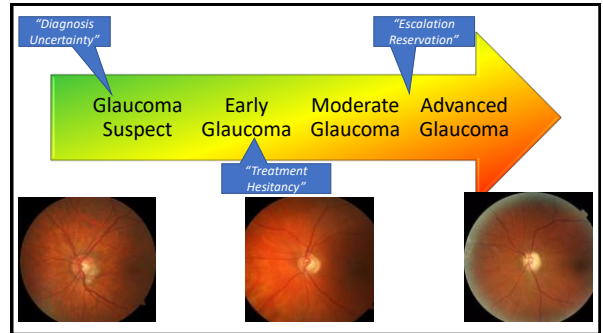


GlaucoMantras

Re-centering our Glaucoma Care

Kentucky Optometric Association
Fall Conference 2024

Center for Sight & Dry Eye Clinic
Diplomate (Glaucoma) AAO
Austin Lifferth OD, FAAO
September 14, 2024



mantra

NOUN

mantra (noun) · *mantras* (plural noun)

1. a statement or slogan repeated frequently.

ORIGIN

late 18th century: Sanskrit, literally 'a thought, thought behind speech or action', from man- 'think', related to mind.



Glaucomantra

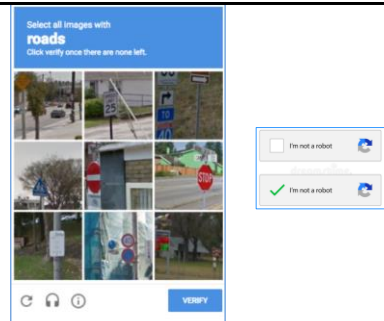
NOUN

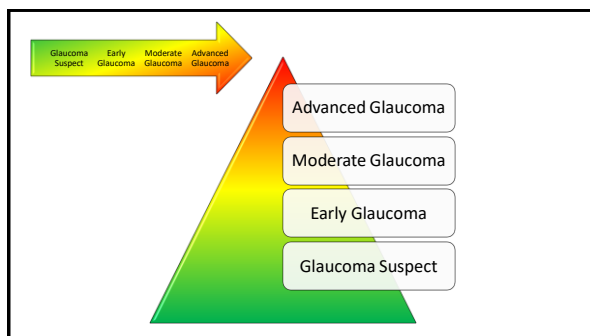
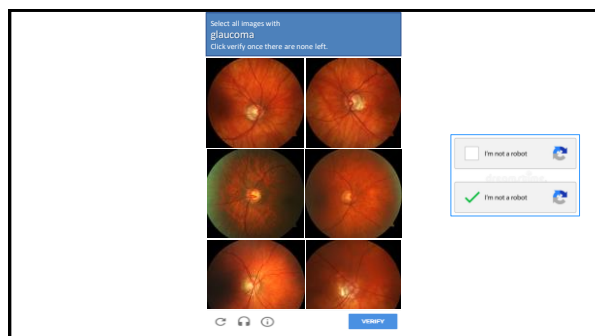
GlaucoMantra (noun) · *GlaucoMantras* (plural noun)

1. a statement or slogan *repeated frequently* to assist with diagnosing glaucoma earlier and treating progression sooner; phrases to *re-center* our glaucoma care

ORIGIN

late 18th century: Sanskrit, literally 'a thought, thought behind speech or action', from man- 'think', related to mind; Spring IOA 2022.





Who is a Glaucoma Suspect?

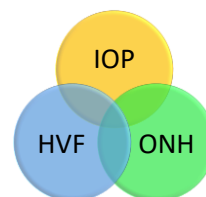
"A glaucoma suspect is an individual with clinical findings and/or a constellation of risk factors that indicate an increased likelihood of developing primary open-angle glaucoma (POAG)."



Phum L, Lim M, Williams R, et al. Preferred practice pattern: Primary Open-Angle Glaucoma Suspect Preferred Practice Pattern® Guidelines. Ophthalmology. January 1, 2016;123:P112-P131.

Who is a Glaucoma Suspect?

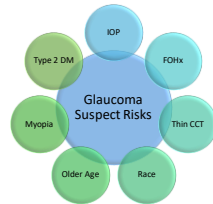
"A diagnosis for primary open-angle glaucoma (POAG) suspect is established by the presence of one of the following conditions: a consistently elevated intraocular pressure (IOP), a suspicious-appearing optic nerve, or abnormal visual field."



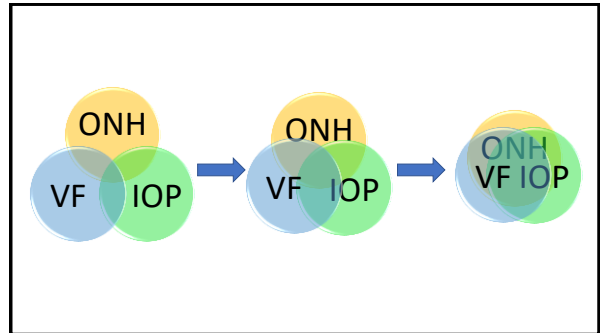
Phum L, Lim M, Williams R, et al. Preferred practice pattern: Primary Open-Angle Glaucoma Suspect Preferred Practice Pattern® Guidelines. Ophthalmology. January 1, 2016;123:P112-P131.

Who is a Glaucoma Suspect?

"Highlights of established risk factors for a POAG suspect diagnosis include an elevated IOP, family history of glaucoma or glaucoma suspect, thin central cornea, race, older age, myopia, and type 2 diabetes."



Pooni L, Lim M, Williams K, et al. Preferred practice pattern: Primary Open-Angle Glaucoma Suspect Preferred Practice Pattern® Guidelines. Ophthalmology. January 1, 2016;123(1):P21-P23.



Case Presentation

- ▶ 41 year-old male returns for *continued* glaucoma suspect evaluation and testing
 - ▶ No relevant ocular history
 - ▶ Review of Systems: unremarkable and noncontributory.
 - ▶ No medications
 - ▶ Non-smoker
 - ▶ Social drinker
 - ▶ Healthy, active
 - ▶ Negative FOHx

Case Presentation

UNCORRECTED VISUAL ACUITY

OD: = 20/20

OS: = 20/20

PUPILS

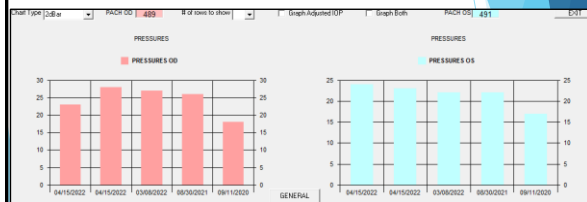
OU: round, reactive; (-) rAPD OD, OS

CONFRONTATION VISUAL FIELDS

OD: Full to FC

OS: Full to FC

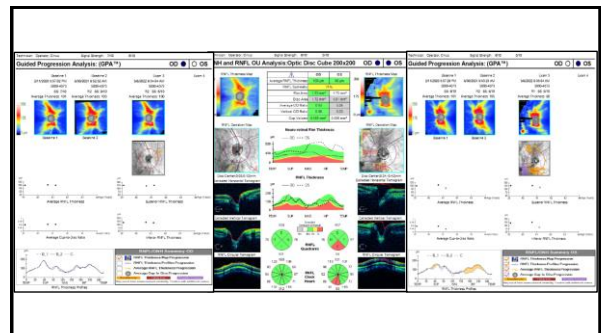
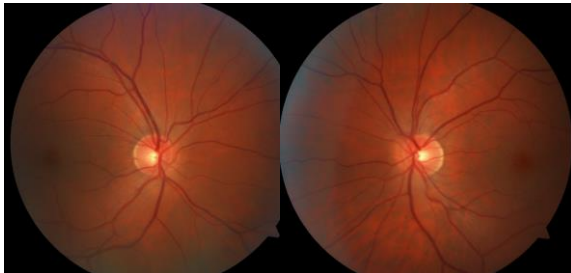
Case Presentation

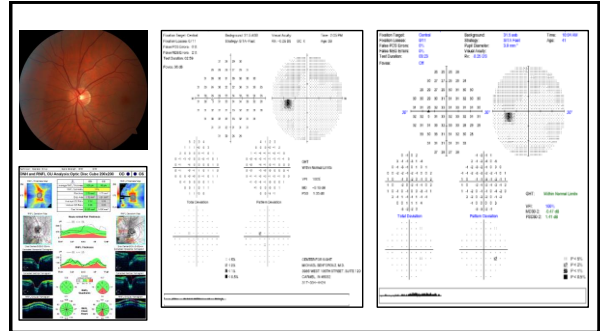
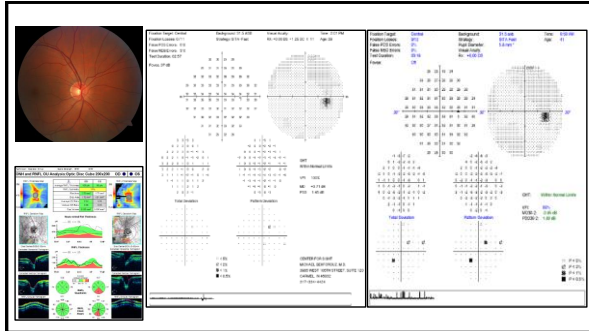


Case Presentation

SLIT LAMP EXAM OU

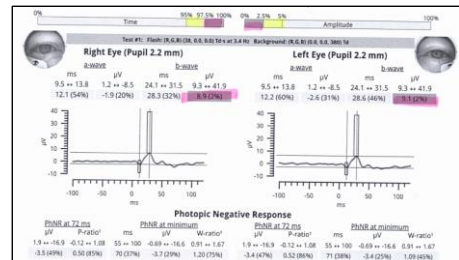
- ▶ Lids/Adnexa: clear OU
- ▶ Conjunctivae: palpebral/bulbar clear
- ▶ Sclera: white
- ▶ Cornea: clear/normal; (-) pigment OU
- ▶ Anterior Chamber: deep and quiet OD, OS; NO cell
- ▶ Iris: clear/normal; NO NVI, synechiae
- ▶ Lens: clear OU; NO PXE OU

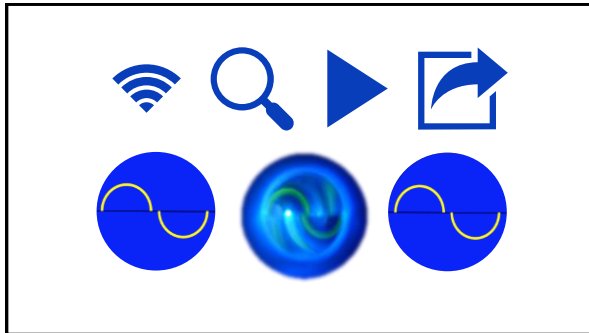




FACTORS	Points for Factors				
	0	1	2	3	4
Age (years)	<input checked="" type="checkbox"/> <45	<input type="checkbox"/> 45 to <55	<input type="checkbox"/> 55 to <65	<input type="checkbox"/> 65 to <75	<input type="checkbox"/> >75
Intraocular Pressure (mm Hg) Mean (3 measurements per eye and average of 2 eyes)	<input type="checkbox"/> <22	<input type="checkbox"/> 22 to <24	<input type="checkbox"/> 24 to <26	<input checked="" type="checkbox"/> 26 to <28	<input type="checkbox"/> ≥28
Central Corneal Thickness (μ) Mean (3 measurements per eye and average of 2 eyes)	<input type="checkbox"/> ≥ 600	<input type="checkbox"/> 576-600	<input type="checkbox"/> 551-575	<input type="checkbox"/> 526-550	<input checked="" type="checkbox"/> ≤525
Vertical Cup/Disc Ratio by Contour Mean (1 measurement per eye and average of 2 eyes)	<input type="checkbox"/> <0.3	<input checked="" type="checkbox"/> 0.3 to <0.4	<input type="checkbox"/> 0.4 to <0.5	<input type="checkbox"/> 0.5 to <0.6	<input type="checkbox"/> ≥0.6
Visual Field: Humphrey Pattern Standard Deviation (sd) Mean (2 measurements per eye and average of 2 eyes)	<input checked="" type="checkbox"/> <1.8	<input type="checkbox"/> 1.8 to <2.0	<input type="checkbox"/> 2.0 to <2.4	<input type="checkbox"/> 2.4 to <2.8	<input type="checkbox"/> ≥2.8
Optic Disc Variance Mean (2 measurements per eye and average of 2 eyes)	<input type="checkbox"/> < 3.24	<input type="checkbox"/> 3.24 to <4.0	<input type="checkbox"/> 4.0 to <5.76	<input type="checkbox"/> 5.76 to <7.84	<input type="checkbox"/> ≥7.84
Sum of Points and Estimated 5-Year Risk of Developing POAG					
Sum of Points	0-4	5	6-8	9-10	11-12
Estimated 5-Year Risk of Developing POAG	<4.0%	10%	15%	20%	≥33%
Total Points: 8	Estimated Risk: 10% The patient's estimated 5-year risk (%) of developing early glaucoma in at least one eye				

Risk Calculator | Ocular Hypertension Treatment Study (OHTS)
[wustl.edu]





Intraocular Pressure Applications

- Mean IOP is the greatest modifiable risk factor in the development^{1,2} and the progression of glaucoma^{3,6}.
- Each mmHg matters¹⁻⁷.
- IOP reduction decreases the risk of development^{1,2} and progression of glaucoma^{3,6}.
 - Even among patients with IOP with a median IOP of < 20mmHg, 30% IOP reduction decreased risk of progression⁷.

1. Kass MA, Heuer DK, Hagan-Johnson H, et al. The 15-year Pigment Dispersion Study: a longitudinal assessment that topical ocular hypotensive medication delays or prevents glaucoma. *Arch Ophthalmol*. 2002;120(6):790-800.
2. Miglior S, Zojan T, Pfeiffer N, et al. Results of the European Glaucoma Prevention Study. *Ophthalmology*. 2008; 112: 366-375.
3. Hogg A, Linds ME, Bhargava R, et al. Reduction of intraocular pressure and glaucoma progression: results from the Early Manifest Glaucoma Trial. *Arch Ophthalmol*. 2002;120(11):2061-2070.
4. Mach DC, Gilliland BW, Nouri LM, Lichter PR, Varma R. C3275 Study Group. Intraocular pressure control and long-term visual field loss in the Collaborative Initial Glaucoma Treatment Study. *Ophthalmology*. 2011;118(7):1366-1373.
5. The Advanced Glaucoma Intervention Study (AGIS) 7. The relationship between control of intraocular pressure and visual field deterioration. The AGIS Investigators. *Ann Ophthalmol*. 2003;35(4):249-444.
6. Sommer A, Tielch JM, Katz J, et al. Relationship between intraocular pressure and primary open angle glaucoma among white and black Americans. The Baltimore Eye Survey. *Arch Ophthalmol*. 1993;111(10):1300-1305.
7. Comparison of glaucoma progression between untreated patients with normal-tension glaucoma and patients with therapeutically reduced intraocular pressures. Collaborative Normal-Tension Glaucoma Study Group. *Ann Ophthalmol*. 1998; 120:487-497.

IOP Implications

"...the most important healthcare implication from this analysis is to avoid being falsely reassured by a lower level of IOP in glaucoma case finding."¹



- IOP is not glaucoma.

1. Chan MPY, Khawaja AP, Broadway DC, Yip J, Luben R, Hayslet S, Peto T, Khaw KT, Foster PJ. Risk factors for previously undiagnosed primary open-angle glaucoma: the EPIC-Norfolk Eye Study. *Br J Ophthalmol*. 2021 Jun 25;95(ophthalmol-2020-317718).

IOP Implications

"Kill the magic number"

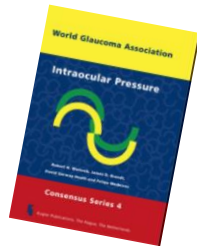
"We have so far failed to eliminate the incorrect notion that the IOP value 21 mmHg is meaningful or represents a benchmark for treatment..."

We should not care if the baseline IOP is 30, 20, or 10mmHg, as we will use whatever is the baseline to set a target lowering."

Guligley HA. 21st century glaucoma care. *Eye (Lond)*. 2019 Feb;33(2):254-260.

IOP Limitations

"Among chronic diseases, glaucoma is remarkable in that its primary risk factor, IOP, is measured only rarely and mostly randomly, perhaps a few time a year in most patients."



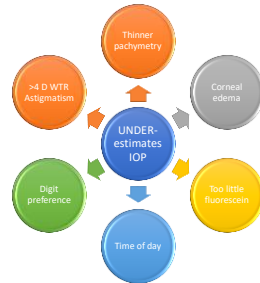
IOP Limitations

"...a single IOP measurement during so-called office hours is a *poor surrogate of the entire IOP profile* of a patient with glaucoma..."

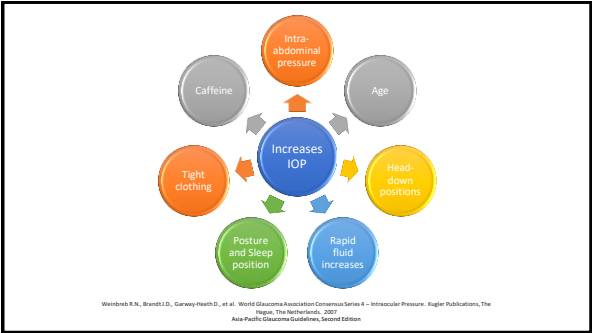
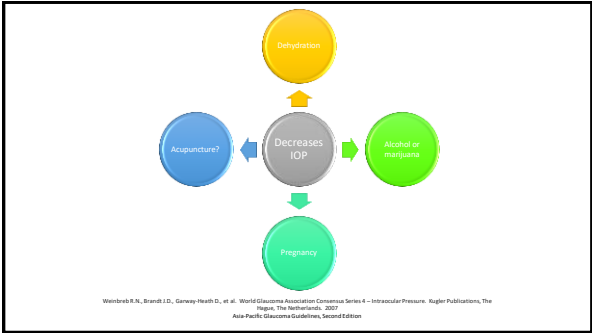
Konrads AG, K. M. (2018). Diurnal and 24-h Intraocular Pressure in Glaucoma: Monitoring Strategies and Impact on Progress and Treatment. *Adv Ther.* 33(12):1775-1804.



Worsham R.N., Brandt J.D., Garway-Heath D., et al. World Glaucoma Association Consensus Series 4 – Intraocular Pressure. Kugler Publications, The Hague, The Netherlands, 2007.
Asia-Pacific Glaucoma Guidelines, Second Edition
Dewanar AM, Gnanasekar SP, Magill RL, Gosh SL. Mask-induced Artificially High Intraocular Pressure Measurement Using Goldmann Applanation Tonometry. J Glaucoma. 2014;23(4):270-272.

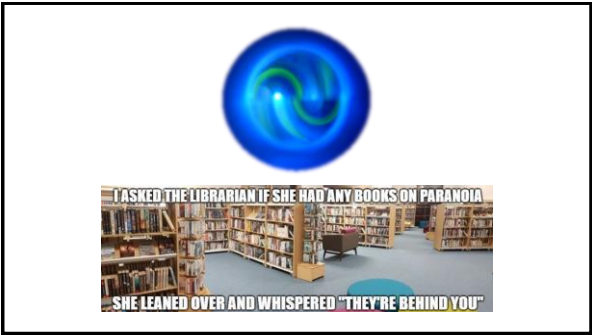


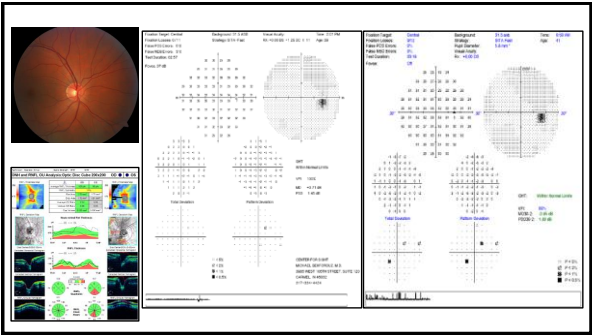
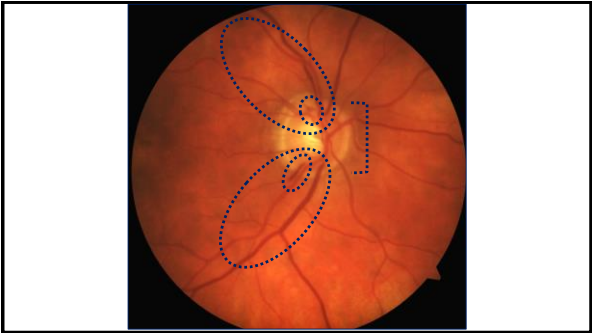
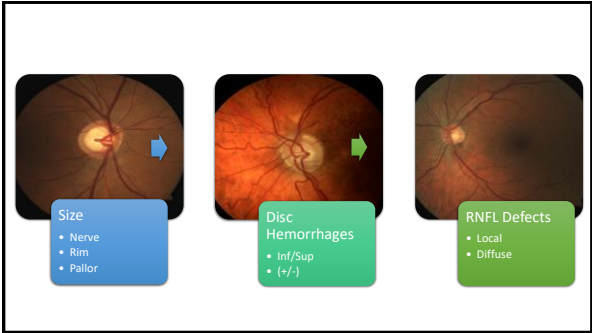
Worsham R.N., Brandt J.D., Garway-Heath D., et al. World Glaucoma Association Consensus Series 4 – Intraocular Pressure. Kugler Publications, The Hague, The Netherlands, 2007.
Asia-Pacific Glaucoma Guidelines, Second Edition

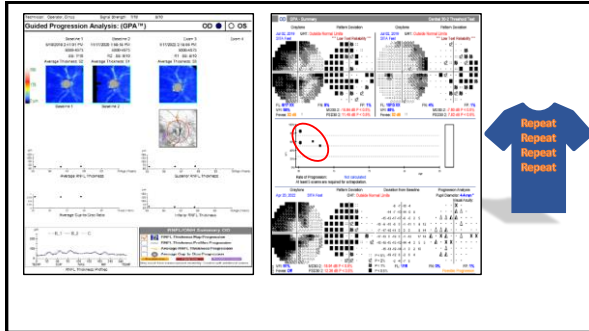


ACTIVITY	INTRACULAR PRESSURE
Light touch through adnexal skin or lids (gentle eye wiping)	Approximately double baseline IOP
Voluntary squeezing of lids (squeinting)	Elevations up to 90 mmHg
Eye Compression (massaging, rubbing, wiping, drying)	Up to 30 mmHg and 40 mmHg
Short duration supine positions	A mean elevation of 4.4 mmHg
The dependent (lower) eye during side sleeping	A mean of 2 mmHg above supine IOP
Long duration prone sleeping	A mean elevation of 40 mmHg
Contact between the eye and bedding surfaces	A mean elevation of 22±5 mmHg
Inverted body position	A mean elevation of 36 mmHg
Wearing swimming goggles	Elevations up to 48 mmHg
Straining involving facial congestion	Elevations of 10-25 mmHg
Playing loud, high pitch notes on a trumpet	Elevations up to 44 mmHg
Wearing a tight necktie	Elevations of 2-4 mmHg

McMonies, C. (2013). An examination of the hypothesis that intraocular pressure elevation episodes can have prognostic significance in glaucoma suspects. *J Glom*, 4(2), 102-116.

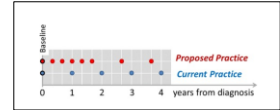




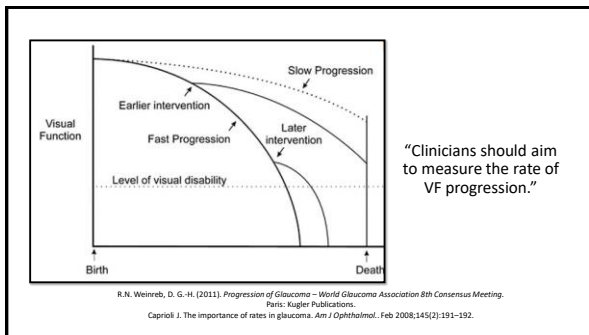


“Perform sufficient examinations to detect change.”

- “A good baseline of reliable VFs is essential to be able to monitor for progression.”
- “Decisions on progression should not be made by comparing only the most recent field with the one before.”
- “Suspected progression should be confirmed by repeating the field.”¹

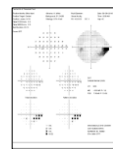
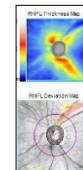


1. R.N. Weinreb, D. G.-H. (2011). *Progression of Glaucoma – World Glaucoma Association 8th Consensus Meeting*. Paris: Kugler Publications.
Boudhva T, Crabb DP. More frequent, more costly? Health economic modelling aspects of monitoring glaucoma patients in England. *BMC Health Serv Res*. 2016;16(1):111.
*See also: Chauhan BC, Darway-Heath CP, Glick FJ, Rossetti L, Bagnasco B, Neveu-Strauss AC, Naj A. Practical recommendations for measuring rates of visual field change in glaucoma. *Br J Ophthalmol*. 2008 Apr;92(4):569-73.



Visual Field Progression - WHERE do we see it? -

- Regional preferential rim loss depending on stage of disease:
 - Early: Look carefully in I.T. and S.T. disc regions
 - Moderate: Temporal horizontal disc region
 - Advanced: Inferior nasal, then superior nasal rim loss
- The sequence of disc sector rim loss correlates with the progression of the VF defects:
 - Early VF loss: Nasal upper or lower quadrant
 - Moderate VF loss: Connecting arcuate
 - Advanced: Island of sensitivity in the inferior-temporal VF

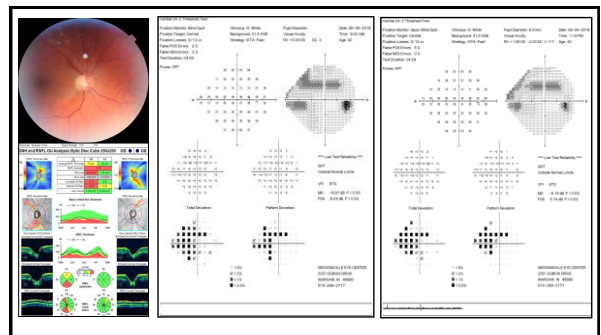
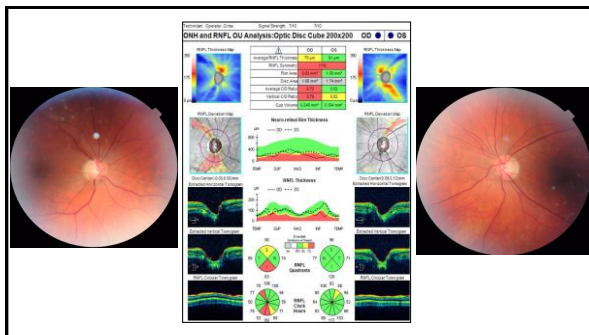
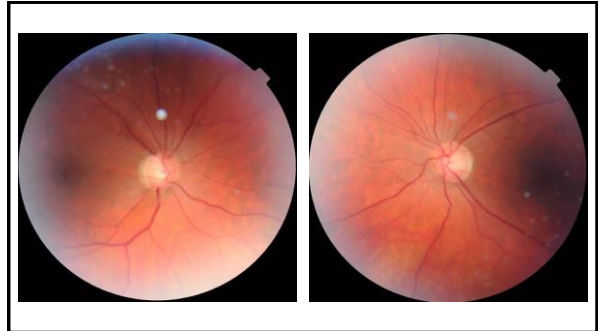


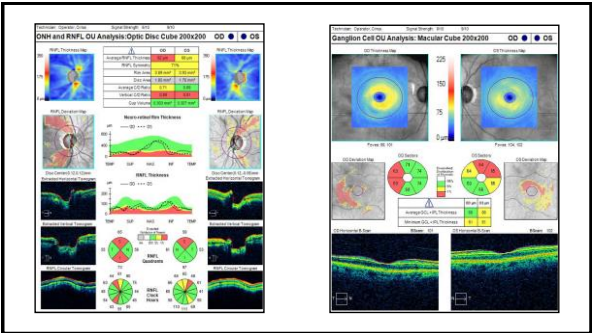
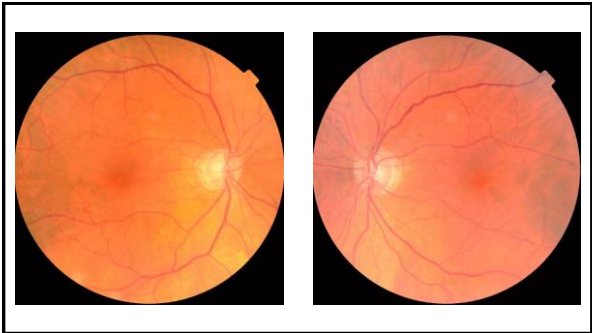
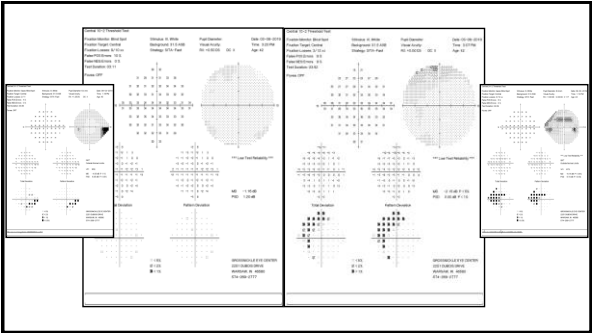
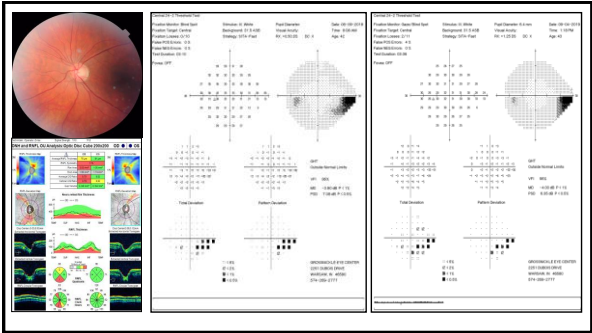
Visual Field Progression

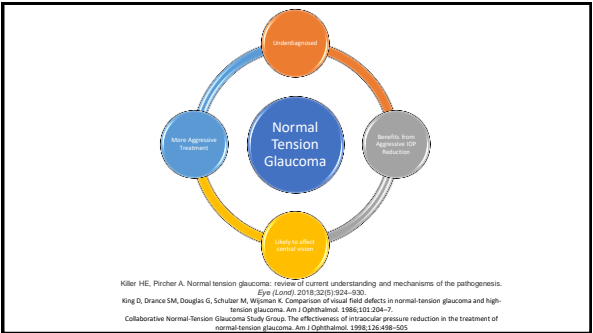
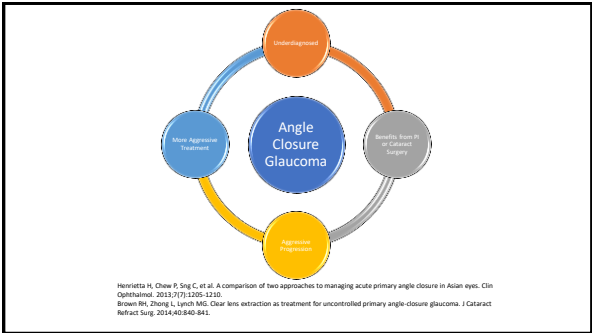
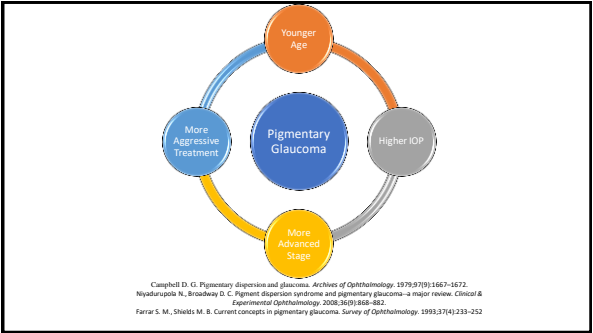
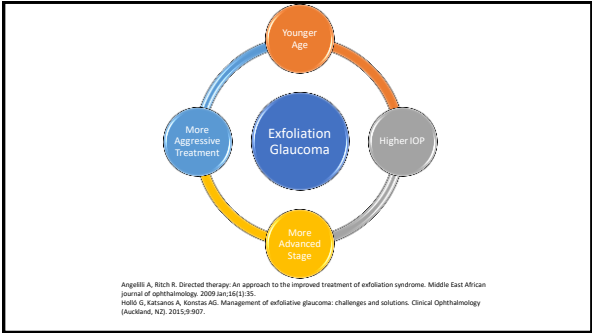
- HOW can we detect it? -

- LOOK for:
 - Deepening of current defects (PSD)
 - Enlargement of current defects (MD)
 - NEW defects
- “Visual field progression may be analyzed by either ‘event-’ or ‘trend-’ based methods”
 - “In general, event-based methods are used early in the follow-up, when few VFs are available for serial analysis.
 - “In general, rate-based analyses are used later in the follow-up, when a greater number of VFs is available over a sufficient period of time to measure the rate of progression.”

R.N. Weinreb, D. G. A. (2015). *Progression of Glaucoma - World Glaucoma Association 8th Consensus Meeting*. Paris: Kugler Publications.







Lower is Better.

- Sufficient IOP reduction
 - Residual life expectancy/Age
- Sufficient treatment
 - Over treatment for older patients
 - Under treatment for younger patients



Target IOP Treatment Principles

“In the end, it will be impossible to know if we overreacted or did too much, but it will be QUITE apparent if we under reacted or did too little.”

(Dr. Darrin M. Peppard
March 20, 2020)

“The decision to initiate glaucoma treatment should be based on the assessment of the risks for development of functional impairment or decrease in vision-related quality of life, taking into account factors such as coexisting ocular conditions, the patient’s life expectancy, and general health status, as well as his/her perception and expectations about treatment.”

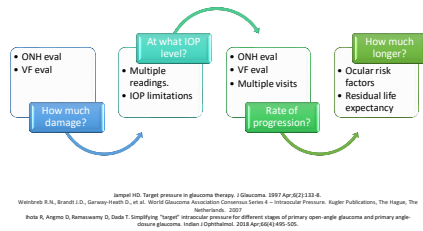
- Age
- Stage

Leiberman J, Weinreb R. World Glaucoma R. Medical Treatment Of Glaucoma: The 7th Consensus Report Of The World Glaucoma Association. Amsterdam: Hoger Publications; 2015. p 3.

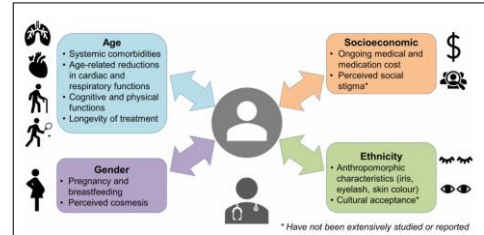
Target IOP Treatment Principles



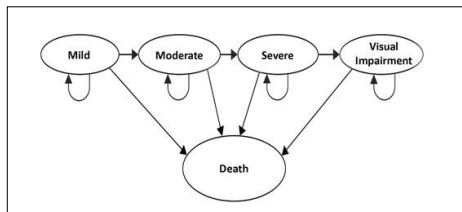
Evaluating Target IOP



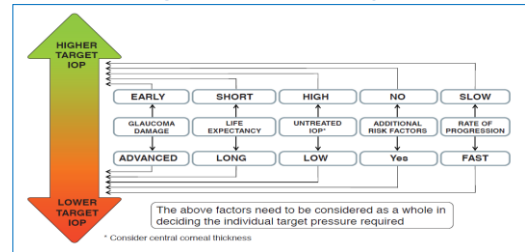
Target IOP Treatment Principles



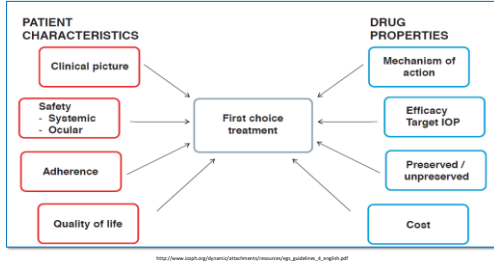
Target IOP Treatment Principles



Target IOP Treatment Principles

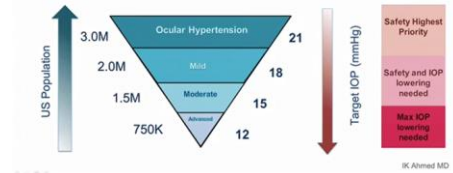


Target IOP Treatment Principles



Target IOP Treatment Principles

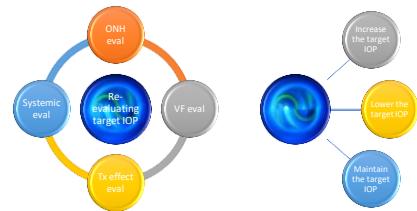
The Glaucoma Pyramid: Target IOP Guides Therapy



"In treated patients, failing to achieve target IOP was associated with more rapid VF worsening. Eyes with moderate glaucoma experienced the greatest VF worsening from failing to achieve target IOP."

Villasana GA, Bradley C, Ramulu P, Unberath M, Yohannan J. The Effect of Achieving Target Intraocular Pressure on Visual Field Worsening. *Ophthalmology*. 2022 Jan;129(1):35-44

Re-evaluating Target IOP



Jampel RD. Target pressure in glaucoma therapy. *J Glaucoma*. 1997 Apr;6(2):122-6.
Weinreb RN, Brubaker RD, Toris JH, et al. World Glaucoma Association Executive Summary Series 4 - IOP Lowering Potentials. Kugler Publications, The Hague, The Netherlands. 2007.
Hsieh R, Argente D, Ramakrishna G, Shih T. Simplifying "target" intraocular pressure for different stages of primary open-angle glaucoma and primary angle-closure glaucoma. *Indian J Ophthalmol*. 2018 Apr;66(4):493-500.



56,000 Ways To Treat Glaucoma

From ignorance our creator flows. The only venturers are the wise.
Matthew Poole (1664-1721)

The addition of new glaucoma medications in the past few years, including several with once daily dosing and improved efficacy, has greatly expanded medical therapy options for glaucoma patients. Our treatment paradigm had long been a stepped algorithm with a reasonably ordered order of drug addition, culminating in what has been labeled, but variably defined, as maximal tolerated medical therapy (MTMT). MTMT can be defined simply as the point at which it makes no sense to add more medications. There are now more steps in the algorithm, corresponding to new classes of medications. There are more choices of individual agents within each class, and no longer in the order of addition class-out. As of the late 1970s, the path to MTMT began with timolol, then progressed through pilocarpine and propine and ended with an oral carbonic anhydrase inhibitor (CAI).

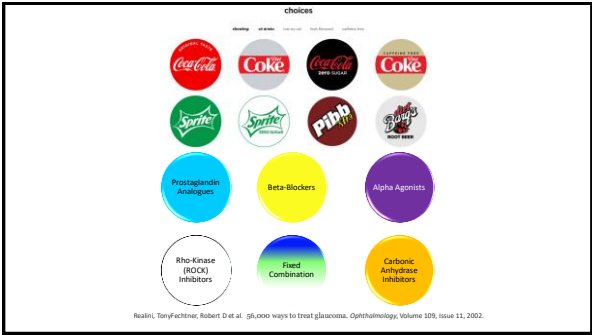
Classes now include beta-blockers (timolol maleate aqueous solution 0.25% or 0.5%, timolol maleate gel-forming solution 0.25% or 0.5%, timolol hemihydrate 0.25 or 0.5%, levobunolol 0.25% or 0.5%, betaxolol 0.25% or 0.5%, metipranolol 0.3%, and carbetolol 1.0%); carbonic anhydrase inhibitors (CAI) topically (dorzolamide 2% and brinzolamide 1%), orally (acetazolamide 125 mg, 250mg, or 500mg sustained-release, acetazolamide 250mg or 500mg, and dichlorophenamide 50mg); prostaglandins (latanoprost 0.005%, unoprostone 0.15%, bimatoprost 0.03%, and travoprost 0.004%); adrenergic agonists (brimonidine 0.2%, brimonidine P 0.15%, apraclonidine 0.5%, dipivefrin 0.1%, and epinephrine 0.5%, 1%, and 2%); and the miotics (pilocarpine solution 0.5%, 1%, 2%, 4%, 6% or 8%, pilocarpine gel 4%, and carbachol 0.75%, 1.5%, 2.25%, or 3%).

As of this writing, ignoring generic equivalents and combination products, glaucoma patients may now be on a regimen consisting of any of 12 beta-blockers, 8 CAIs, 4 prostaglandins, 7 adrenergic agonists and 11 miotics. For each class of drugs, a patient is either on one of the available agents or not in that class, so that the number of choices per class is one more than the number of agents in that class. Additionally, as one choice is zero from all classes, we must subtract one from the total. Considering all possible regimens from monotherapy to MTMT with an agent from each of the five pharmacologic classes, there are some 56,159 different ways to reach MTMT, $(12 \text{ (beta-blockers)} \times 9 \text{ (CAI)} \times 5 \text{ (prostaglandins)} \times 8 \text{ (adrenergic agonists)} \times 12 \text{ (miotics)}) - 1 = 56,159$.

Latanoprostene Bunod 1 Rho-Kinase Inhibitor (Netarsudil)

67,391!

Realini, TonyFecthner, Robert D et al. 56,000 ways to treat glaucoma. Ophthalmology, Volume 109, Issue 11, 2002.



Prostaglandin Analogues

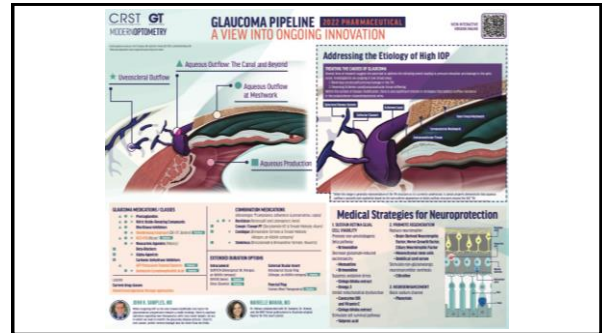
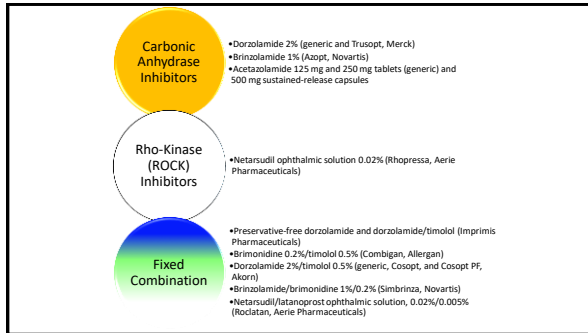
- Latanoprost 0.005% (generic and Xalatan, Pfizer)
- Latanoprost ophthalmic emulsion 0.005% (Xelpros, Sun Pharma)
- Bimatoprost 0.01% (Lumigan, Allergan); Bimatoprost 0.03% (generic)
- Travoprost 0.004% (generic and Travatan Z, Novartis)
- Tafluprost 0.0015% (Zioptan, Akorn)
- Latanoprostene bunod 0.024% (Vyzulta, Bausch + Lomb)

Beta-Blockers

- Timolol maleate 0.25% and 0.5% (Timoptic, Timoptic Ocudose and Timoptic XE, Valeant Ophthalmic) and 0.5% (Istalol, Bausch + Lomb, and generic)
- Timolol hemihydrate 0.25% and 0.5% (Betimol, Akorn)
- Beta-1 selective betaxolol hydrochloride 0.5% (generic) and 0.25% (Betoptic-S, Novartis)
- Levobunolol hydrochloride 0.5% (Betagan, Allergan) and 0.25% and 0.5% (generic)

Alpha Agonists

- Apraclonidine (Iopidine 0.5% and 1%, Novartis and generic)
- Brimonidine: 0.2% preserved with BAK and 0.15% non-BAK preservative and 0.1% (Alphagan P, Allergan, preserved with Purite)



Lower IS More

- Sufficient IOP reduction
 - Residual life expectancy/Age
- Sufficient treatment
 - Over treatment for older patients
 - Under treatment for younger patients



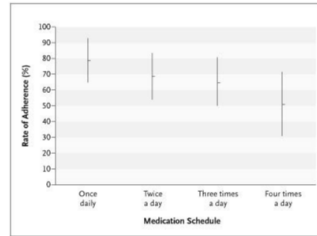
"The goal of glaucoma treatment is the preservation of vision *and* vision-related quality of life throughout the patient's lifetime."

Corbode Consultation in Glaucoma: 40 Clinical Questions Dale K., Phum, Richard A., Lewis, Steve J., Gedde, 2008. SLACK Incorporated, Thorofore, New Jersey, PA.

“Patient adherence is the wild card in the deck for controlling glaucoma progression.”



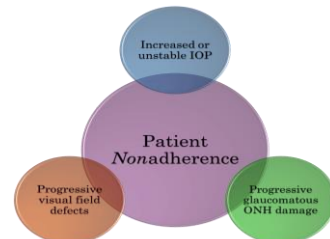
Figueret M, Dickerson J. The Role of Minimally Invasive Glaucoma Surgery Devices in the Management of Glaucoma. *Optometry And Vision Science: Official Publication Of The American Academy Of Optometry* [serial online]. February 2018;95(2):155-162.



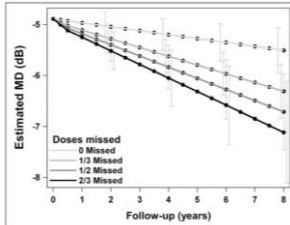
Osterberg L, Blaschke T. Adherence to medication. *N Engl J Med*. 2005 Aug 4;353(5):487-97.

“Ultimately,... nonadherent patients...fail to achieve the intended or full effect of the treatment.”

Budenz D. A Clinician's Guide to the Assessment and Management of Nonadherence in Glaucoma. *Ophthalmology* January 1, 2009;116:543-547.



1. Savath B, Baloch S, Rubin A, et al. Original article: The Relationship between Glaucoma Medication Adherence, Eye Drop Technique, and Visual Field Defect Severity. *Ophthalmology* January 1, 2012;118:2398-2402.
2. Schwartz G, Gagny H. Adherence and Persistence with Glaucoma Therapy. *Survey Of Ophthalmology* January 1, 2008;53(Supplement):557-568.
3. O'Brien C, Schuster L, van der Horst E, Walters C. Original article: Nonadherence with Ocular Hypertension Treatment in Patients with Glaucoma or Ocular Hypertension: An Evidence-Based Review. *Ophthalmology*. January 1, 2005;112:953-961.e7.
4. Stewart WC, Chouk RP, Hunt RH, Sethuraman G. Factors associated with visual loss in patients with advanced glaucomatous changes in the optic nerve head. *Am J Ophthalmol*. 1995;119:176-183.



Newman Casey PR, Nizol LM, Gillespie BW, Jantz NK, Lichter PR, Musch DC. The Association between Medication Adherence and Visual Field Progression in the Collaborative Initial Glaucoma Treatment Study. *Ophthalmology*. 2020 Apr;127(4):477-483.

“To increase the effectiveness of our current glaucoma treatments, there is a *critical need* to focus on helping support patients in *improving their glaucoma medication adherence.*”

“...increasing the effectiveness of adherence interventions may have a far greater impact on the health of the population than any improvement in specific medical treatments.”

Haynes RB, McDonald H, Garg AX MR. World Health Organization. http://www.who.int/ctcp/ncpeledge/publications/adherence_full_report.pdf

“[Eye care providers]...do a poor job of detecting nonadherence in their patients.”

Bailey D. A Clinician's Guide to the Assessment and Management of Nonadherence in Glaucoma. *Ophthalmology* January 1, 2008;116:543-547.

“Physician *attitude* has been shown to play a large role in patient adherence...”

Bailey D. A Clinician's Guide to the Assessment and Management of Nonadherence in Glaucoma. *Ophthalmology* January 1, 2008;116:543-547.

“...addressing adherence issues involves changing physician behavior, which may result in changes in patient behavior.”

Budenz D. A Clinician's Guide to the Assessment and Management of Nonadherence in Glaucoma. *Ophthalmology* January 1, 2008;116:550-561.

I know it must be difficult to take all your medications regularly. How often do you miss taking them?²³
Of the medications prescribed to you, which ones are you taking?
Of the medications you listed, which ones are you taking?
Have you had to stop any of your medications for any reason?
How often do you not take medication X? (address each medication individually)
When was the last time you took medication X? (address each medication individually)
Have you noticed any adverse effects from your medications?

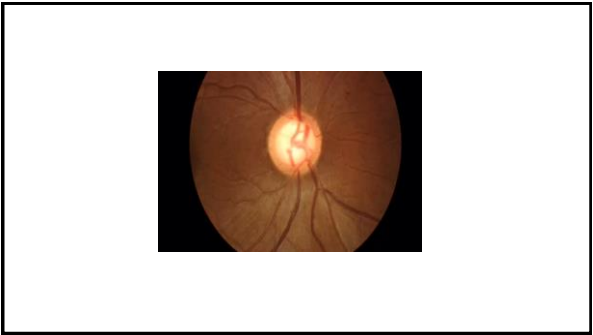
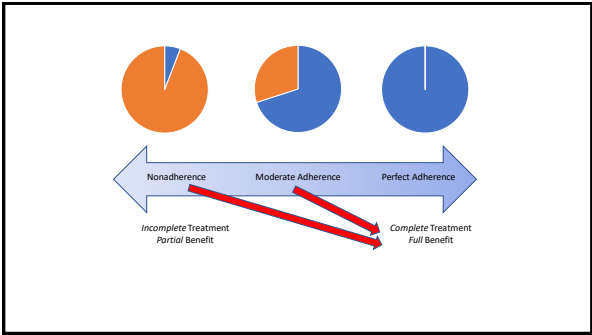
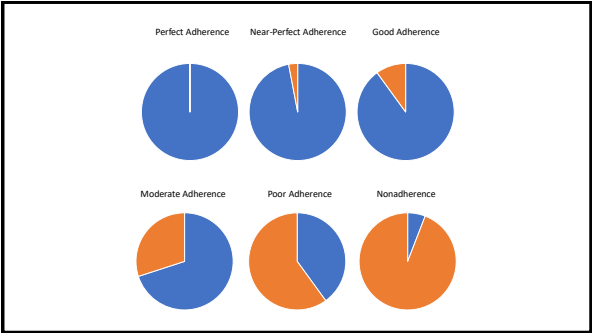
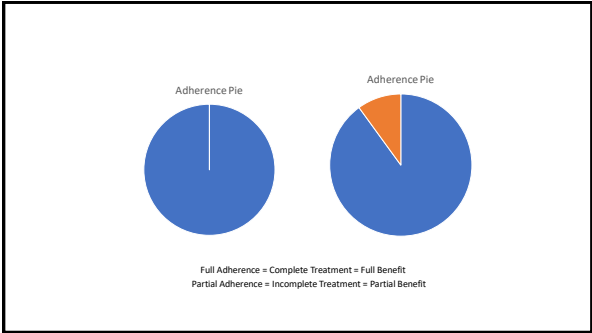
Brown MT, Bussell JK. Medication adherence: WHO cares?. *Mayo Clin Proc.* 2011;86(4):304-314. doi:10.4065/mcp.2010.0575

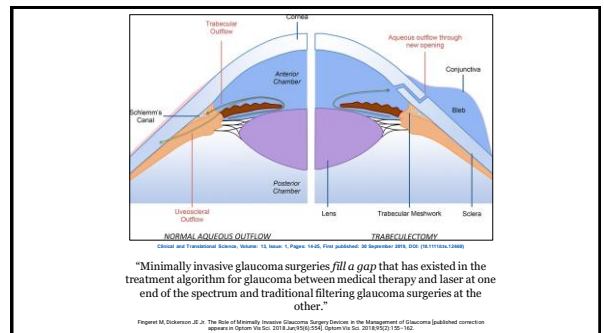
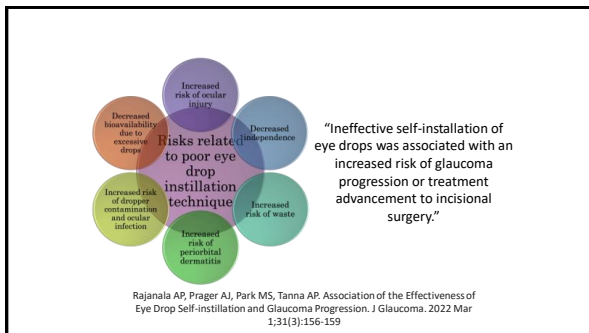
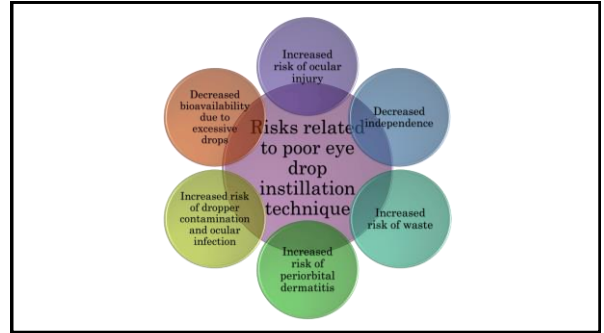
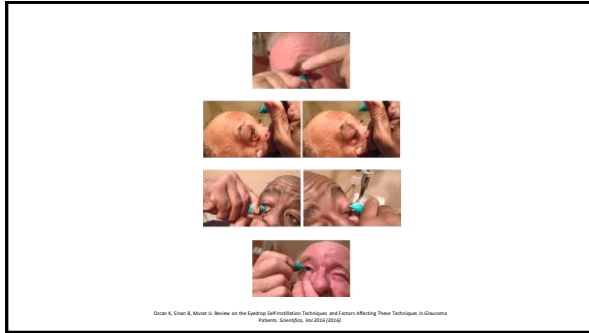
“Managing glaucoma...is influenced by a person's *perceived susceptibility* to the disease, the *perceived severity* of the disease, the *perceived benefits to treatment* and the *perceived barriers* to the recommended behavior change.”

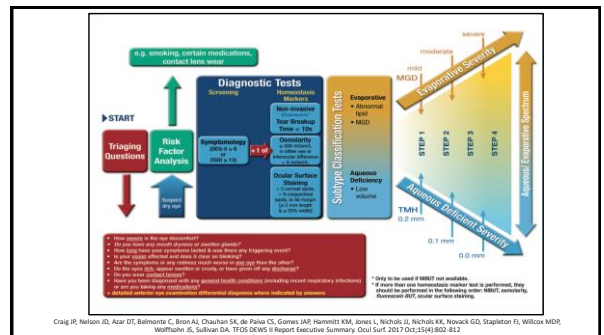
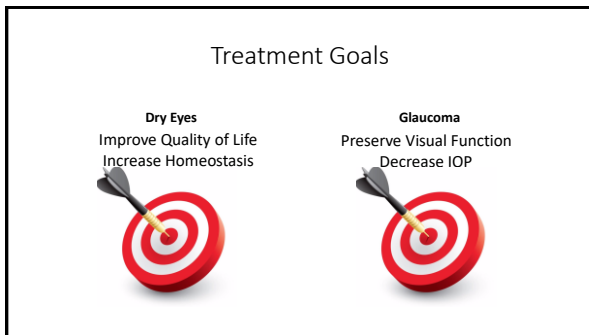
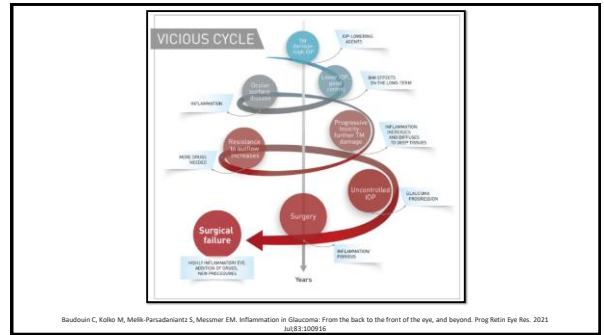
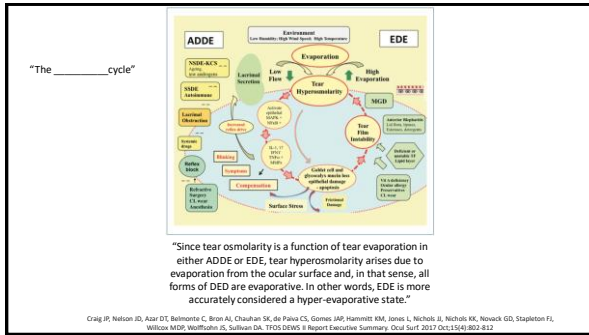
Newman-Costy PA, Steiner RM, Coleman AL, Hendricks L, Lee SP. Why Patients With Glaucoma Lose Vision: The Patient Perspective. *J Glaucoma.* 2016;25(7):e688-e675.

“For a glaucoma patient, this would mean that the person would only take their medication and return for their follow-up appointments *if* they believed that glaucoma would cause undesirable vision loss, the treatments offered by their doctor could mitigate this effect, and the barriers to following their physician's recommendation were not so difficult to overcome that they outweighed the perceived benefit of treatment.”

Newman-Costy PA, Steiner RM, Coleman AL, Hendricks L, Lee SP. Why Patients With Glaucoma Lose Vision: The Patient Perspective. *J Glaucoma.* 2016;25(7):e688-e675.









Better providers...Better care



Better providers...Better care

<u>Start</u>	<u>Stop</u>	<u>Continue</u>
<i>"What should we/I start doing?"</i>	<i>"What should we/I stop doing?"</i>	<i>"What should we/I continue doing?"</i>
List ideas/items: • Things that are not being done, but should be done • Things to begin doing to get better results • Things worth experimenting with for better results	List ideas/items: • Things that are not working or helping • Things that impede or are not practical • Not delivering desired results • We or others dislike	List ideas/items: • Things that are working well • Things that we want to keep • Worth continuing to see if they're worthwhile • We like or need

Questions??

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Instagram: [glaucomaqd](https://www.instagram.com/glaucomaqd)

Website: www.glaucomaqd.org

