

# Glaucoma Imaging and Analysis

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# Financial disclosures

- None

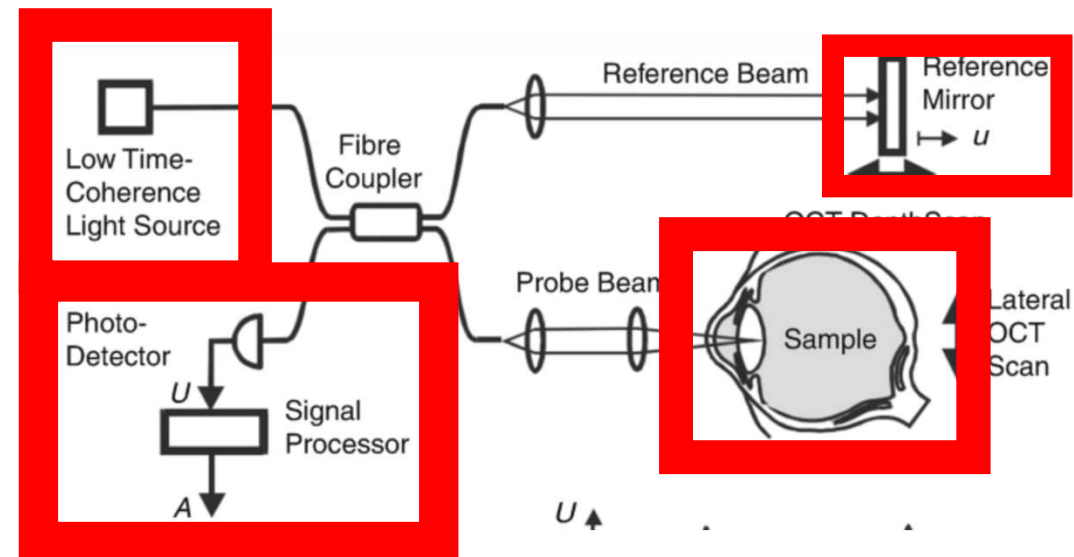
# Learning objectives

- How to determine if an OCT or HVF is reliable
- What are common artifacts skewing OCTs
- Identify green and red disease
- Define the global indices of HVF
- Define trend-based and event-based progression
- Note slow progressors vs. fast progressors on HVF

# Optical Coherence Tomography

# OCT

- Light source
  - Split to reference arm and a media
- Reference arm
  - Mirror reflects light back
- Media
  - eg, the eye
- Detector
  - Returning light from both arms combines to form interference pattern generating cross-sectional image



# OCT: distinguishing normal from glaucoma

- Most reliable RNFL parameters:
  - Average RNFL
  - Inferior quadrant RNFL
  - IT clock hour (7/5 OD/OS, respectively)
  - ST clock hour (11/1)
  - Clock hour 6
- Excellent ability
  - AUC 0.923– 0.957
  - AUC 1 = perfect test w/ 100% sensitivity and specificity
- Regardless of disease severity

# OCT: distinguishing normal from glaucoma

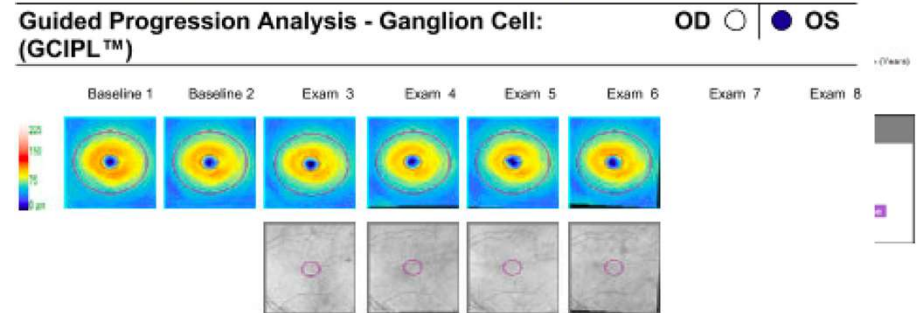
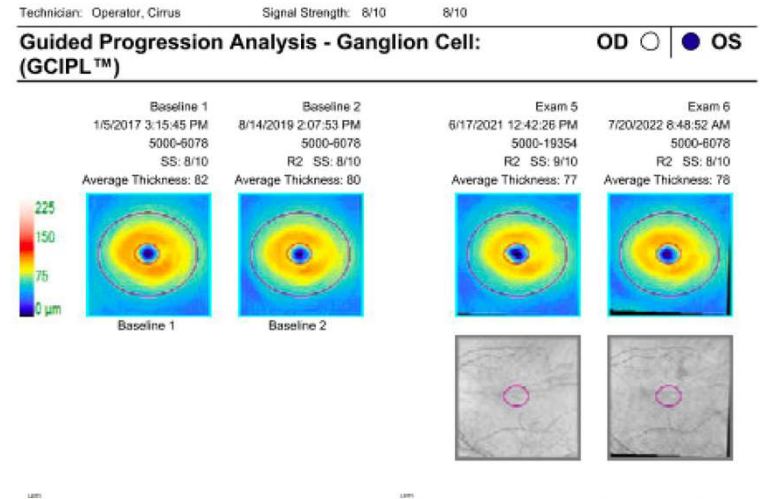
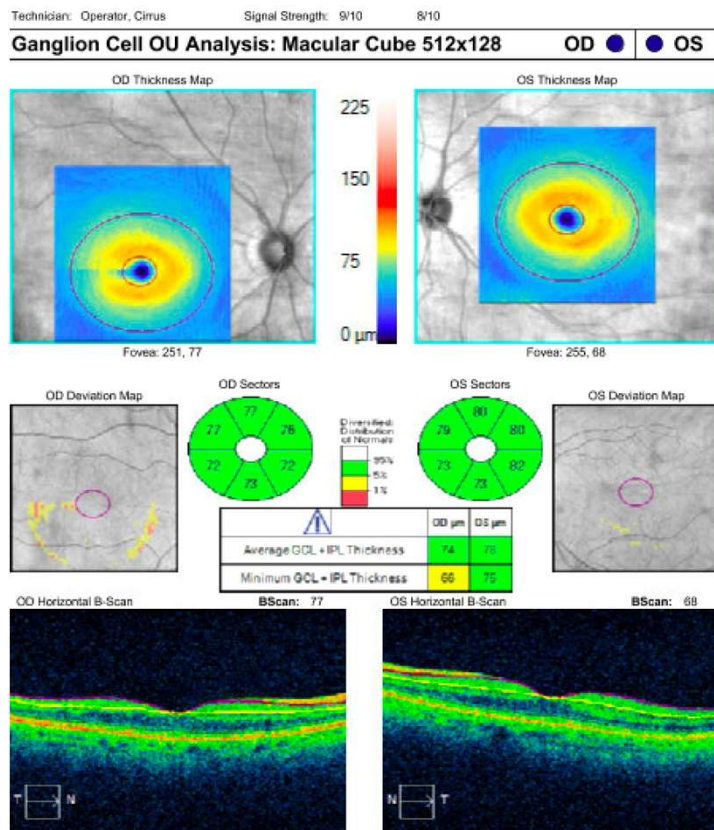
- Most reliable ONH parameters
  - VRT
    - Vertical rim thickness
    - Total rim thickness in vertical meridians
  - Rim area
  - VCDR
  - CDR
  - HRT
    - Horizontal rim thickness
    - Total rim thickness in horizontal meridians
      - AUC 0.901 and 0.963

# OCT: distinguishing normal from glaucoma

- Macular ganglion cell analysis
  - Higher concentration of ganglion cells
  - Less effected by peripheral chorioretinal disease
  - Useful in early and advanced glaucoma
  - Useful when RNFL reaches 'floor'
  - Skewed by macular disease



# OCT: ganglion cell analysis



**GCL + IPL Summary Parameters**

	Exam Date/Time	Serial Number	Registration Method	SS	Average Thickness:	Total Superior Thickness ( $\mu\text{m}$ )	Total Inferior Thickness ( $\mu\text{m}$ )
Baseline 1:	1 1/5/2017 3:15:45 PM	5000-6078		8/10	82	83	81
Baseline 2:	2 8/14/2019 2:07:53 PM	5000-6078	R2	8/10	80	81	80
	3 2/11/2020 5:05:23 PM	5000-6138	R1	10/10	78	79	77
	4 5/25/2021 1:15:08 PM	5000-21009	R2	9/10	78	79	77
	5 6/17/2021 12:42:26 PM	5000-19354	R2	9/10	77	78	76
Current:	6 7/20/2022 8:48:52 AM	5000-6078	R2	8/10	78	79	77

# OCT: age-related loss

- Histologically, we lose approx. 5,000 axons/year
- OCT-measured average RNFL loss varies: 0.16um/year - 0.44um/year
- 0.365um/year
  - Most consistent approximation
  - Remember approximately 1/3<sup>rd</sup> micron RNFL loss per year

# OCT: reliable test

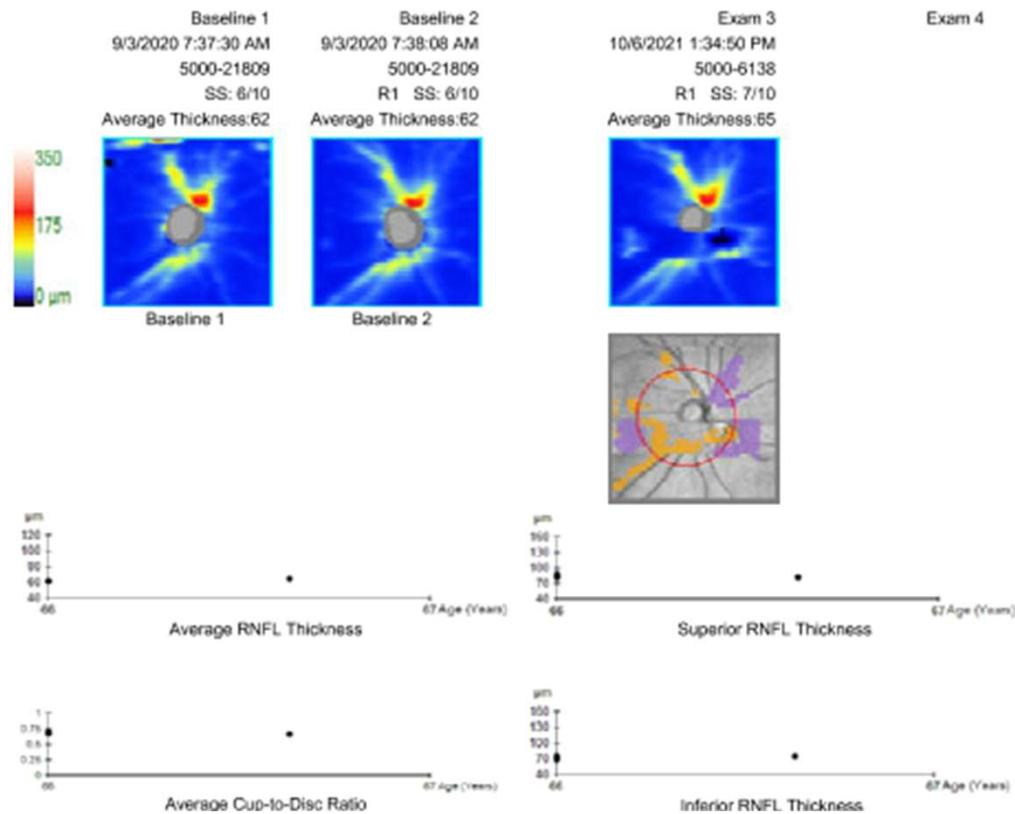
- Signal strength
  - $\geq 6$
  - Decreasing SS erroneously measures a thinner RNFL
- Artifact
  - Blocking
  - Movement
- Improper identification of ON head
- Segmentation error
  - Improperly identifying RNFL borders

# OCT: artifact

- Saccade Artifact

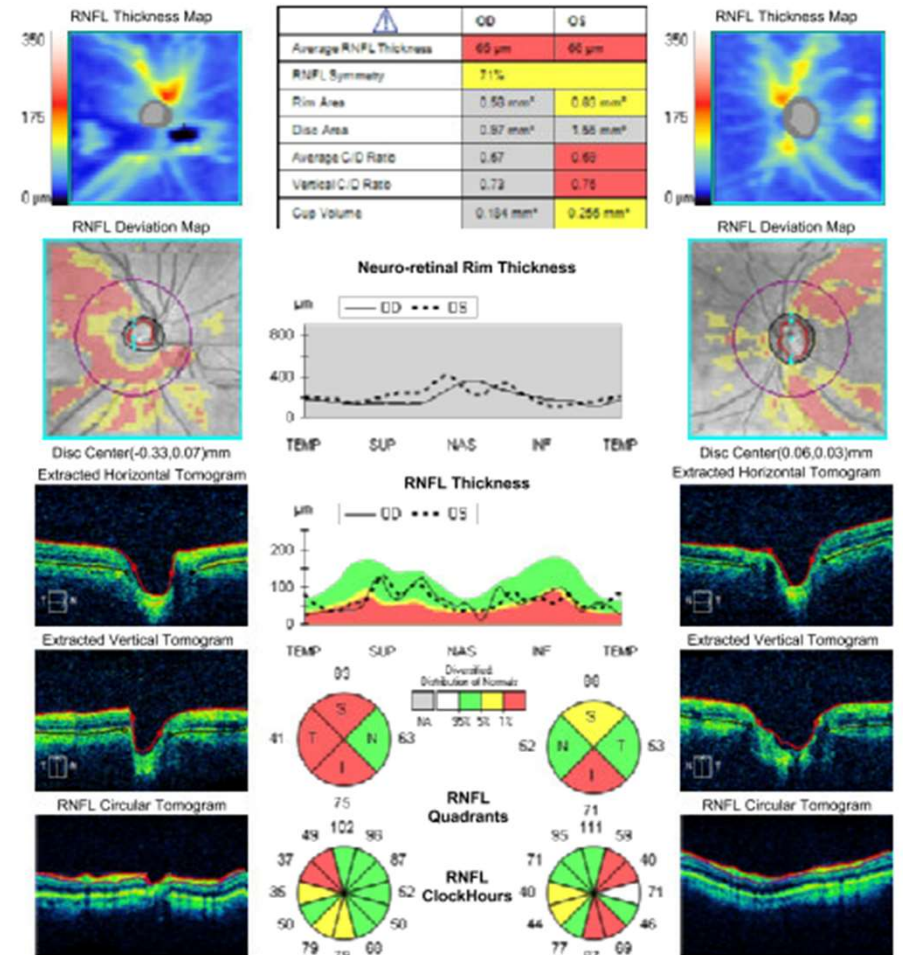
## Guided Progression Analysis: (GPA™)

OD ● ○ OS



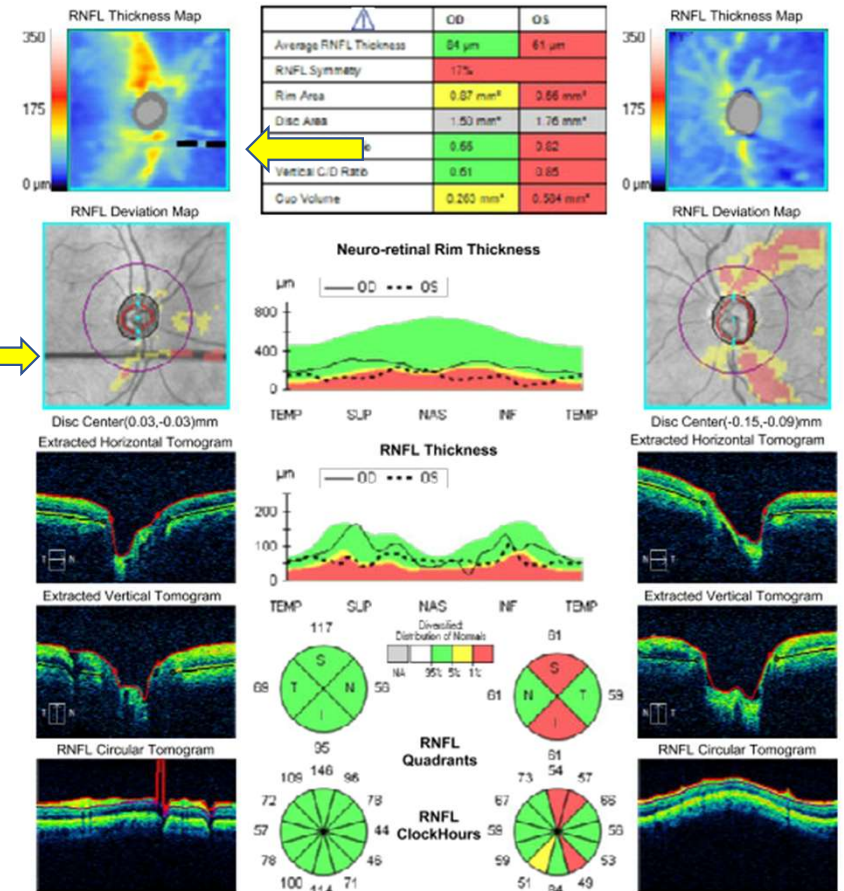
## ONH and RNFL OU Analysis: Optic Disc Cube 200x200

OD ● ● OS

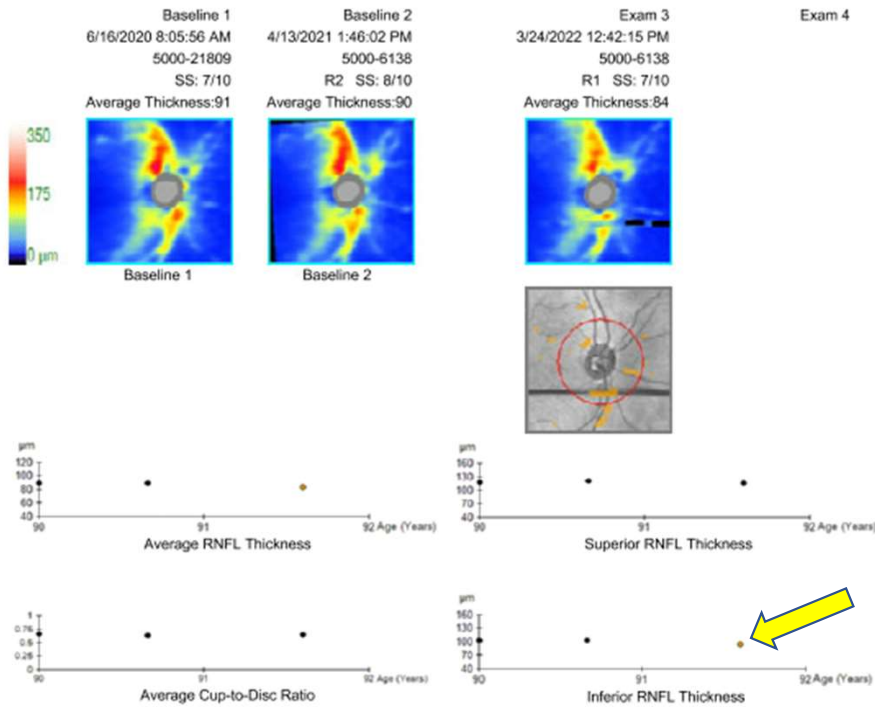


# OCT artifact

- Blink artifact



**Guided Progression Analysis: (GPA™)** OD ● OS ○

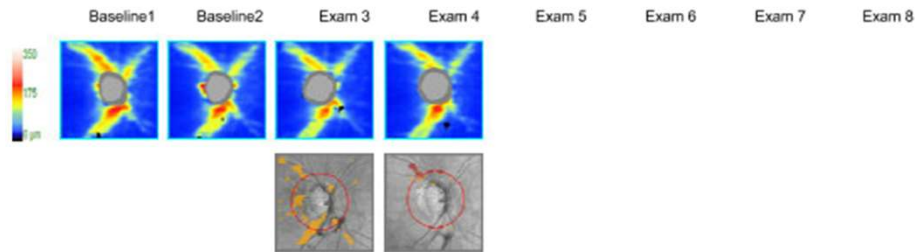




# OCT: artifact

- Blocking artifact
- Vitreous

## Guided Progression Analysis: (GPA™) OD ● OS ○

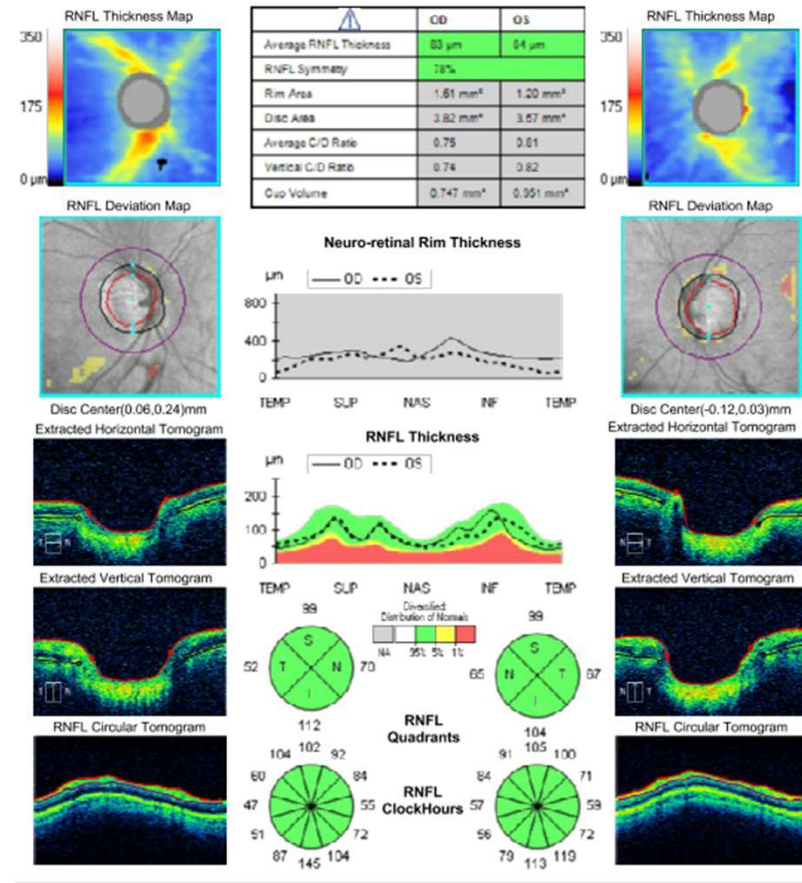


RNFL and ONH Summary Parameters

	Exam Date/Time	Serial Number	Registration Method	SS	Avg RNFL Thickness (μm)	Inf Quadrant RNFL (μm)	Sup Quadrant RNFL (μm)	Rim Area (mm²)	Average Cup-to-Disc Ratio	Vertical Cup-to-Disc Ratio	Cup Volume (mm³)
Baseline 1:	1 4/23/2019 1:14:36 PM	5000-6138		6/10	88	115	108	1.42	0.73	0.71	0.570
Baseline 2:	2 9/18/2020 8:35:02 AM	5000-19354	R1	8/10	86	114	100	1.08	0.78	0.75	0.607
	3 9/29/2021 9:37:28 AM	5000-6078	R1	7/10	81	103	104	1.36	0.77	0.76	0.756
Current:	4 12/8/2022 1:25:54 PM	5000-19354	R1	6/10	83	112	99	1.61	0.75	0.74	0.747

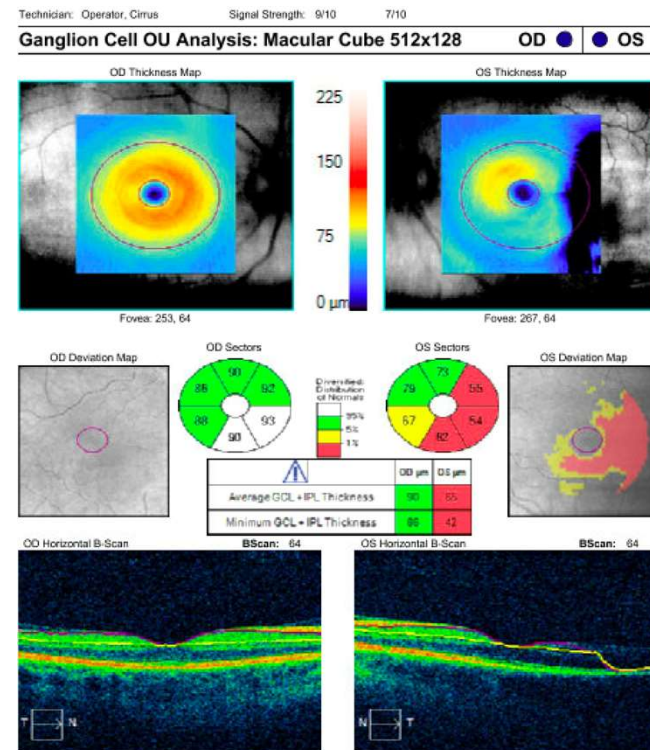


## ONH and RNFL OU Analysis: Optic Disc Cube 200x200 OD ● OS ●



# OCT: artifact

- Segmentation failure
  - Improper identification of limits of
    - RNFL, GCIPL, GCC
  - Result in
    - Augmented or exaggerated RNFL measurements
- Occur when
  - ONH, inner retinal layers, and/or outer retinal layers improperly identified
  - Boundaries of inner and outer layers not completely identified
  - "Stretching" these segmentation lines up or down



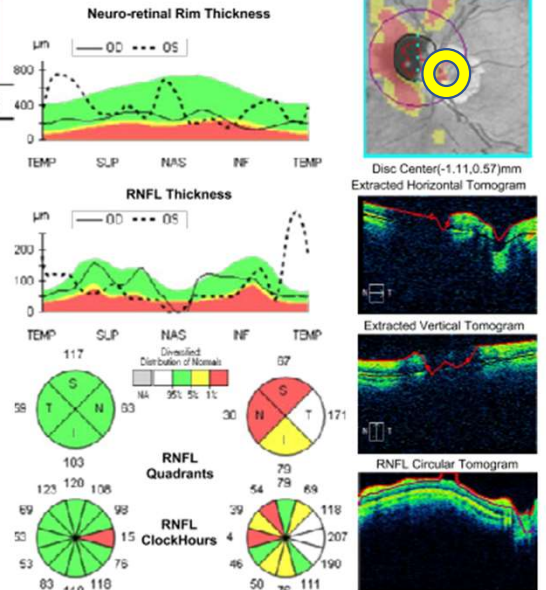
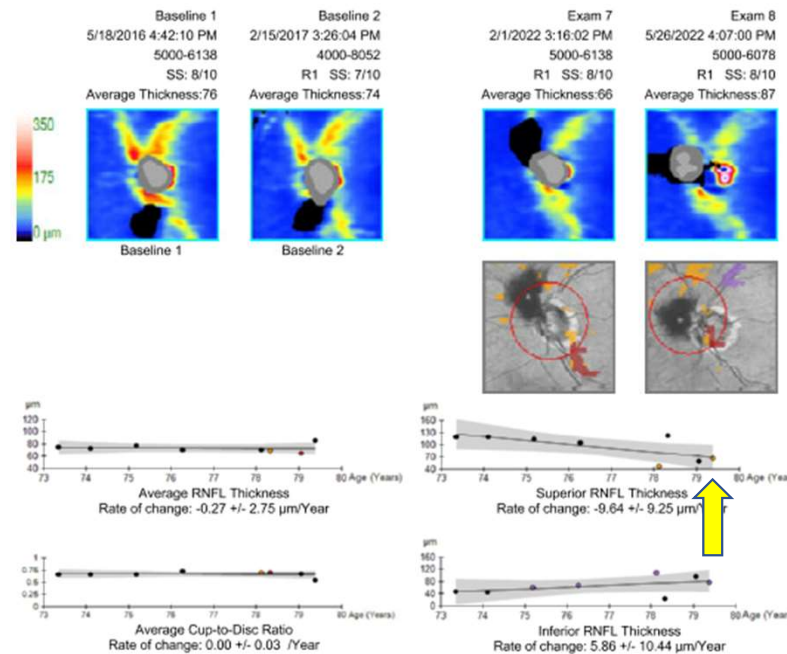
# OCT: artifact

## • Failure of ONH recognition algorithm

- Decentration
- PPA
- Improper structure
  - Macula
  - PVD



## Guided Progression Analysis: (GPA™)





# OCT: red and green disease

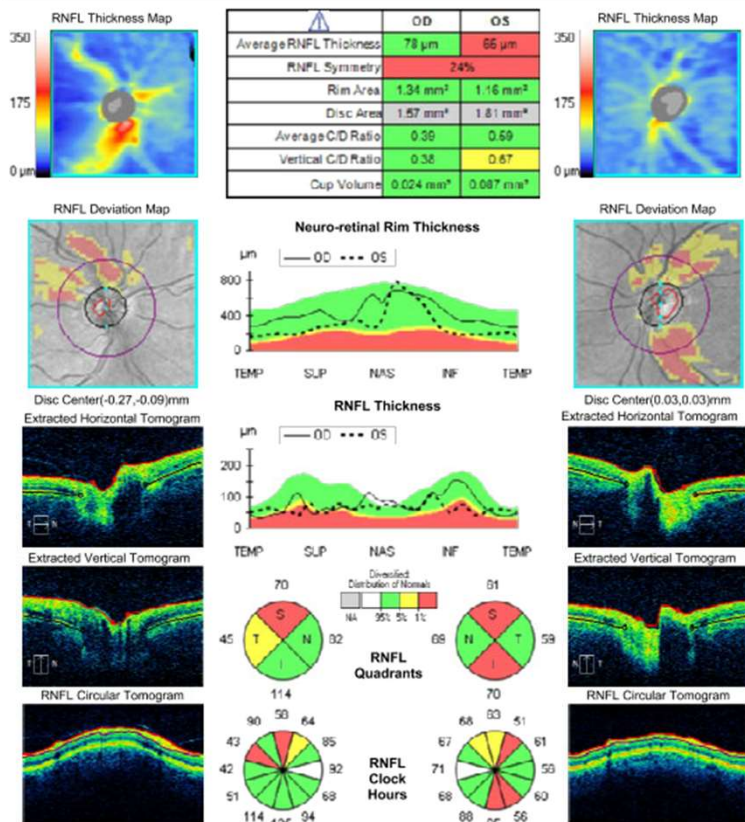
- Remember:
  - Color coding parameters are based on RNFL measurements compared with age-matched controls
  - Green
    - 'within normal limits'
    - 95%
  - Yellow
    - 'borderline'
    - <5%
  - Red
    - 'outside normal limits'
    - <1%
- Thinner than average RNFL exists without pathology, and
- Pathology can exist with seemingly 'normal' RNFL

# OCT: red disease

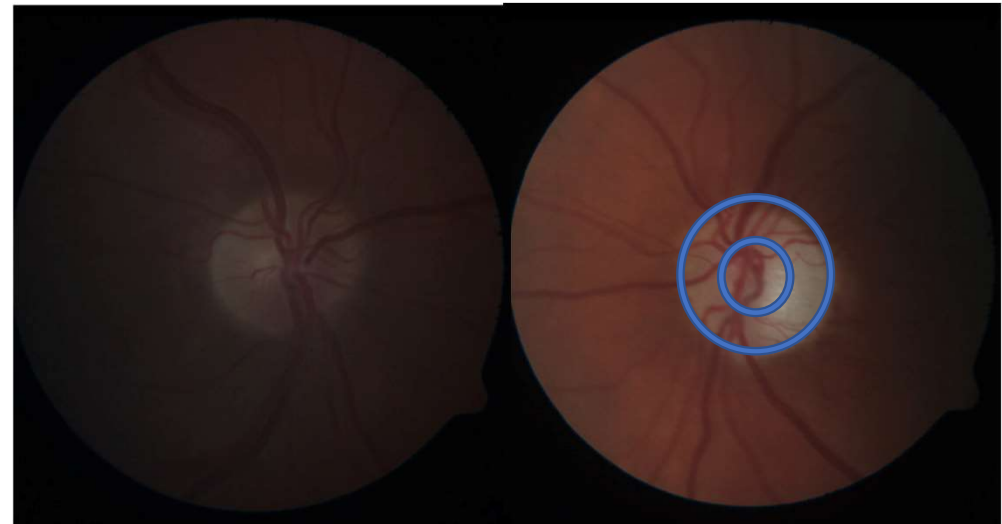
- Abnormal OCT w/ no glaucoma, i.e. a false positive
  - ON drusen
  - ON ischemia
  - ON atrophy
  - ON hypoplasia
  - Small ON
  - Tilted ON
  - Congenital anomalies
  - High myopia
  - Poor SS
  - Artifact
  - Peripheral retinal disease
  - PRP/chorioretinal scarring

# OCT: Red disease

Technician: Operator, Cirrus Signal Strength: 9/10 9/10  
**ONH and RNFL OU Analysis: Optic Disc Cube 200x200** OD ● ● OS

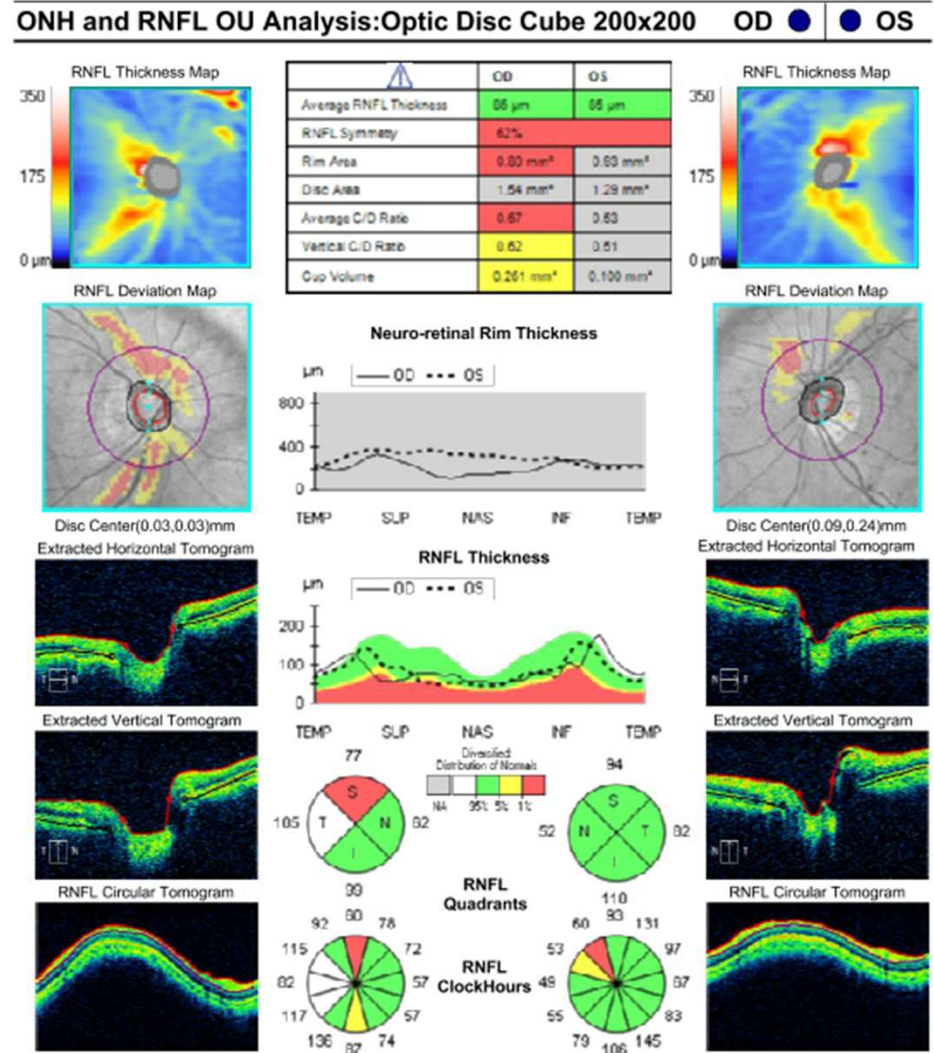


- Patient with h/o OHT
- AND bilateral sequential NAION following cataract surgery



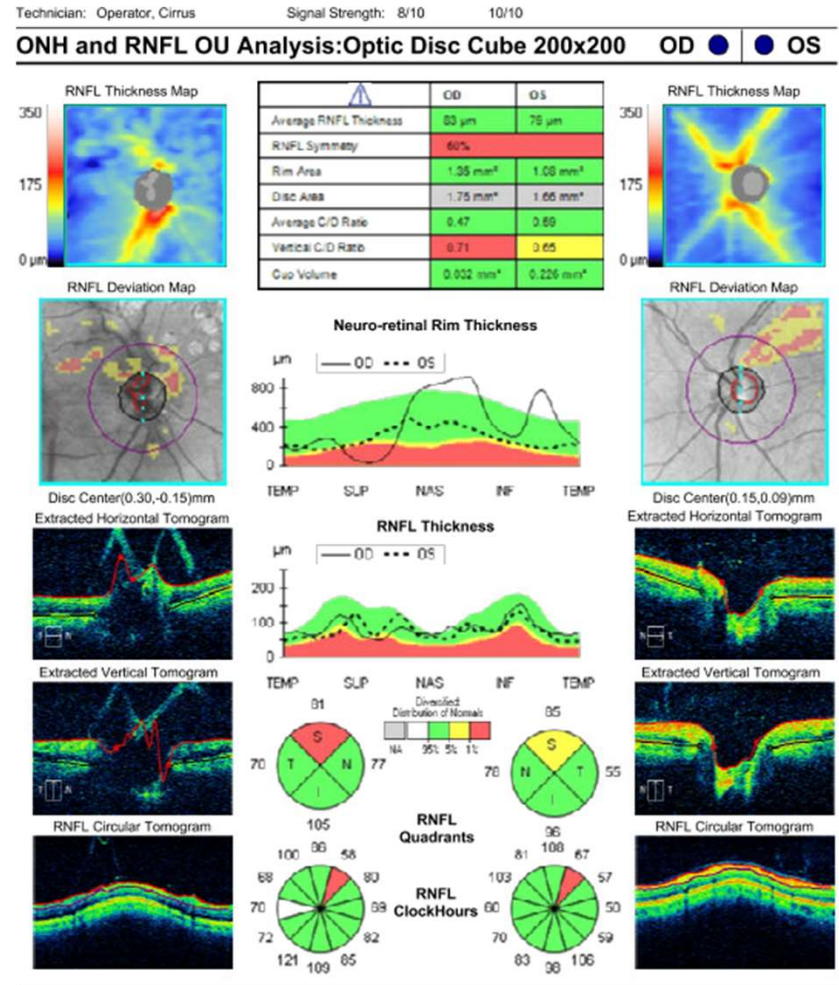
# OCT: red disease

- High myopia
  - RNFL thickness decreases w/ increasing AL
  - False positive (red disease) seen more often in
    - RNFL (~50% FP)
    - Macular GCL-IPL (~25% FP)
    - ONH measurements perform best (only 7% FP)
  - Temporal deviation of RNFL 'butterfly'



# OCT: red disease

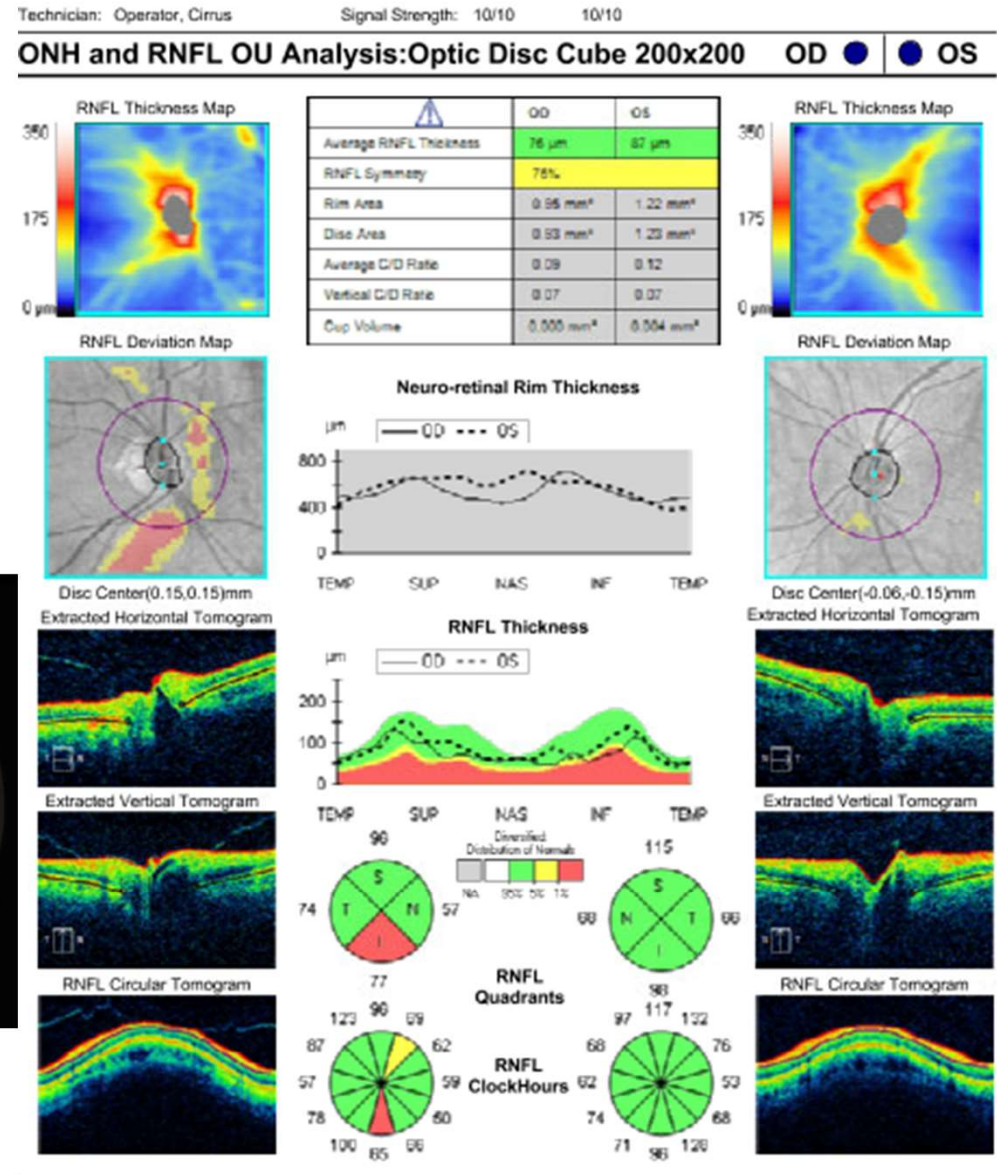
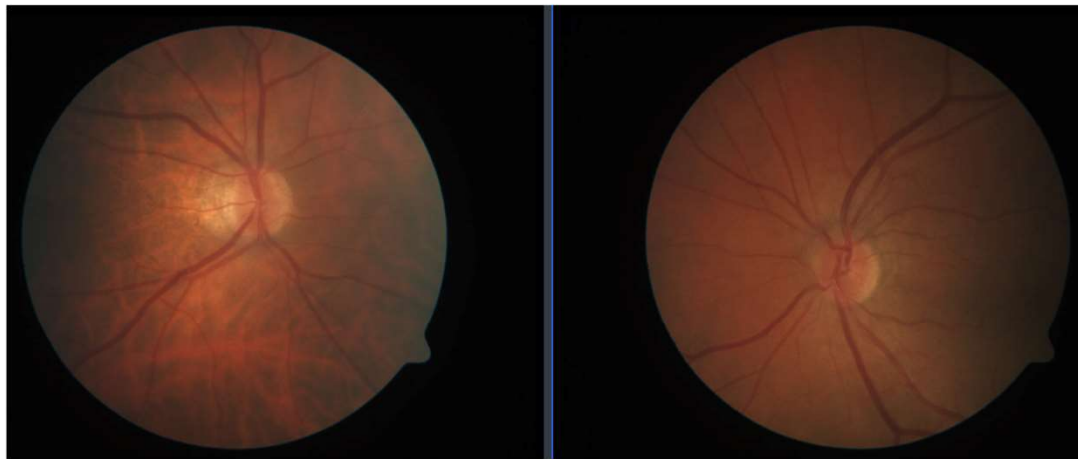
- Red dz 2/2 BRVO S/P PRP





# OCT: red disease

- ON hypoplasia or small ON

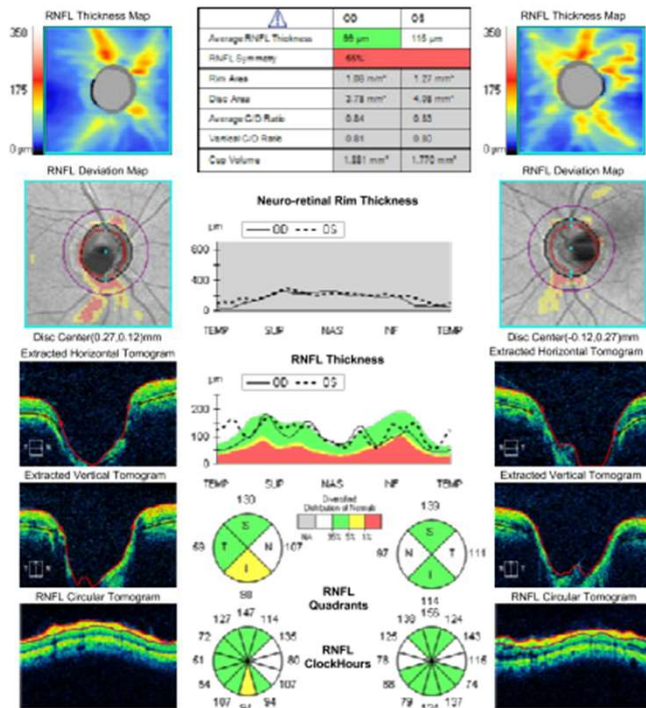


# OCT: green disease

- 'Normal' OCT with glaucomatous pathology, i.e. a false negative
  - Secondary to:
    - Other ocular pathology augmenting RNFL measurement in setting of glaucoma
      - Thickened/edematous RNFL
        - ERM
        - Macular edema
        - DME
        - RVO
        - Uveitis
      - ON edema
        - Acute NAION
        - Uveitis
        - Papilledema
  - OR
    - Pathologic, glaucomatous thinning of RNFL greater than age-related change while still measuring 'within normal limits.'

# OCT: green disease?

ONH and RNFL OU Analysis: Optic Disc Cube 200x200 OD OS

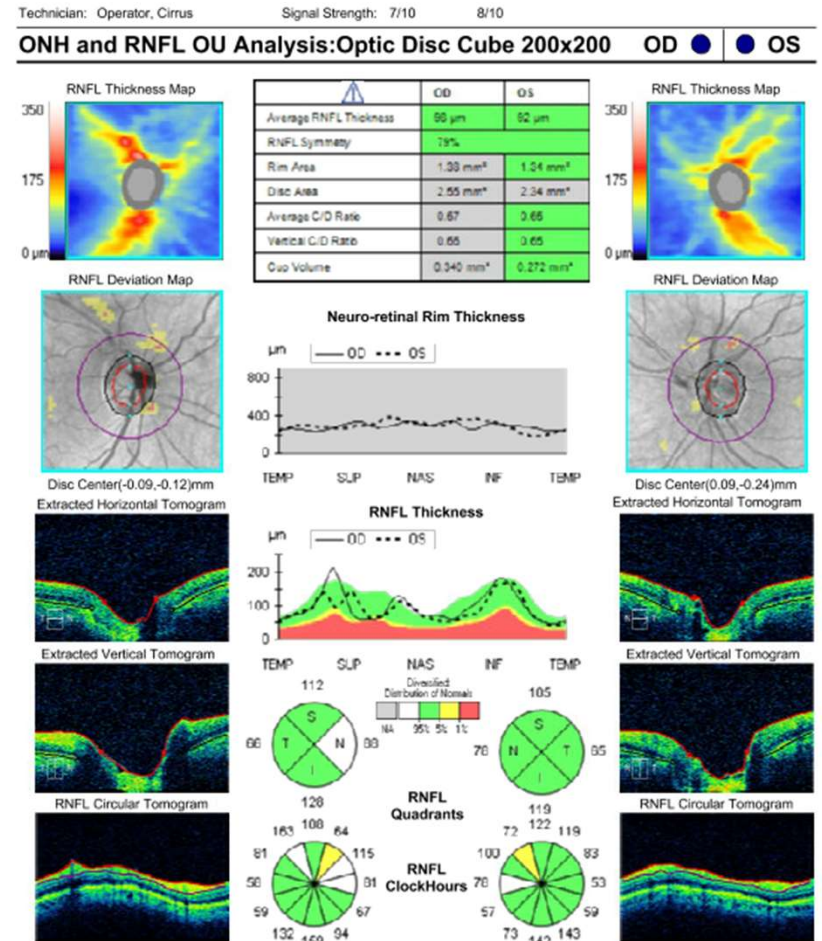


- No
- ON pit



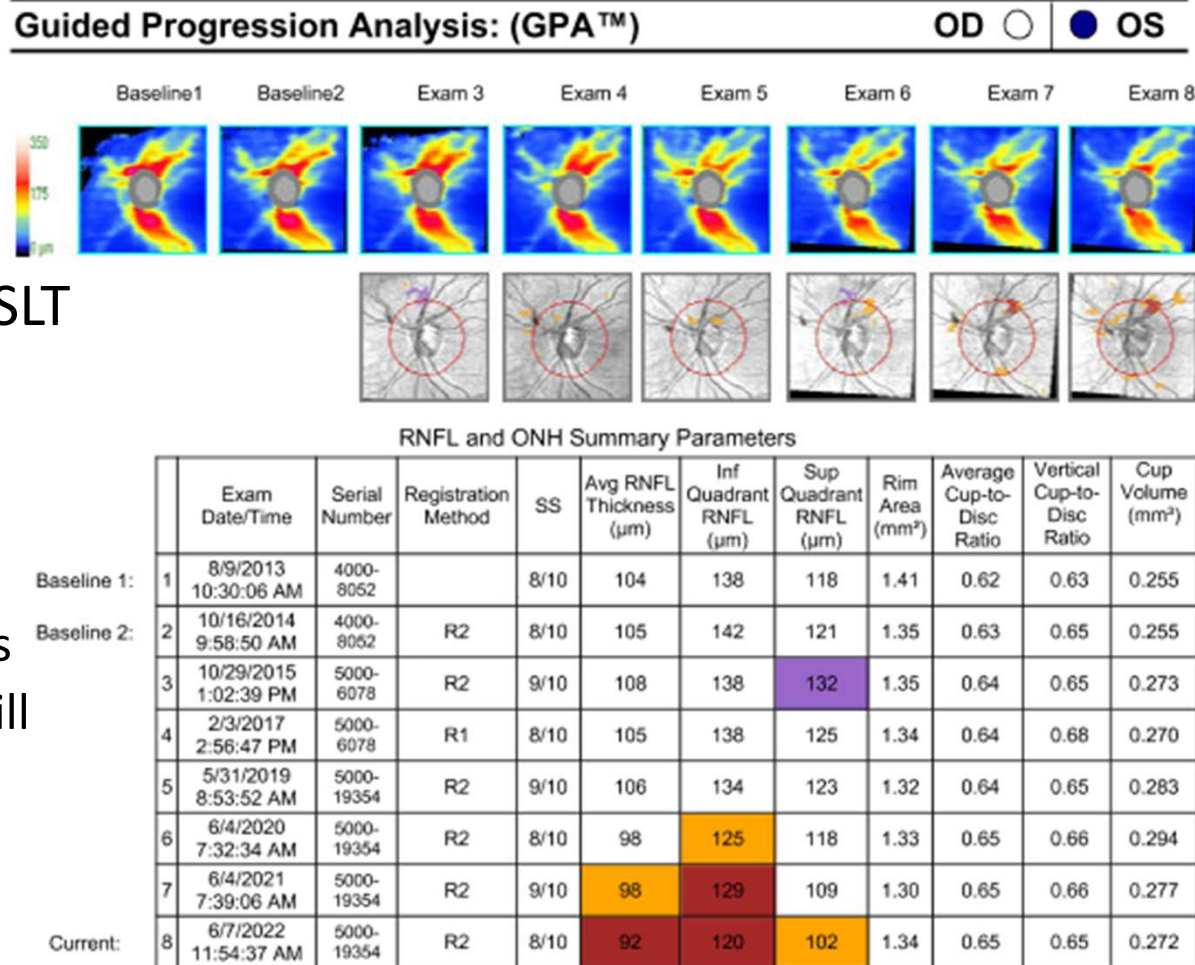
# OCT: green disease

- 66 yo WF
- Tmax 30s
- Originally treated as OHT w/ PGa lowering IOP to low 20s



# OCT: green disease

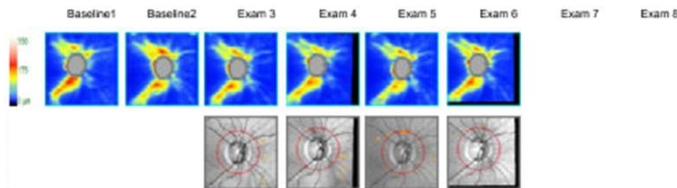
- Subsequent scans revealed progressive loss of RNFL
- Treatment augmented with SLT
- IOP maintained in  $\leq 18$
- What next?
  - Reset the baseline OCT exams
  - Or every subsequent exam will appear as progressing



# OCT: green disease

- Originally followed as GS
- Repeat OCT show progressive RNFL loss

## Guided Progression Analysis: (GPA™) OD ● ○ OS



RNFL and ONH Summary Parameters

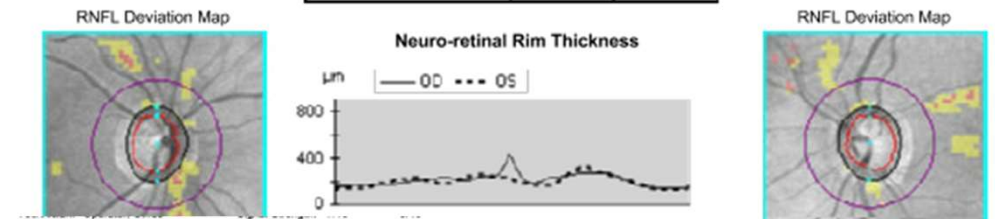
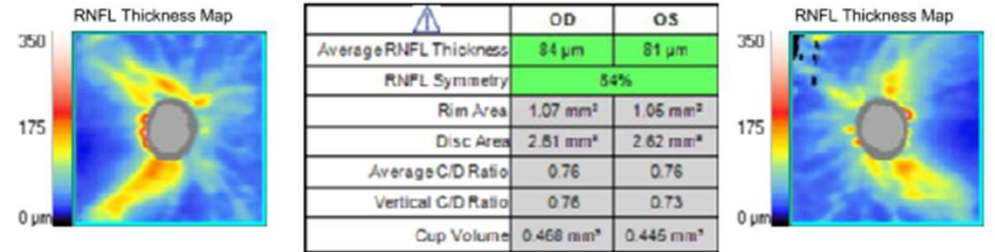
	Exam Date/Time	Serial Number	Registration Method	SS	Avg RNFL Thickness (μm)	Inf Quadrant RNFL (μm)	Sup Quadrant RNFL (μm)	Rim Area (mm²)	Average Cup-to-Disc Ratio	Vertical Cup-to-Disc Ratio	Cup Volume (mm³)
Baseline 1:	3/13/2015 9:40:23 AM	4000-3411		6/10	87	112	107	1.12	0.75	0.74	0.506
Baseline 2:	7/28/2017 9:57:44 AM	5000-6138	R1	7/10	88	114	107	1.09	0.76	0.76	0.471
3:	3/1/2019 8:34:10 AM	5000-6138	R1	8/10	83	110	103	1.11	0.74	0.75	0.420
4:	2/9/2021 6:00:30 AM	5000-19354	R2	7/10	80	104	91	1.09	0.74	0.77	0.448
5:	11/23/2021 7:46:32 AM	5000-6138	R1	7/10	77	104	91	1.08	0.73	0.74	0.419
Current:	11/9/2022 8:12:09 AM	5000-21809	R2	7/10	82	107	94	1.10	0.75	0.75	0.458

Technician: Operator, Cirrus

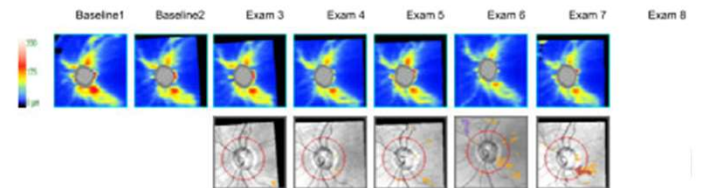
Signal Strength: 7/10

8/10

## ONH and RNFL OU Analysis: Optic Disc Cube 200x200 OD ● ● OS

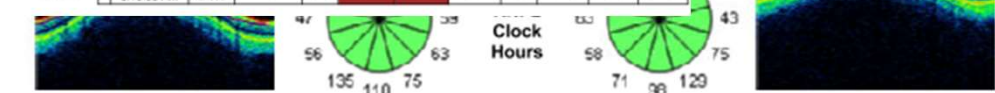


## Guided Progression Analysis: (GPA™) OD ○ ● OS



RNFL and ONH Summary Parameters

	Exam Date/Time	Serial Number	Registration Method	SS	Avg RNFL Thickness (μm)	Inf Quadrant RNFL (μm)	Sup Quadrant RNFL (μm)	Rim Area (mm²)	Average Cup-to-Disc Ratio	Vertical Cup-to-Disc Ratio	Cup Volume (mm³)
Baseline 1:	9/30/2011 9:01:30 AM	4000-3411		7/10	87	107	98	1.14	0.72	0.67	0.373
Baseline 2:	3/13/2015 9:41:09 AM	4000-3411	R2	7/10	84	106	97	1.03	0.76	0.71	0.448
3:	7/28/2017 9:58:56 AM	5000-6138	R2	9/10	86	108	96	1.05	0.76	0.74	0.442
4:	3/1/2019 8:34:51 AM	5000-6138	R2	7/10	81	100	96	1.02	0.76	0.73	0.420
5:	2/9/2021 6:01:09 AM	5000-19354	R2	10/10	82	97	94	1.03	0.77	0.73	0.426
6:	11/23/2021 7:46:59 AM	5000-6138	R1	7/10	74	96	87	1.00	0.76	0.75	0.401
Current:	11/9/2022 8:13:03 AM	5000-21809	R2	8/10	81	98	95	1.05	0.76	0.73	0.445



# Humphrey Visual Fields

# HVF

- How it works
- Reliability indices
- Global parameters
- Structure-function correlation
- Trend-based analysis
- Event-based analysis

# HVF, what are we measuring?

- Apostilb (asb)
  - Measurement of the intensity of light (AKA luminance ( $L$ ) =  $\text{cd}/\text{m}^2$ )
    - Cd = candelas
    - $\text{Cd}/\text{m}^2 = \text{asb}$
  - BUT, our eyes can see large ranges of luminance (3-4 orders of magnitude),
  - Visual function does not have a linear relationship with luminance,
    - eg, luminance increase from 0-100 asb more likely to be appreciated than 1000-1100
  - Luminance is inversely correlated to retinal light sensitivity,
  - VF loss difficult to display w/ luminance levels,
  - THEREFORE, we use decibels (dB) instead
- Decibel
  - Sensitivity threshold of the retina
    - $\text{dB} = 10 \cdot \log(L_{\text{max}}/L)$
  - Foveal range: approximately 0-32 dB
  - Brightest stimulus required = 0 dB, dimmer stimulus required = 32 dB
- Threshold
  - “The intensity of light stimulus, which, when presented at a particular location, “n” number of times is detected by the corresponding retinal point at least 50% of the time.”

# HVF

- HVF 24-2 SITA STANDARD
  - Most commonly used
  - SITA = Swedish Interactive Thresholding Algorithm
    - Optimized to reduce time and errors from fatigue
  - 54 points tested
  - 3 degrees from each point
  - 24:
    - central 30 degrees around fixation (fovea)
    - eliminating the outermost points save for the nasal area
  - 2: equidistant points on either side of the vertical and horizontal meridians
    - 6 degrees from point on other side of meridian
    - Vertical: neuro
    - Horizontal: glaucoma

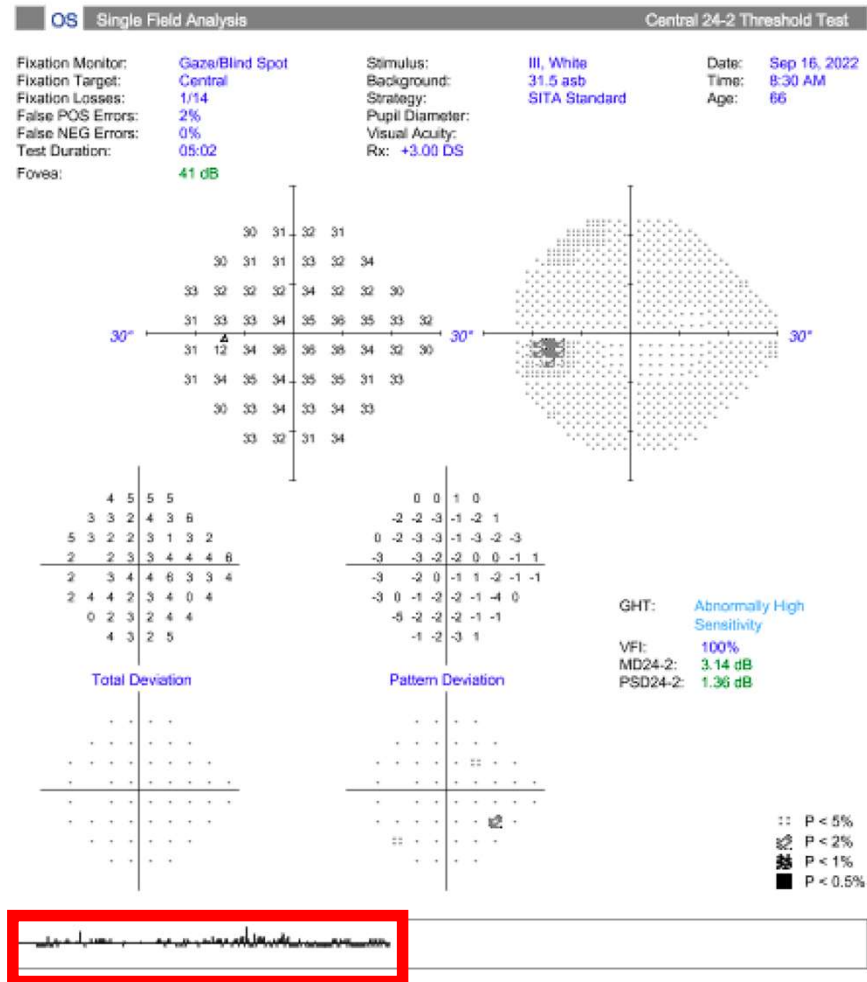
# HVF: reliability indices

- Fixation Losses
  - Responding to stimulus presented in the physiologic blind spot
  - $\leq 20\%$
- False Positives
  - Responding although stimulus has not been presented
  - $\leq 20\%$
- False Negatives
  - Lack of response to stimulus presented at a previously seen location at a lower stimulus
  - $\leq 33\%$



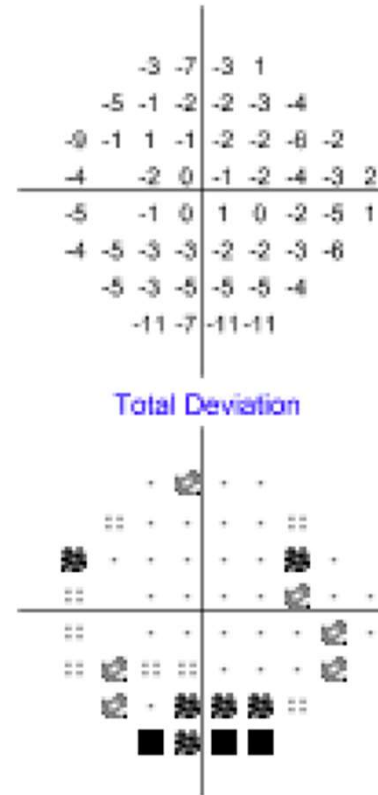
# HVF: Gaze tracking

- Upward deflection
  - eye movement
- Downward deflection
  - blink



# HVF: Total deviation

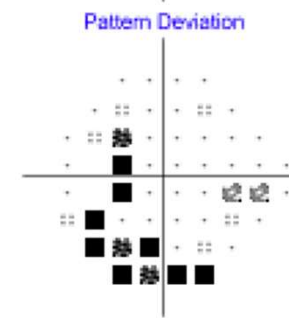
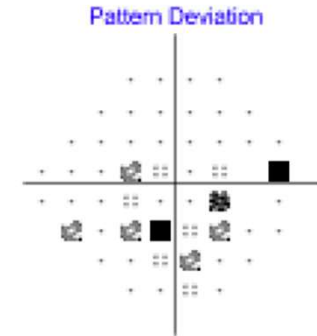
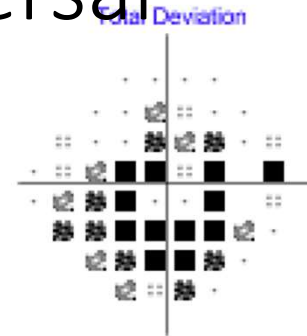
- Total deviation
  - Sensitivity (dB) at all 54 plot locations
  - Compared w/ age-matched control
  - <5% sensitivity considered abnormal
  - Probability plot color-coded from <5% to <0.5%
  - Used to calculate MD and PSD





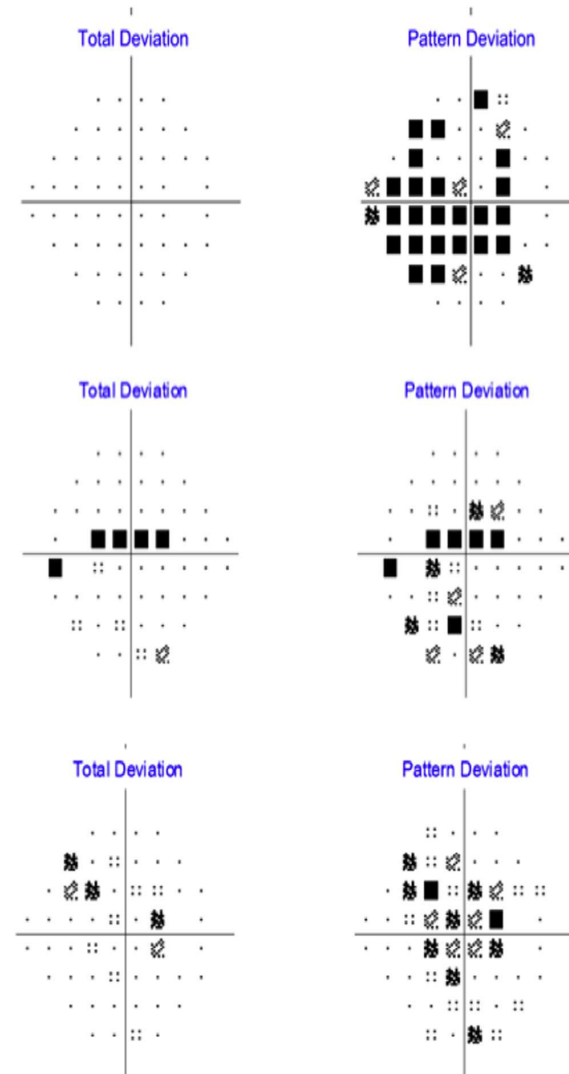
# HVF: TD, PD, pattern reversal

- Total deviation
  - Age-matched sensitivities at every location
- Pattern deviation
  - Total deviation adjusted to correct for generalized depressions or 'shifts in field sensitivities'
- Pattern reversal
  - Pattern deviation with more depressed locations than total deviation
  - The 'reverse' of what would be expected in a generalized depression from a media opacity like cataract



# HVF: Pattern reversal

- Pattern reversal
  - Most often to be considered an unreliable test
    - Trigger happy
    - Higher false positives
  - BUT, be careful. . .
  - This can represent true paracentral scotomas
  - Scotomas that may only present on pattern deviation map up to 20% of the time

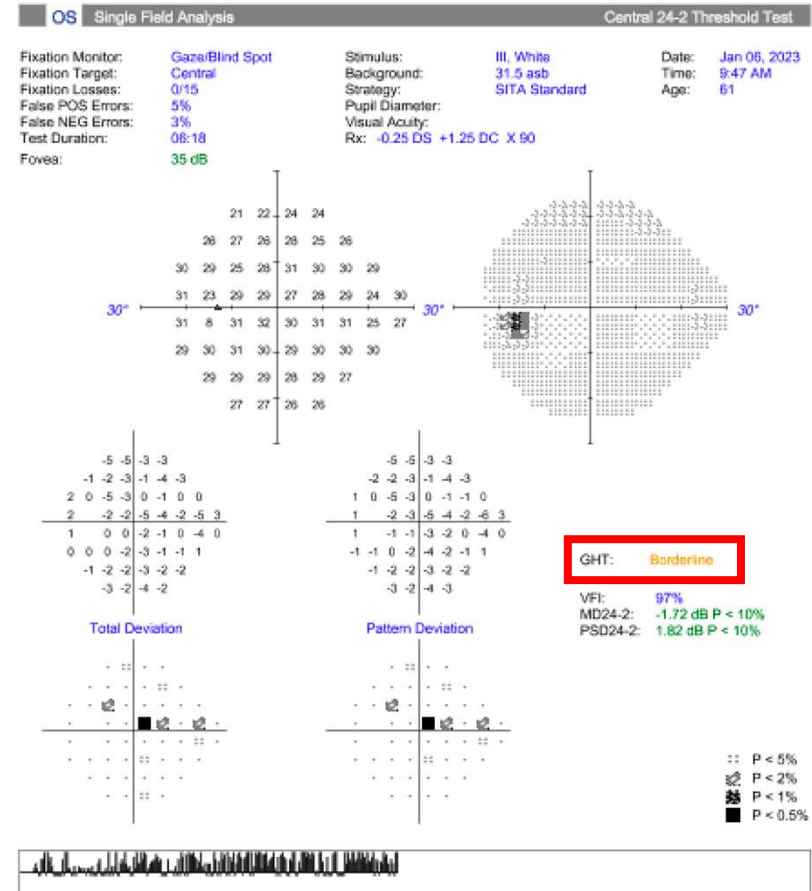


# HVF: Global indices

- Mean deviation (MD)
  - Average of all points of total deviation
    - Remember, total deviation is retinal sensitivity (dB) at all points in the VF
    - Positive value
      - Dimmer stimuli seen c/w age-matched controls
    - Negative value indicates defects
      - Brighter stimuli required c/w age-matched controls
- Pattern standard deviation (PSD)
  - Glaucoma defects are usually non uniform
  - Reflects the island or hill of vision
  - Low value (0)
    - Normal hill of vision OR
    - Severely depressed visual field
  - Higher value
    - Irregular shape of hill
    - Severely depressed focal points
    - Helpful in early glaucoma
    - With greater depression of hill of vision (e.g. advanced glaucoma), PSD becomes less useful

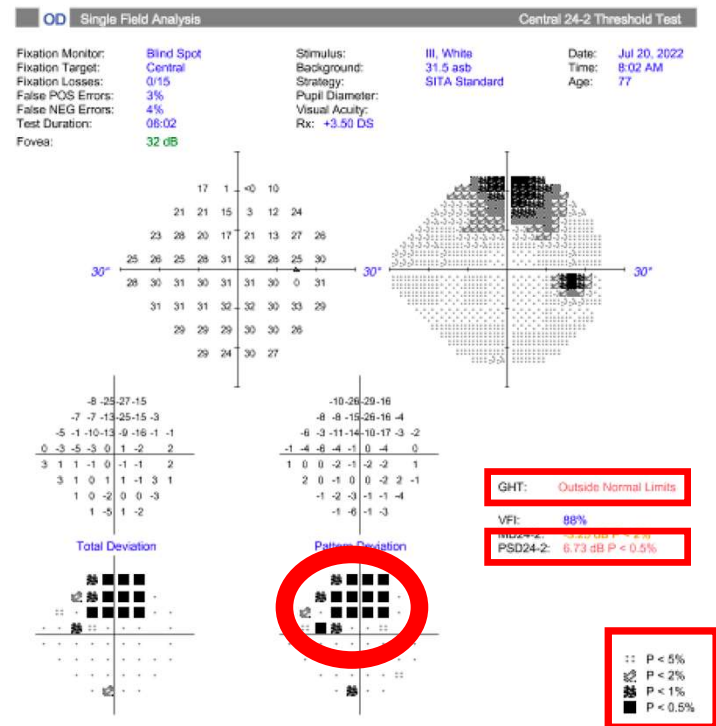
# HVF: Global indices

- Glaucoma Hemifield Test (GHT)
  - Compares retinal sensitivity across horizontal meridian
  - Assumes symmetry of upper and lower hemifields
  - Abnormal GHT is early indicator of glaucoma



# HVF: Glaucomatous findings

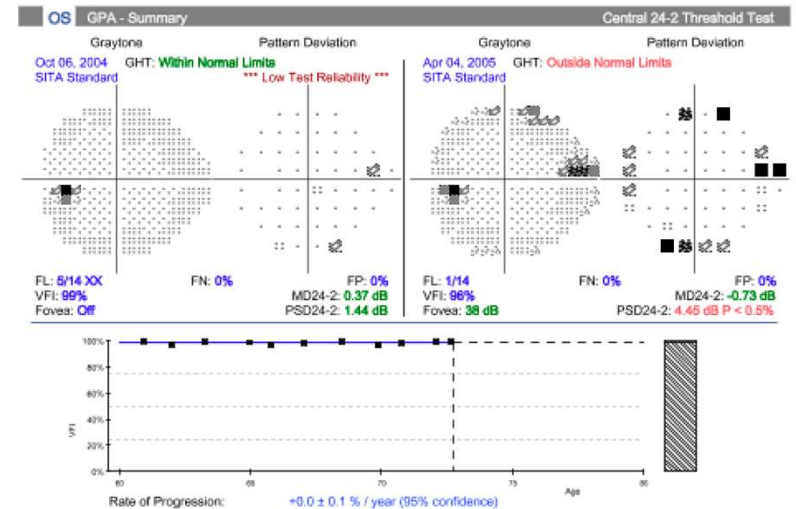
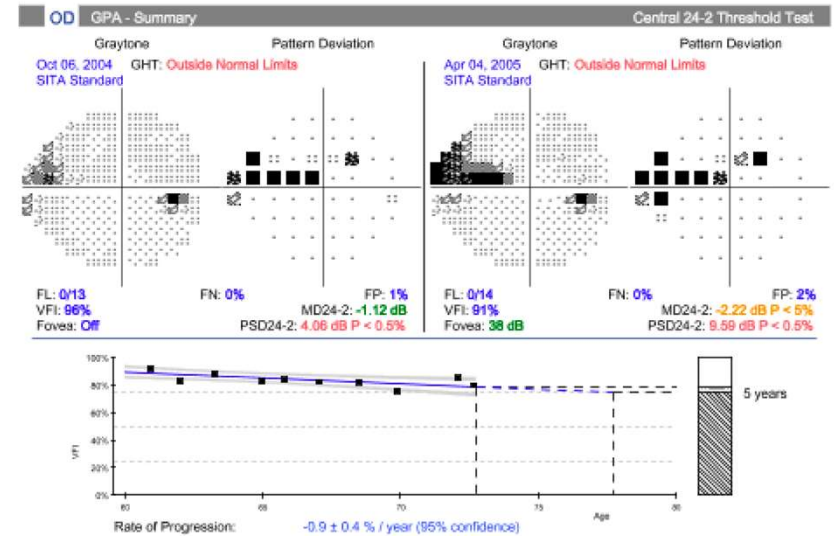
- Anderson criteria:
  - Abnormal GHT
  - Abnormal PSD
    - p-value < 5%
  - 3 contiguous, non-edge defects
    - One w/ p-value < 5%
    - One w/ p-value < 0.5%





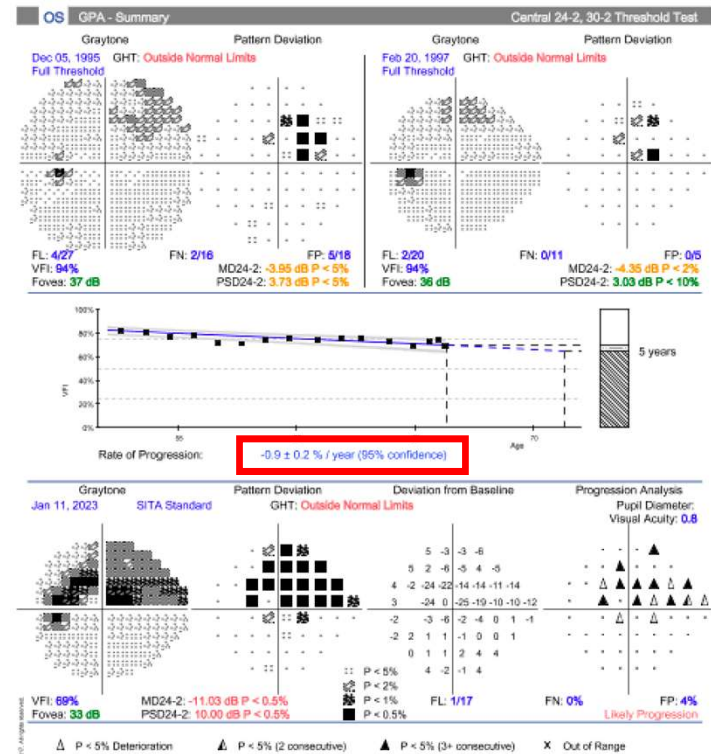
# HVF: Trend-based analysis

- Based on Visual Field Index
- VFI
  - Direct, linear correlation w/ MD
    - Therefore, affected by anything that reduces retinal sensitivity
  - Central locations weighted more heavily than peripheral points
  - Represents the entire VF as a numerical percentage
- 100%
  - Normal, full, unaffected VF
- 0%
  - Perimetrically blind
- Slope is extrapolated over time to predict future progression



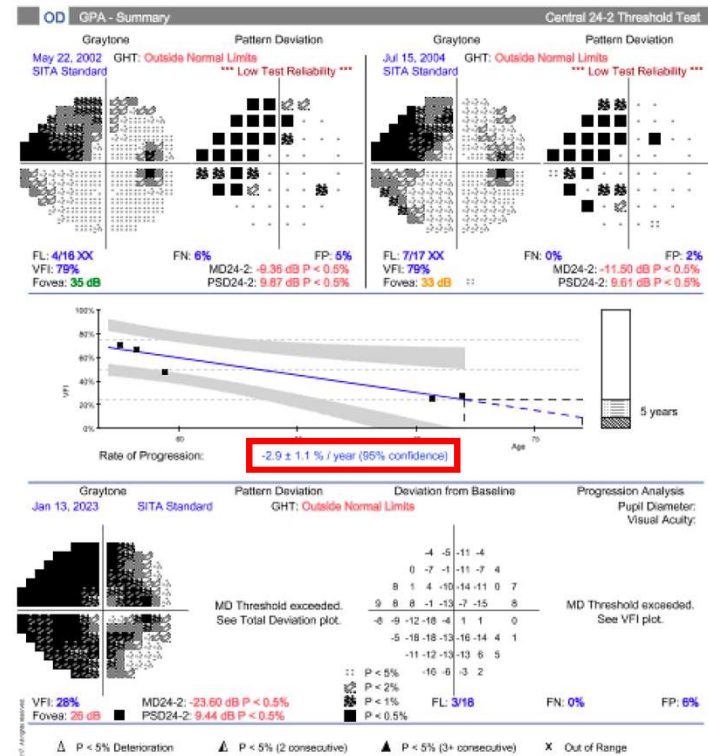
# HVF: Trend-based analysis

- VFI slope
  - Expressed as %/year
  - $< -0.5\%/year$ 
    - Slow progressor
  - $< -1.0\%/year$ 
    - Fast progressor
  
- Slow progressor



# HVF: Trend-based analysis

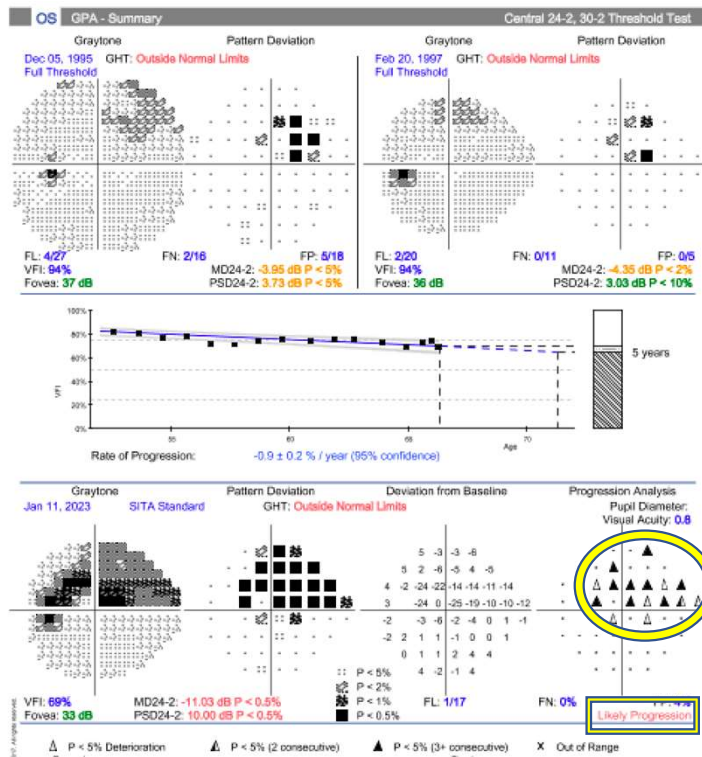
- Fast progressor



# HVF: Event-based analysis

- Based on EMGT method of detecting VF progression
- Each point on pattern deviation probability plots evaluated
- Most recent test(s) compared with 2, reliable baseline tests
- Open triangle:
  - 1 location significantly deteriorates c/w baseline
  - Significant if location degrades more so than is expected in < 5% of stable glaucoma patients
- Half-filled triangle:
  - 2 consecutive tests with significant deterioration at same location
- Closed triangle:
  - 3 consecutive tests with significant deterioration at same location
- Possible progression:
  - If  $\geq 3$  same points show significant degradation ( $p < 0.05$ ) on 2 consecutive tests
  - Points do not have to be clustered together
- Likely progression:
  - If  $\geq 3$  same points show significant degradation ( $p < 0.05$ ) on 3 consecutive tests

# HVF: Event-based analysis



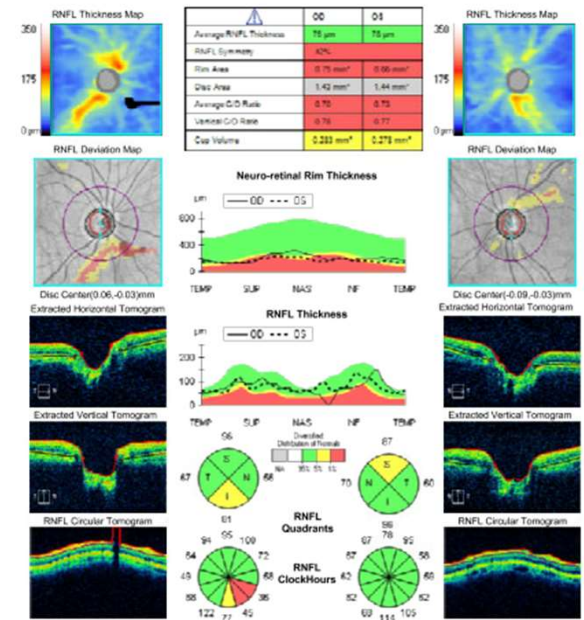
# Is my patient progressing????

Every TRRR participant's favorite game

# Case 1:

- 78 yo black female
- h/o POAG
- Tmax 30s
- IOP 14/15 on latanoprost and timolol OU
- OCT reveals . . .

## ONH and RNFL OU Analysis: Optic Disc Cube 200x200 OD ● OS ○



## Guided Progression Analysis: (GPA™) OD ● OS ○

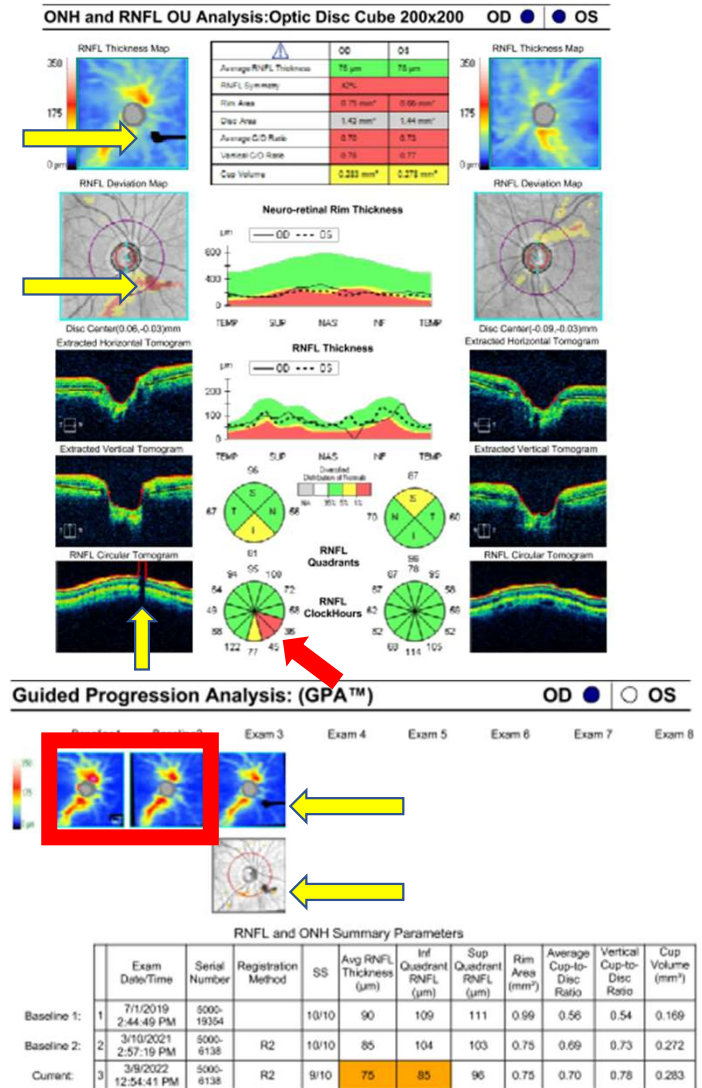


RNFL and ONH Summary Parameters

	Exam Date/Time	Serial Number	Registration Method	SS	Avg RNFL Thickness ( $\mu$ m)	Inf Quadrant RNFL ( $\mu$ m)	Sup Quadrant RNFL ( $\mu$ m)	Rim Area (mm <sup>2</sup> )	Average Cup-to-Disc Ratio	Vertical Cup-to-Disc Ratio	Cup Volume (mm <sup>3</sup> )
Baseline 1:	7/1/2019 2:44:49 PM	5000-19304		10/10	90	109	111	0.99	0.56	0.54	0.169
Baseline 2:	3/10/2021 2:57:19 PM	5000-6138	R2	10/10	85	104	103	0.75	0.69	0.73	0.272
Current:	3/9/2022 12:54:41 PM	5000-6138	R2	9/10	75	85	96	0.75	0.70	0.78	0.283

# Case 1, cont:

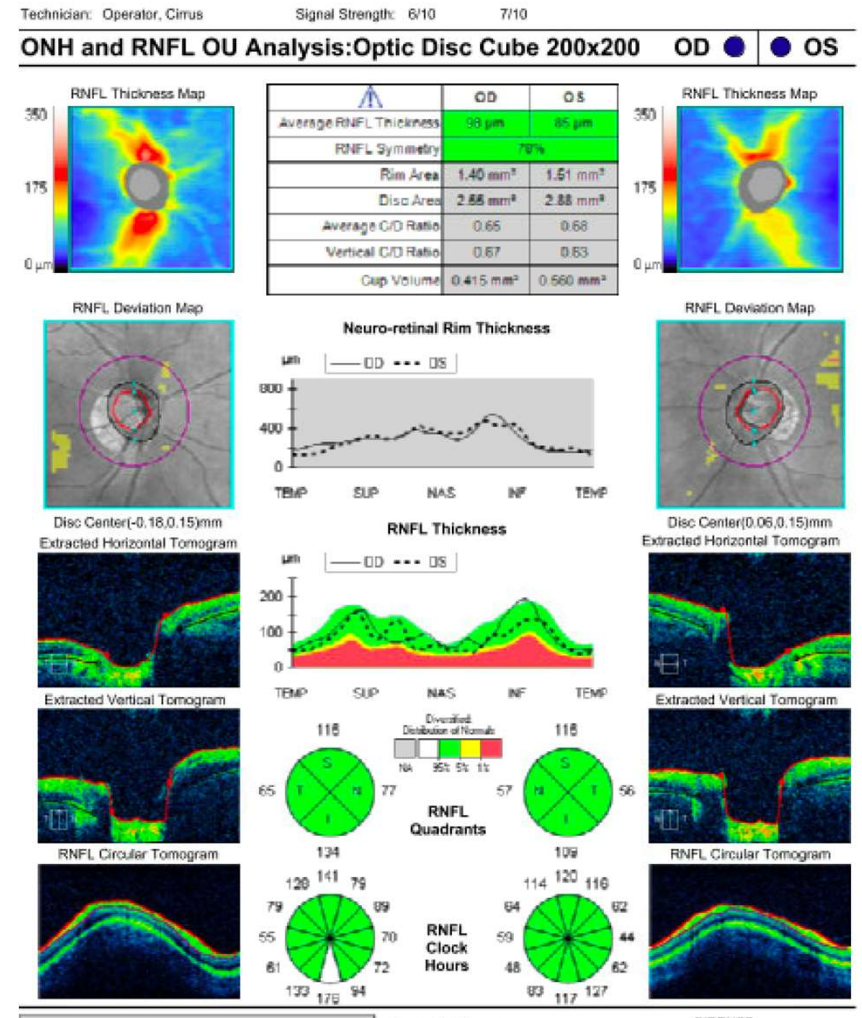
- Is my patient progressing?
  - NO
- PVD artifact OD
  - Blocking artifact
  - Atypical nasal 'loss' of RNFL
  - Note the absence of vitreous artifact in the baseline scans





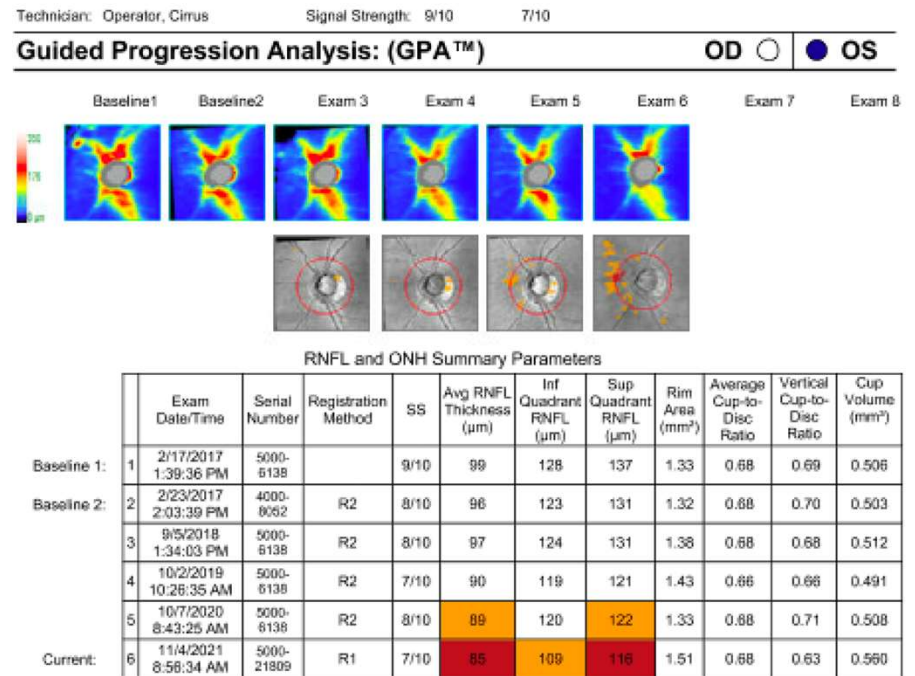
# Case 2:

- 59 yo WF
- Followed as glaucoma suspect
- No fam hx
- Nml Ks
- Nml IOP
- Full fields
- OCT reveals . . .



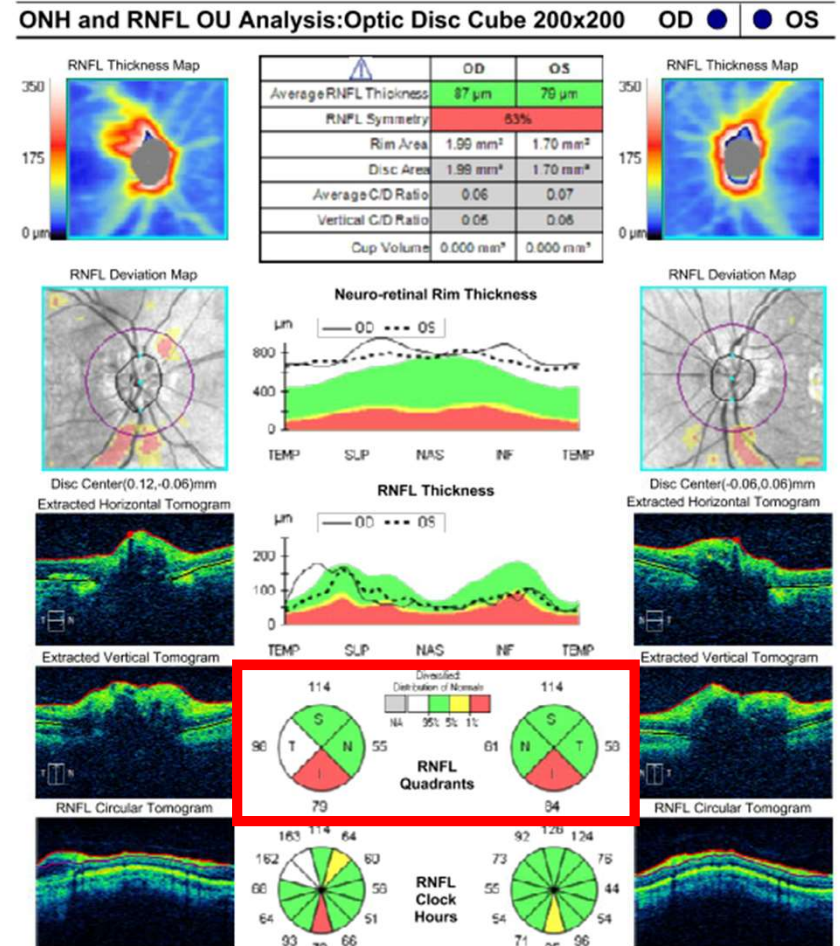
# Case 2, cont:

- Is my patient progressing?
- Yes
- Progression in the green
  - Green disease
- Pt diagnosed w/ NTG
- Began treatment w/ PGa



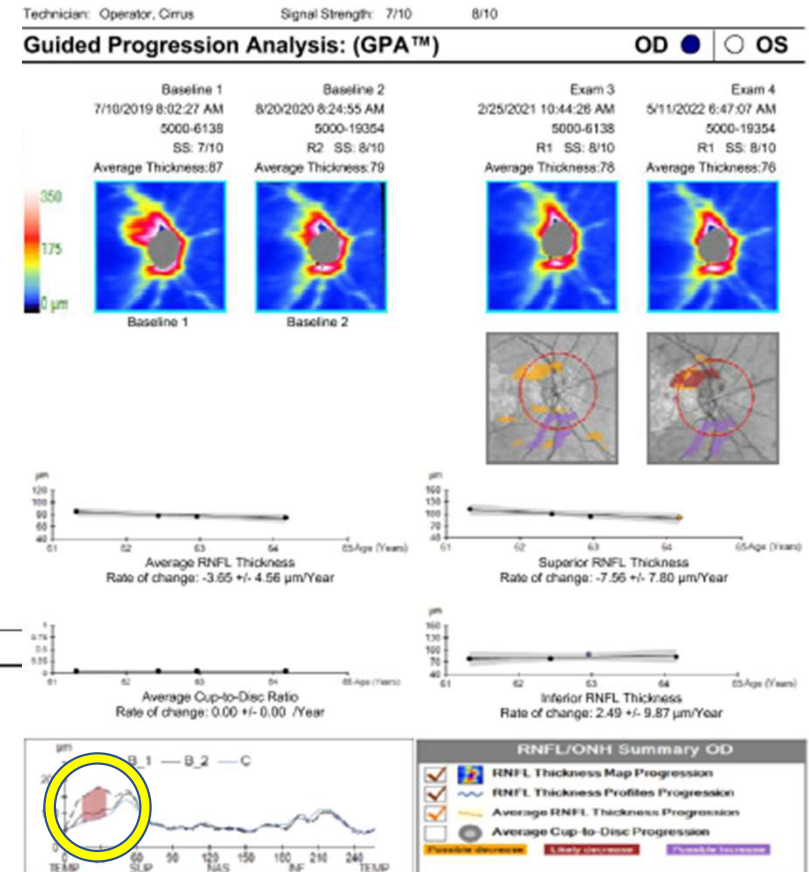
# Case 3:

- 64 yo WF
- Referred for glaucoma eval
- h/o AMD
- IOP 24/24
- OCT reveals . . .
- Is my patient progressing? (from glaucoma suspect to fulminant glaucoma)
  - NO,
    - OHT + ON drusen
    - Red disease

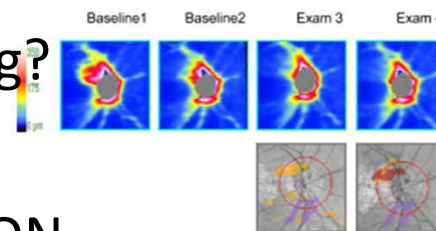


# Case 3, cont:

- OHT treated with aqueous suppressants, IOP lowered, followed
- Repeat OCT reveals . . .
- Is my patient progressing?
- Do we have co-existing ON drusen AND glaucoma?



### Guided Progression Analysis: (GPA™)



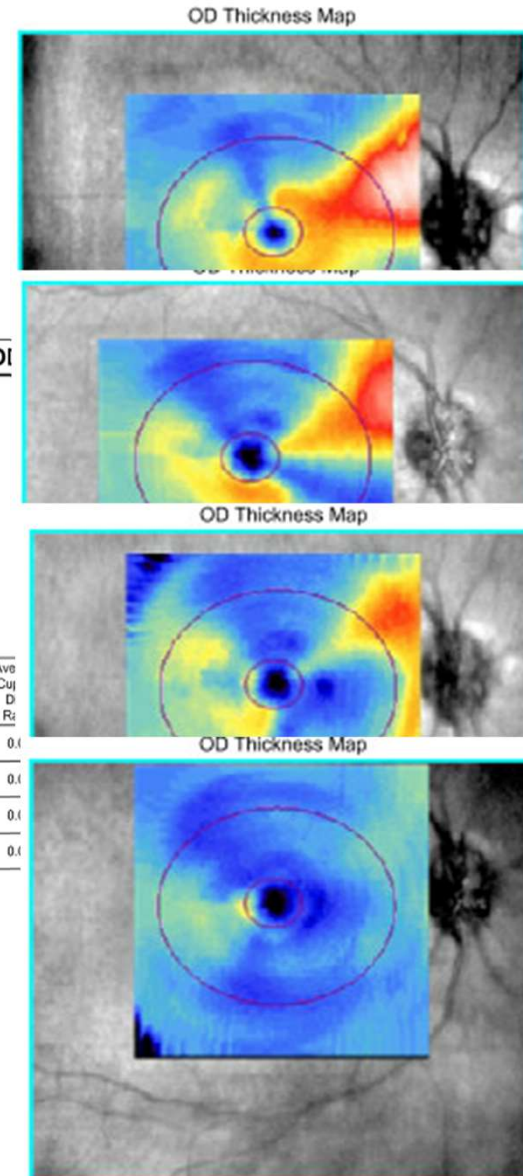
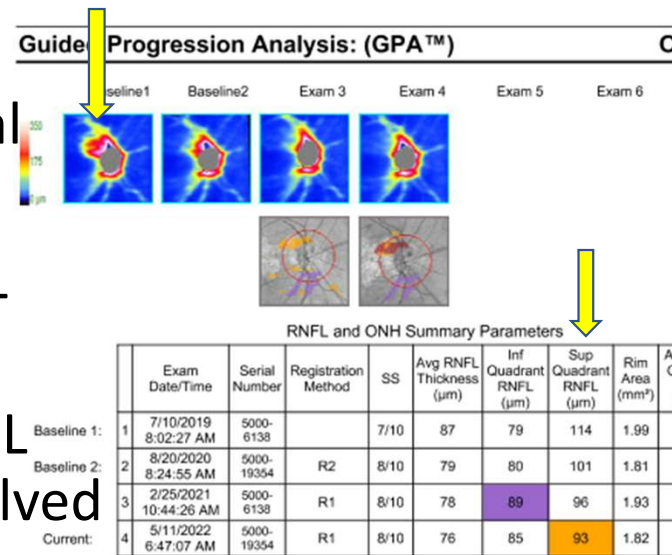
RNFL and ONH Summary Parameters

	Exam Date/Time	Serial Number	Registration Method	SS	Avg RNFL Thickness (µm)	Inf Quadrant RNFL (µm)	Sup Quadrant RNFL (µm)	Rim Area (mm²)	Average Cup-to-Disc Ratio	Vertical Cup-to-Disc Ratio	Cup Volume (mm³)	
Baseline 1:	1	7/10/2019 8:02:27 AM	5000-6138		7/10	87	79	114	1.99	0.06	0.05	0.000
Baseline 2:	2	8/20/2020 8:24:55 AM	5000-19354	R2	8/10	79	80	101	1.81	0.07	0.06	0.000
	3	2/25/2021 10:44:26 AM	5000-6138	R1	8/10	78	89	96	1.93	0.06	0.05	0.000
Current:	4	5/11/2022 6:47:07 AM	5000-19354	R1	8/10	76	85	93	1.82	0.07	0.06	0.000



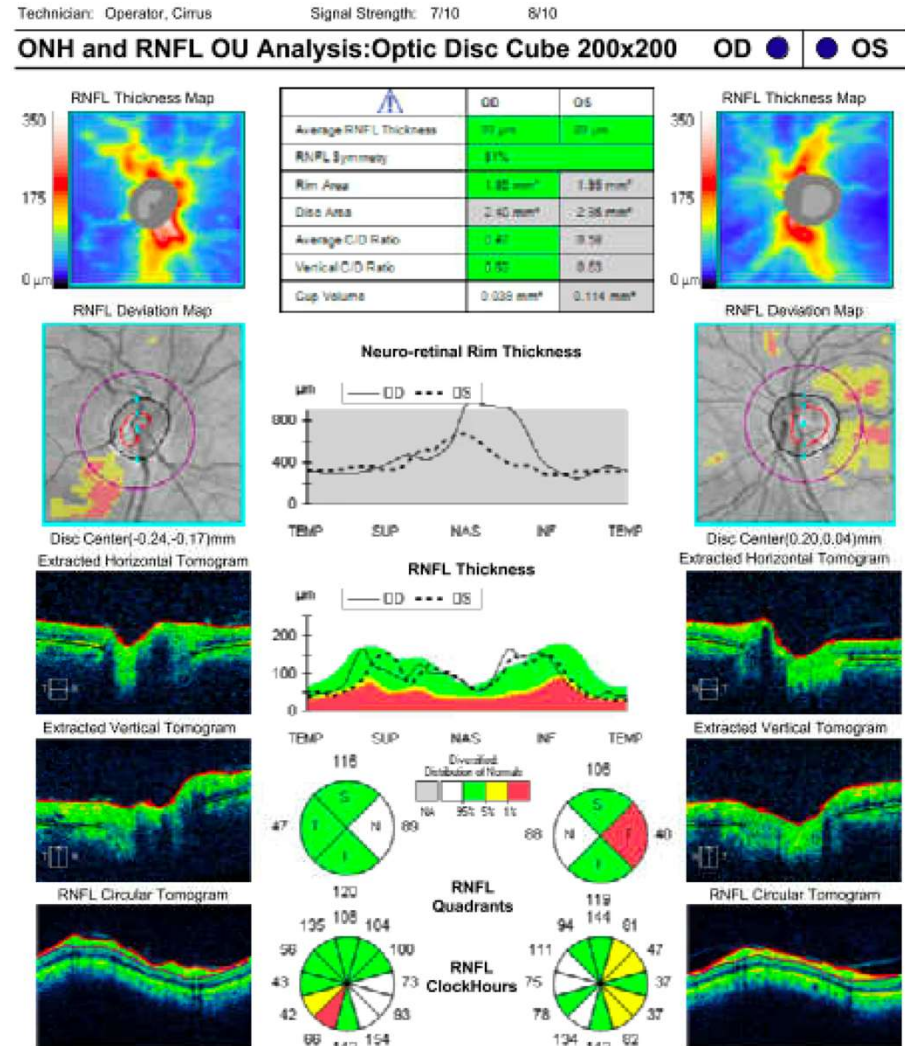
# Case 3, cont:

- NO
- Peripapillary CNVM on initial scan resolved over time
- Artificially augmenting RNFL measurement (2019)
- Artificially exaggerating RNFL “loss” once completely resolved (2022)

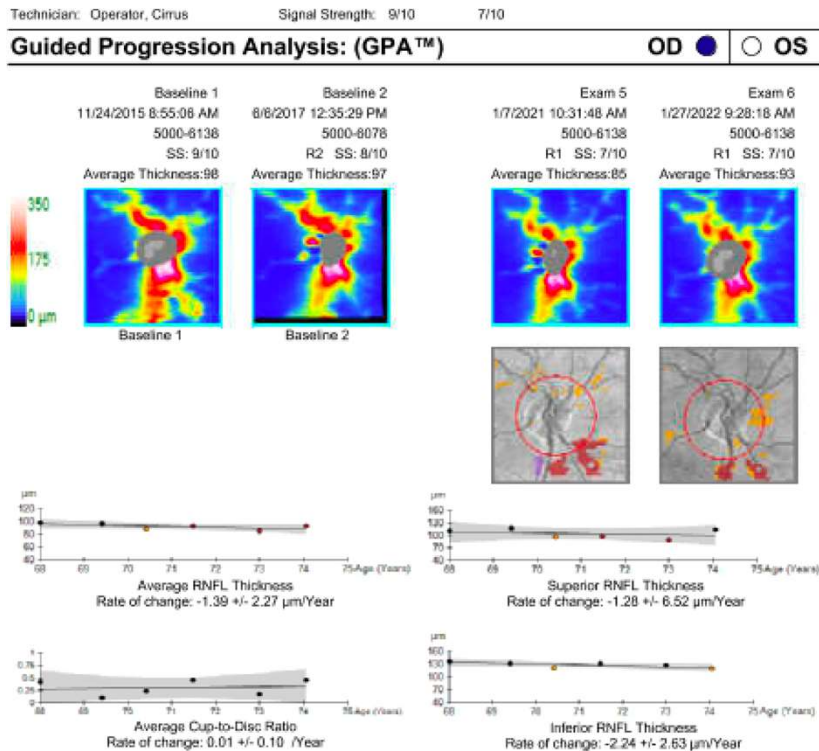


# Case 4:

- 75 yo F w/ h/o OHT
- IOP high teens, low 20s
- s/p SLT OU
- On topical CAI OU BID
- Presents for f/u w/ OCT



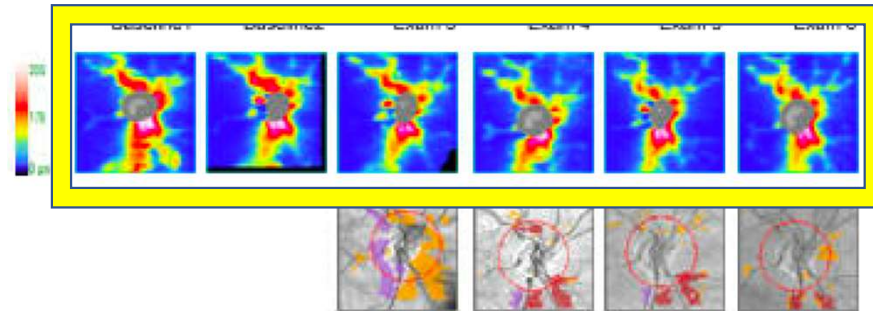
# Case 4, cont:



- GPA reveals . . .
- Is my patient progressing?
- Is this POAG instead of OHT?
  - NO
  - Improper identification of ON head

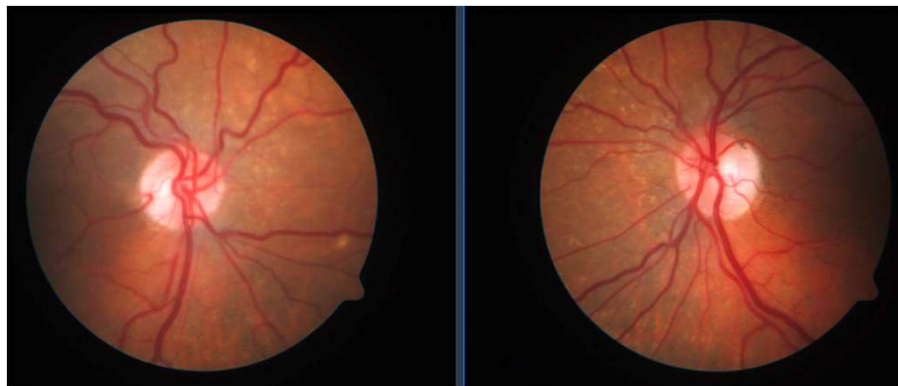
# Case 4, cont:

- Failure of ON head recognition



Exam 7

Exam 8



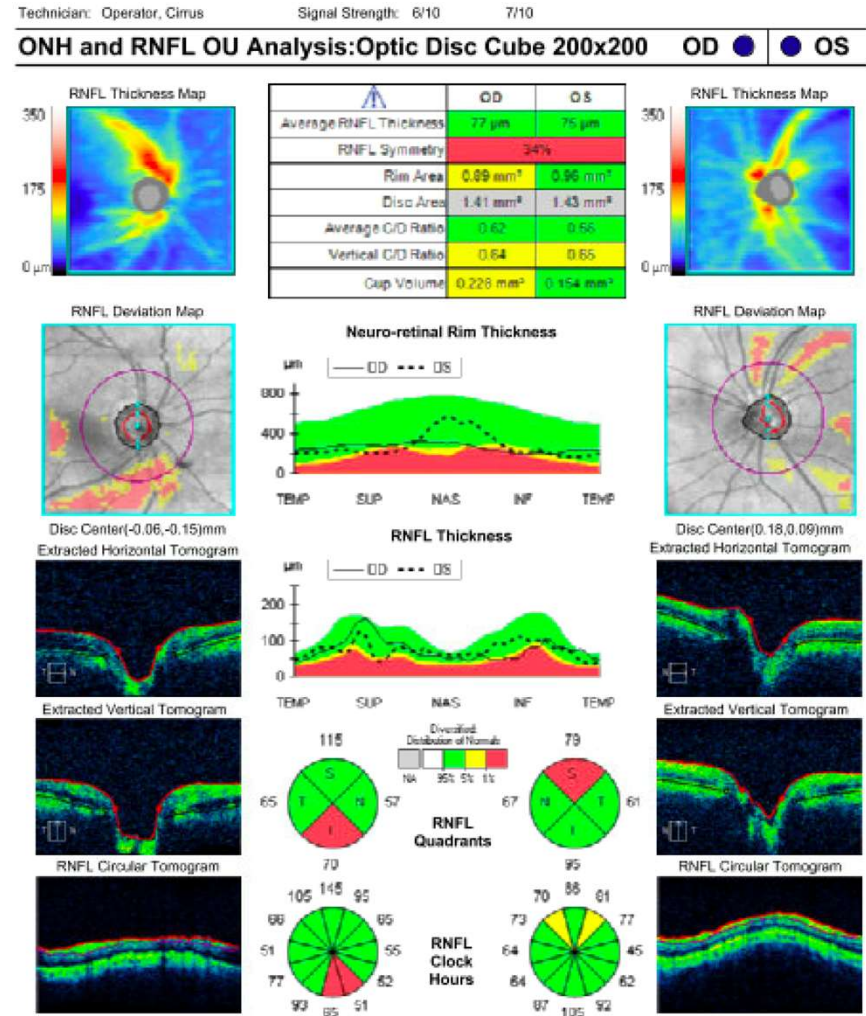
RNFL and ONH Summary Parameters

	Exam Date/Time	Serial Number	Registration Method	SS	Avg RNFL Thickness (µm)	Inf Quadrant RNFL (µm)	Sup Quadrant RNFL (µm)	Rim Area (mm²)	Average Cup-to-Disc Ratio	Vertical Cup-to-Disc Ratio	Cup Volume (mm³)
Baseline 1:	1 11/24/2015 8:55:06 AM	5000-6138		9/10	98	138	111	1.79	0.44	0.50	0.030
Baseline 2:	2 6/6/2017 12:35:29 PM	5000-6078	R2	8/10	97	133	118	1.41	0.11	0.10	0.008
	3 6/13/2018 12:43:58 PM	5000-6138	R1	7/10	89	123	97	1.39	0.24	0.26	0.009
	4 7/2/2019 3:11:27 PM	5000-6078	R1	8/10	84	133	99	1.79	0.46	0.53	0.044
	5 1/7/2021 10:31:48 AM	5000-6138	R1	7/10	85	127	91	1.31	0.18	0.20	0.008
Current:	6 1/27/2022 9:28:18 AM	5000-6138	R1	7/10	83	120	116	1.85	0.47	0.53	0.039



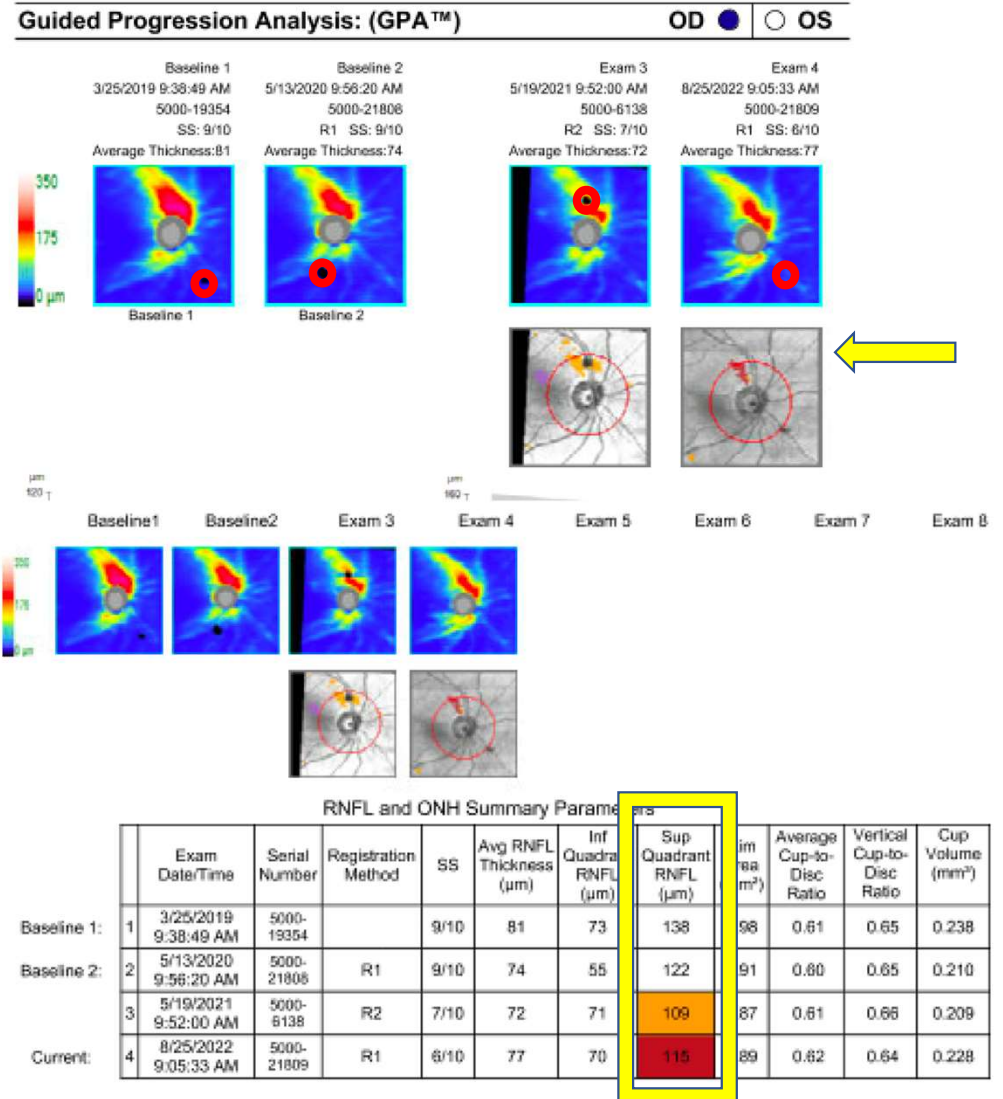
# Case 5:

- 75 yo WM
- NTG, mild OD, moderate OS
- Tmax 21
- IOP at target (mid teens) on PGa
- OCT reveals. . .



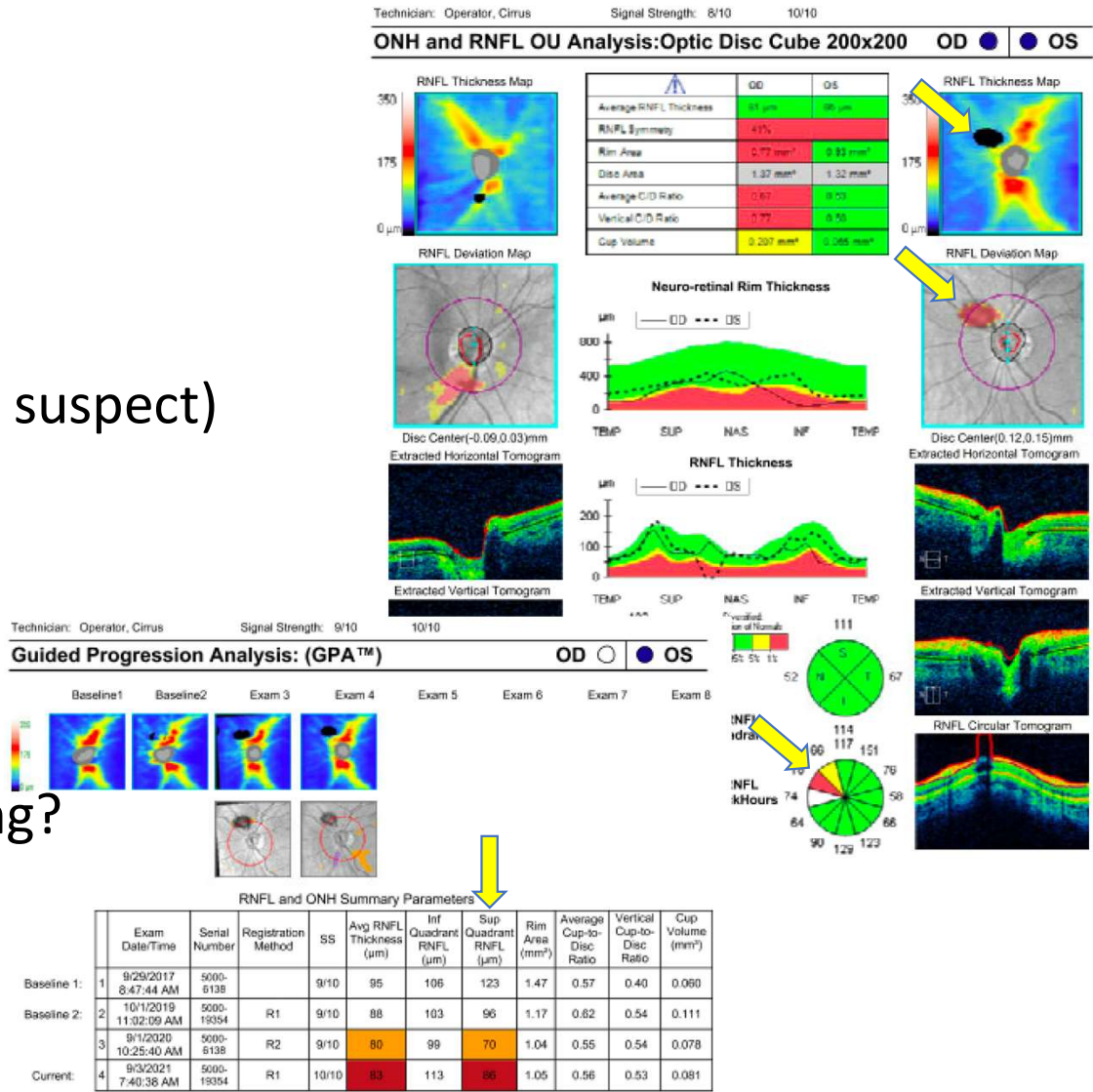
# Case 5, cont:

- GPA:
- Is my patient progressing?
  - Maybe
- Roaming vitreous artifact
- Once PVD moves from obscuring superior quadrant, RNFL rebounds
  - But not fully to baseline
  - Does superior saccade artifact play a role?
  - Watch closely

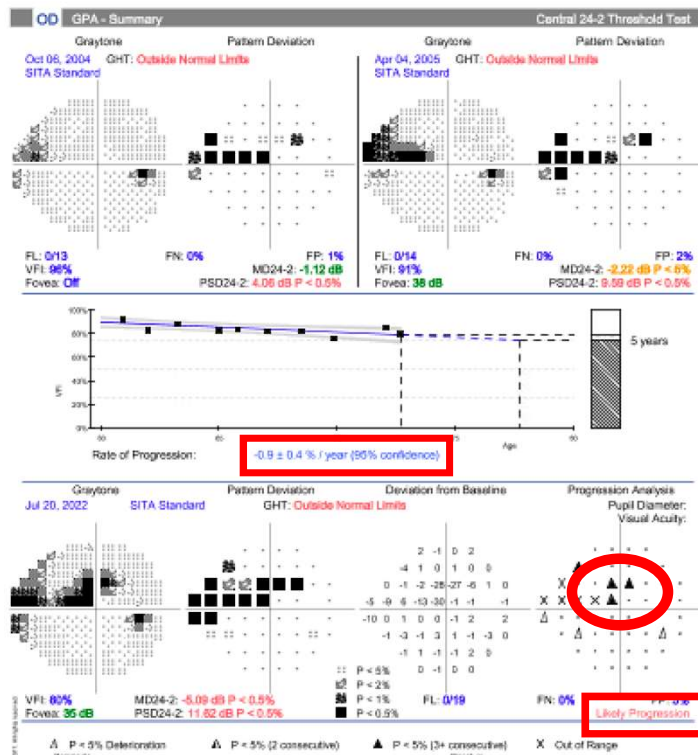


# Case 6:

- 37 yo WF
- NTG, severe OD, mild (v suspect) OS
- Target IOP: low teens
- IOP controlled on PGa
- OCT reveals. . .
- Is my patient progressing?
- No,
  - PVD artifact



# Case 6, cont:

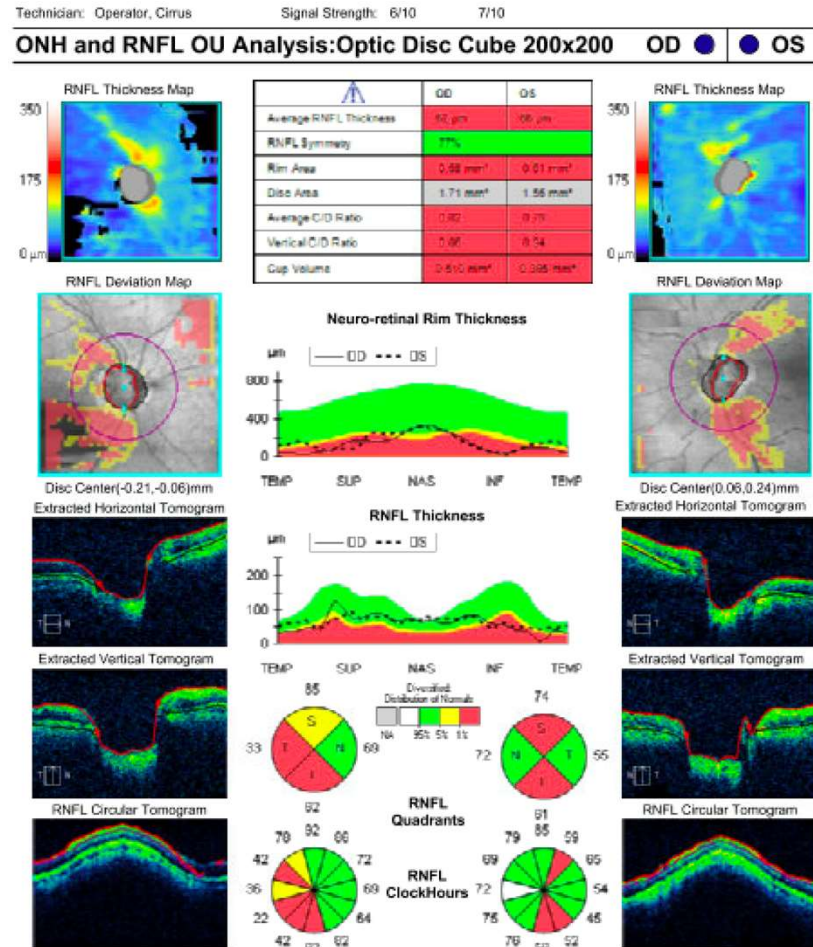


- However, HVF reveals . . .
- ‘Slow progression’ on VFI
- ‘Likely progression’ on GPA
- **IMPORTANTLY**
  - Event-based analysis revealing extension of scotoma into fixation
- So, is my patient progressing?
  - Structurally, no (OCT GPA was stable OD as well)
  - Functionally, yes



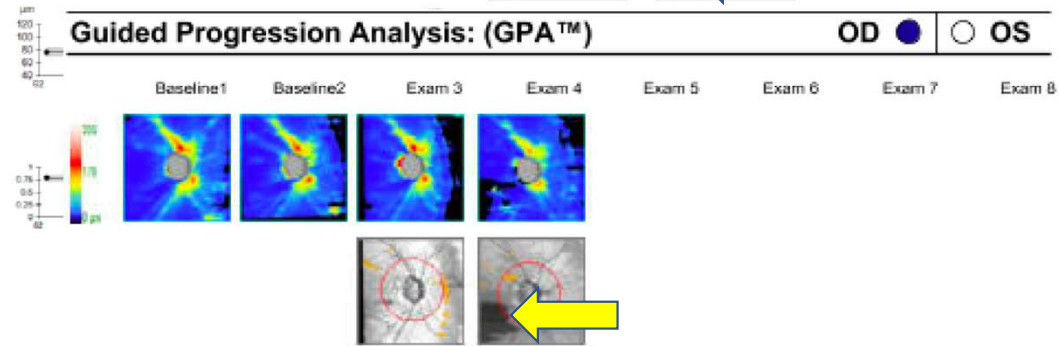
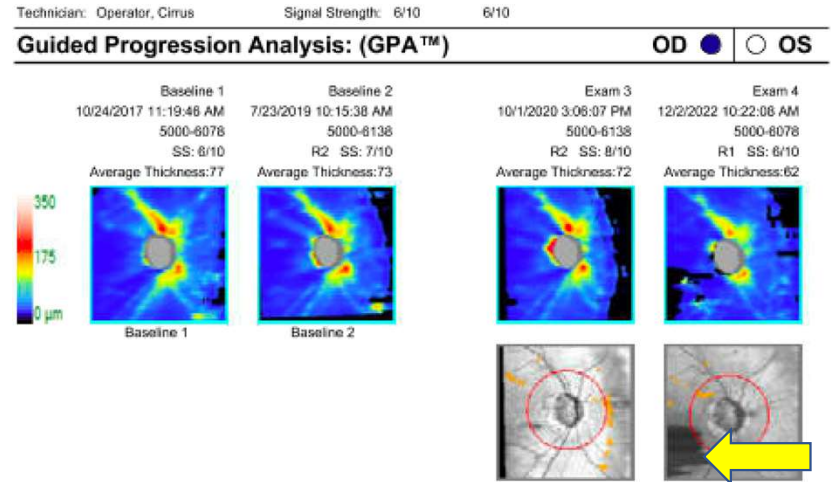
# Case 7:

- 67 yo, phakic, WM
- h/o severe NTG
- Tmax 17, tentative target 11-12
- IOP at target on PGa and fixed-combination timolol-brimonidine
- OCT reveals . . .



# Case 7, cont:

- OCT GPA reveals. . .
- Is my patient progressing?
- No
- Blocking artifact

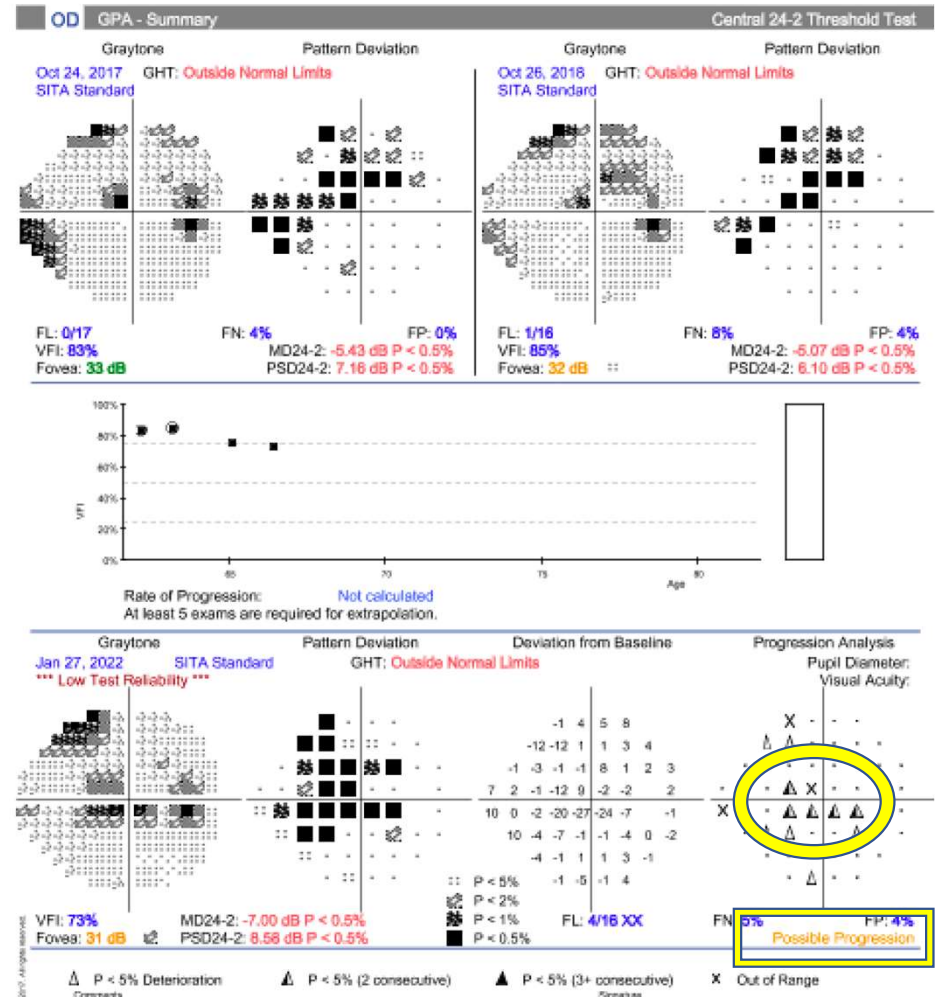


RNFL and ONH Summary Parameters

	Exam	Exam Date/Time	Serial Number	Registration Method	SS	Avg RNFL Thickness (μm)	Inf Quadrant RNFL (μm)	Sup Quadrant RNFL (μm)	Rim Area (mm <sup>2</sup> )	Average Cup-to-Disc Ratio	Vertical Cup-to-Disc Ratio	Cup Volume (mm <sup>3</sup> )
Baseline 1:	1	10/24/2017 11:19:46 AM	5000-6078		6/10	77	83	96	0.66	0.80	0.88	0.527
Baseline 2:	2	7/23/2019 10:15:38 AM	5000-6138	R2	7/10	73	78	89	0.67	0.77	0.86	0.434
	3	10/1/2020 3:08:07 PM	5000-6138	R2	8/10	72	82	90	0.58	0.82	0.88	0.504
Current:	4	12/2/2022 10:22:08 AM	5000-6078	R1	6/10	62	62	85	0.58	0.82	0.86	0.510

# Case 7, cont:

- BUT
- HVF reveals. . .
- Is my patient progressing?
- Maybe
- But patient has symptomatic cataract which we believe is affecting VF performance,
- So watch closely



# Conclusions

- Know your instruments
  - How they work
  - What they are measuring
  - How the measurements can be skewed
- Know your diseases
  - Structure/function correlation
  - How they progress
  - What can mimic them
- Know your patients
  - What do these tests mean when applied individually



# REFERENCES

- Budenz DL. Atlas of Optical Coherence Tomography for Glaucoma. Cham, Switzerland. Springer Nature Switzerland AG. 04 Jul 2020.
- Aumann, S., Donner, S., Fischer, J., Müller, F. (2019). Optical Coherence Tomography (OCT): Principle and Technical Realization. In: Bille, J. (eds) High Resolution Imaging in Microscopy and Ophthalmology. Springer, Cham. [https://doi.org/10.1007/978-3-030-16638-0\\_3](https://doi.org/10.1007/978-3-030-16638-0_3)
- Mwanza JC, Oakley JD, Budenz DL, Anderson DR; Cirrus Optical Coherence Tomography Normative Database Study Group. Ability of cirrus HD-OCT optic nerve head parameters to discriminate normal from glaucomatous eyes. *Ophthalmology*. 2011 Feb;118(2):241-8.e1. doi: 10.1016/j.ophtha.2010.06.036. Epub 2010 Oct 28. PMID: 20920824; PMCID: PMC3017237.
- Ali Riza Cenk Celebi, G. Ertugrul Mirza; Age-Related Change in Retinal Nerve Fiber Layer Thickness Measured With Spectral Domain Optical Coherence Tomography. *Invest. Ophthalmol. Vis. Sci.* 2013;54(13):8095-8103. doi: <https://doi.org/10.1167/iovs.13-12634>.
- Zhang X, Dastiridou A, Francis BA, Tan O, Varma R, Greenfield DS, Schuman JS, Huang D; Advanced Imaging for Glaucoma Study Group. Comparison of Glaucoma Progression Detection by Optical Coherence Tomography and Visual Field. *Am J Ophthalmol*. 2017 Dec;184:63-74. doi: 10.1016/j.ajo.2017.09.020. Epub 2017 Sep 28. PMID: 28964806; PMCID: PMC5894829.
- Shin JW, Sung KR, Lee GC, Durbin MK, Cheng D. Ganglion Cell-Inner Plexiform Layer Change Detected by Optical Coherence Tomography Indicates Progression in Advanced Glaucoma. *Ophthalmology*. 2017 Oct;124(10):1466-1474. doi: 10.1016/j.ophtha.2017.04.023. Epub 2017 May 23. PMID: 28549518.
- Savini G, Barboni P, Parisi V, Carbonelli M. The influence of axial length on retinal nerve fibre layer thickness and optic-disc size measurements by spectral-domain OCT. *Br J Ophthalmol*. 2012 Jan;96(1):57-61. doi: 10.1136/bjo.2010.196782. Epub 2011 Feb 24. PMID: 21349942.
- Aref AA, Sayyad FE, Mwanza JC, Feuer WJ, Budenz DL. Diagnostic specificities of retinal nerve fiber layer, optic nerve head, and macular ganglion cell-inner plexiform layer measurements in myopic eyes. *J Glaucoma*. 2014 Oct-Nov;23(8):487-93. doi: 10.1097/IJG.0b013e31827b155b. PMID: 23221911; PMCID: PMC3986352.
- Scuderi G, Fragiotta S, Scuderi L, Iodice CM, Perdicchi A. Ganglion Cell Complex Analysis in Glaucoma Patients: What Can It Tell Us? *Eye Brain*. 2020 Jan 31;12:33-44. doi: 10.2147/EB.S226319. PMID: 32099501; PMCID: PMC6999543.
- Racette, Lyne, et al. Visual Field Digest, A Guide to Perimetry and the Octopus Perimeter, 7<sup>th</sup> Ed. Koniz, Switzerland. Haag-Streit AG. 2018.
- Ruia S, Tripathy K. Humphrey Visual Field. [Updated 2022 Aug 22]. In: StatPearls [Internet]. Treasure Island (FL): StatPearls Publishing; 2022 Jan-. Available from: <https://www.ncbi.nlm.nih.gov/books/NBK585112/>
- Aimee C. Chang, Andrew S. Camp, Vincent M. Patella, Robert N. Weinreb, Association of Visual Field Pattern Reversal with Paracentral Visual Field Loss, *Ophthalmology Glaucoma*, Volume 5, Issue 3, 2022, Pages 353-358, ISSN 2589-4196, <https://doi.org/10.1016/j.ogla.2021.10.009>.
- Bengtsson B, Heijl A. A visual field index for calculation of glaucoma rate of progression. *Am J Ophthalmol*. 2008 Feb;145(2):343-53. doi: 10.1016/j.ajo.2007.09.038. PMID: 18078852.
- Iutaka NA, Grochowski RA, Kasahara N. Correlation between Visual Field Index and Other Functional and Structural Measures in Glaucoma Patients and Suspects. *J Ophthalmic Vis Res*. 2017 Jan-Mar;12(1):53-57. doi: 10.4103/jovr.jovr\_98\_16. PMID: 28299007; PMCID: PMC5340064.
- Zhang X, Parrish RK 2nd, Greenfield DS, Francis BA, Varma R, Schuman JS, Tan O, Huang D; Advanced Imaging for Glaucoma Study Group. Predictive Factors for the Rate of Visual Field Progression in the Advanced Imaging for Glaucoma Study. *Am J Ophthalmol*. 2019 Jun;202:62-71. doi: 10.1016/j.ajo.2019.02.015. Epub 2019 Feb 20. PMID: 30794787; PMCID: PMC6548618.
- Arnalich-Montiel F, Casas-Llera P, Muñoz-Negrete FJ, Rebolleda G. Performance of glaucoma progression analysis software in a glaucoma population. *Graefes Arch Clin Exp Ophthalmol*. 2009 Mar;247(3):391-7. doi: 10.1007/s00417-008-0986-1. Epub 2008 Nov 4. PMID: 18982343.