

# Memphis, TN



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## Ground Rules...

- References/sources available if you want them...
- 139 articles used for this presentation...
- I'm not perfect...
- Please email me with questions:
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# OCT & Retrograde Degeneration

Retrograde vs. Anterograde degeneration?				
<u>Retrograde:</u> Cell bodies → RGC's in retina Synapse → LGN				
LGN $\rightarrow$ RGC's = retrograde				
Ex: Compression, TON, MS, etc.				
<u>Anterograde:</u> Cell bodies: RGC's in retina Synapse: LGN	$\longrightarrow$			
RGC's → LGN = anterograde				
Ex: Glaucoma				

<u>(</u>	<u>Case #1:</u>		
		- Courtesy of Rena Lepine, OD	

## Case history...

- 19 yo AAF
- · CC: "blurry vision"
- HPI: OD only, constant, onset 2 months (?), (+)glare noted, (-)trauma, (-) relief with Visine drops, (+)frontal HA's started about same time, (-)relief with ibuprofen
- · POH: glasses only
- · PMH: unremarkable
- · Meds: Ibuprofen PRN
- Allergies: nickel
- · SH: denies all, (-)pregnant
- BP: 128/78

- BCVA: 20/25+ OD, 20/20 OS
- Pupils: PERRL, (+)APD OD
- · EOM's: Full OU
- $\cdot$  CVF: sup temp defect OD, FTFC OS
- · Cover Test: ortho @ D & N
- · Color: very reduced OD, normal OS
- SLE: unremarkable OU
- DFE: see pics
- · Tonometry: 16 mmHg OU with Goldman





































- 55 yo female
- CC: "glaucoma suspect repeat testing"
- HPI: OD>OS, onset 5-6 years ago, no Tx at this time, (-)FHx glaucoma, vision stable OU per patient
- $\cdot$  POH: glaucoma suspect OD>OS
- $\cdot$  PMH: nasopharyngeal carcinoma at age 44, s/p chemo/radiation/resection
- · Meds: multivitamin
- ALL: NKDA

- VA: 20/20 OD, 20/20 OS
- Pupils: PERRL, (-)APD
- · CVF: FTFC OU
- · EOM: FROM OU
- SLE: normal OU
- $\cdot$  DFE: see photos of nerves
- $\cdot$  IOP: 12 OD, 15 OS via iCare





















## Final Diagnosis...

- $\cdot \ Dx$  = Collateral radiation damage to optic tract
- $\cdot$  None of the MRI scans showed carcinoma/tumor close enough to optic tract to cause compression...
- $\cdot$  Occam's Razor  $\rightarrow$  the most likely explanation is most often the correct one
- · Radiation Toxicity to Optic Tract

## HEAT CHECK!

Quick Case Topic...

### Case...

- · 46 yo AAF
- · CC: "reduced vision"
- $\cdot$  HPI: OS only, slowly getting worse, central vision>>peripheral vision, spot just off of central vision OS, (-)trauma
- · POH: unremarkable; first eye exam "in a long time"
- PMH: T2DM x 16 yrs, unknown A1c, last BG was 500 but doesn't check regularly, was in diabetic ketoacidosis 3 weeks ago, PCP recommended eye exam at last PCP visit 2 weeks ago; HTN x 16 yrs, unknown last BP
- · Meds: metformin, insulin, Onglyza, metoprolol, losartan
- Allergies: NKDA
- SH: unremarkable

- $\circ$  VAcc: 20/20 OD, 20/25+ OS
- Pupils: PERRL, (-)APD
- $\cdot$  CVF: FTFC peripherally OU, central depressions OU?
- EOMs: FROM OU
- SLE: unremarkable OU
- $\cdot \mbox{ DFE:} \ see \ photos$
- IOP: 20 OD, 22 OS via Goldman

















# **OCT & Ocular Torsion**

#### Case

- 58 yo AAM
- · CC: "double vision"
- $\cdot$  HPI: started 4 days ago, vertical/oblique diplopia, (-)trauma, diplopia goes away when covering either eye
- POH: unremarkable
- · PMH: hypercholesterolemia
- Meds: rosuvastatin
- Allergies: NKDA

#### Exam...

- · VAsc: 20/20 OD, 20/20 OS
- · Pupils: PERRL, (-)APD
- · CVF: FTFC OU
- $\cdot\,$  EOM: FROM OU, OS overshoot in attempted right gaze
- · Cover Test: 4^ LHT primary gaze
- · SLE: normal OU · DFE: normal OU
- · IOP: 16 OD, 16 OS with Goldman

Notice Right Head Tilt















• 60 yo AAM

- $\cdot$  CC: double vision since stroke
- $\cdot$  HPI: vertical diplopia x 1 month, brainstem CVA x 1 month ago
- $\cdot$  POH: mild cataracts OU, LEE x 1 year ago
- PMH: HTN, unknown BP ("it was high"); T2DM, unknown BG ranges/levels ("it was high too")
- · Meds: metformin, glipizide, atenolol, lisinopril
- Allergies: NKDA
- SH: unremarkable

- VAcc: 20/20 OD, 20/20 OS
- Pupils: PERRL, (-)APD
- CVF: FTFC OU
- EOMs: FROM OU but clear right hypertropia noted
- $\cdot$  Cover Test: 6^ RHT in primary gaze
- $\cdot$  SLE: grossly normal OU, mild cataracts OU
- DFE: unremarkable
- IOP: 12 OD, 12 OS via Goldman















- · 63 yo AAM
- $\cdot\,$  CC: "routine exam; do I need glasses?"
- HPI: adequate vision OU per patient, maybe some mild blur?, never had eye exam before
- · POH: unremarkable
- · PMH: HTN well controlled
- · Meds: lisinopril
- Allergies: NKDA
- · SH: unremarkable

- $\cdot$  VA: 20/30 OD, 20/30 OS
- Pupils: PERRL, (-)APD
- $\cdot$  EOM: FROM OU
- CVF: FTFC OU
- SLE: normal OU
- $\circ\,\mathrm{DFE:}\,$  see photos/OCTs
- $\cdot$  IOP: 18 OD, 18 OS via iCare











## How did this happen?

- Upon further questioning to patient...
- Patient: "Man, when I was II years old I saw the original Superman movie – and Superman got his strength from the sun – so I stared at the sun about 5 times for as long as I could in hopes that I dg eX r-ay vision like Superman did!...is that what caused what you're seeing??"

• Me: "Yes!"

Final Dx  $\rightarrow$  Solar Retinopathy OU

## How about this OCT scan?

• Is this Solar Retinopathy also?



















### Case:

- $\cdot$  Best corrected VA: 20/30 OD, 20/40 OS
- Pupils: PERRL, (-)APD
- CVF: FTFC OU
- $\cdot$  EOM: FROM OU
- $\cdot$  CT: or tho distance/near
- SLE: irregular astigmatism, EBMD  $\rightarrow\,$  RCE in past
- DFE: see photos
- $\cdot$  IOP: 18 OD, 18 OS











## Alport Syndrome

- Inherited/genetic X-linked collagen IV disorder
- · Pathophysiology: defective or absent collagen IV in basement membranes throughout body
- · Associated with:
- Renal failure (99%)
- $^\circ$  Hearing loss (82.5%)
- Ocular findings (44-81%)

ru K, Nakanishi K, Abe Y, et al. A review of clinical characteristics and rige J, Colville D. Ocular features aid the diagnosis of Alport syndrome. rige J, Sheth S, Leys A, et al. Ocular features in Alport syndrome: path and genetic backgrounds a ne. Nat Rev Nephrol 2009 athogenesis and clinical si Clin Exp Nephrol 2019; 23: 158-68 lin J Am Soc Nephrol 2015; 10: 703

#### Ocular findings... • 70% males vs. 15% females

- \*\*Temporal retinal thinning (81%)
- Dot-and-fleck retinopathy (13-70%)
- · Lozenge's sign/dull macular reflex (19%)
- · Lenticonus (2-50%)
- · Other ocular findings possible:
- Keratoconus, RCE, posterior polymorphous K dystrophy
- Iris atrophy
   Cataracts

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## Temporal retinal thinning

\*\*Most common ocular finding in X-linked Alport syndrome • AKA: "stair-case" thinning

- Temporal macula is ~59 μm thinner than nasal macula
- · No significant effect on vision in general
- · Predominantly affects superficial layers of retina • Temporal retinal thinning  $\rightarrow$  How? Why?
- Idiopathic/unknown at this time
  Believed to be related to underlying collagen IV dysfunction
- Abnormal vitreoretinal traction?
- Abnormal Muller cell adhesion? Presumed congenital? Acquired seems unlikely...



## Enlarged vs. Absent FAZ's on OCT-A?

 Apparently both are possible in Alport Syndrome!

st MWM

Pfau M, Wi









\* 44 yo AAF

- $\cdot$  CC: new onset diplopia  $\rightarrow$  final diagnosis was CN VI palsy  $\cdot$  Ultimately diagnosed with MS!
- · However, incidental retinal finding on DFE.....let's take a look!





















## **Chorioretinal Folds**

First documented by Nettles in 1884

- $\cdot\,$  Striae/grooves/lines in posterior pole with numerous patterns possible
- $\label{eq:seudo-sinusoidal undulations of inner choroid, Bruch's membrane and RPE = choroidal folds \\ \cdot \ If neurosensory retina involved then proper name <math>\Rightarrow$  choroiretinal folds
- · Alternating light-dark streaks usually present
- $\cdot\,$  Underlying autoimmune condition has been identified in about 28% of cases
- Unilateral: 44% to 58% cases
- Variable visual acuity but usually near normal acuity

· Patient's usually asymptomatic

· MOA: largely idiopathic; not completely elucidated at this time

## **Common Associations**

HTN (47.5%)

- Hyperopia (45%)
- · Known systemic autoimmune disease (28%)
- Idiopathic/Unknown (11-17%)
- Other reported assocations include:
   Papilledema, optic neuritis, thyroid eye disease, CNVM, CSCR, choroidal nevi, orbital/choridal turnos, surgical procedures, autoimmune diseases (RA, PMR, MS, VKH, etc), infectious causes (ie. Lyme disease)

al. Chorioretinal folds: Associated disorders and a related maculonathy. Am J Onhthalmol. 2014;157:1038-1

## Best Tools to Identify Choroidal Folds

- OCT\*\*\*
- $\cdot$  OCT-A
- FAF
- IVFA
- Retinal Photos

Del Turco C, et al. Optical coherence tomography angiography fastures of chorioretinal folds: A case aeries, Eur J Ophthalmol. 2017;27:a35-a38. Giuffa G et al. Optical coherence tomography of chorioretinal and choriskal folds. Acto Ophthalmol. Scinid. 2007;85:333-336. Domendial H, et al. Zies Angiojes aerietal domain optical coherence tomography angiography technical aspects. Em Ophthalmol. 2016;66:18-29

## OCT & Narrow Angle Glaucoma

#### Case

- $\cdot$  65 yo AAF
- + CC: routine clinical exam"
- $\cdot\,$  Upon SLE, noted pretty severe narrow angles per Von Herrick
- Gonio: no structures seen  $\operatorname{OU}$
- Additional testing possible?
   Anterior segment OCT
- Anterior segment OCT
   Anterior segment ultrasound
- · Is this pupillary block/narrow angles? Or plateau iris configuration?
- + Is it time to refer for cataract surgery and/or LPI?





## Plateau Iris Configuration (PIC)

"Classic teaching is that PIC <u>cannot</u> be diagnosed without first performing an iridatomy or iridectomy to rule out PB...there is no definitive criteria for diagnosing PIC on gonioscopy <u>without</u> a patent iridotomy or iridectomy present."
 Crowell et al. J Glaucoma. 2020

· Can we use anterior segment OCT to diagnose PIC without iridotomy/iridectomy???

Crowell EL, et al. Using anterior segment optical coherence tomography (ASOCT) parameters to determine pupillary block versu

### OCT and the Iridocorneal Angle in Glaucoma

- · Gonioscopy is still the gold standard! · OCT is a supplement, <u>not</u> a replacement!
- $\label{eq:specific/best} Specific/best measurements of anterior segment OCT to determine pupil block (PB; ie. narrow angles) vs. plateau iris configuration (PIC) are still being worked out$
- Crowell et al. (2020) suggest two best parameters of PB vs. PIC to be: 1. Lens/pupil size: • Horizontal> 4.1 mm = PIC • Vertical>3.8 mm = PIC

  - . Pupillary margin-center scleral spur length angles (PM-CSSL)  $\cdot$  Nasal  $\rightarrow$  <13.76° = PIC

  - $\begin{array}{l} \text{Nasar} \rightarrow <15.76^\circ \text{PIC} \\ \text{Superior} \rightarrow <16.45^\circ = \text{PIC} \\ \text{Temporal} \rightarrow <18.39^\circ = \text{PIC} \\ \text{Inferior} \rightarrow <18.58^\circ = \text{PIC} \end{array}$

Med. 2021;10:1-1 EL, et al ntical coh anhy (ASOCT) no munillary block ye

Pupillary margin-center scleral spur length angles  $(PM-CSSL) \rightarrow$  yellow angle

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ers to determine pupillary block

ell EL, et al. Using anterior segment optical cohe

## Conclusion: OCT for Pupil Block vs. Plateau Iris

· Need more studies...

- · OCT might be key to differentiate these in the future before iridotomy/iridectomy performed
- · I don't see gonio being replaced any time soon. · But OCT is a very good adjunct to gonio; I use it all the time!
- · Stay tuned for future research!

