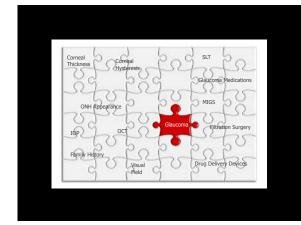




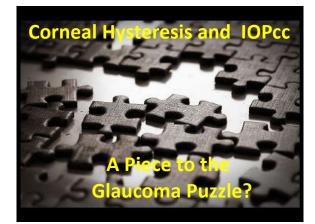


Aerie Pharmaceuticals, Inc.	CL	
Alcon Laboratories. Inc.		
Alderya Therapeutics, Inc.		
Alimera Sciences, Inc.		
Allergan, Inc.	CLS	
Avelino Labs		
Bausch + Lomb		
Beaver-Visitec International, Inc.		
BELKIN Laser Ltd		
Carl Zeiss Meditec		
Dompe		
Elios Vision		
Ellex		
EyePoint Pharmaceuticals		
Glaukos Corporation		
ImprimisRx		
IRIDEX		
IrisVision		
Neric Bio		
Kala Pharmaceuticals, Inc.		
Lumenis Vision		
New World Medical Inc		
Novartis Pharma AG		
Ocular Therapeutix		
Omeros Corporation		
Orasis Pharmaceuticals		
Quantel Medical		
Rayner		
Reichert, Inc.		
Santen, Inc.		
Shire		
Spyglass Pharma	C,SO	
Tarsus Pharmaceuticals		
TearClear	C,SO	
THEA		
Walase Inc		

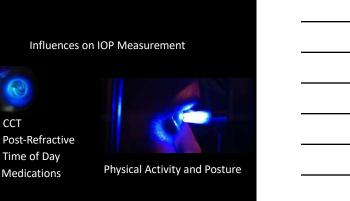








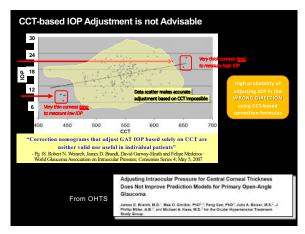




CCT

Time of Day

Medications



When gold standards change: time to move on from Goldmann tonometry?

Gus Gazzard,^{1,2} Hari Jayaram ⁽¹⁾,^{2,3} Ana M Roldan ⁽¹⁾,⁴ David S Friedman⁵

Br J Ophthalmol: 10.1136/bjophthalmol-2020-317112 24 September 2020

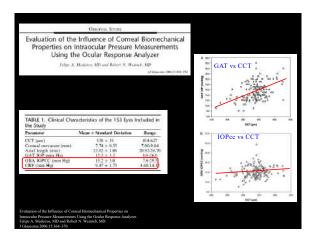
"Why are we persisting in using GAT clinically? The test itself is relatively time consuming, physicians often repeat the measurement because they cannot fully trust a technician, it slows down the clinic requiring technical staff to have slit lamps and place drops in patient's eyes and worse, it may be giving us a false sense of security.

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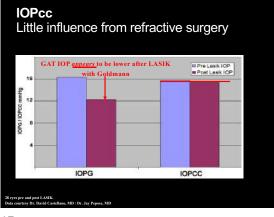
Reducing the Corneal effect on Measured IOP ORA's Patented IOPcc

ORA derived Corneal biomechanical information, which gives us Corneal Hysteresis, is also used to quantify (and reduce) the impact of these properties on the IOP measurement.

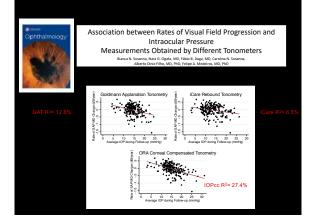
IOPCC: a pressure measurement that is less affected by corneal properties than other methods of tonometery, such as Goldmann (GAT).



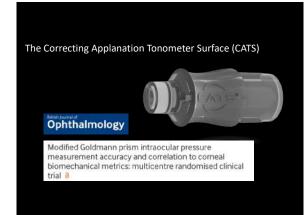


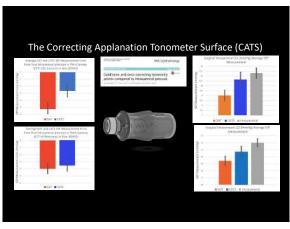




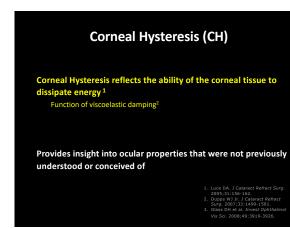












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Intro to Corneal Hysteresis

Viscoelastic tissue with complex, interconnected microstructure

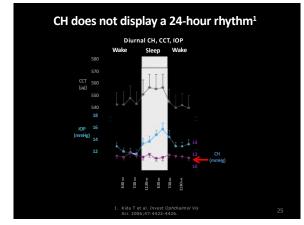
Geometrical attributes are not a surrogate for biomechanical properties

The eye appears to be a mechanical structural continuum

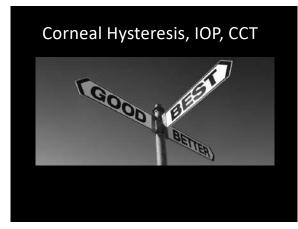
More than 13,000 + papers published on hysteresis

Average CH in Normal Subjects					
	Brazil	105	10.1 +/- 1.8		
	UK	272	10.2 +/- 1.2		
	China	125	10.9 +/- 1.5		
	Japan	204	10.2 +/- 1.3		
	Spain	88	10.8 +/- 1.5		
	USA	44	10.5 +/- 1.2		







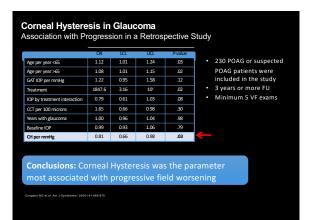






Clinical Evidence Why is CH relevant in Glaucoma?

(Low) CH has been consistently shown to be independently and strongly associated with or predictive of glaucoma progression



2013 Aug:120(6):1533-40. doi: 10.1016).opithe.2013.01.032. Epub-2013 May 1. Corneal hysteresis as a risk factor for glaucoma progression: a prospective longitudinal tudy. na FA¹, Meira-Freitas D, Lisboa R, Kuang TM, Zangwill LM, Weimeb RN-Author information

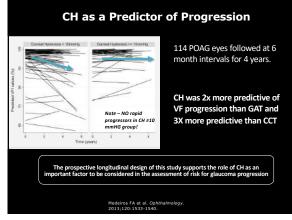
befract UNPOSE To evaluate the role of owneal hysteresis (CH) as a risk factor for the rate of visual field progression in a othor of patients with glaucoma followed prospectively over time. IESIGN: Proceedive schematicces choice study.

PARTICIPANTS: The study group included 114 eyes of 68 patients with glaucoma followed for an average of 4.0 ± 1.1 years. Vasual fields were obtained with standard automated parimetry. Included eyes had a median number of 7 (range, 5-12) tests during follow-up.

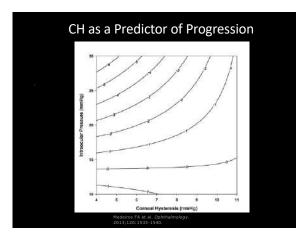
5-r2) relia during store-up. MENTODES: The CA measurements were acquired at baseline using the Ocular Response Analyzer (Reichert Instruments, Depex, NY). Exeluation of naise of visual field change during bolow-up was performed using the visual field index (VFI). Linear mixed models were used to investigate the reliatorship between rates of visual field loss and baseline CH, baseline intraocdar pressure (ICP), and central correlations (CCT), while adjusting for colornality confluenting factors. An interaction item between (ICP and CH was included in the model to investigate whether the effect of CP on rates of progression depended on the level of CH. **NAIN OUTCOME MEASURES:** Effects of CH, ICP, and CCT on rates of VFI loss over time:

NANN OUTCOME MEASURES: Interds of CH, I/O, and CL I on rates of VH loss over time. RESULTS: The CH had a significant filterial role and set visual field progression over time. The universitie model including only CH as a predictive factor along with time and their instanction, each 1 mm/s] ower CH was associated into 325% years taker rete of VH outcine over time (PC 0001). The multivatisatie model along with the chart of the ch

CONCLUSIONS: The CH measurements were significantly associated with risk of glaucoma progression. Eyes with lower CH had taster rates of visual field loss than those with higher CH. The prospective long/had raid design of this study supports the cloid CH as an important factor to be considered in the assessment of the risk of progression in patients with glaucoma.



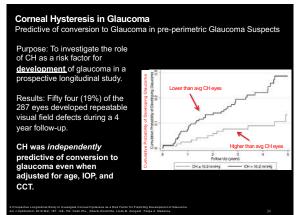


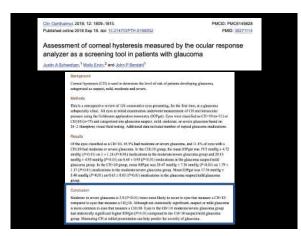




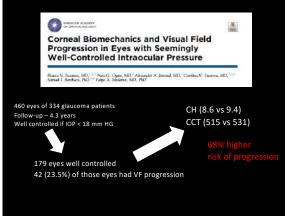
Corneal Hysteresis in Glaucoma Association with Normal Tension Glaucoma 82 progressing eyes of NTG patients under treatment Eyes were split into two groups: higher & lower than average CH b (95% c) Practice 1.18 (0.96 to -1.44) 0.12 0.99 (0.97 to 1.01) 0.35 ne VF MD (dB) • Of the 39 eyes with low CH, 26 (66.7%) CCT (µm) ubfoveal choroidal thickness 0.99 (0.98 to 1.00) 0.08 showed progression NFL thickness (average) 0.96 (0.92 to 0.99) 0.04 • Of the 43 eyes with high CH, 15 (34.9%) 0.97 (0.94 to 1.01) 0.09 RNFL thic ness (temporal) 0.98 (0.96 to 1.01) 0.13 0.32 (0.17 to 0.62) <0.01 NFL thickness (inferior) showed progression Corneal Hysteresis (mmHg) These findings suggest that CH can be used as one of the prognostic factors for progression, independent of corneal thickness or IOP Park Et. Al Br J Ophthalmol. 2015 Jan 2. pit bjophth 305962. doi: 10.1136/bjophthalmol-2014-305962.

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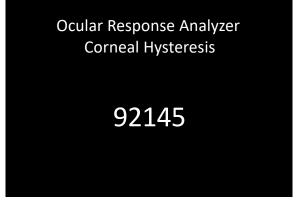


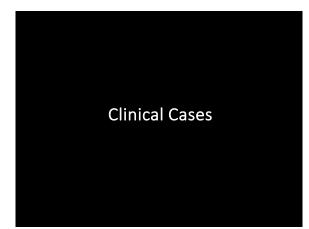














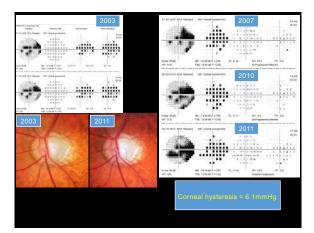
Slow Progression Despite IOP control

62 yr old, male, with diagnosis of POAG

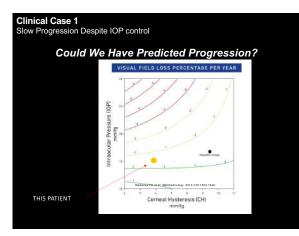
Currently on maximum tolerated medical therapy and having undergone 2 sessions of laser trabeculoplasty

- BCVA: 20/20 OU
- Biomicroscopy: normal
- GAT IOP: 13mmHg to 15mmHg on maximum tolerated medical therapy
- Corneal thickness: 545μm OD 541μm OS

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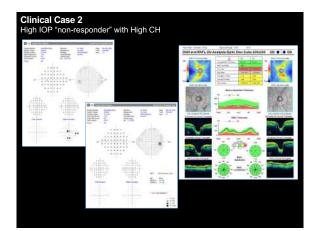


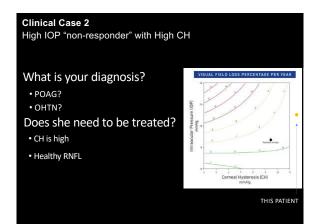




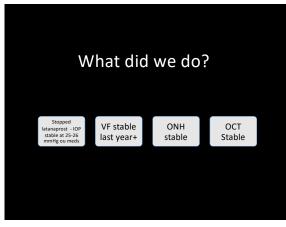
Clinical Case 2 High IOP "non-responder" with High CH 73 y/o Caucasian Female diagnosed with OHTN <u>3 years prior by outside provider</u> Sister also being followed for glaucoma but not being treated Meds: Bystolic, Pravastatin, MVI Ocular Meds: Latanaprost qhs Tmax: 26 mmHg OU Medicated IOP: 21 mmHg ou on multiple visits CCT: 558 OD 562 OS CH: 11.6 OD 12.3 OS Heathy RNFL, C/D 0.65 OD 0.6 OS, no disc heme or beta zone atrophy

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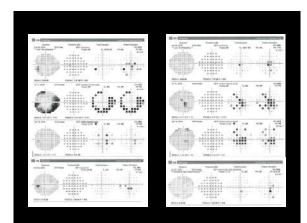


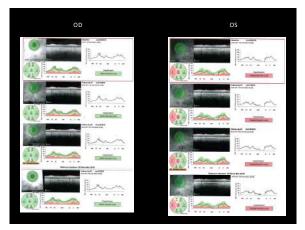
Case 3: Patient CT (65 YO Caucasian male)

CC: Glaucoma Followup PmHx: Hyperlipidemia FeHx: Unremarkable Medications: Lipitor Topical Medications: latanoprost 0.005% qhs OU

Tmax IOP: 28 mmHg OU Current IOP: 22 mm HG OD, 23 mm HG OS Corneal Hysteresis: 10.1 OD, 11.3 OS Pach: 545 OU Gonioscopy: Open to CB, no pigment present in TM SLE: Unremarkable, except for well centered IOL's

ONH: C/D OD: 0.70/0.70 C/D OS: 0.75/0.75







Would you treat SK?

Case Data:

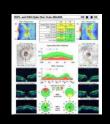
- Age: 70 year old man presents
- IOPs (GAT): 28 mmHg OU

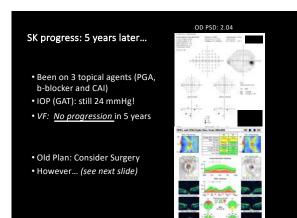
• CCT: 545 microns

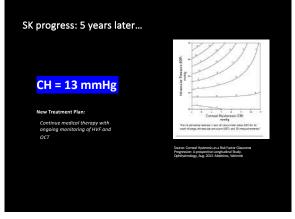
- VF: Full (PSD 1.4)
- OCT: borderline, some thinning

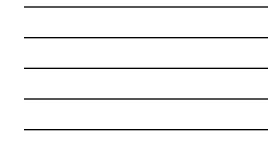


Corneal Hysteresis: not available









Summary & Considerations of SK Case

SUMMARY

- High risk OHTN, IOP: 28 mmHg
 CCT average: 545 microns
 Patient's IOP not much lower with
 treatment
- No progression in 5 years
 High Corneal Hysteresis may have predicted this

CONSIDERATIONS

What might have been done differently if Corneal Hysteresis was known 5 years ago? How might knowing Corneal Hysteresis today change management going forward?



