

INTERPRETING RETINAL OCT'S & INTRODUCING OCT-ANGIOGRAPHY

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1

Disclosures

- | | |
|-------------------------|------------------|
| ▣ Aerie Pharmaceuticals | ▣ Nova Ocular |
| ▣ Biotissue | ▣ Novartis |
| ▣ Diopsys | ▣ Optovue |
| ▣ Ellex | ▣ Quantel |
| ▣ EyePromise | ▣ Reichert |
| ▣ Ivantis | ▣ RevolutionEHR |
| ▣ Lumenis | ▣ Sight Sciences |
| ▣ Maculogix | ▣ Shire |
| ▣ Nidek | ▣ Sun Pharma |

2

Overview

- ▣ Beyond (Retina) First
 - History/principles of the OCT
 - What does the normal retinal OCT look like
 - Vitreal disorders
 - Retinal/RPE disorders
 - Choroidal disorders
- ▣ Glaucoma
 - What does the normal ONH OCT look like
 - ▣ rNFL
 - ▣ GCA
 - ONH disorders

3

History of OCT

- ▣ 1991: 1st scientific description of the OCT
 - Huang et al, Science. 1991; 254 (5035): 1178-1181.
- ▣ Original Founders:
 - David Huang, M.D., PhD
 - Dr. James Fujimoto, PhD
 - Eric Swanson, MS
 - Carmen Puliafito, M.D.
 - Joel Schulman, M.D.
- ▣ Introduced commercially in the mid-1990's

4

The Beginning OCT

- ▣ 1995 OCT1 debuted at 100 axial scans per second with a resolution of 20 microns

5

Evolving the OCT

- ▣ Stratus OCT – 2002
 - “Time domain”
 - 500 axial scans/second
 - 10 micron resolution

6

Evolving the OCT

- ▣ "Spectral-Domain" OCT – 2007
 - "Fourier-Domain"
 - 27,000 – 40,000 axial scans/second
 - ▣ Analyzes data using a spectrometer
 - ▣ Does not use a moving mirror
 - ▣ Very fast acquisition speed
 - 65x greater acquisition speed
 - ▣ 3-D imaging

*** 3.5 – 6 micron resolution ***

7

Understanding and Interpreting the Retina OCT

- ▣ Choroid
- ▣ 10 layers of the retina
 - RPE
 - Photoreceptors
 - ELM
 - Outer nuclear layer
 - Outer plexiform layer
 - Inner nuclear layer
 - Inner plexiform layer
 - Ganglion cell layer
 - Nerve fiber layer
 - ILM
- ▣ Vitreous

8

Posterior Vitreous Detachment (PVD)

9

Vitreo-macular adhesion/traction (VMT)

10

Vitreomacular traction

11

Vitreomacular traction

12

Macular hole

- ▣ Unilateral, decreased vision
 - Often in 60-80 year old women
 - Anyone w/ a history of trauma
- ▣ Symptoms:
 - Decreased vision, metamorphopsia
 - 20/200 for full thickness holes
- ▣ Signs:
 - Red hole in the macula
 - (+) Watzke-Allen sign

13

Macular hole

- ▣ Stages
 - Stage 1a -> impending hole. Normal foveal depression with yellow spot/dot in fovea.
 - Stage 1b -> Abnormal foveal depression with yellow ring.

Stage 1b macular hole

14

Macular hole

- ▣ Stages
 - Stage 2 -> Small full-thickness hole. 20/80 - 20/400.
 - Stage 3 -> Full-thickness hole w/ cuff of SRF. No PVD
 - Stage 4 -> Full-thickness hole with cuff of SRF, with complete PVD.

Stage 2 macular hole

15

Macular hole

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 - Stage 4 -> Full-thickness hole with cuff of SRF, with complete PVD.

Stage 3
Macular
hole

Stage 4 macular hole →

16

Macular Hole

- ▣ Treatment:
 - Stage 2 holes or beyond (full thickness macular holes)
 - Vision 20/40 or worse
 - How long has the hole been there???
 - Vitrectomy & membrane peel
 - Face down???
- ▣ Prognosis:
 - 20/40 or better in up to 65% of cases

17

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18

Lamellar Macular hole

- ▣ "Partial thickness macular holes"
- ▣ Aborted macular holes
- ▣ "Upside down anvil" "anvil-like"
- ▣ VA -> usually 20/40 or better
- ▣ 4 characteristics
 1. Irregular foveal contour
 2. Break in inner fovea
 3. Intraretinal split
 4. Intact foveal photoreceptors

19

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20

Pseudohole

- ▣ "False hole"
- ▣ Simulates macular hole w/o actual tissue dehiscence
- ▣ Full thickness retinal tissue is still present
 - Not an anvil
- ▣ VA
 - Usually 20/20 - 20/30 unless significant ERM is present

21

Cystoid Macular Edema (CME)

22

Cystoid Macular Edema (CME)

23

Diabetic macular edema (DME)

24

CSME vs. CME???

25

Central Serous Chorioretinopathy (CSR)

- ▣ Demographics
 - 25-50 year old men, stressed/Type A personalities
- ▣ Symptoms
 - Unilateral, blurred vision
 - VA -> usually 20/20 - 20/80
 - Metamorphopsia
- ▣ Signs
 - Localized serous detachment of the neurosensory retina in the macula

26

Central Serous Chorioretinopathy

- ▣ DDx:
 - Optic disc pit
 - CNVM

27

Central Serous Chorioretinopathy

- ▣ Med associations:
 - Steroids
 - Nasal sprays, steroid creams, oral, injectable
 - Ephedra
 - Ephedrine & pseudoephedrine
- ▣ Treatment:
 - Observation/lifestyle change
 - D/C steroid if possible
 - Possible laser therapy

28

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29

Plaquenil Toxicity

- ▣ Antimalarials:
 - Chloroquine
 - Hydroxychloroquine (Plaquenil)
- ▣ Now used for RA, SLE, Sjogren's, etc.
- ▣ Toxicity risk is low, but....
- ▣ Lots of different screening recommendations have been proposed

30

Plaquenil Toxicity

- ▣ Risk Factors:
 - Cumulative dose**
 - 1000 gram cumulative dose for Plaquenil
 - 6.85 years to reach that
 - Daily dose
 - Age
 - Liver or kidney dysfunction
 - Pre-existing retinal disease or maculopathy

31

Plaquenil Toxicity

- ▣ Symptoms:
 - Asymptomatic early
 - Paracentral visual field defects affecting reading
 - Color vision changes
- ▣ Signs:

32

Plaquenil Toxicity

- ▣ Recommended Screening Guidelines:
 1. Baseline exam within the first year of starting Plaquenil
 - Biomicroscopy exam, 10-2 VF, Fundus photos, OCT
 - After 5 years, annual screening exams
 - SD-OCT or
 - mfERG or
 - Fundus autofluorescence

33

Plaquenil Toxicity

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 1. Baseline exam within the first year of starting Plaquenil
 - Biomicroscopy exam, 10-2 VF, Fundus photos
 - SD-OCT or mfERG or fundus autofluorescence
 - After 5 years, annual screening exams
 - Biomicroscopy exam along with 10-2 VF and SD-OCT or
 - mfERG or
 - Fundus autofluorescence

34

Plaquenil toxicity “saucer sign”

35

Plaquenil Toxicity

- ▣ Tests not recommended for screening
 - Fundus photography
 - Time-domain OCT
 - FA
 - Full-field ERG
 - EOG
 - Color vision testing
 - Amsler grid

36

Plaquenil Toxicity

- ▣ Treatment:
 - No medical therapy is available to treat/cure the toxicity
 - D/C the med if possible
 - Work with the PCP

37

Dry Age-related Macular Degeneration (AMD)

38

Dry Age-related Macular Degeneration (AMD)

39

Dry Age-related Macular Degeneration (AMD)

40

Dry Age-related Macular Degeneration (AMD)

41

Dry Age-related Macular Degeneration (AMD)

42

Wet Age-related Macular Degeneration (AMD)

43

Wet Age-related Macular Degeneration (AMD)

44

OCT-ANGIOGRAPHY

45

OCT Angiography: the Next Chapter in Posterior Imaging

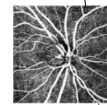
- Images retinal microvasculature without dye injection
- Displays structure and function from a single imaging system



2002: Time Domain OCT



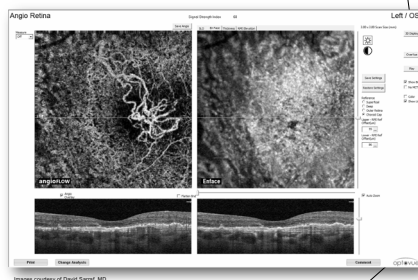
2006: Spectral Domain OCT



2014: OCTA

46

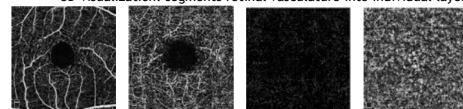
Structure & Function from One System



47

A New Approach to Visualizing Blood Flow

- Patient Benefits
 - Reduces patient burden to allow more frequent imaging
 - Avoid potential side-effects of fluorescein injection
- Clinical Benefits
 - Faster than a dye-based procedure
 - Ultra-high resolution imaging of retinal microvasculature
 - 3D visualization: segments retinal vasculature into individual layers



48

OCT-Angiography

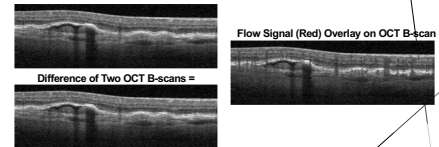
How Does it Work?

49

Principles of OCTA

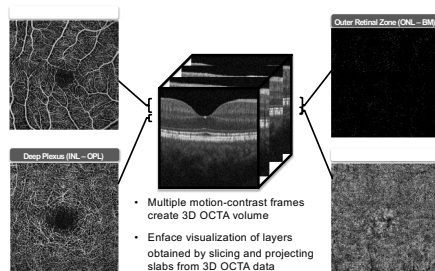
OCTA uses motion contrast to detect flow from OCT data

- Rapidly acquires multiple cross-sectional images from a single location on the retina
- Flow is the difference in signal between two sequential B-scans

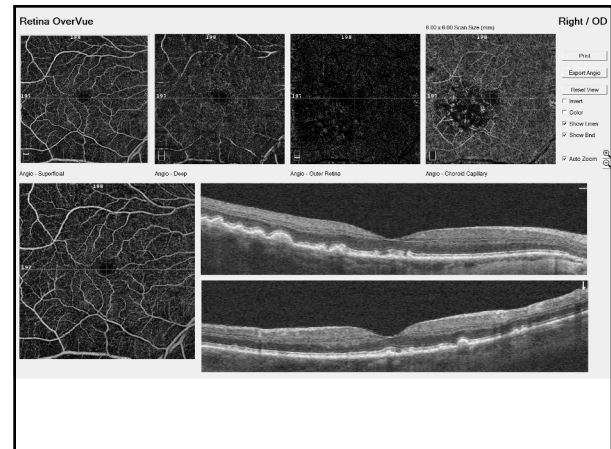


50

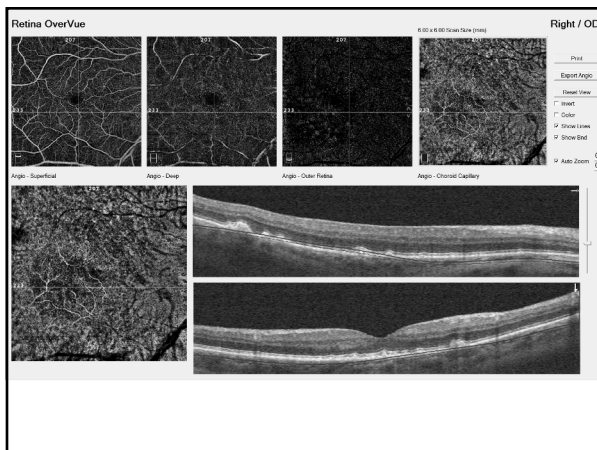
Enface OCTA Generated from OCTA Volume Data



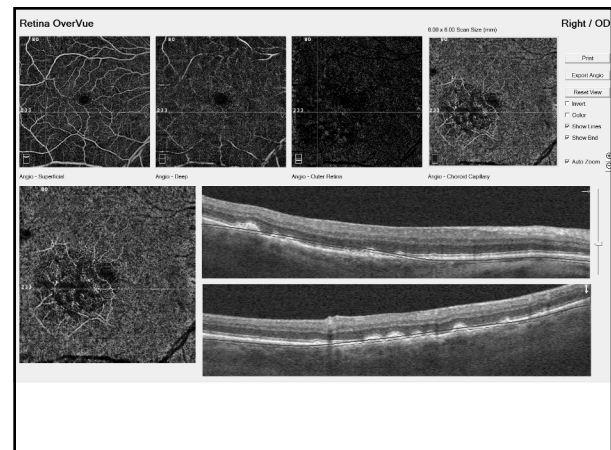
51



52



53

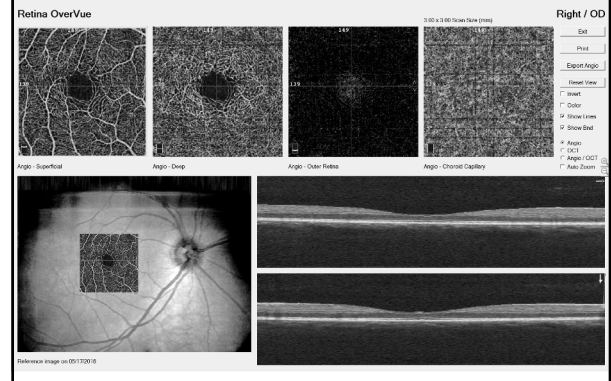


54

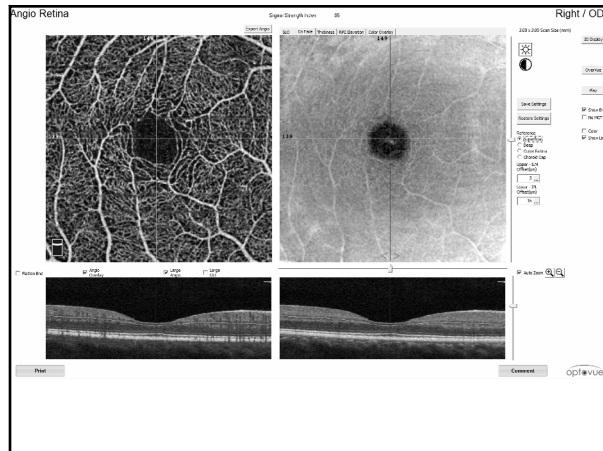
OCT-ANGIOGRAPHY

55

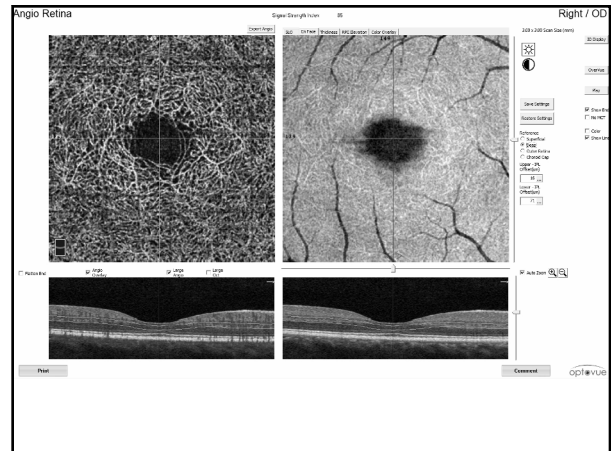
What is normal?



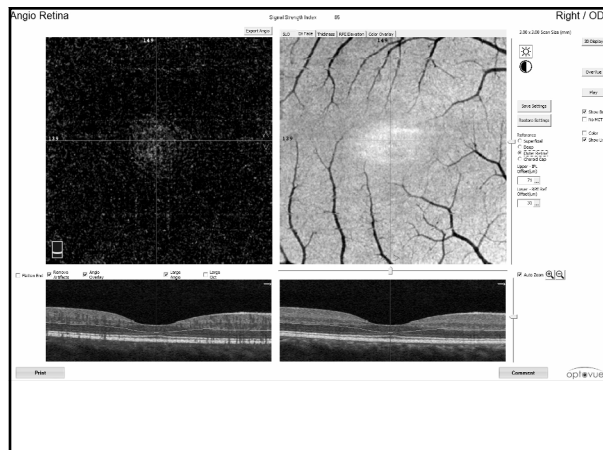
56



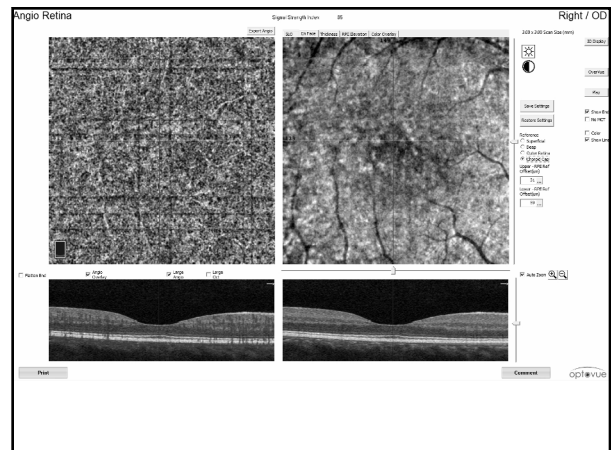
57



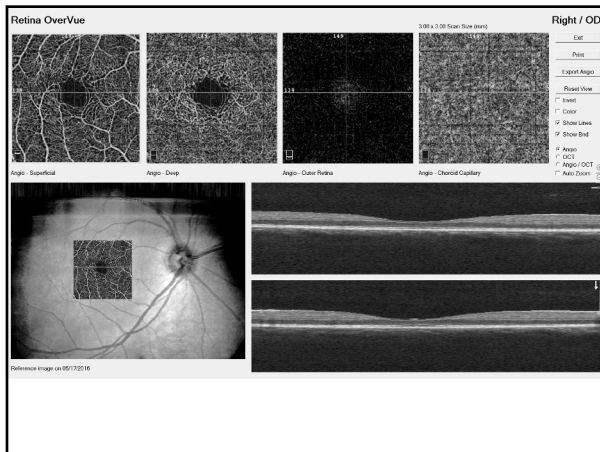
58



59



60



61

OCT-A in our clinic

Indications:

AMD - dry vs. wet

Diabetics -

is there neo?

is there non-perfusion (capillary dropout)?

Vein Occlusions

Glaucoma patients

nerve perfusion?

62

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63