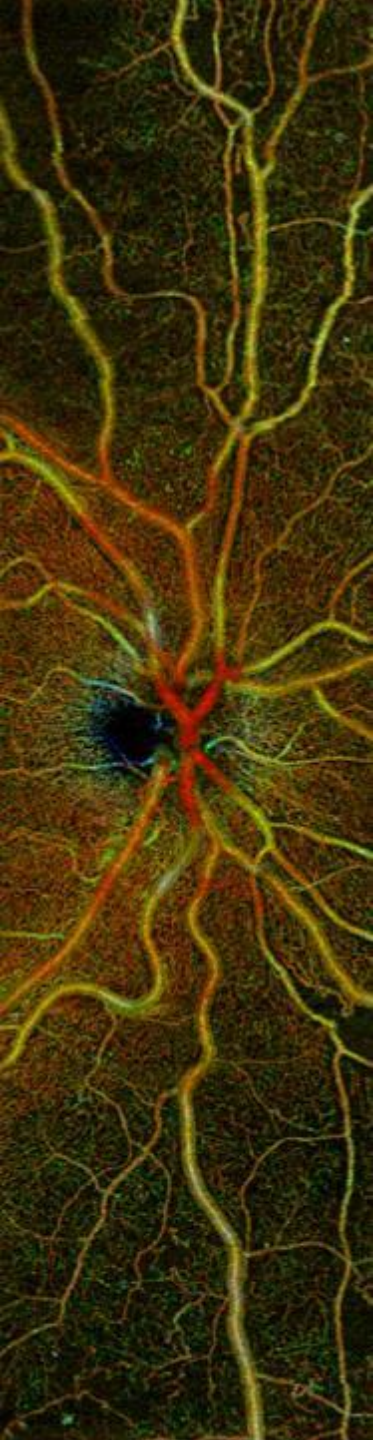
# DIABETIC RETINOPATHY



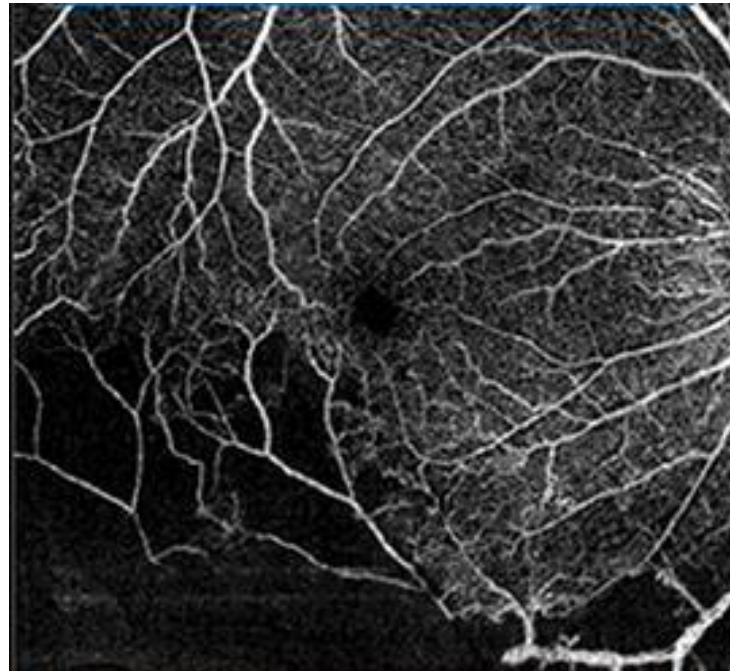
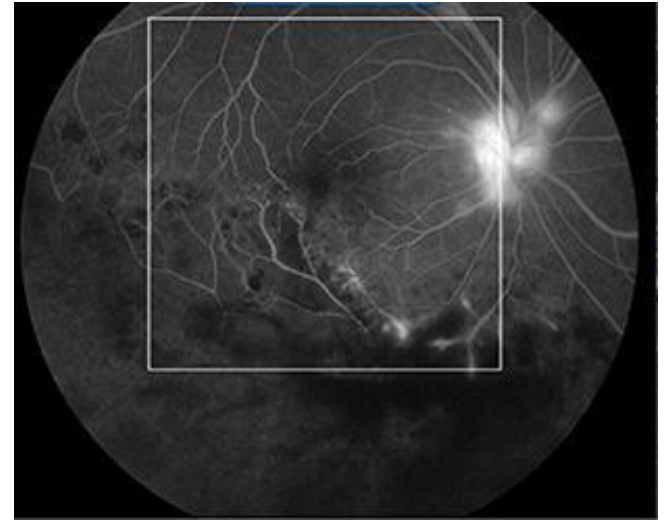
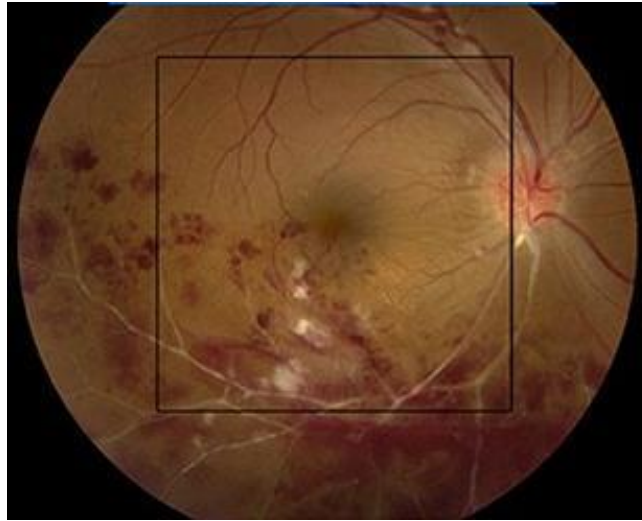
# VENOUS OCCLUSION

## OCTA Clinical Applications

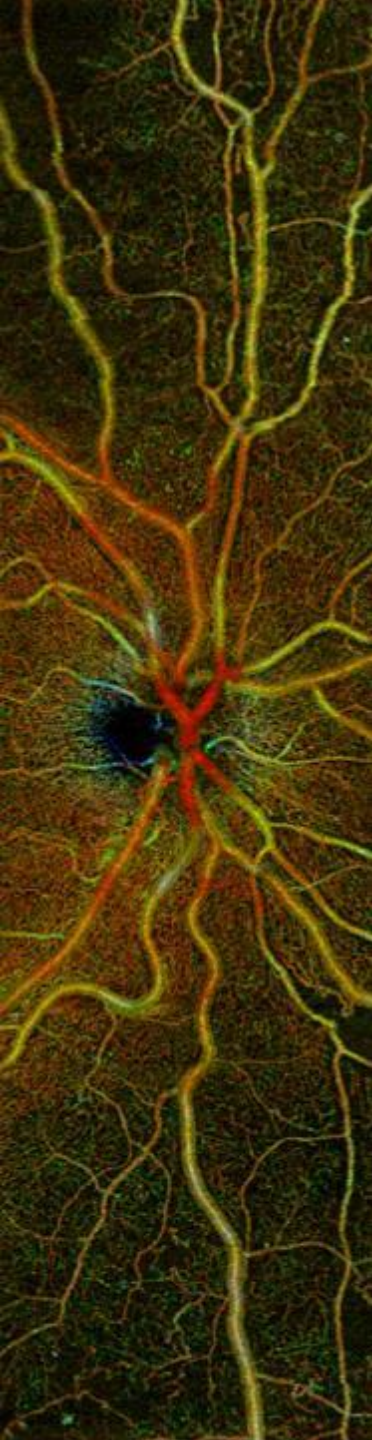
- Quantify non-perfusion
  - Ischemic vs non-ischemic?
  - BRVO: Risk of post segment neo
  - CRVO: Risk of ant segment neo and NVG
- Visualization of macular ischemia
- Earlier detection of posterior segment neo



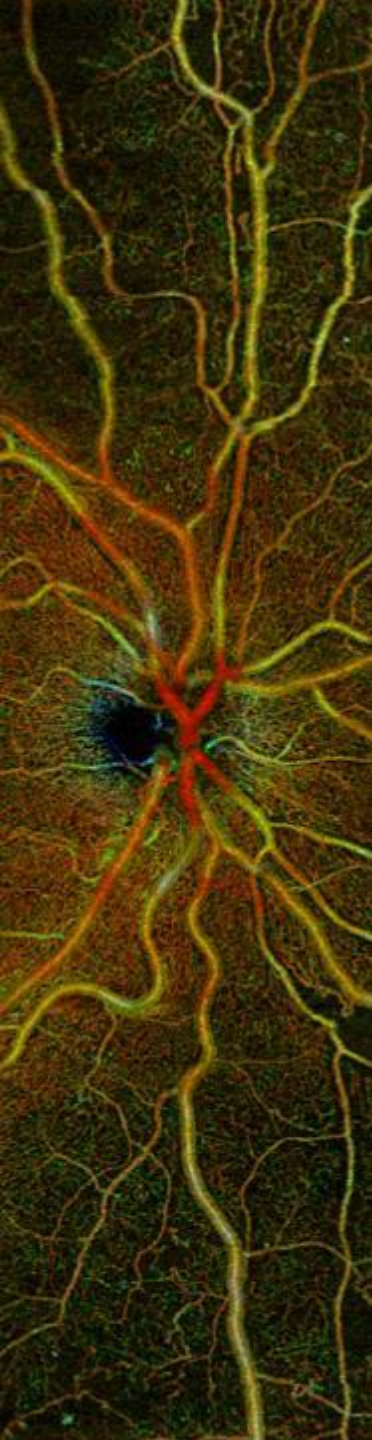
# BRVO - Ischemic

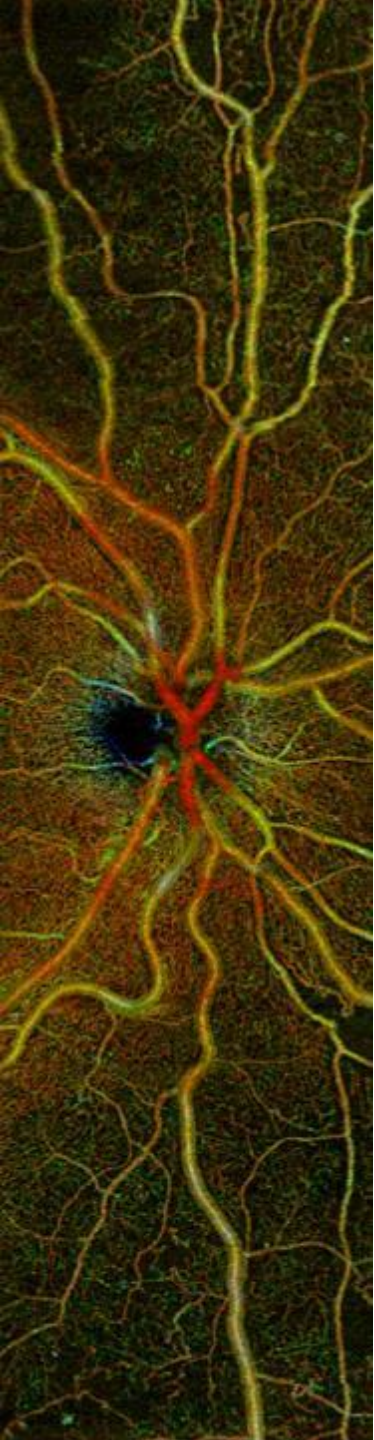


# BRVO – Nonischemic



# BRVO – Nonischemic





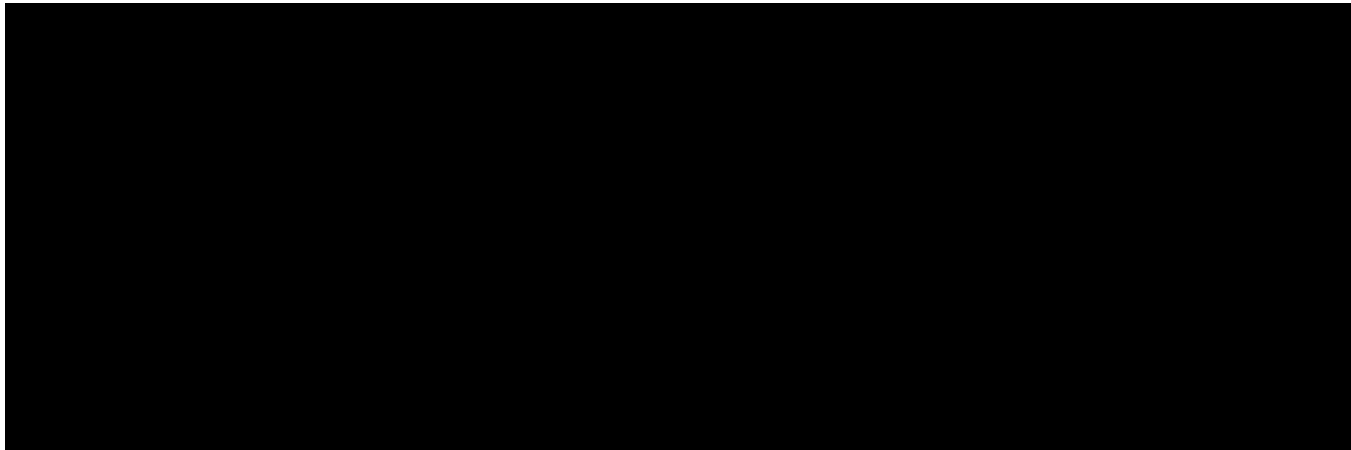
# Retinal Vein Occlusion

## ***CASE REPORT***

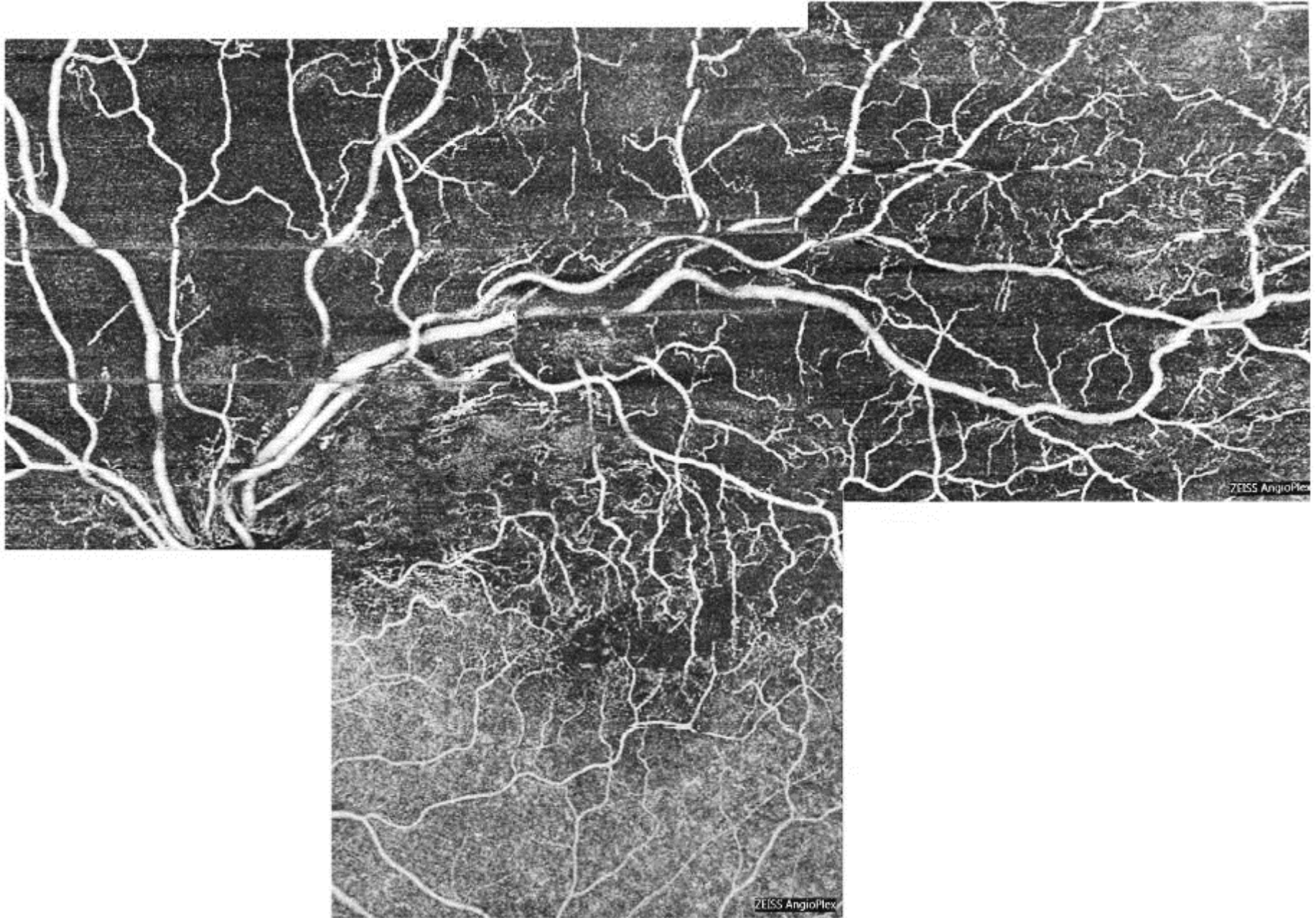
52yo HM

- Referred to our retinal service for evaluation of RVO OS
- Pt c/o blur OS x 5 days – getting worse
- BCVA: 20/25 OD, 20/200 OS
- Ta 19/14
- SLE: WNL OU, No NVI

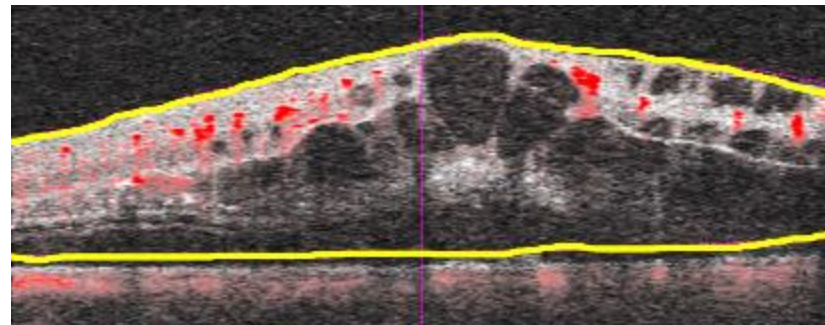
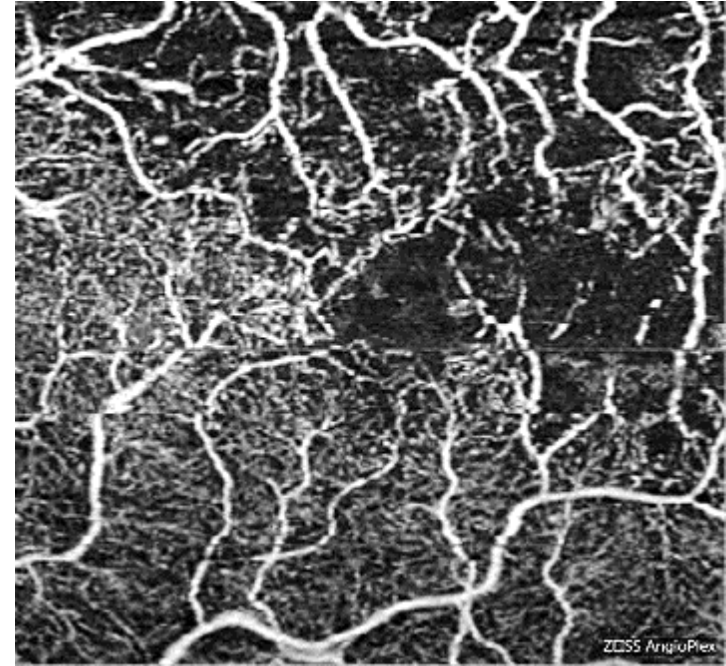
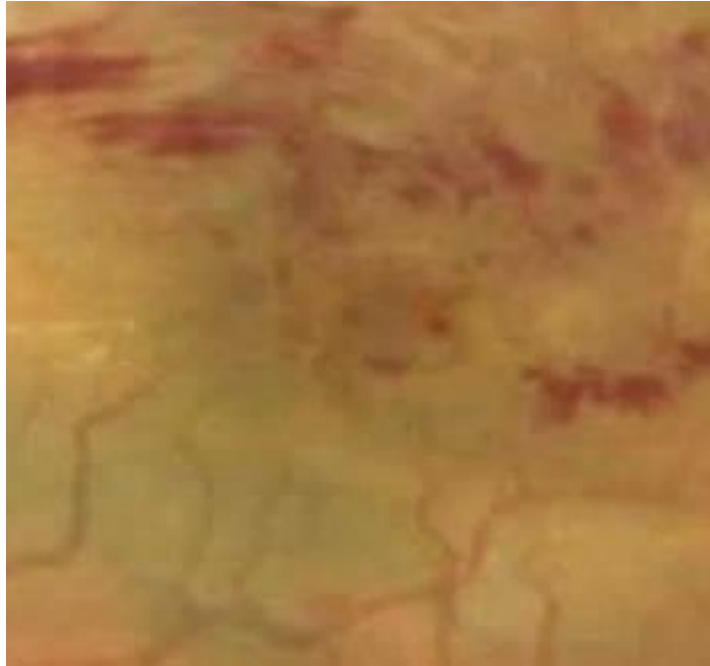
# Hemicentral Vein Occlusion



# Hemicentral Vein Occlusion



# Hemicentral Vein Occlusion



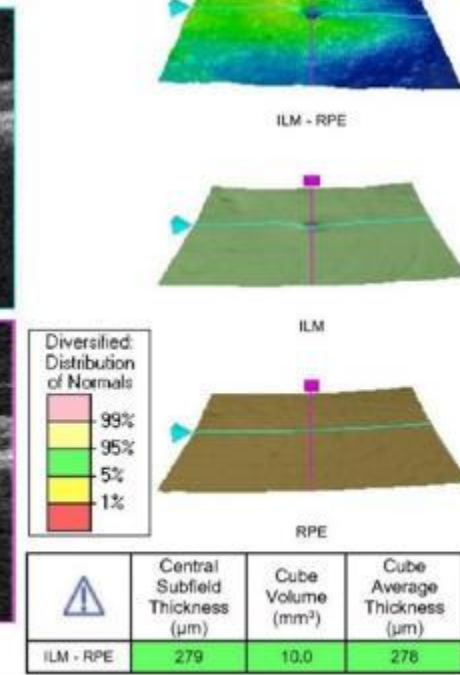
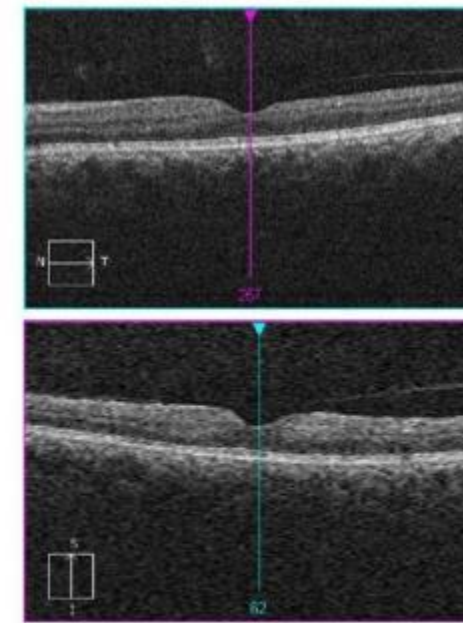
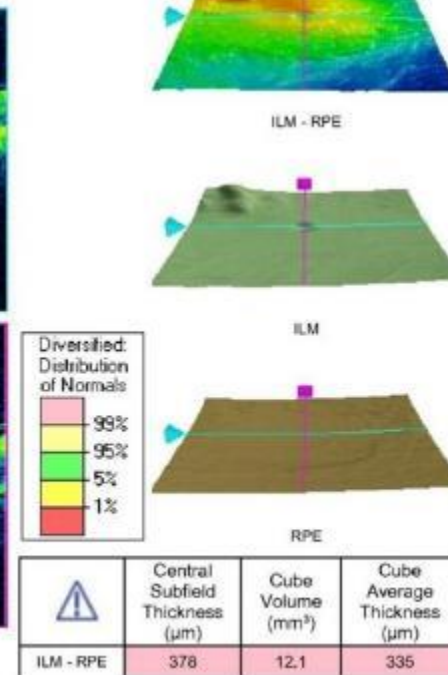
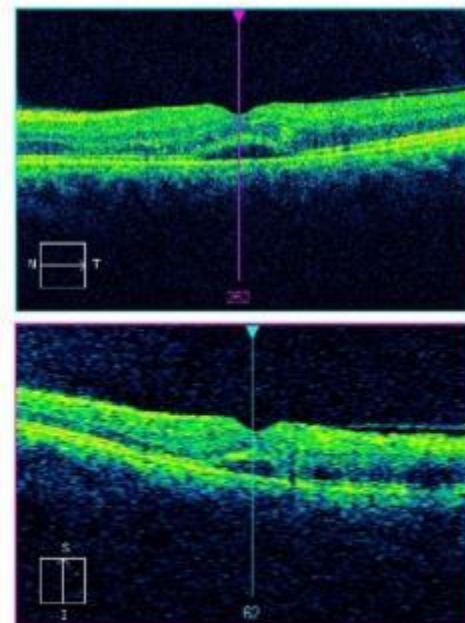
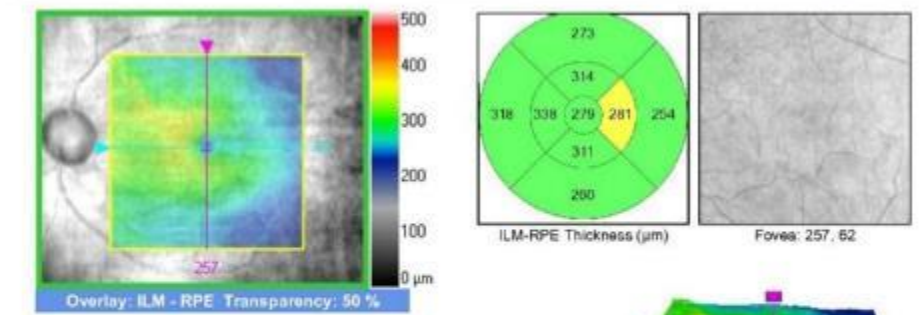
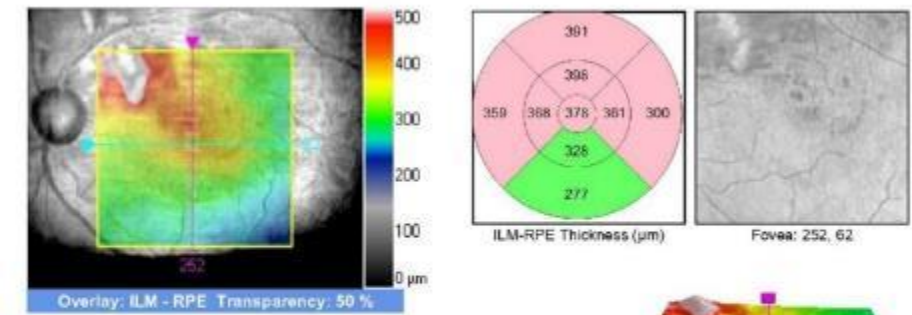
# Hemicentral Vein Occlusion

S/P IVB x 1

S/P IVB x 3 (5 mos after presentation)

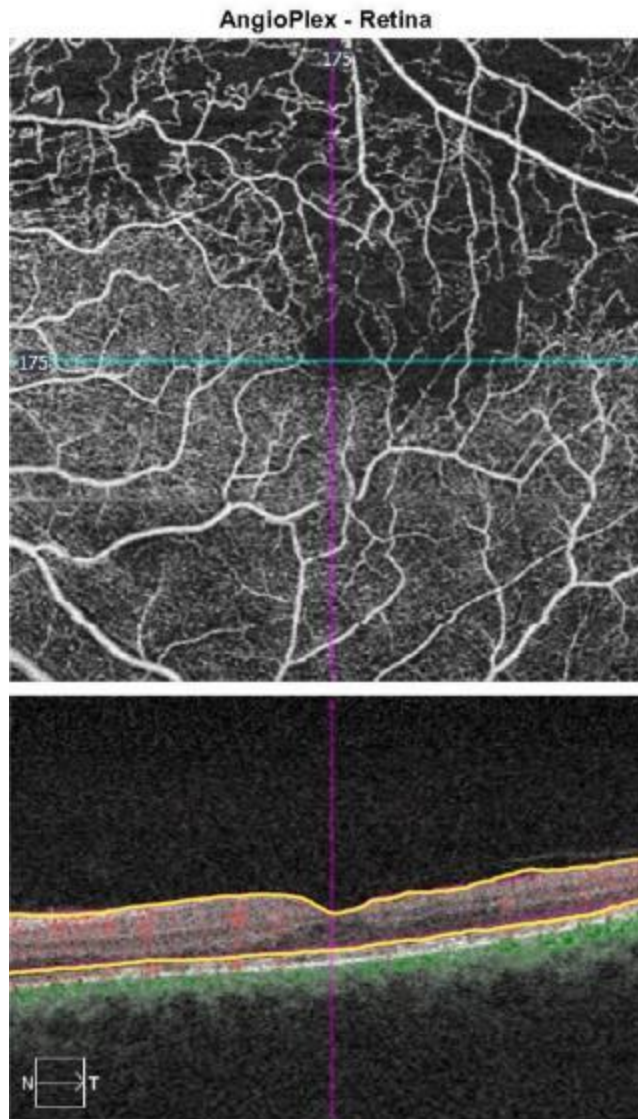
Macula Thickness : Macular Cube 512x128 OD ☐ OS ☒

Macula Thickness : Macular Cube 512x128 OD ☐ OS ☒



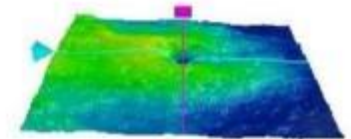
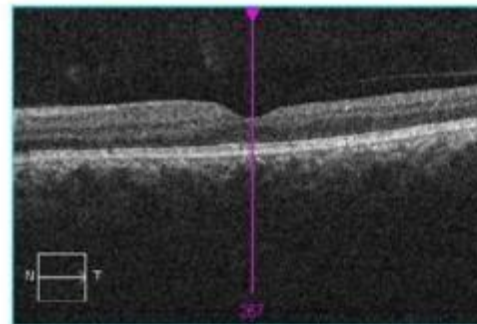
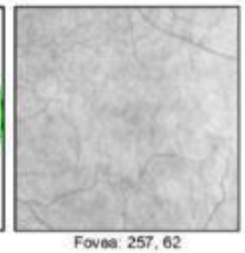
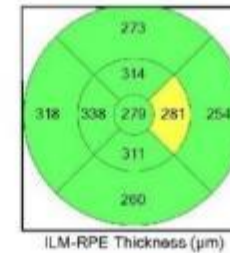
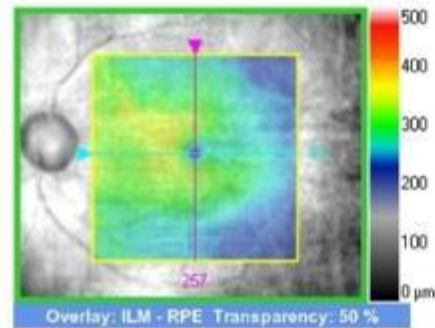
# Hemicentral Vein Occlusion

BCVA 20/40

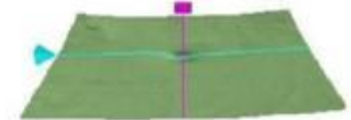


Macula Thickness : Macular Cube 512x128

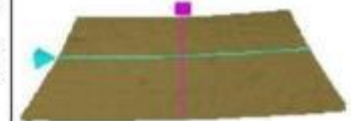
OD ☐ OS ☒



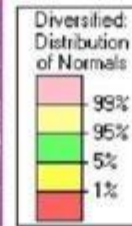
ILM - RPE



ILM

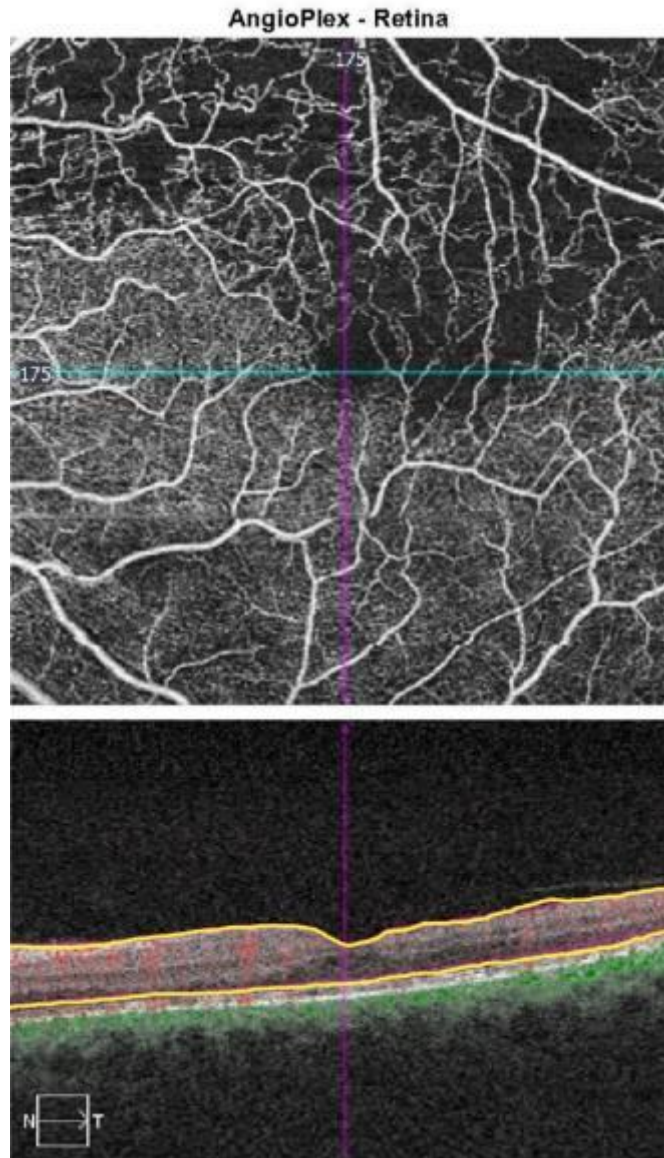


RPE



|           | Central Subfield Thickness ( $\mu\text{m}$ ) | Cube Volume ( $\text{mm}^3$ ) | Cube Average Thickness ( $\mu\text{m}$ ) |
|-----------|----------------------------------------------|-------------------------------|------------------------------------------|
| ILM - RPE | 279                                          | 10.0                          | 276                                      |

# Hemicentral Vein Occlusion



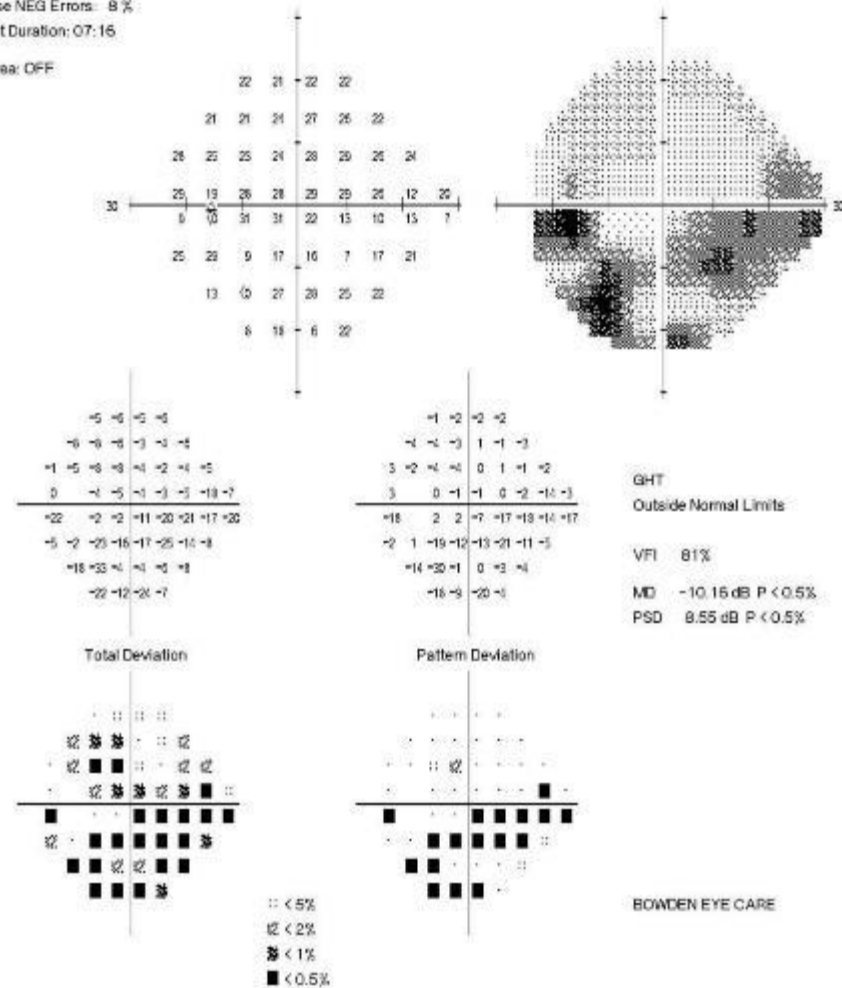
### Central 24-2 Threshold Test

Fixation Monitor: Gaze/Blind Spot  
Fixation Target: Central  
Fixation Losses: 0/19  
False POS Errors: 2 %  
False NEG Errors: 8 %  
Test Duration: 07:16

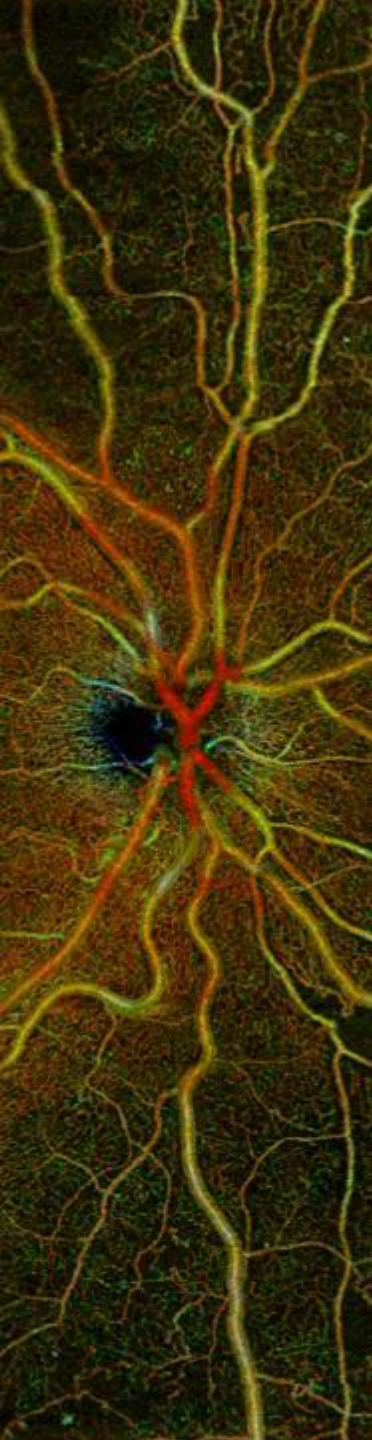
Fovea: OFF

Stimulus: III. White  
Background: 31.5 ASB  
Strategy: SITA-Standard

Pupil Diameter:  
Visual Acuity:  
RX: +2.00 DS      DC X



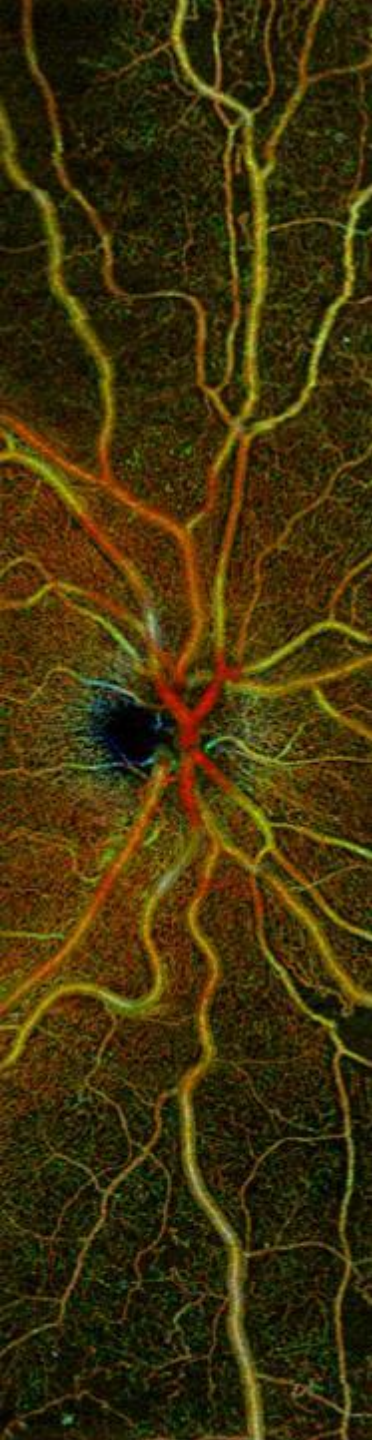
BOWDEN EYE CARE



# Retinal Vein Occlusion

## ***KEY POINTS***

- Vision loss from RVO may be a consequence of macular ischemia and/or edema
- More extensive retinal ischemia increases the risk of neovascular complications
- OCTA is an excellent tool for evaluating retinal perfusion and detecting preretinal neovascularization following RVO

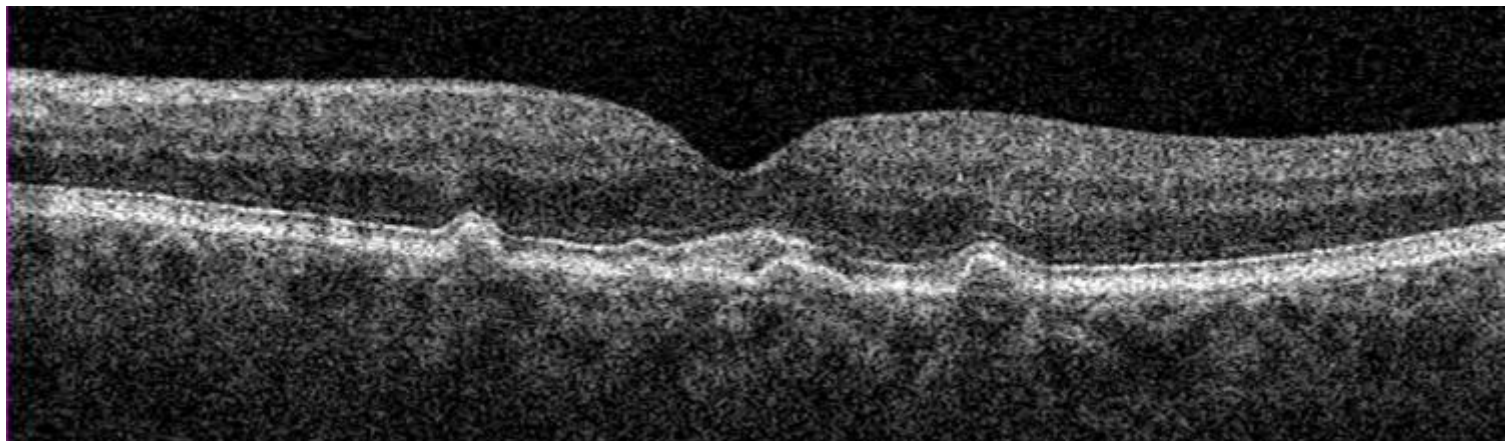
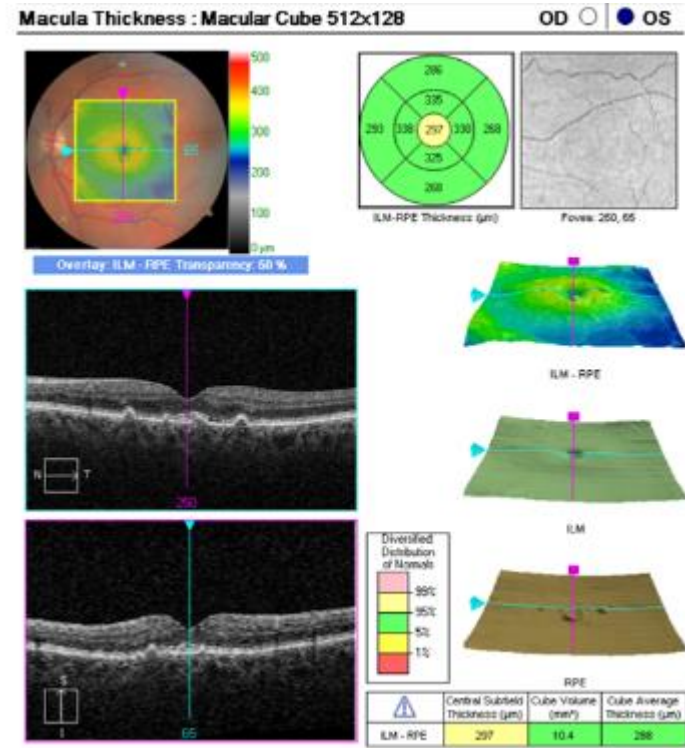
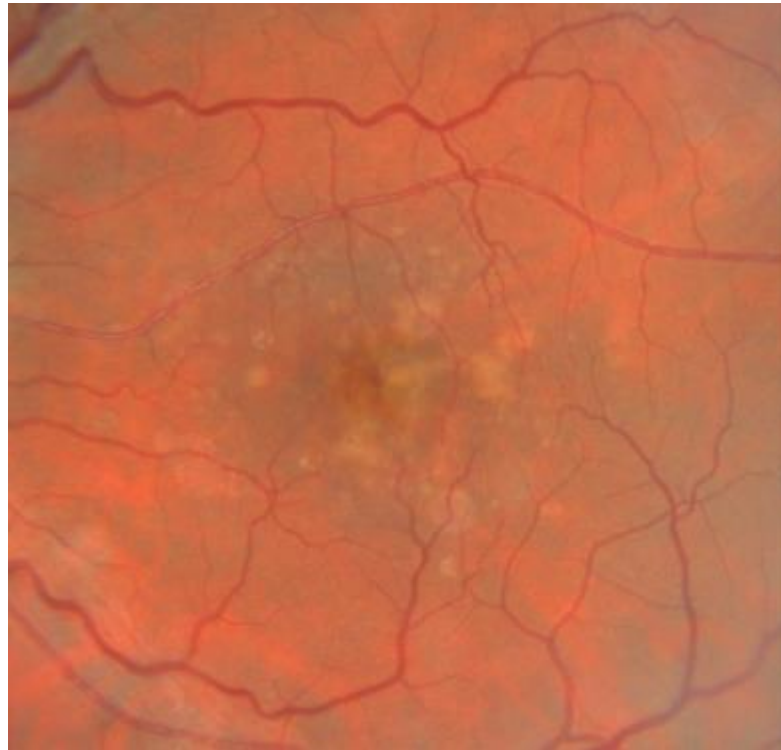


# AMD & OTHER CAUSES OF CNV

## OCTA Clinical Applications

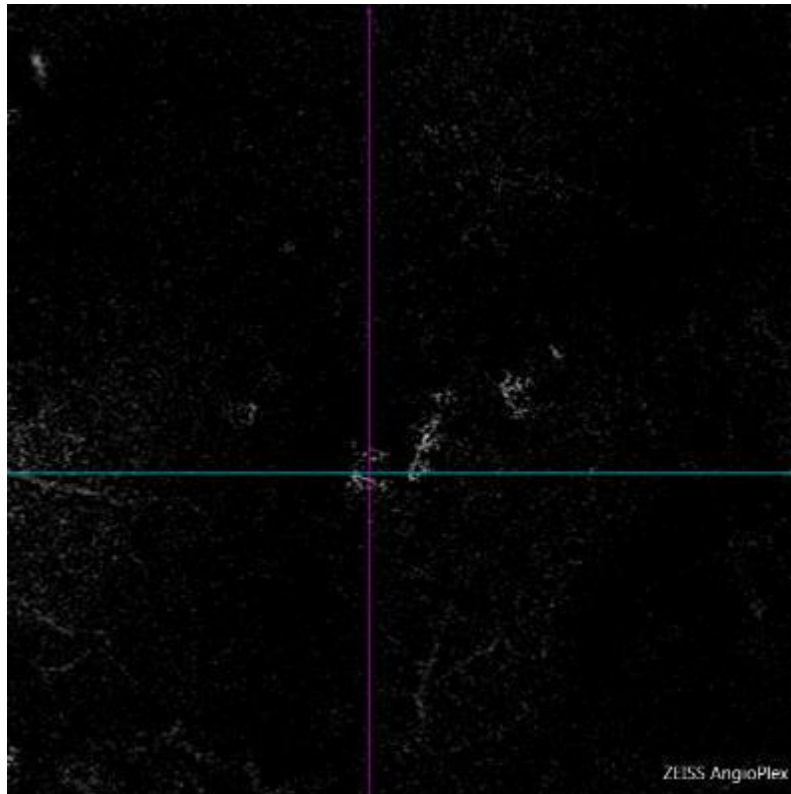
- CNV detection, classification and quantification
- Assess responsiveness to treatment
- Assess need for retreatment
- Detect quiescent (inactive) CNV membranes

# AMD

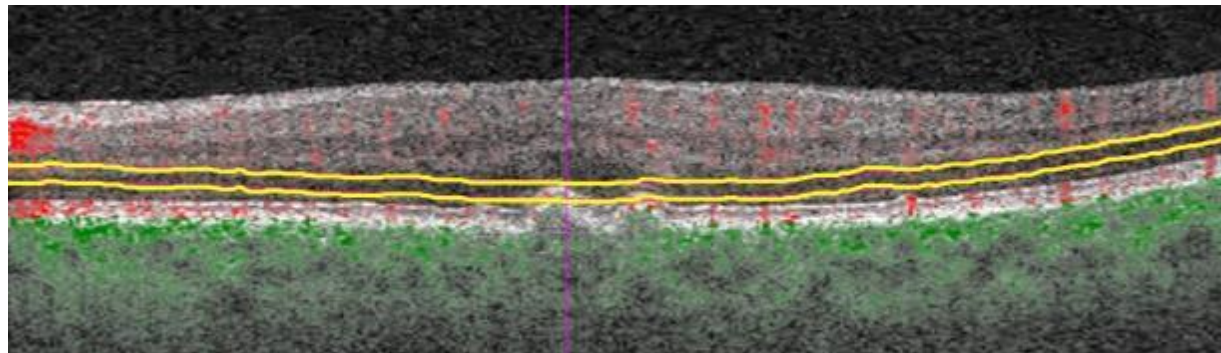
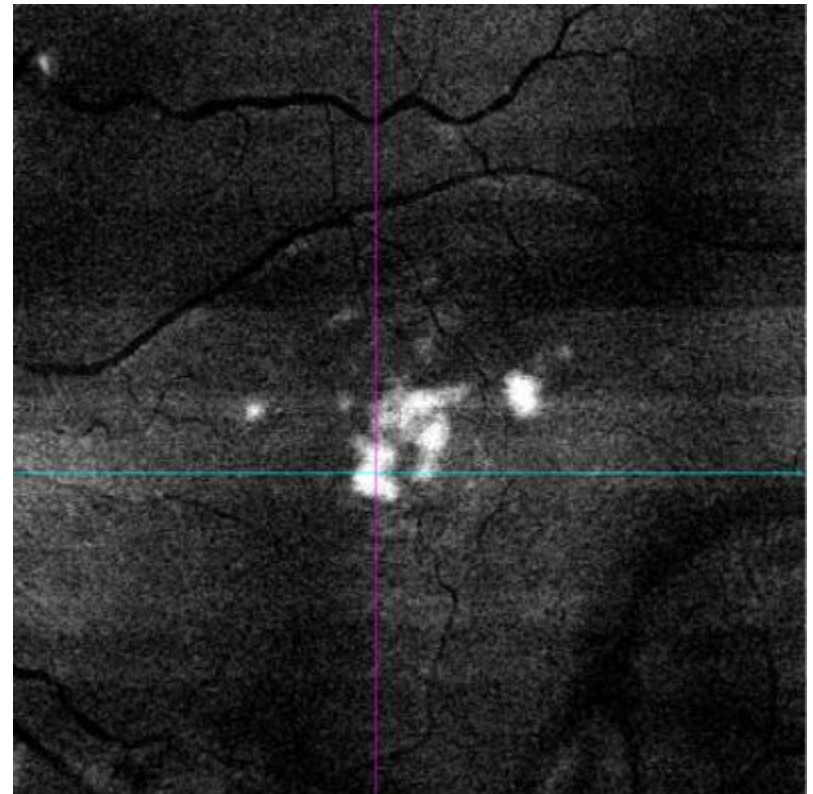


# AMD - Drusen

Avascular Angio



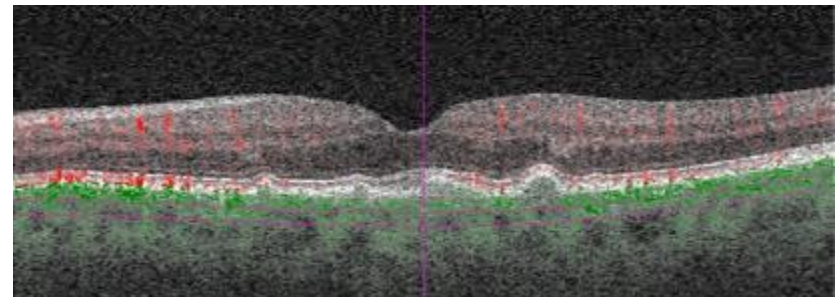
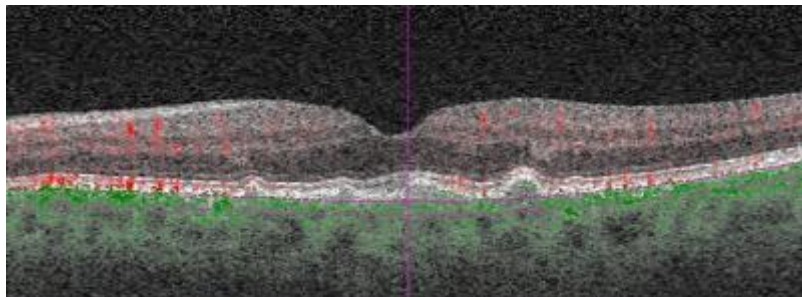
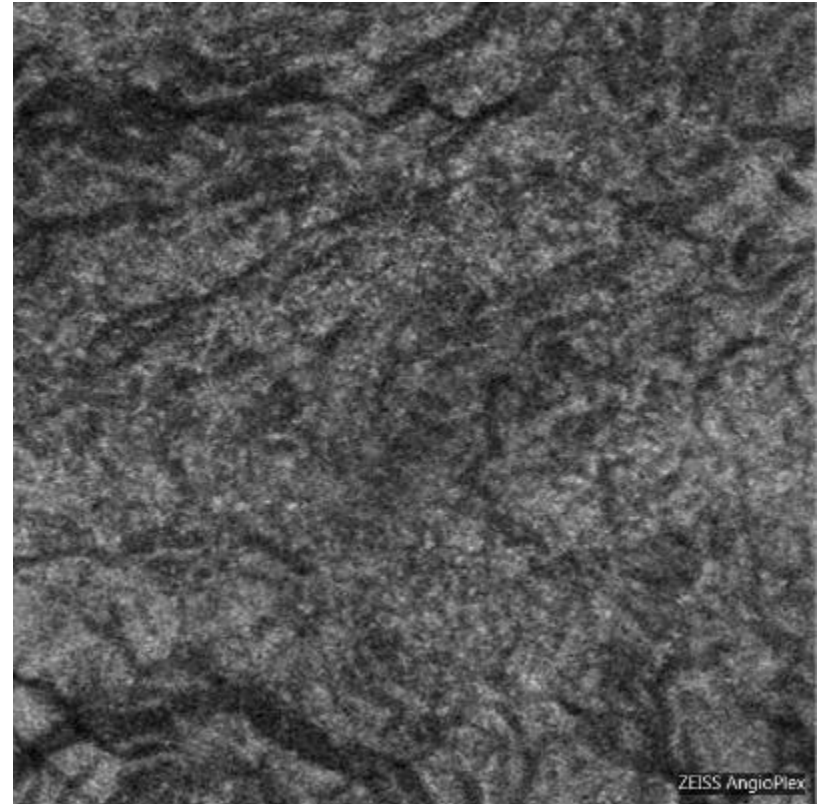
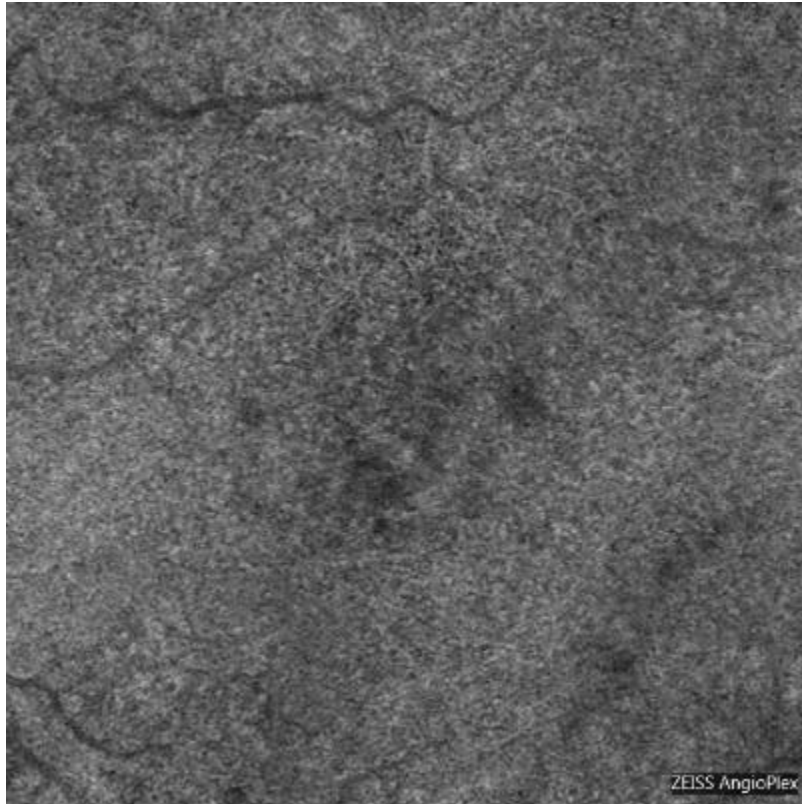
Avascular Structural



# AMD - Drusen

Choriocapillaris

Choroid



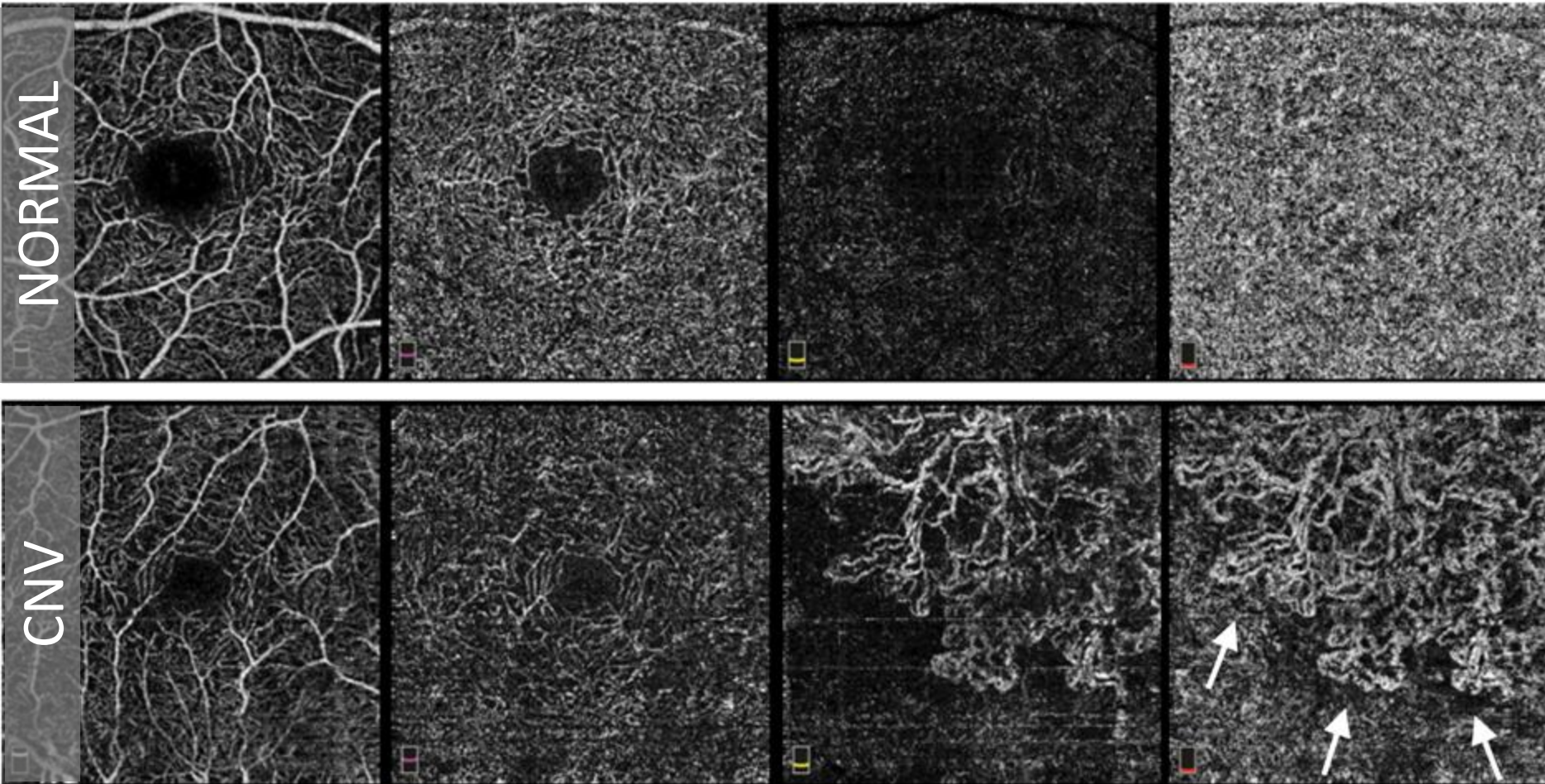
# AMD- CNV

SCP

DCP

Avascular

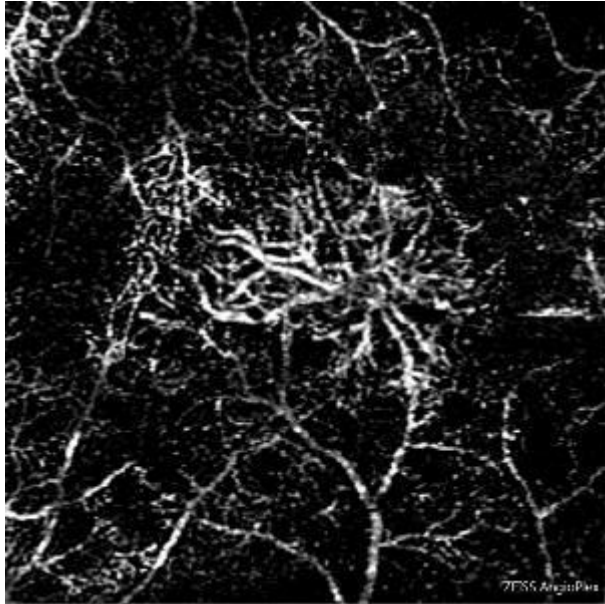
Choriocapillaris



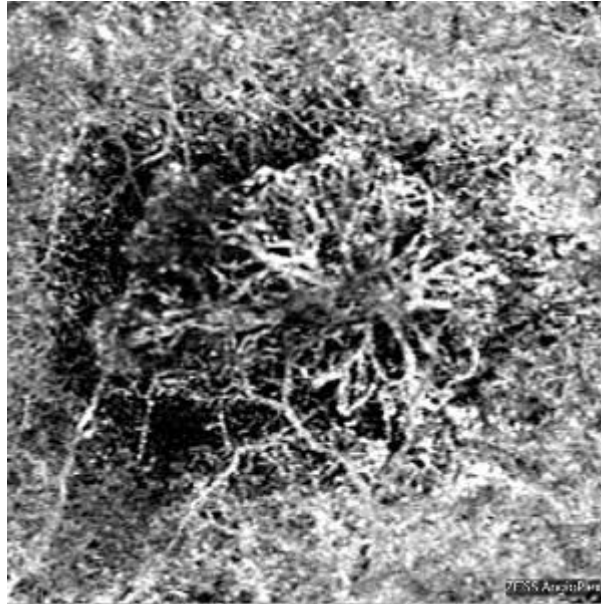
PMID: 32826223

# AMD- Exudative

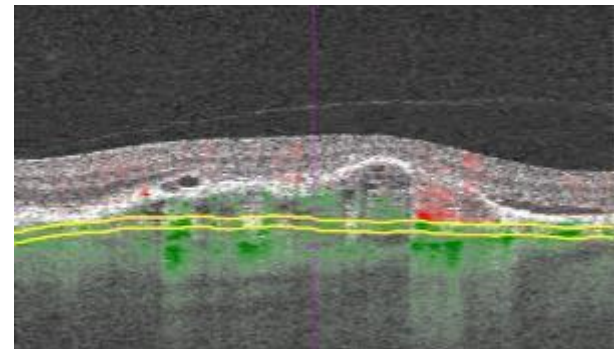
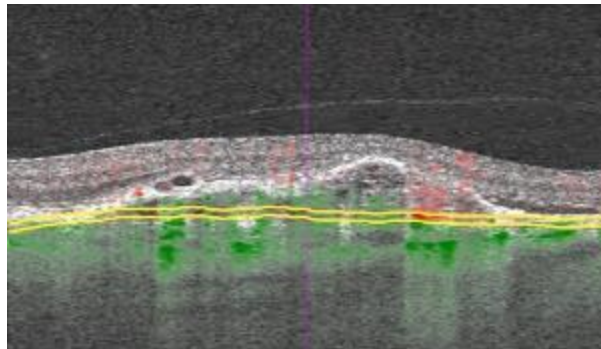
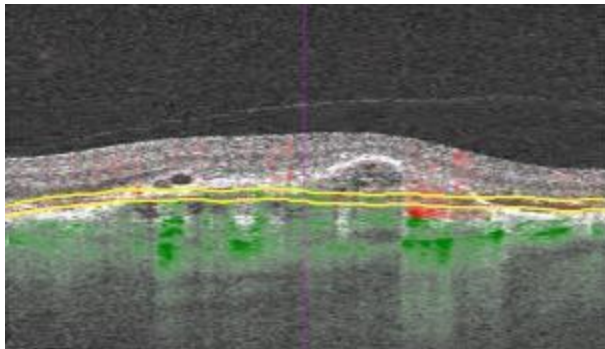
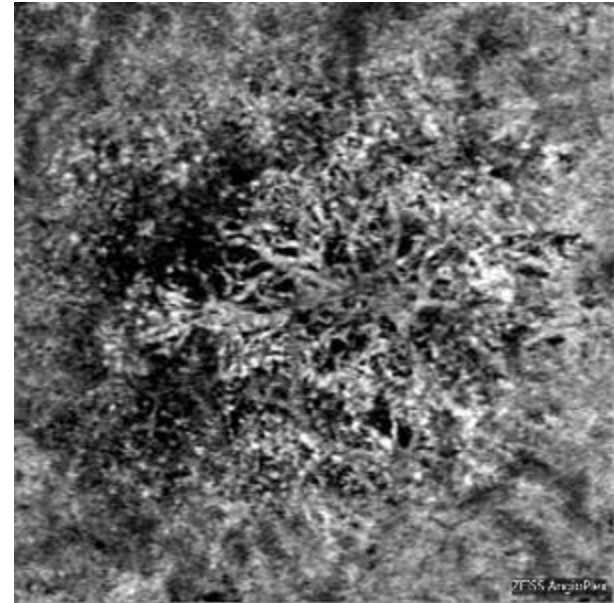
Avascular



Choriocapillaris

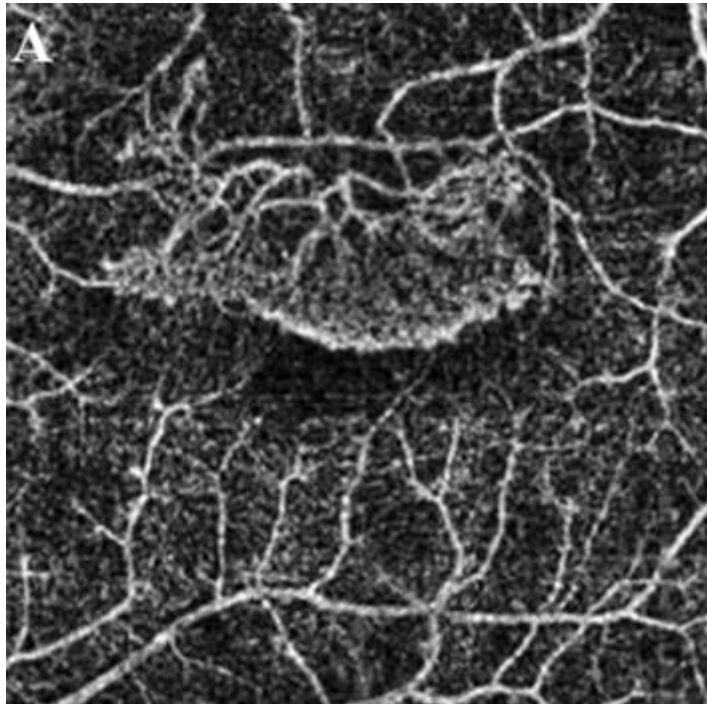


Choroid



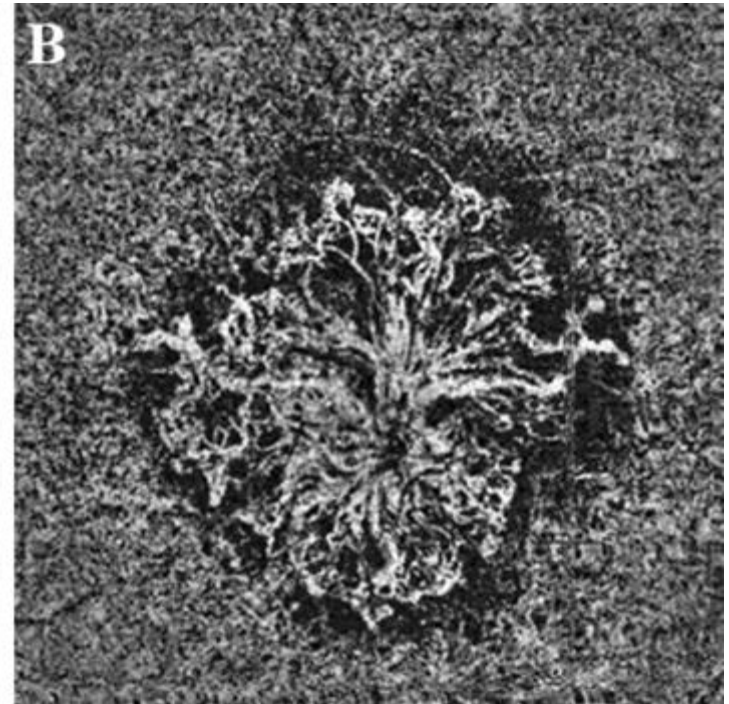
# AMD- Exudative

Seafan Pattern



Vessels radiate from one side of the lesion

Medusa Pattern



Vessels branch in all directions from the center of the lesion

# AMD- Exudative

Clinical science

## Evaluation of the clinical utility of optical coherence tomography angiography in age-related macular degeneration

Melina Cavichini,<sup>1,2</sup> Kunny C Dans,<sup>1</sup> Mahima Jhingan,<sup>1,3</sup> Manuel J Amador-Patarroyo,<sup>1,4</sup> Shyamanga Borooah,<sup>5,6</sup> Dirk-Uwe Bartsch,<sup>1</sup> Eric Nudleman,<sup>1</sup> William R Freeman <sup>7</sup>

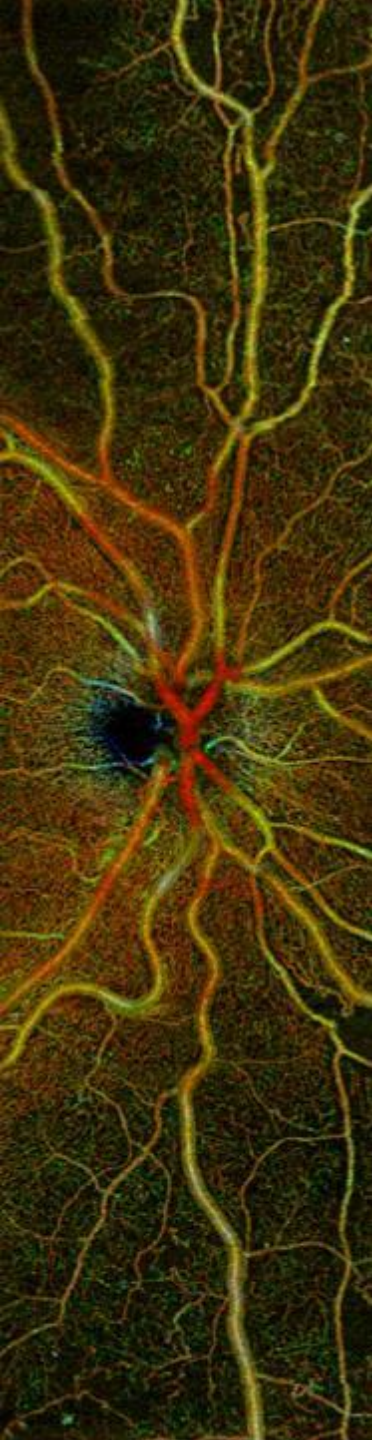
|                           |     |                                     |
|---------------------------|-----|-------------------------------------|
| Sensitivity               | 80% | Correctly identify presence of CNV  |
| Specificity               | 85% | Correctly identify absence of CNV   |
| Positive Predictive Value | 60% | CNV found after positive result     |
| Negative Predictive Value | 94% | CNV not found after negative result |

PMID 32826223

# AMD- Exudative

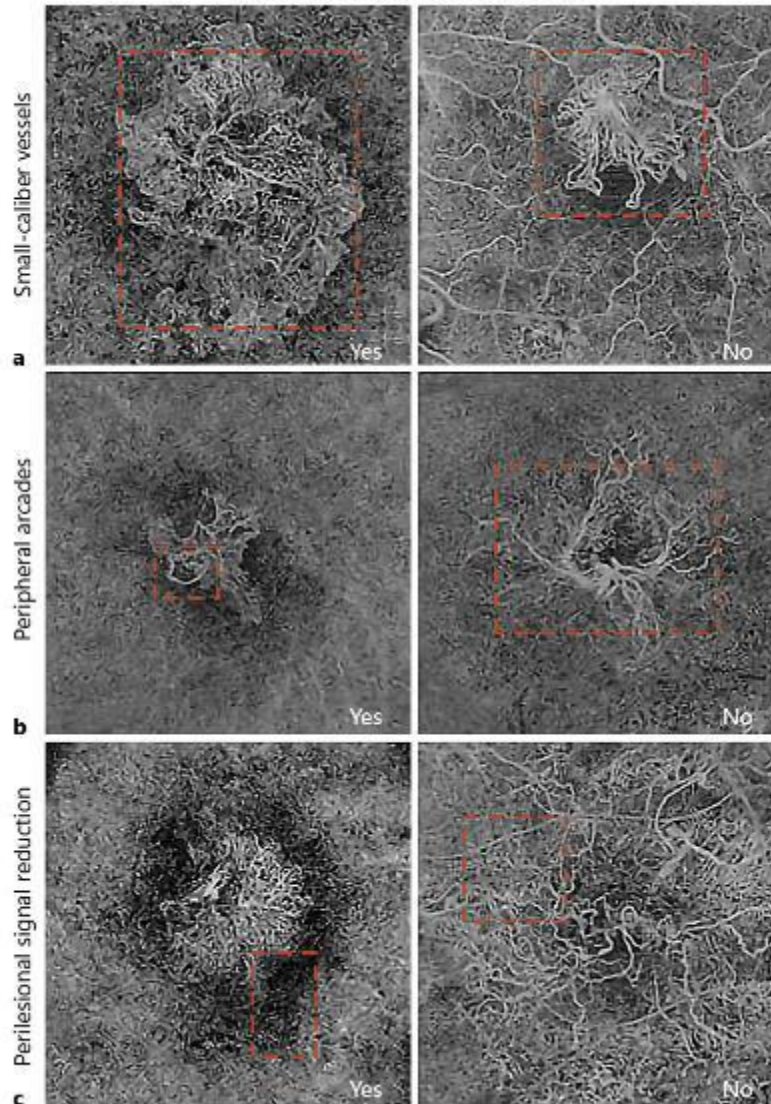
## THREE CLINICAL CNV VARIANTS

1. Active lesions: Symptomatic, signs of exudation on OCT (subretinal fluid)
2. Inactive lesions: Status following successful anti-VEGF therapy. No signs of exudation on OCT
3. Subclinical lesions: Asymptomatic, no signs of exudation on OCT. Often found in fellow eyes of active CNV. High risk of becoming active



# AMD- Exudative

Presence of qualitative OCTA criteria



## OCTA SIGNS OF ACTIVE CNV

Active lesions contain smaller caliber vessels

Active lesions have looping vessels at margins.  
Inactive lesions have a “dead tree” appearance

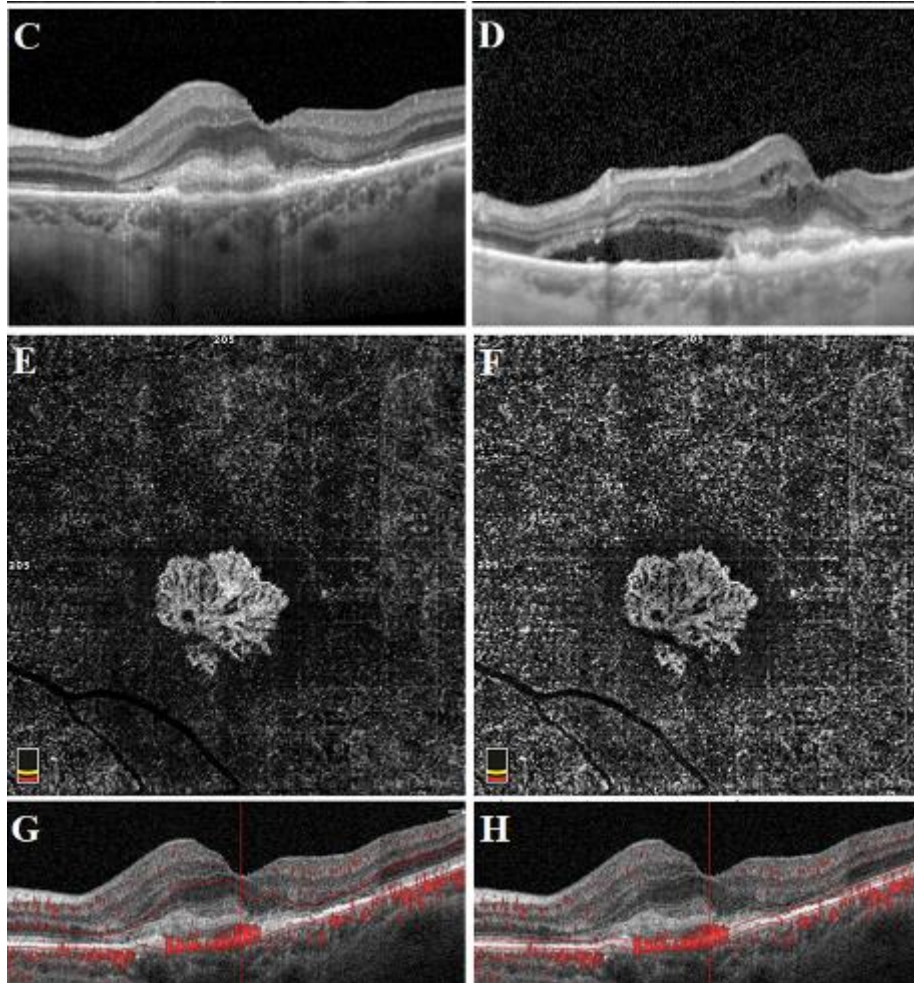
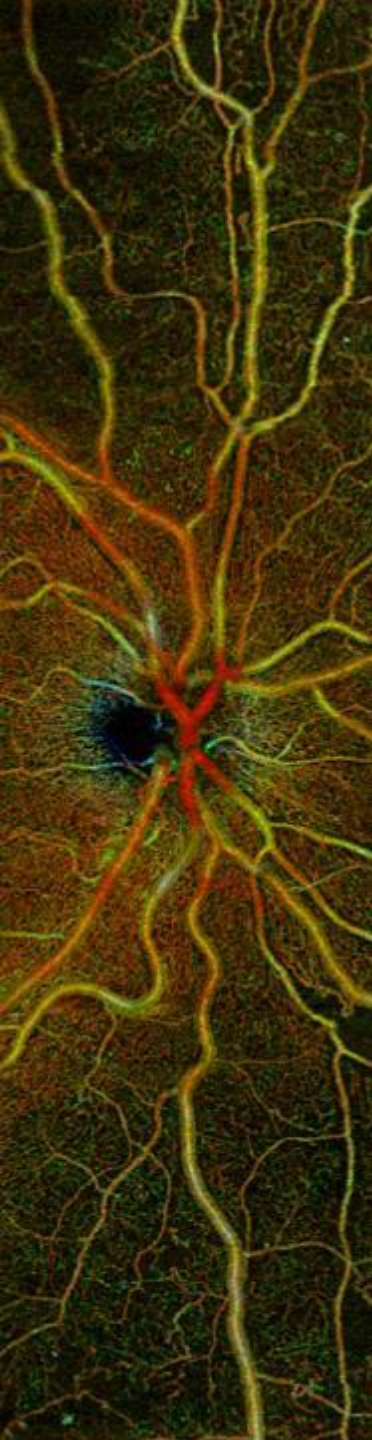
Active lesions are surrounded by a zone of decreased signal intensity (“halo sign”)

**ACTIVE**

**INACTIVE**

PMID: 31665719

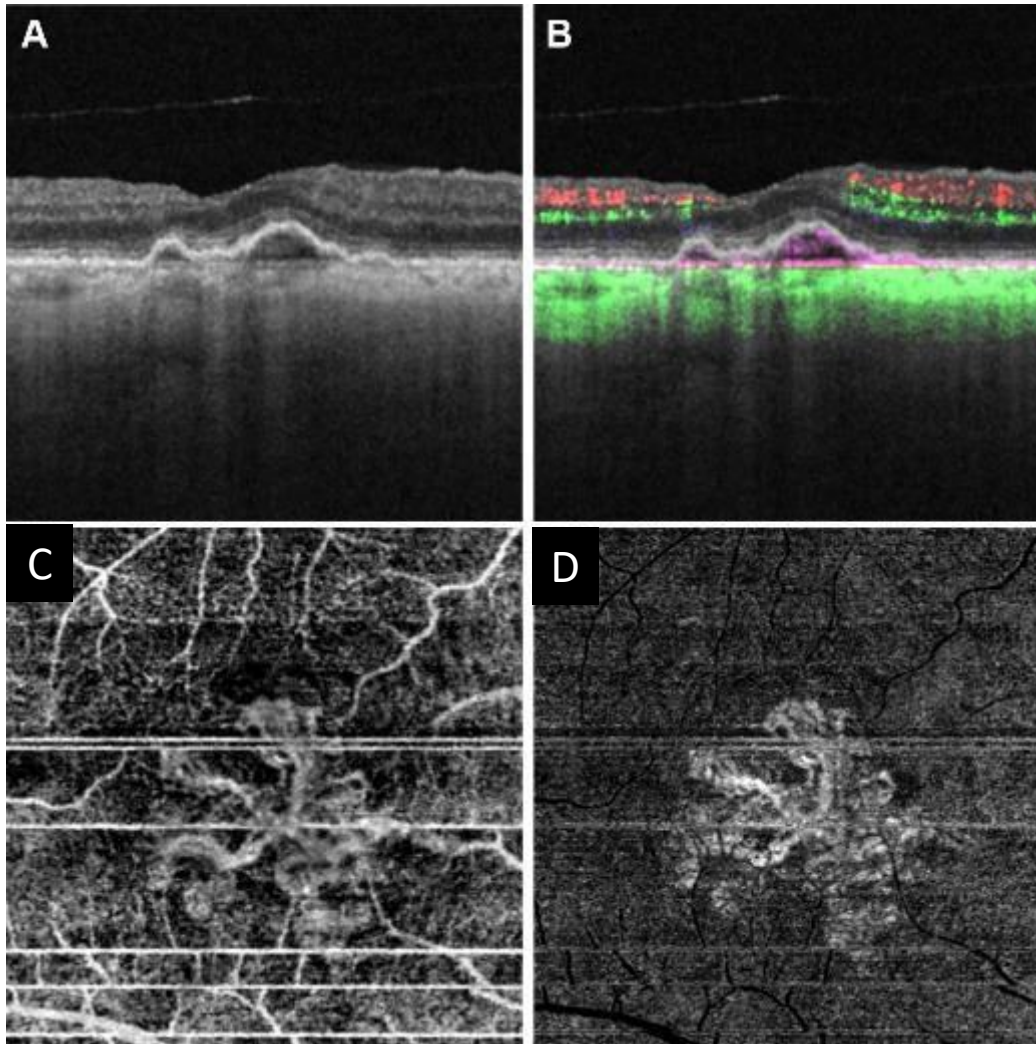
# AMD- Exudative



Subretinal lesion  
with fluid leakage

OCTA reveals sea-  
fan CNV with OCTA  
signs that it is active  
Left: Outer retina  
Right: Choroid

# AMD- Nonexudative (subclinical) CNV

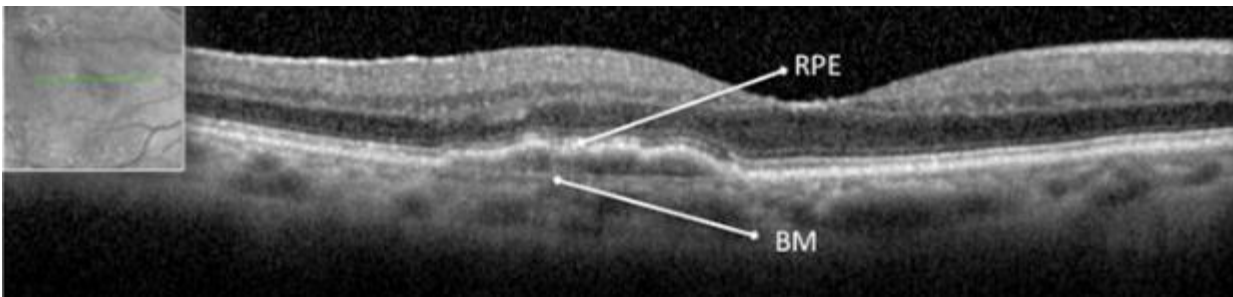


- A. OCT shows PED w/o SRF in right eye of 78yo asymptomatic man with dry AMD. VA 20/20
- B. OCTA B-scan shows flow signal within PED
- C. En-face OCTA reveals CNV within PED. Note projection artifact of retinal vascular tree
- D. Same image with projection artifact removed

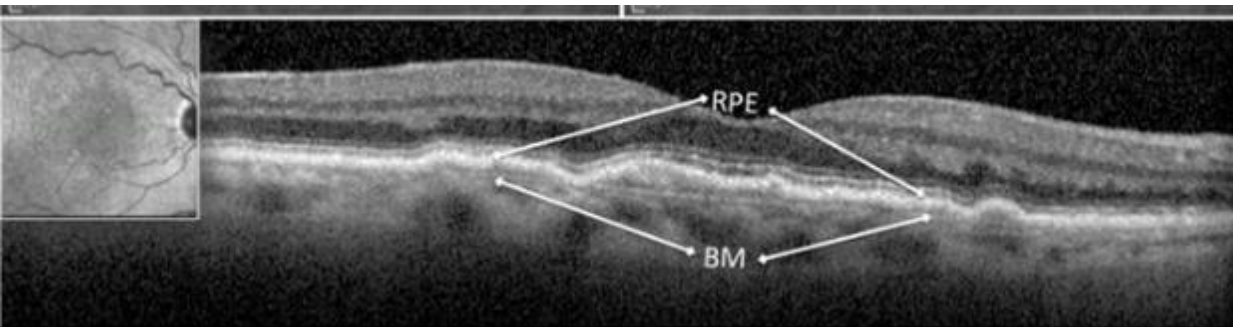
PMID 26876696

**These lesions are asymptomatic but carry a high risk of exudation.  
These lesions may remain stable for years but require close monitoring**

# AMD- Nonexudative (subclinical) CNV

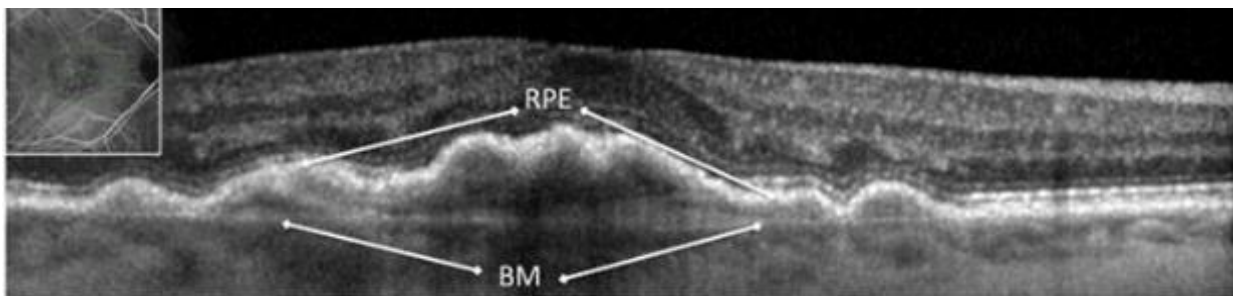


**“DOUBLE LAYER”  
SIGN TO IDENTIFY  
SUBCLINICAL CNV  
IN EYES WITH  
“DRY” AMD**



Top layer = RPE

Bottom layer = Bruch's

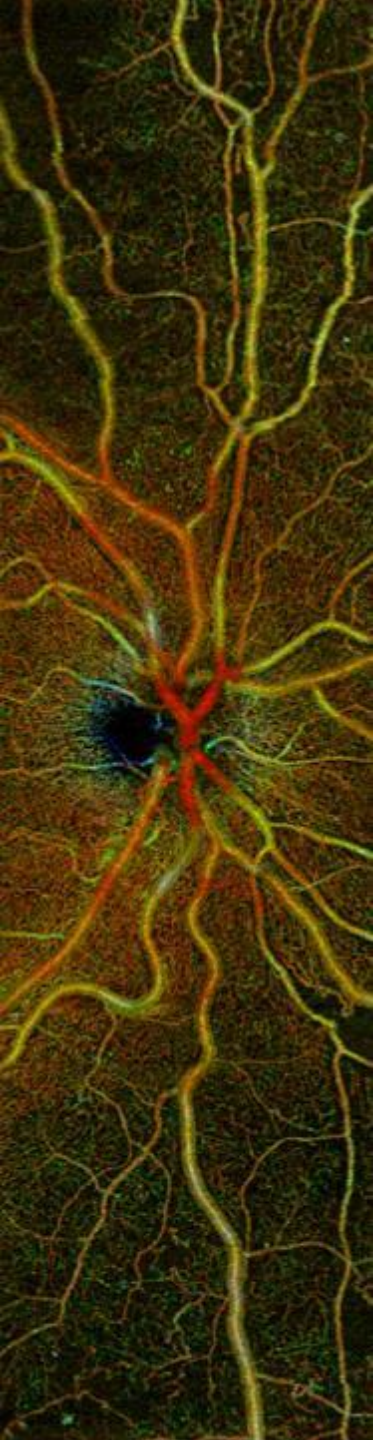


Absence of SRF, blood,  
and other signs of  
exudation indicates  
nonexudative nature of  
the lesion

**Sensitivity: 88%**  
**Specificity: 87%**

**PPV: 76%**  
**NPV: 94%**

PMID: 31014697



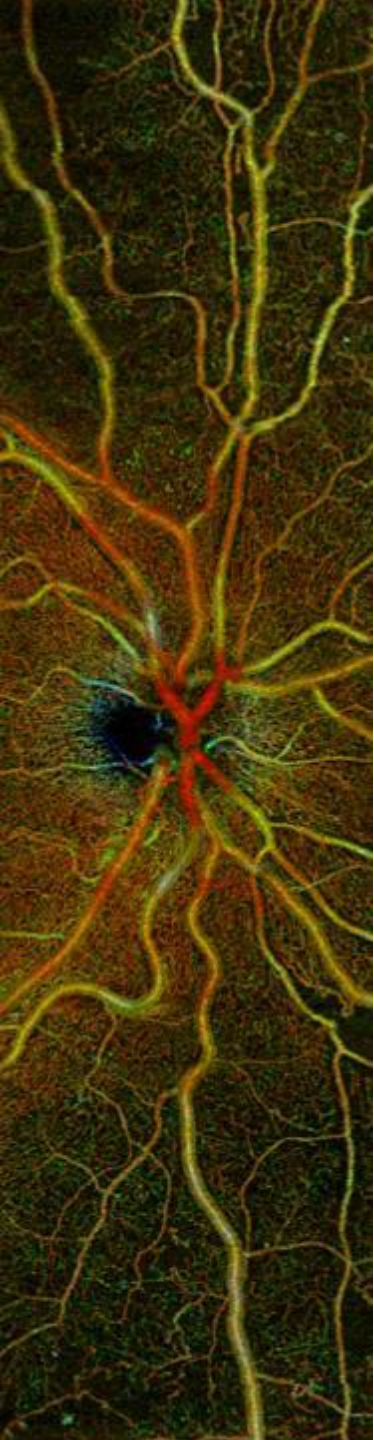
# Macular Telangiectasia Type 2

## ***WHAT IS IT?***

MacTel type 2 is an idiopathic bilateral neurodegenerative disease with characteristic alterations of the macular capillary network and neurosensory atrophy

## ***CLINICAL FEATURES INCLUDE:***

- Loss of macular pigment
- Retinal hyporeflective cavities on OCT
- Telangiectatic capillaries (early)
- Retinal pigment plaques, foveal atrophy, and subretinal neovascularization (late)

A fluorescein angiogram of the retina. The image shows a dense network of retinal vessels. The macula, located in the center-left, exhibits telangiectatic changes, characterized by dilated, tortuous, and hyperfluorescent vessels. The surrounding retinal vasculature appears relatively normal.

# Macular Telangiectasia Type 2

## ***CASE REPORT***

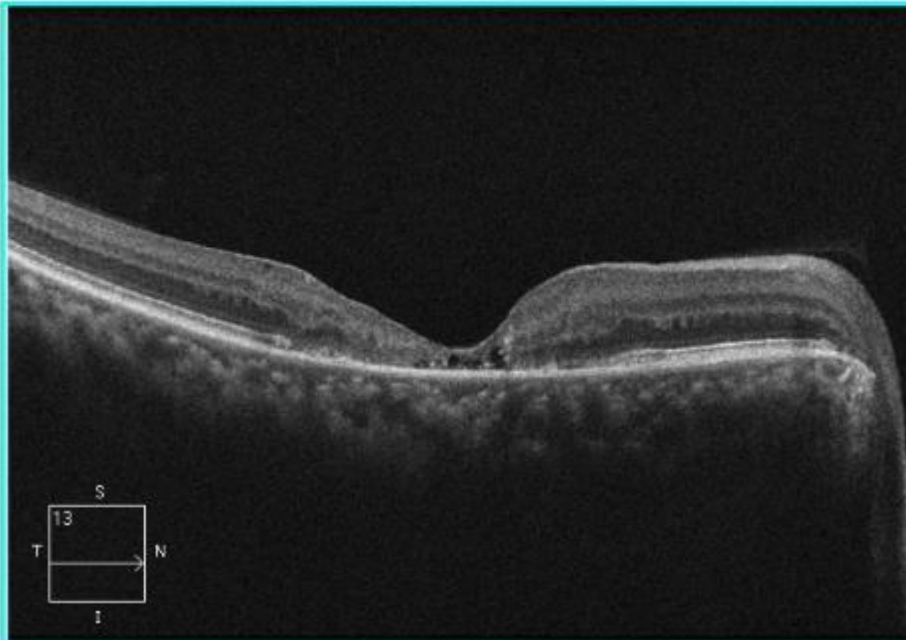
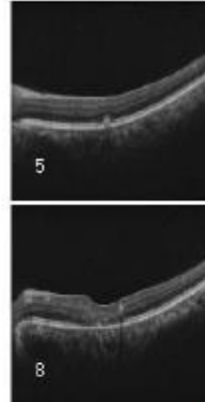
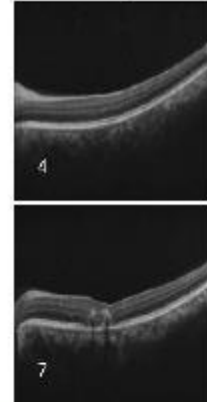
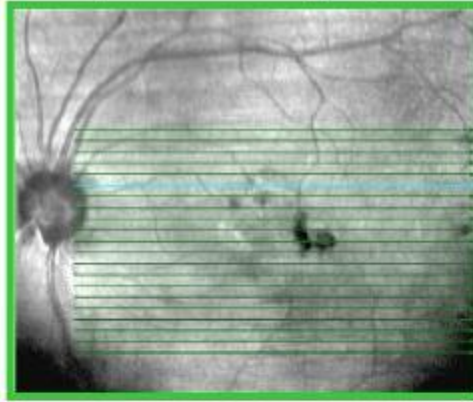
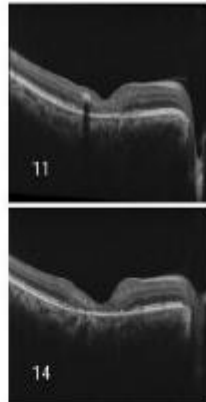
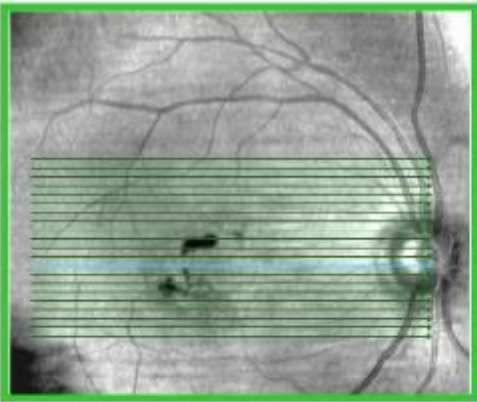
56yo HF

- Consultation for evaluation of maculopathy OU
- C/O bilateral progressive decrease in vision x 2 yrs
- MH: Good health. No meds
- POH: Unremarkable
- BCVA: 20/100 OD, 20/200 OS
- Ta 25/25 @ 2pm
- SLE: WNL OU

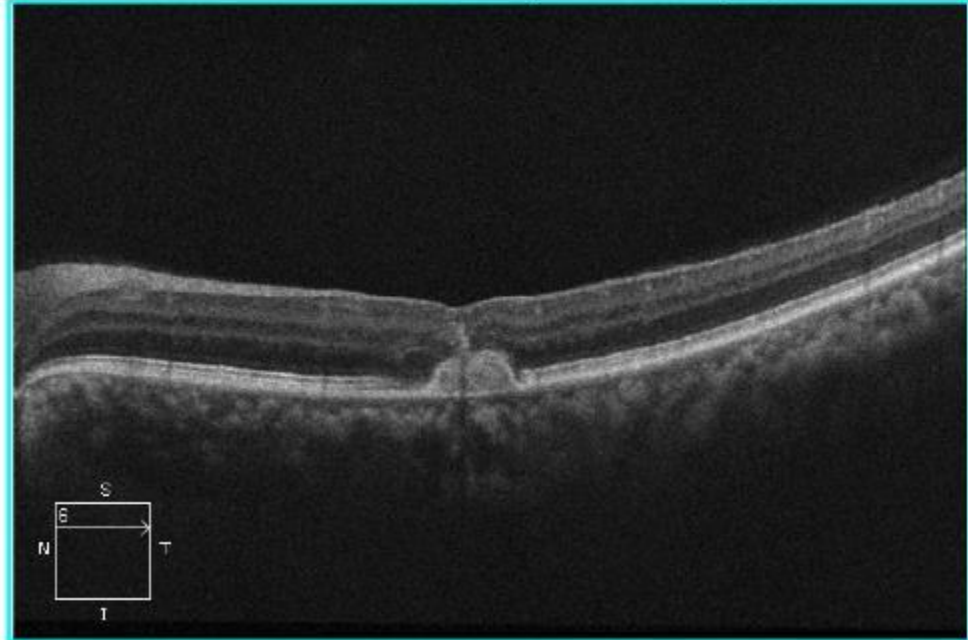
# Macular Telangiectasia Type 2



# Macular Telangiectasia Type 2



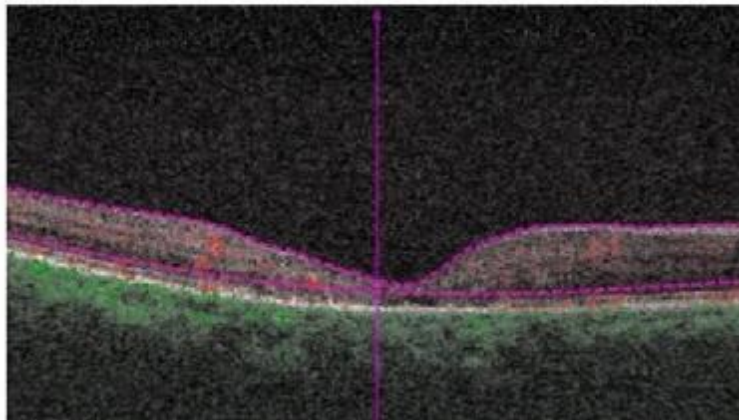
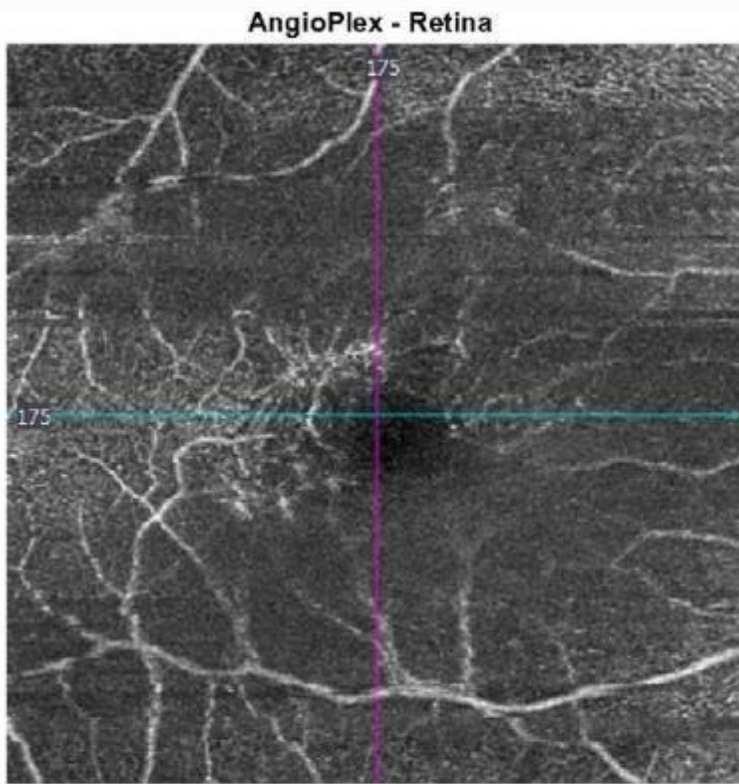
OD



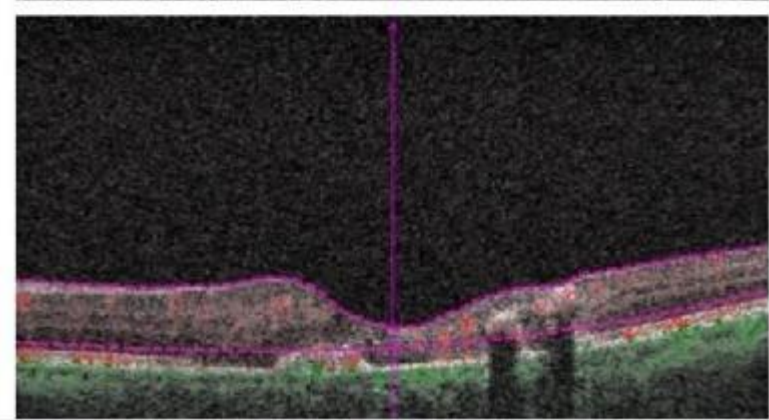
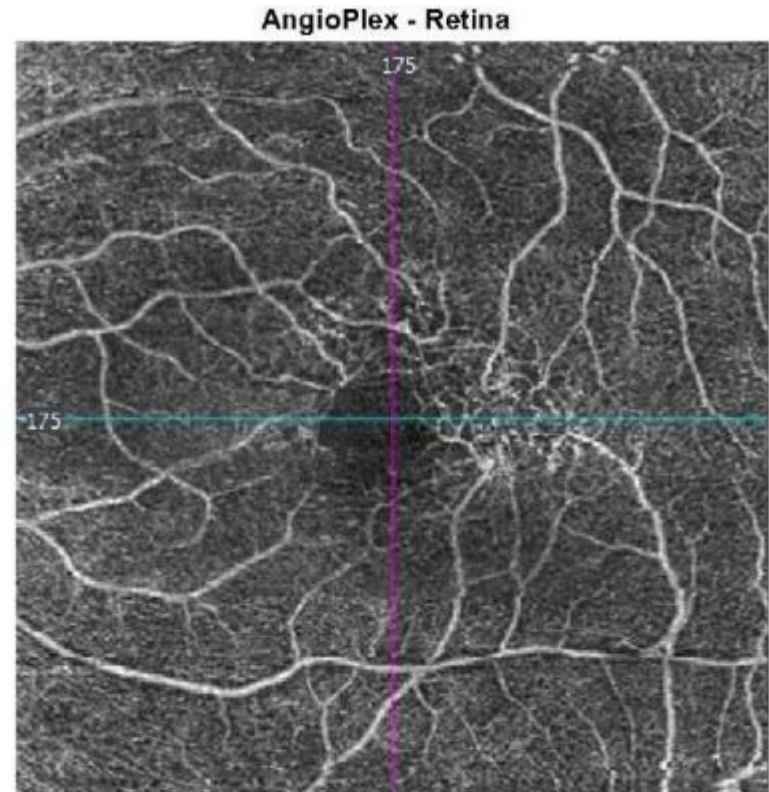
OS

# Macular Telangiectasia Type 2

OD



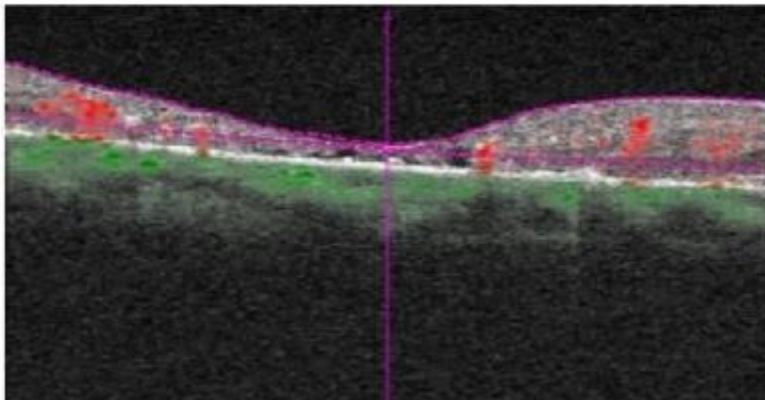
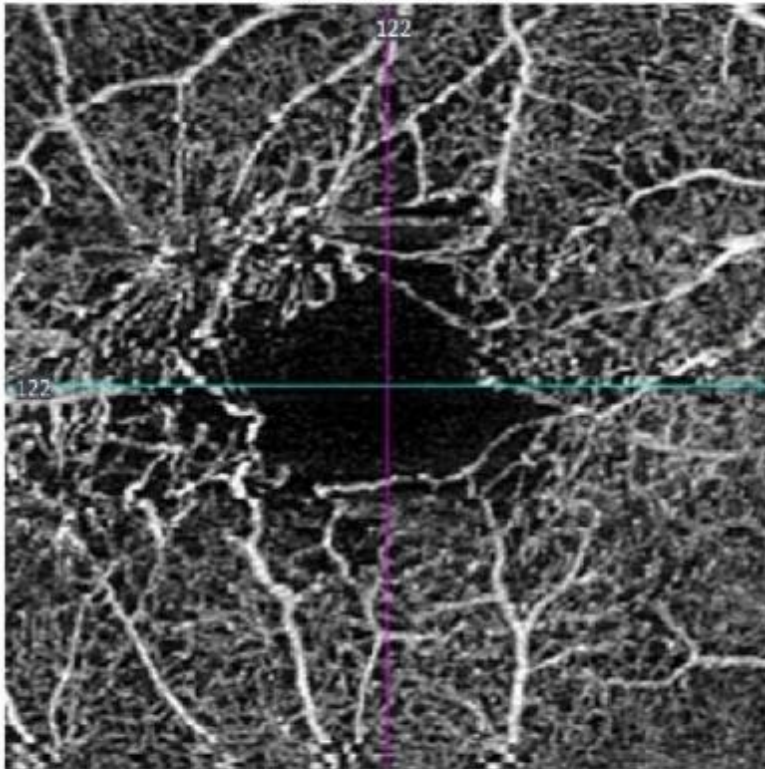
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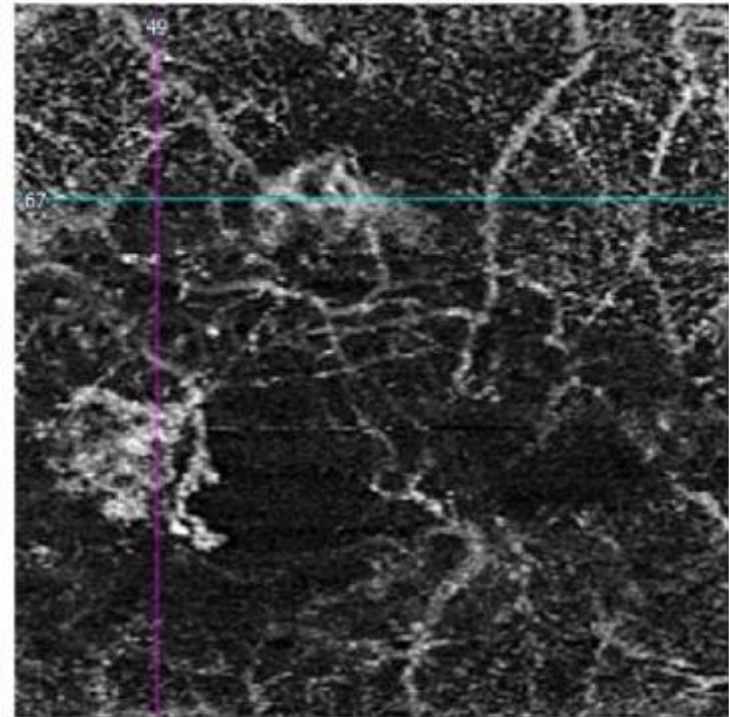
# Macular Telangiectasia Type 2

OD

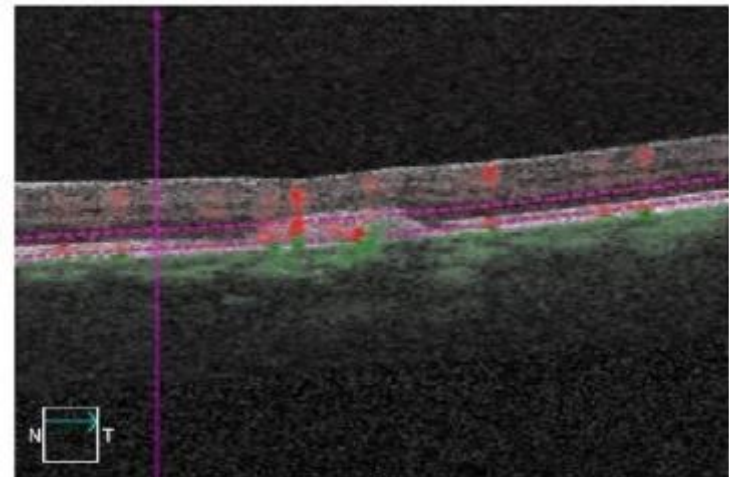
AngioPlex - Retina



AngioPlex - Custom



OS



# Macular Telangiectasia Type 2

## Single Field Analysis

Eye: Left

Name: ID:  
DOB: 10-12-1960

## Central 30-2 Threshold Test

Fixation Monitor: OFF

Fixation Target: Central

Fixation Losses: 0/0

False POS Errors: 4 %

False NEG Errors: 15 %

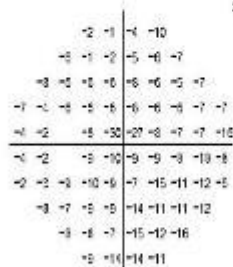
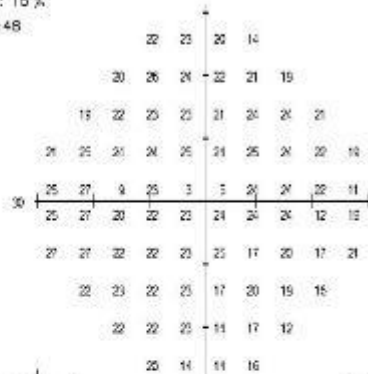
Test Duration: 12:48

Fovea: OFF

Stimulus: III, White  
Background: 31.5 ASB  
Strategy: SITA-Standard

Pupil Diameter:  
Visual Acuity:  
RX: DS DC X

Date: 05-04-2017  
Time: 11:04 AM  
Age: 56



Total Deviation



Pattern Deviation

GHT  
Outside Normal Limits

VFI 74%

MD -9.15 dB P < 0.5%

PSD 6.22 dB P < 0.5%

BOWDEN EYE CARE

□ < 5%  
□ < 2%  
□ < 1%  
■ < 0.5%

## Single Field Analysis

Eye: Right

Name: ID:  
DOB: 10-12-1960

## Central 30-2 Threshold Test

Fixation Monitor: Gaze/Blind Spot

Fixation Target: Central

Fixation Losses: 10/23 xv

False POS Errors: 5 %

False NEG Errors: 15 %

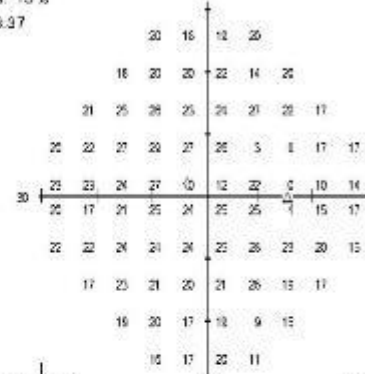
Test Duration: 13:37

Fovea: OFF

Stimulus: III, White  
Background: 31.5 ASB  
Strategy: SITA-Standard

Pupil Diameter:  
Visual Acuity:  
RX: DS DC X

Date: 06-04-2017  
Time: 10:48 AM  
Age: 56



Total Deviation



Pattern Deviation

\*\*\* Low Test Reliability \*\*\*

GHT  
Outside Normal Limits

VFI 73%

MD -9.67 dB P < 0.5%

PSD 7.22 dB P < 0.5%

BOWDEN EYE CARE

□ < 5%  
□ < 2%  
□ < 1%  
■ < 0.5%

# Macular Telangiectasia Type 2

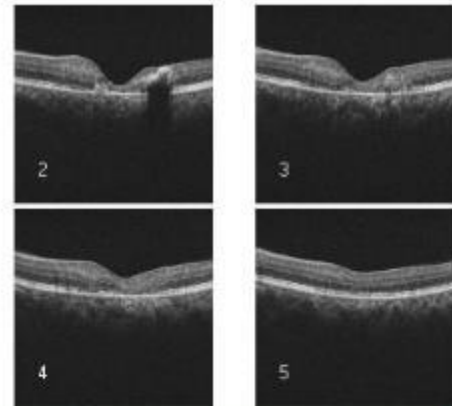
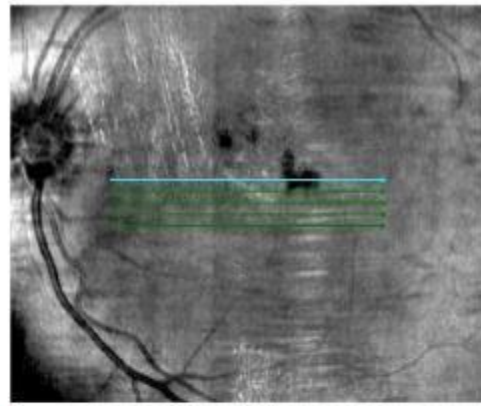
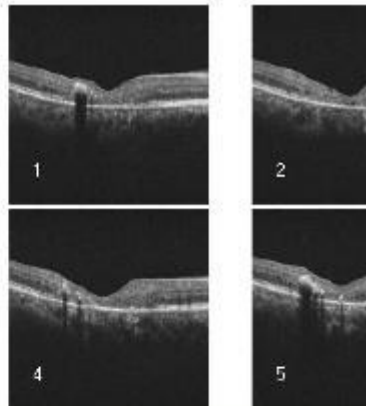
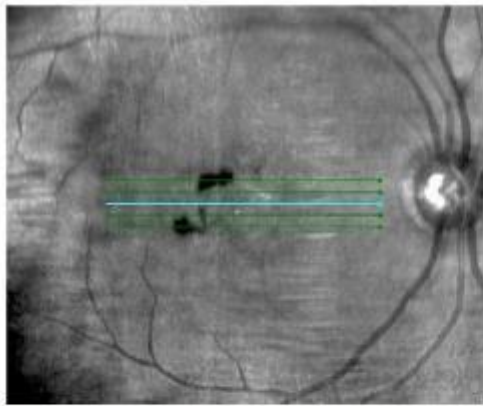
## ***OUR PLAN:***

(1) IVB OS, (2) Glaucoma eval, (3) Low vision

## ***PATIENT'S PLAN:***

- Return 8 mos later
- C/O vision getting worse OS.
- BCVA: 20/100 OU
- Ta 24/24 @ 4pm
- SLE: WNL OU

# Macular Telangiectasia Type 2



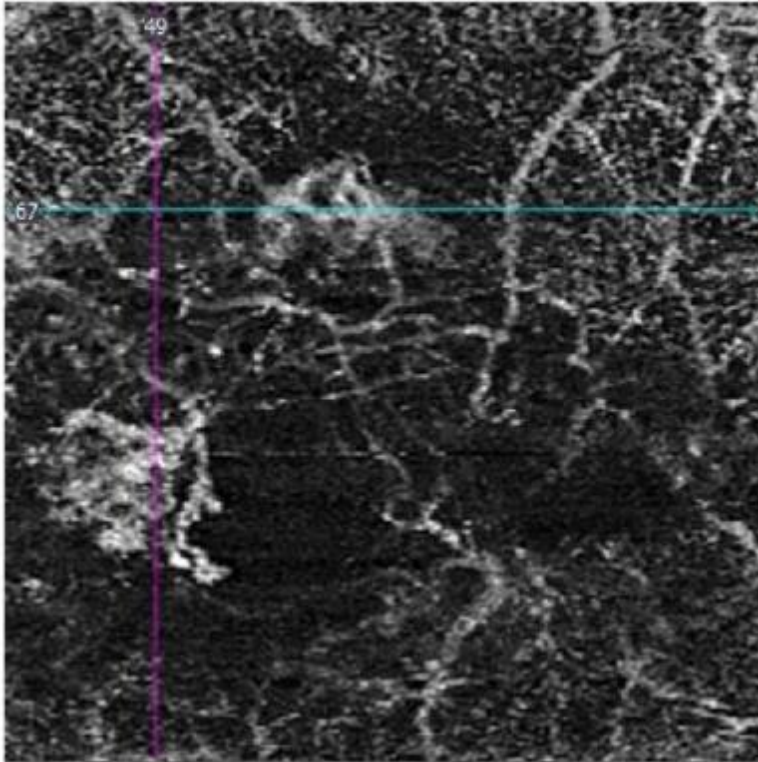
OD



OS

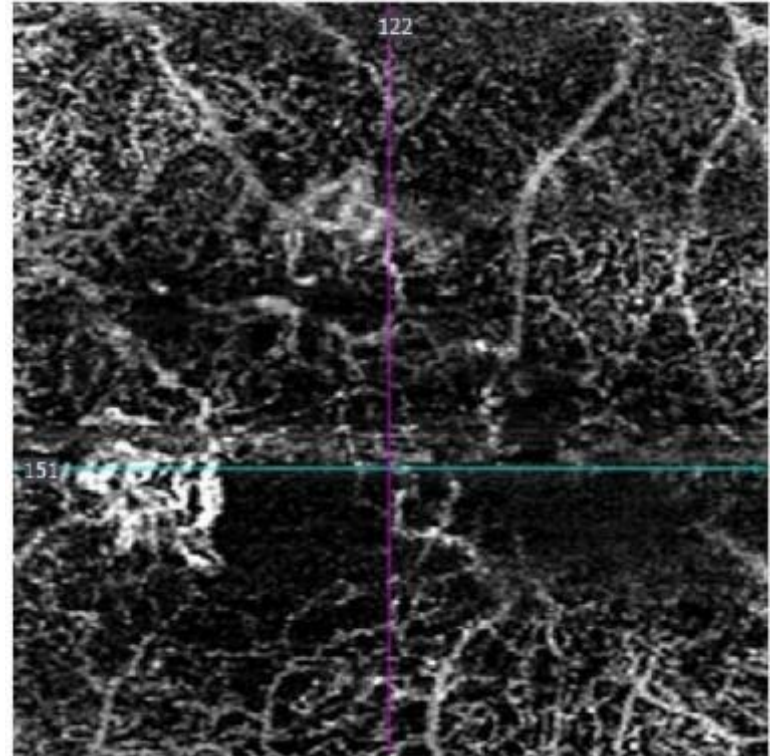
# Macular Telangiectasia Type 2

AngioPlex - Custom



2017

AngioPlex - Custom



2018

Registration : Automatic

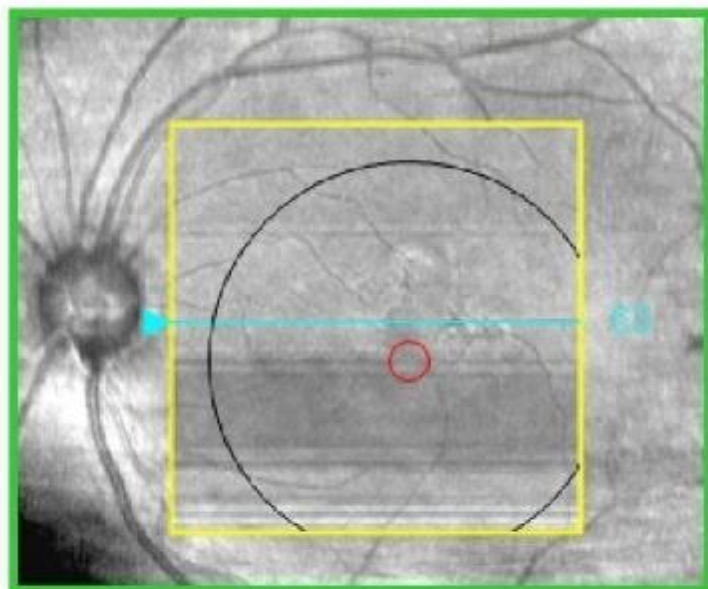
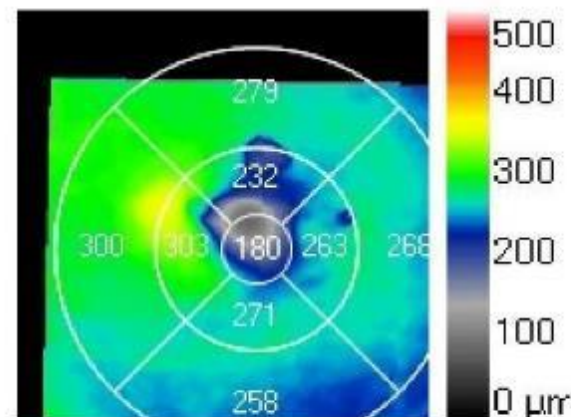
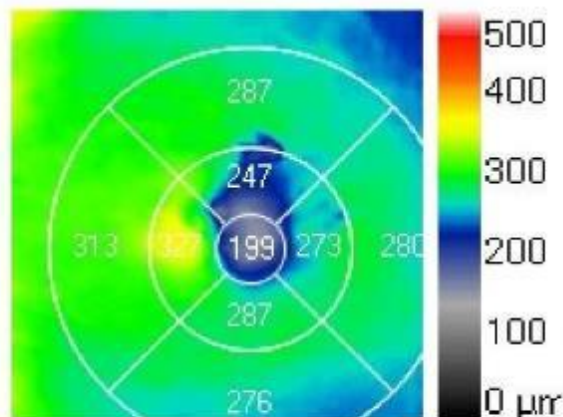
Registration succeeded

Exam from 8/4/2017 1:02:18 PM

Exam from 4/13/2018 4:21:07 PM

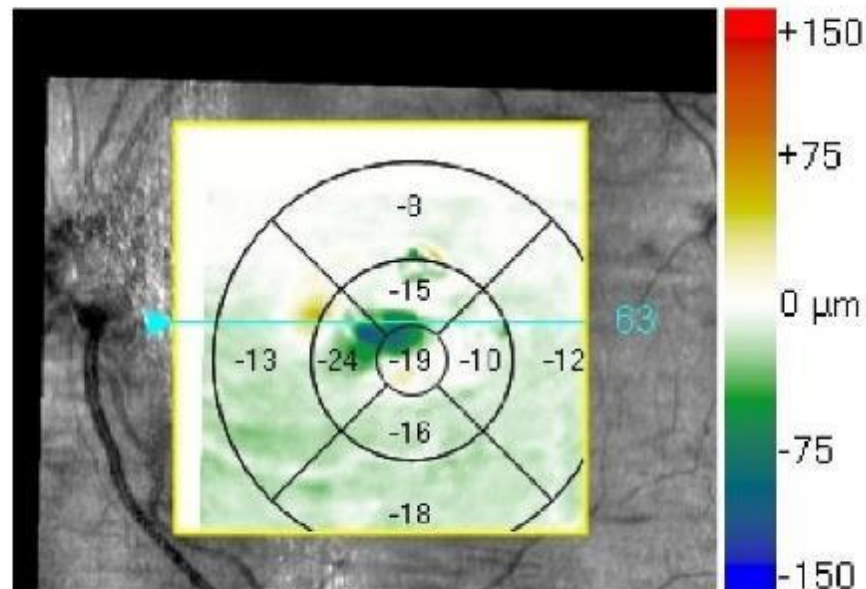
Fovea: 297, 75

Fovea: 297, 75



Overlay: OCT Fundus

Transparency: 0 %



Overlay: ILM-RPE Difference

Transparency: 0 %

# Macular Telangiectasia Type 2

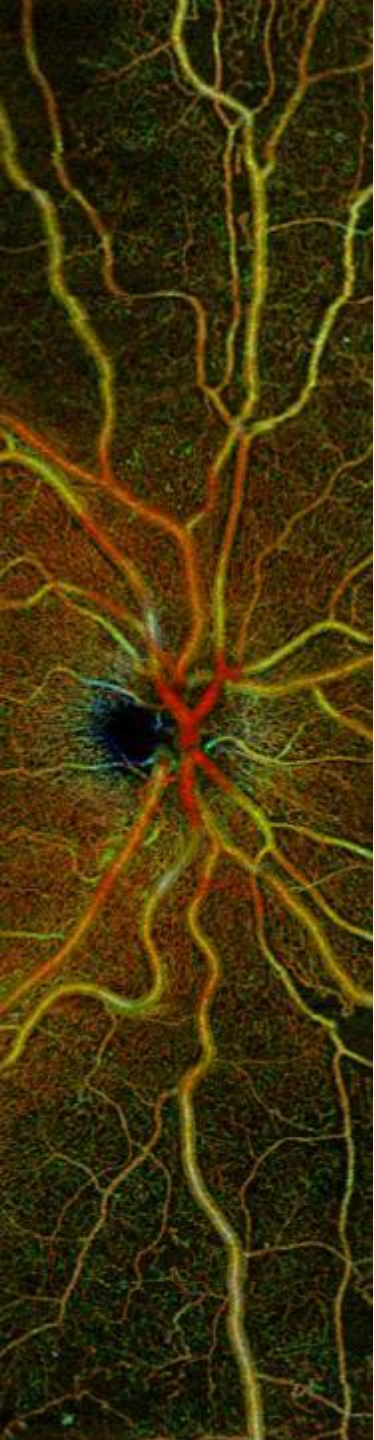
## ***KEY POINTS***

- No effective treatment for nonproliferative disease (macular atrophy)
- Treatment of subretinal neovascularization can prevent vision loss
- OCTA is well suited to visualizing the retinal vascular abnormalities associated with MacTel2, particularly with monitoring subretinal neovascularization

# GLAUCOMA

## OCTA Clinical Applications

- Screening/diagnosing
- Staging
- Monitoring progression
  - High myopia



# GLAUCOMA



AMERICAN ACADEMY  
OF OPHTHALMOLOGY®

## Ophthalmic Technology Assessment



### OCT Angiography for the Diagnosis of Glaucoma

*A Report by the American Academy of Ophthalmology*

Darrell WuDunn, MD, PhD,<sup>1</sup> Hana L. Takusagawa, MD,<sup>2</sup> Arthur J. Sit, MD,<sup>3</sup> Julia A. Rosdahl, MD, PhD,<sup>4</sup> Sunita Radhakrishnan, MD,<sup>5</sup> Ambika Hoguet, MD,<sup>6</sup> Ying Han, MD, PhD,<sup>7</sup> Teresa C. Chen, MD<sup>8</sup>

**Purpose:** To review the current published literature on the use of OCT angiography (OCTA) to help detect changes associated with the diagnosis of primary open-angle glaucoma.

**Methods:** Searches of the peer-reviewed literature were conducted in March 2018, June 2018, April 2019, December 2019, and June 2020 in the PubMed and Cochrane Library databases. Abstracts of 459 articles were examined to exclude reviews and non-English articles. After inclusion and exclusion criteria were applied, 75 articles were selected and the panel methodologist rated them for strength of evidence. Three articles were rated level I and 57 articles were rated level II. The 15 level III articles were excluded.

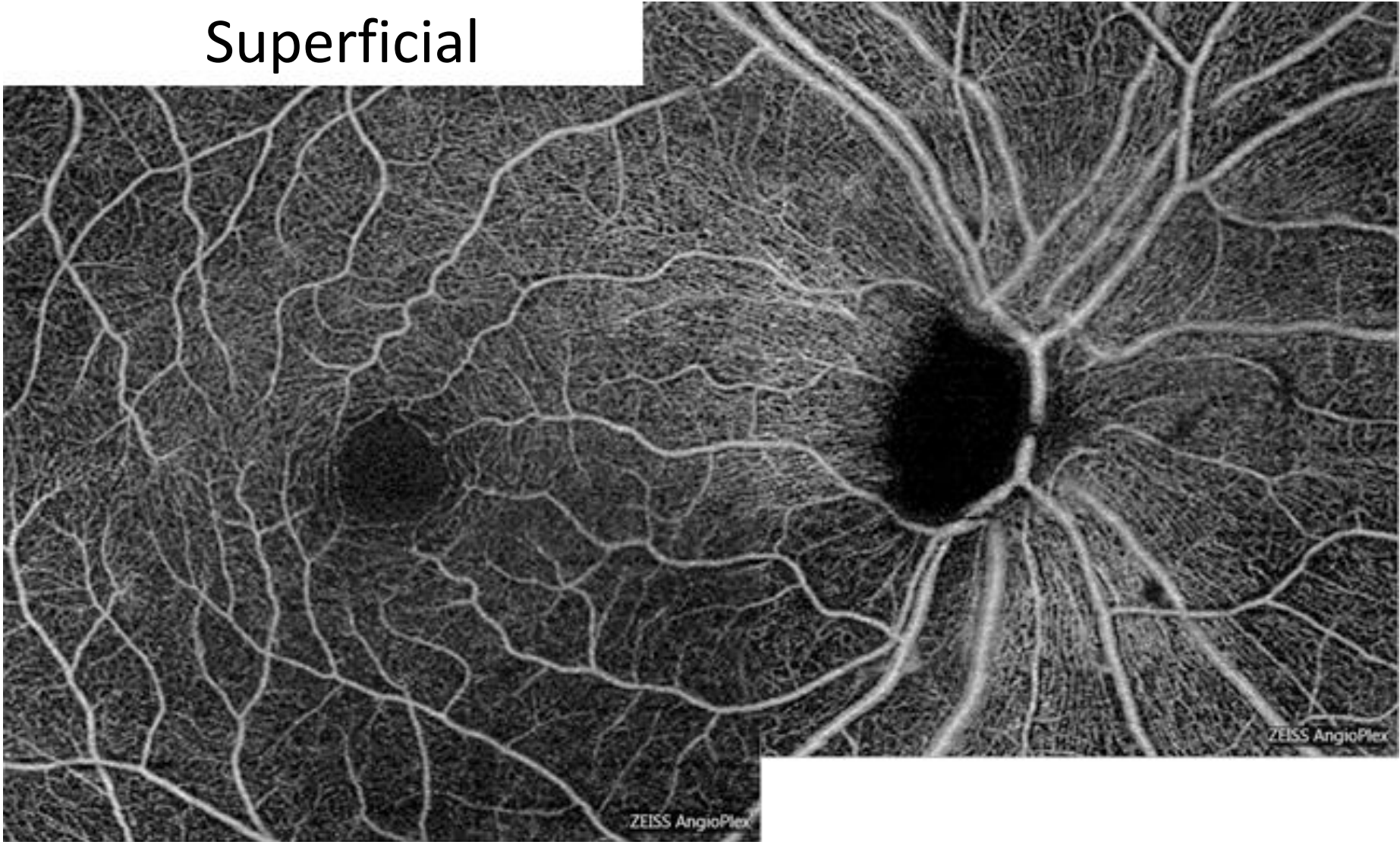
**Results:** OCT angiography can detect decreased capillary vessel density within the peripapillary nerve fiber layer (level II) and macula (level I and II) in patients with suspected glaucoma, preperimetric glaucoma, and perimetric glaucoma. The degree of vessel density loss correlates significantly with glaucoma severity both overall and topographically (level II) as well as longitudinally (level I). For differentiating glaucomatous from healthy

“Vessel density loss associated with glaucoma can be detected by OCTA.”

PMID 33632585

# Glaucoma

Superficial



# A Review of OCT Angiography in Glaucoma

Astrid C. Werner and Lucy Q. Shen 

*Department of Ophthalmology, Massachusetts Eye and Ear Infirmary, Boston, USA*

## ABSTRACT

There is growing evidence that vascular dysfunction plays a role in the pathogenesis of glaucoma. The details of this relationship have remained elusive partially due to limitations in our ability to assess blood flow in the optic nerve. Optical coherence tomography angiography (OCTA) has emerged as a promising new technology well positioned to become the first clinically suitable test of optic nerve perfusion. OCTA uses the motion of red

“There is early evidence that OCTA may be of particular use in very early or very late stage disease where our current functional or structural diagnostic modalities fall short, however, its superiority to existing technology has not been confirmed.”

PMID 31158045

# CONCLUSION

- Many advantages over IVFA
- Better detection of non-perfusion in diabetes and vein occlusion
- Earlier detection of pre-retinal neovascularization
- Improved visualization of CNV and precursors
- Potential for early glaucoma detection and end-stage management

