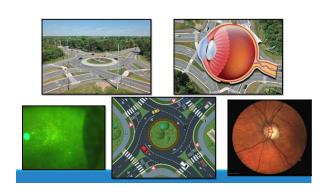




Glaucoma and Dry Eyes *"Frenemies" for Life*

OPTOMETRIC MANAGEMENT SYMPOSIUM 2022 CENTER FOR SIGHT & DRY EYE INSTITUTE AUSTIN LIFFERTH OD, FAAO DIPLOMATE, GLAUCOMA AAO NOVEMBER 4, 2022



Prevalence and Association

80 million people worldwide will have glaucoma this year.¹

Prevalence of DED in studies up to 75% in certain populations.²

5 million people in U.S. older than age 50 who have dry eyes, 11% have glaucoma.³

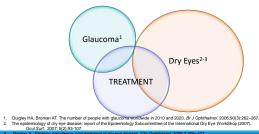
Nearly 60% of patients on topical glaucoma therapy have DED.³



Than YC, Li X, Wong TY, Quigley HA, Aung T, Cheng CY Global provisions of glascons and projections of glascons burden through 2040; a systematic review.
 Casq. JP, Nelson 3D, Aus TD, et al. 1706 DEWIS B Report Executive Summay. The Qualit Surface (2017), <u>Neurile Surface Surfa</u>

Prevalence and Association



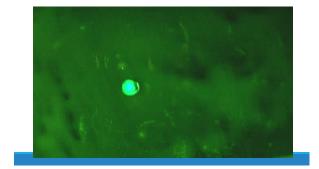






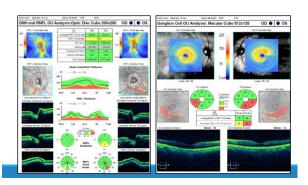


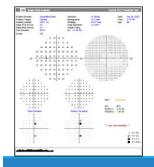


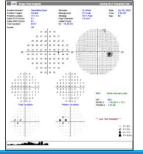




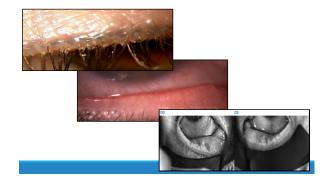


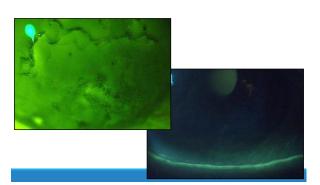


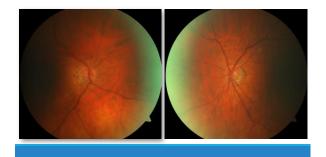


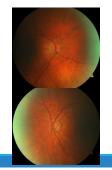


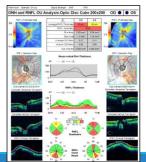


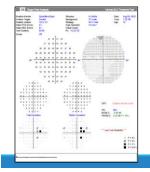


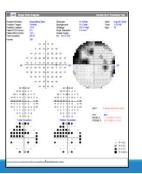














Glaucoma and Dry Eyes "Frenemies for Life"

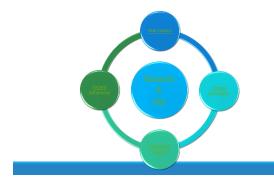
- To increase awareness of both chronic, progressive, and unyielding conditions.
- To increase awareness of the strong association between both chronic, progressive, and unyielding conditions.

 They love to be together.
 Inseparable and Inevitable
- To increase understanding of the similarities of both chronic, progressive, and unyielding conditions.
 They have a lot in common.
- To increase understanding of the collateral damage between both chronic, progressive, and unyielding conditions when together.

 They are worse when together.

 It does not happen overnight.





Risk Factors

MODIFIABLE AND NONMODIFIABLE

#1 Modifiable Risk Factor



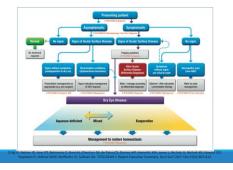


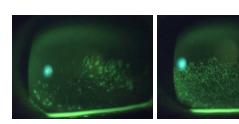
"The cycle"

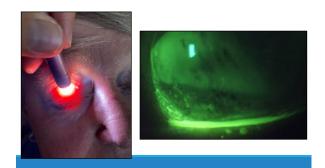


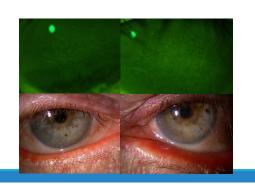
"Since tear osmolarity is a function of tear evaporation in either ADDE or EDE, tear hyperosmolarity arises due to evaporation from the ocular surface and, in that sense, all forms of EDE are evaporative. In other words, EDE is more accurately considered a hyper-evaporative state."

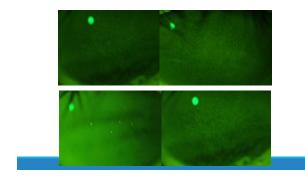
Craig JP, Nelson JD, Azar DT, Belmonte C, Bron AJ, Chauhan SK, de Paiva CS, Gomes JAP, Hammitt KM, Jones L, Nichols JJ, Nichols KK, Novack GD, Stapleton FJ, William MMP, Modificable IS, Sullians DA, TCOS DSMS II Report Supration Suprempts On a Surf 2017 Onto SERIA 1973, 913



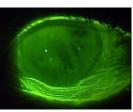


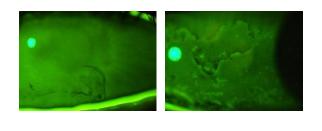


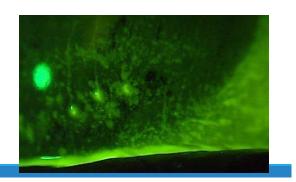


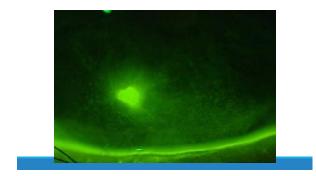


















Intraocular Pressure

IOP is the greatest modifiable risk factor in the development and progression of glaucoma^{1,2}.

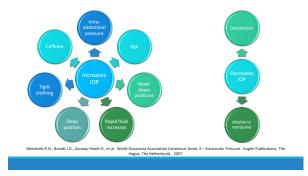
- Each mmHg of matters^{1,4}.

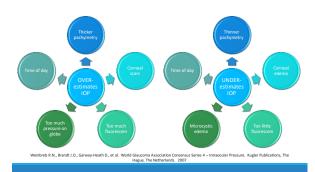
- Corneal biomechanical properties (central corneal thickness, corneal hysteresis) and applanation technique (decentration, thick/thin tear film, etc.) affect IOP measurement accuracy^{1,4}.

 $Thinner\ central\ corneal\ thickness\ and\ lower\ corneal\ hysteres is\ associated\ with\ the\ development\ and\ progression\ of\ glaucoma^7.$

Advanced glaucoma at time of diagnosis is associated with poorer prognosis and greater risk of progression^{8,9}. Poor adherence to prescribed topical medications is associated with visual field progression $^{10}.\,$

NON-MODIFIABLE MODIFIABLE O HEALTH LITERACY
O BMI
O DISEASE MITTIGATION (HTN,
DM, SLEEP APNEA)
LIFESTYLE
O NUTRITION IOP¹⁻⁷





Over-reliance on IOP levels?



"...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

"Kill the magic number"²

- IOP is not glaucoma.
- How would you monitor for glaucoma if we did not measure IOP?

Clinical Evaluation

PREFERENTIAL STRUCTURAL LOSS AND RED FLAGS

Preferential Loss

DRY EYES



QUESTIONS TO CONSIDER...

Do you have any family history of glaucoma? Have you ever had any eye injury or eye surgery? Do your eyes ever feel burning? Tired? Foreign body sensation?

Does your vision fluctuate after prolonged reading or other tasks?

Have you ever been diagnosed with dry eyes?

Do you currently use any artificial tears or prescription dry eye treatment?

SIGNS TO LOOK FOR...

Obstructed meibomian glands
Telangiectatic lid vessels
Frothy and/or low tear lake
Rapid TBUT

Corneal/conjunctival staining Endothelial pigment/keratic precipitates Anterior chamber reaction

Transillumination iris defects
Posterior synechiae

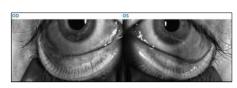
Narrow angles Elevated/asymmetric IOPs

Vertical neuroretinal rim thinning

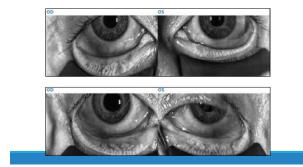


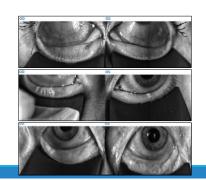


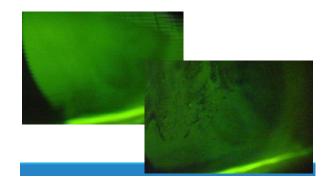


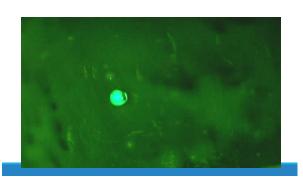








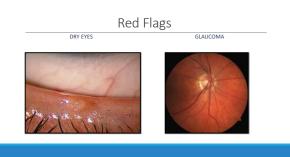






"Clinical diagnosis of glaucoma is predicated on the detection of a thinned retinal nerve fiber layer (RNFL) and narrowed neuroretinal rim. These features often are accompanied by deformation of the optic nerve head (ONH) (cupping). These features often appear first in the supero- or inferotemporal quadrants. Although these features are characteristic of POAG, it is important to exclude non-glaucomatous optic neuropathies. Detecting progressive glaucomatous RNFL thinning and neuroretinal rim narrowing are the best currently available gold standards for glaucoma diagnosis."





"...its presence should be an unfavorable prognostic event."1

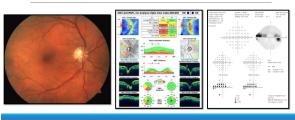


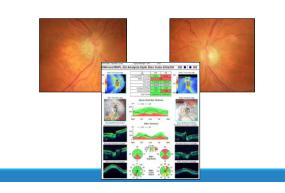
Increased risk of disc hemorrhages was comparable to...

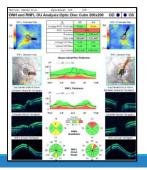
Increasing age by 10.5 years Increasing IOP by 11.4 mm Hg Decreasing CCT by 23.1 microns Worsening PSD by 1.3-dB Increasing vertical CDR by 0.1 unit.

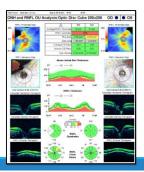


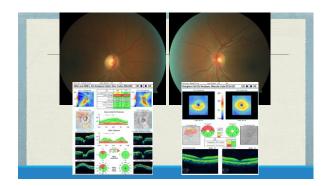
Preferential Loss
- Structural and Functional Correlation -

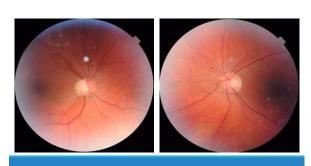


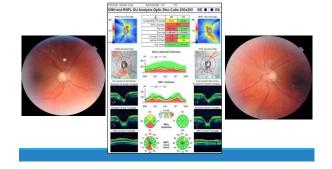


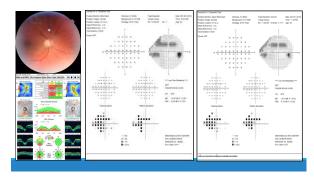


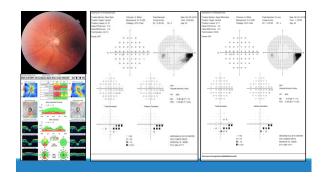


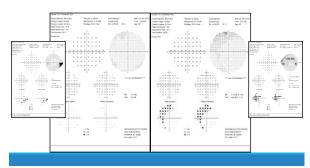


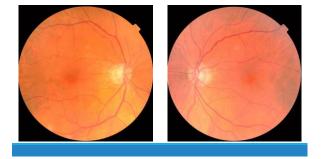


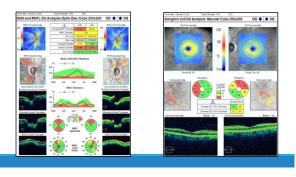


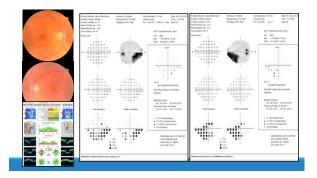


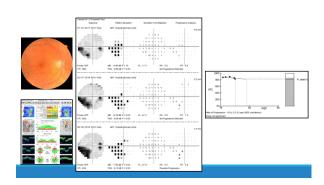


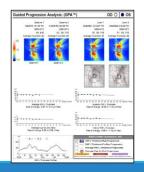


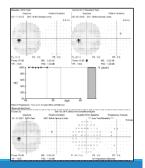


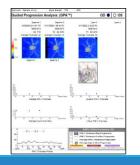


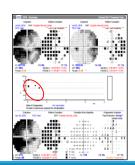












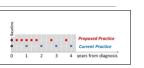
"Perform sufficient examinations to detect change."

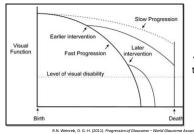
- "A good baseline of reliable VFs is essential to be able to monitor for progression."
- "Decisions on progression should not be made by comparing only the most recent field with the one hefore"
- "Suspected progression should be confirmed by repeating the field."1

1. B.N. Weimen, D. G. H. (2011). Proprising of Genomes - New College - According to the Contract Meeting Parks Right Publications.

Boodina T. Crabb DP. More Stropert more costly? Heart according aspects of motioning discovera patients in England. BMC Health Save Res. 2016;16(1):811.

Contract Contract





"Clinicians should aim to measure the rate of VF progression."

Weinreb, D. G.-H. (2011). Progression of Glaucoma – World Glaucoma Association 8th Consensus Meetir Paris: Kugler Publications.

Caprioli J. The importance of rates in glaucoma. Am J Ophthalmol.. Feb 2008;145(2):191–192.

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:
- 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 inferiortemporal VF



- 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"
- $^\circ$ "In general, event-based methods are used early in the follow-up, when few VFs are available for serial analysis.

Visual Field Progression

 "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."

Treatment Goals

HOME THERAPY VS IN-OFFICE THERAPY

ADHERENCE – INDEPENDENT OSMOLARITY REDUCTION

ADHERENCE – INDEPENDENT IOP REDUCTION

Treatment Goals

DRY EYES

Improve Quality of Life Increase Homeostasis



GLAUCOMA

Preserve Visual Function Decrease IOP



Treatment Principles

"The management of DED is complicated, due to its multifactorial etiology... This aspect of determining the major causative factors behind the DED is critical to appropriate management. The sufficience is of DED management is to restore the homeostasis of the ocular surface and tear film, through breaking the vidous cycle of the disease.

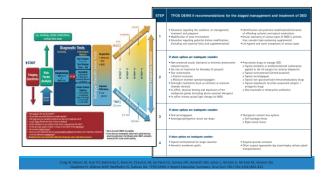
"Overall, the treatment of DED remains something of an ort, not easily lending itself to a rigid, evidence-based algorithm that accommodates all patients with DED symptoms signs. All eye care providers who treadge the significance of each of the varied pathogenic processes that may amaltest similar subjective compliants and similar signs of ocular surface dysfunction."

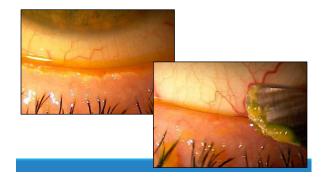
"In general, treatment is indicated for patients with glaucoma or glaucoma suspects who are at risk for patients of glaucoma suspects who are at risk for in vision-related quality of life from the disease. Treatment is generally indicated when the risks of progressive disease outweigh the risks and potential side effects of treatment.

All treatment decisions should take into account the presence of coesting outor conditions (50s). The patients life expectancy and general health status, or self-such as the processing outor and expectations about the treatment.

Treatment analic include IDP visual function and

Treatment goals include IOP, visual function and structural (optic disc,RNFL) outcomes and QOL."2









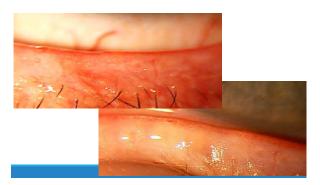


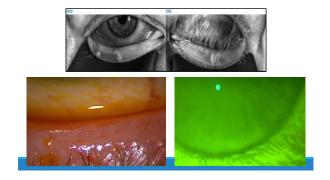


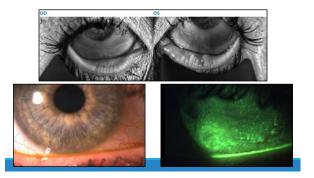












- Causes

 Health conditions

 Diabetes

 Auto-immune conditions

 Anthritis

 Spigren's Syndrome

 Medications

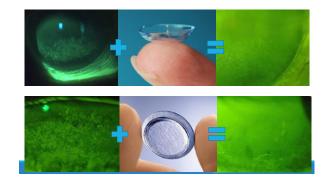
 Allegg, Blood pressure, Depression

Anti-inflammatory medications
• Eysuvis, Flarex, Lotemax etc
• Nutritional supplements

- Immunomodulator medications
 Cequa, Restasis, Xiidra, etc

Cequa, restasts, xarura, ecc
 Additional therapy
 Autologous serum (Ex: Vital tears)
 Oral secretagogues (Ex: pilocarpine and cevimeline)
 Neurostimulation (Ex: Tyrvaya, Tear 100)





Topical Therapy

DRY EYE



GLAUCOMA 56,000 Ways To Treat Glauce

Matters Part (India), visualization of sever glasscome medications at part fore years, visualizing seroual with once opposed efflower, has greatly expanded student flowarys for glasscome particles. Our new particles are searched to the student for the particles are searched to the student flowarys of the particles are searched to the student flowarys of the particles are searched to the particles are resulted to the particles are partic

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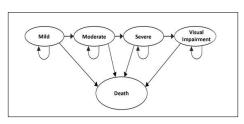
Realini, TonyFechtner, Robert D et al. 56,000 ways to treat glaucoma. Ophtholmology, Volume

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



Boodina T, Crabb DP. More frequent, more costly? Health economic modelling aspects of monitoring glaucoma patients in England. BMC Health San Alex. 2014;16(1):581.

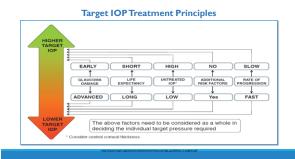
Target IOP Treatment Principles

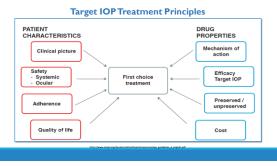
"The decision to initiate glaucoma treatment should be based on the assessment of the risks for development of functional impairment or decrease in vision-related quality of life, taking into account factors such as coexisting ocular conditions, the patient's life expectancy, and general health status,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

Age Stage

nann), Weinreb R, World Glaucoma A. Medical Treatment Of Glaucoma : The 7th Consensus Report Of the World Glaucoma Association. Amsterdam: Kugler Publications; 2010. p. 1.

Target IOP Treatment Principles Age Systemic concribitions Congoing medical and respiratory functions Congoing medical and response of the congoing medical and respiratory functions Congoing medical and response of the cong





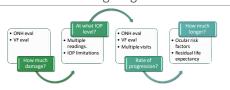


Target IOP Treatment Principles



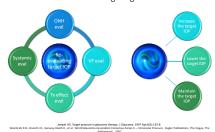


Evaluating Target IOP



Imagel Ho. 1 Earget pressure in glucoms therapy. J Glucoms. 1997 App.(5):1311-8.
Welchrich R.N., Brandt J.D., Garway-Heisth Welf of Glucoms Association Consensus Series 4 – Intracociar Pressure. Kingler Publications, The Hages, The Heistenich. 2007
Philip Research Discounter On Ind. 1 Gentlifered "Years" ("Part 1): 1997
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Re-evaluating Target IOP





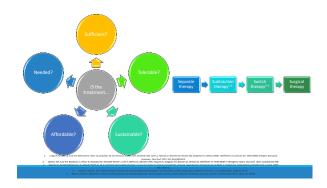
"In treated patients, failing to achieve target IOP was associated with more rapid VF worsening. Eyes with moderate glaucoma experienced the greatest VF worsening from failing to achieve target IOP."

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

Target IOP Treatment Principles

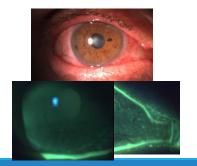
"In the end, it will be impossible to know if we overreacted or did too much, but it will be QUITE apparent if we under reacted or did too little."

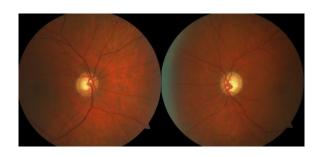
(Dr. Darrin M. Peppard March 20, 2020)

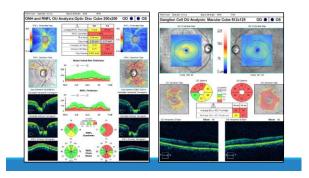


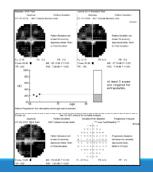


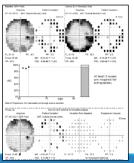


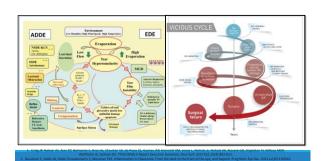


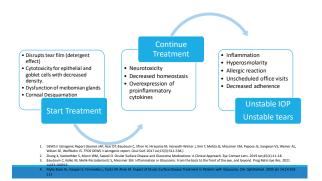














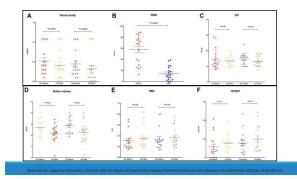


"Patients had a diagnosis of glaucoma for 9.82 ± 7.92 years and had been on antiglaucoma topical medication since then. The mean number of IOPlowering drugs used by the patients was 3.05 ± 0.91, among which 2.58 ±. 1.17 were BAK-preserved drops. The mean number of instilled drops was 5.21 ±1.90 per day." "All patients underwent a complete OSD treatment, consisting of eyelid hygiene using a gel twice a day, fluorometholone acetate 0.1% one drop at night, preservative-free lubricant every 2 hrs, oral free-acid supplementation (omega 3 and flaxseed oil capsule 2g a day) and oral tetracyclin derivate (doxycycline hydrochloride 100 milligram per day, during 30 days). Anti-glaucoma treatment was not modified, and a second appointment was scheduled for the same evaluation 1 to 3 months after treatment."

					Table 2 Comparison (Control Group)	between the tr		, m and story	aroup board oc	DEF SOFTEE TICES	THE REAL PROPERTY.	ay susjec
					Parameter	Pretreatment			Control			
						MeantSD	95% CI	Median	MeantSD	95% CI	Median	P value
					BCVA OD (LogMAR)	0.52±0.35	0.34-0.69	0.40	0.83±0.24	0.71-0.95	1	0.02
Table I OSD Class	fication A	ccordings	to Each Param	neter Studied	BCVA OS (LigMAR)	0.44±0.35	0.26-0.62	0.30	0.85±0.13	0.79-0.92	0.9	0.0018
	0.	t.	2.	3.	OSDI	57.86±25.63	45.51-70.21	68.18	11.64±12.69	5.11-18.16	10.41	<0.0001
	None	Mild	Moderate	Severe	TMH OD	0.46±0.65	0.12-0.79	0.25	0.24±0.03	022-026	0.24	0.82
CISIDI Score	<03	13-22	23-32	>33	TMH OS	0.4810.65	0.15-0.82	0.30	0.26±0.08	0.22-0.30	0.28	0.49
FBUT (seconds)	8-15	7-5	4-1	Immediate	Schirmer OD	16.76±11.33	10.94-22.59	14.00	14.72±9.57	9.96-19.48	11.00	0.65
Represent staining	0-1	2-4	5-9	10-15	Schirmer OS	17.82±10.11	12.62-23.02	20.00	16.28±8.89	11.85-20.70	14.50	0.58
	-	-	-		NITBUT OD	5.74±6.20	2.54-8.92	3.44	14.42±5.05	11.98-16.85	12.48	0.0002
Lissamine green	0-1	2-3	4-5	6-9	NITBUT OS	7.75±5.90	4.60-10.90	6.98	12.78±5.272	10.24-15.32	13.02	0.0063
_	-	_	_	-	FBUT OD	4.00±3.57	1.25-6.74	4.00	9.18±4.37	6.85-11.52	7.50	0.008
Schirmer test (mm)	>10	10-5	S+I	0	FBUT OS	1.77±1.20	0.85-2.70	4.00	9.87±4.91	7.25-12.49	9.50	0.0001
Meboscore	0	1	2	3	Bulbar Redness OD	2.70±0.72	2.31-3.08	2.60	1.55±0.35	1.38-1.72	1.50	<0.0001
Tear meriscus	>0.3	0.3-0.2	0.2-0.1	0.1-0	Bulbar Redness OS	2.84±0.69	2.47-3.21	2.90	1.57±0.40	1.378-1.769	1.50	<0.0001
height (mm)					Fluorescein OD	6.29±4.61	4.02-8.56	6.00	0.31±0.74	(-0.04)-0.67	0.0	<0.0001
Note: Criteria used for dry eye disease severity classification. Abbreviations: OSES: Onder Surface Disease Index: FBLIT Fluorescein Bresina.			Fluorescein OS	5.64±4.09	3.54-7.75	5.00	0.52±0.77	0.15-0.89	0.0	<0.0001		
Time mrs. rellimeters.	nder harbo	e Disease 9	stee; FBLIT, Fluo	rescen Brinkup	Lissamine OD	1.35±0.86	0.9099-1.796	1.00	0.78±1.31	0.15-1.42	0.0	0.03
					Lissamine OS	1.37±0.95	0.86-1.88	1.00	0.63±1.01	0.14-1.11	0.0	0.009
					Meiboscore	1.76±0.9	1.3-3.22	2.00	1±0.6	0.67-1.32	1.00	0.01
					Age	66.74±9.79	62.02-71.46	70.00	62.3±4.97	59.92-64.72	62.00	0.05

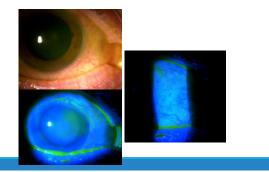
Mylla Boso AL, Gasperi E, Fernandes L, Costa VP, Alves M. Impact of Ocular Surface Disease Treatment in Patients

Parameter	Pre-tre	atment				Post-Tr	reatment					
	Mean	Median	PD	Min	Max	Mean	Median	PD	Min	Max	Mean A	Р
BCVA OD (LogMAR)	0.52	0.40	0.35	1.08	0.10	0.41	0.30	0.37	1.08	0.00	(-0.13)	0.0031
BCVA OS (LogMAR)	0.44	0.30	0.35	1.08	0.10	0.32	0.15	0.38	1.08	0.00	(-0.12)	0.0003
OSDI	57.86	68.18	25.63	12.50	88.88	14.60	11.36	12.13	0.00	38.90	(-43.3)	<0.0001
TMH OD	0.46	0.25	0.65	0.15	2.90	0.35	0.32	0.20	0.17	0.83	(-0.11)	0.2834
TMH OS	0.49	0.30	0.65	0.14	2.90	0.37	0.30	0.19	0.15	0.75	(-0.12)	0.7851
Schirmer OD	16.76	14.00	11.33	3.00	35.00	19.12	17.00	9.33	6.00	35.00	2.35	0.1228
Schirmer OS	17.82	20.00	10.11	2.00	35.00	17.53	14.00	10.63	4.00	35.00	(-0.29)	0.7505
NITBUT OD	5.74	3.44	6.20	0.00	20.84	7.69	7.75	4.84	0.00	19.12	1.87	0.2117
NITBUT OS	7.76	6.98	5.91	1.72	23.52	9.92	6.31	7.12	3.25	24.00	3.22	0.1742
FBUT OD	4.00	4.00	3.57	0.00	9.00	5.11	5.00	2.37	2.00	9.00	1.29	0.2894
FBUT OS	4.75	4.00	3.28	1.00	10.00	6.75	6.00	3.88	2.00	12.00	2.00	0.0797
Bulbar Redness OD	2.70	2.60	0.73	1.70	4.00	2.26	2.40	0.95	0.24	4.00	(-0.45)	0.0414
Bulbar Redness OS	2.84	2.90	0.70	1.90	4.00	2.18	2.25	0.76	0.15	3.40	(-0.64)	0.0196
Fluorescein OD	6.29	6.00	4.41	1.00	15.00	2.59	1.00	3.57	0.00	15.00	(-3.71)	<0.0001
Fluorescein OS	5.65	5.00	4.09	0.00	14.00	1.88	1.00	2.26	0.00	7.00	(-3.76)	<0.0001
Lissamine OD	1.35	1.00	0.86	0.00	3.00	0.94	1.00	0.65	0.00	2.00	0.13	0.131
Lissamine OS	1.37	1.00	0.95	1.00	3.00	0.88	1.00	0.99	0.00	3.00	0.10	0.101
IOP OD	14.65	12.00	6.23	8.00	29.00	13.44	13.00	3.16	9.00	20.00	(-1.38)	0.9471
IOP OS	14.65	16.00	3.60	10.00	24.00	13.06	13.00	3.07	8.00	18.00	(-1.59)	0.0510



How would you treat the patient if it was...you?

MMT MTMT MRMT



Less IS More

Less is:

- More understanding
- · Less confusion...drop sheets?
- More adherence
- More disease control
- More comfort
- Decreased side effects
- Decreased office visits
- More quality of life
- More ocular surface health
 Decreased ocular surface disease





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

Patient Adherence

EDUCATION AND ADHERENCE

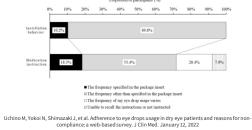
Treatment Goals – Adherence

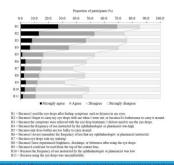
DRY EYES

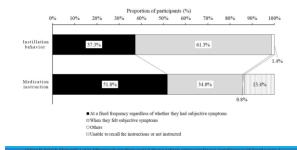












"We showed that most participants with DED did not instill the DED eye drops at the specified frequency..."

Decreased Tear Quality

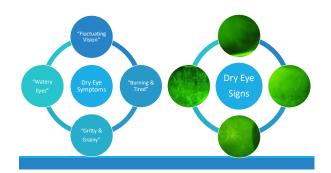
Decreased Tear Quantity

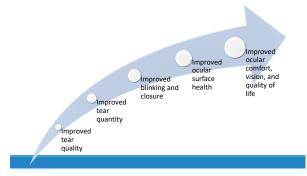
Decreased Blinking/Eyelid Closure

Decreased Surface Health

Decreased Comfort, Vision, Quality of Life

33



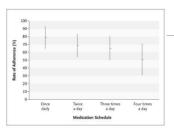


"Patient adherence is the wild card in the deck for controlling glaucoma progression."



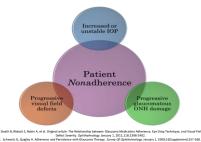
Fingeret M, Dickerson J. The Role of Minimally Invasive Glaucoma Surgery Devices in the Management of Glaucoma.

Optomator And Union Sciences Official Bublication Of The American Academy Of Cetemator Legical police. Solvens

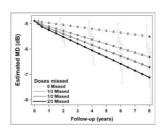




Osterberg L, Blaschke T. Adherence to medication. N Engl J Med. 2005 Aug







"To increase the effectiveness of our current glaucoma treatments, there is acritical need to focus on helping support patients in improving their glaucoma medication adherence."

Newman-Casey PA, Niziol LM, Gillespie BW, Janz NK, Lichter PR, Musch DC. The Association betw

"...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."

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

"Physician attitude has been shown to play a large role in patient adherence..."

Budenz D. A Clinician's Guide to the Assessment and Management of Nonadherence in Glaucoma.

"...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 Glaucomo

I know it must be difficult to take all your medications regularly. How often do you miss taking them?³

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)

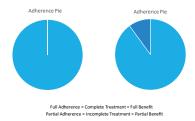
Have you noticed any adverse effects from your medications?

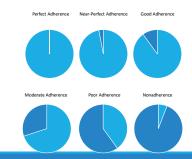
Brown MT, Bussell JK. Medication adherence: WHO cares?. *Mayo Clin Proc.* 2011;86(4):304-314, doi:10.4065/mcp.2010.0575 "Managing glaucoma...is influenced by a person's perceived susceptibility to the disease, the perceived severity of the disease, the perceived benefits to treatment and the perceived barriers to the recommended behavior change."

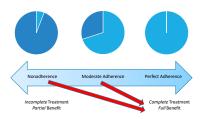
Newman-Casey PA, Shtein RM, Coleman AL, Herndon L, Lee PP. Why Patients With Glaucoma Lose Vision: The Patient Perspective. J Glaucoma. 2016;25(7):e668-e675. "For a glaucoma patient, this would mean that the person would only take their medication and return for their follow-up appointments if they believed 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-Casey PA, Shtein RM, Coleman AL, Herndon L, Lee PP. Why Patients With Glaucoma Lose Vision The Patient Perspective. J Glaucoma. 2016;25(7):e668-e675. "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 Glaucom Ophtholmology January 1, 2009;116:S43-S47.









What are some barriers to adherence?

Provider Factors

- °Dissatisfaction: "I quit taking my drops because I was dissatisfied with my doctor's care."
- Communication: "I stopped taking my drops because I didn't understand initially that I need to take them forever."



Tsai JC, McClure CA, Ramos SE, Schlundt DG, Pichert JW. Compliance barriers in glaucoma: a systematic classification

What are some barriers to adherence?

Situational/environmental Factors

- Major life events: "Two years ago when my wife died I had a hard time taking my drops."
- Travel/away from home: "When I am on vacation it is more difficult to take my drops"
- ° Competing Activities: "I miss my drops on Sunday mornings when I go to church.

Tsai JC, McClure CA, Ramos SE, Schlundt DG, Pichert JW. Compliance barriers in glaucoma: a systematic classification. J Glaucomo. 2003;12(5): 393-398.

What are some barriers to adherence?

"Major reasons cited for non-adherence include:

medication side effects (both local and systemic),

the inability to notice a visual benefit from the medications (at least short term),

and difficulty in administering the medication."



What are some barriers to adherence?

Table 1. Barriers to Glaucoma Medication Adherence					
Barriers to Glaucoma Medication Adherence	Literature Sources				
Beliefs about glaucoma, skepticism that glaucoma will cause vision loss	Friedman et al (2008), ²² Lucey et al (2009), ²³ Tsai et al (2003), ²² Slexth 2010 ²⁶				
beliefs about glaucotus medications, skepticism that glaucotus medications will mitigate vision loss Foot self-efficacy	Friedman et al (2008), ³³ Lucey et al (2009), ³³ Strykov et al (2010), ³³ Tsui et al (2003), ³³ Steath et al (2010), ³³ Skoth et al (2012), ³⁴ Steath et al (2010), ³⁴				
Foor knowledge about gloucoma	Friedman et al (2008), ³² Lacey et al (2009), ³³ Snyker et al (2010), ¹⁴ Lunnela et al (2010), ³⁵ Tsai et al (2003) ³⁷				
Mismst of physician Difficulty with eye drop administration	Stryker et al. (2010), ¹⁵ Lunnela et al. (2010), ¹⁶ Tsai et al. (2003), ¹⁷ Taylor et al. (2002), ¹⁸ Luces et al. (2009), ¹⁷ Tsai et al. (2003), ¹⁷ Tsalor et al. (2010), ¹⁸ Sleath et al. (2010), ¹⁸				
Medication cost	Friedman et al (2008). Tun et al (2003). Turior et al (2002).				
Medication-induced side effects	Friedman et al (2008), 7 Tsai et al (2003), 7 Taylor et al (2002) 7				
Forgetfylness	Lacev et al (2009),13 Stroker et al (2010),23 Tsai et al (2003),12 Taxlor et al (2002)33				
Difficulties with the medication schedule	Lacey et al. (2009), 17 Soi et al. (2003), 17 Taylor et al. (2002) 17				
Life stress	Hall (2014), 7 Keetchy (2014), 2 Cohen et al (1983) 7				

What are some barriers to adherence?

Barter'	Univariate Analysis, Olds Bario (99% Confidence Interval)	P Valer	Biogriste Analysis, Odds Ratio (95% Confidence Interval)	P Value
Officially with shop administration here self-efficient have been self-efficient fallets about administrative that stood from the self-efficient shot officers that officers Officially with the medication wheelide disease of physician here of been self-efficient here of physician he of the rison.	2.3 (10-4.5) 47 (23-9.7) 1.5 (03-2.9) 1.0 (03-2.1) 0.9 (03-1.1) 0.9 (03-1.9) 2.2 (13-4.4) 3.5 (28-11.4) 1.8 (09-5.6) 2.8 (14-6.7) 1.0 (03-2.7) 1.0 (03-2.7)	0.04 <0.0001 0.3 0.9 0.01 <0.0001 0.01 0.1 0.000 0.9 0.000	2.5 (1.1—4.9) 4.7 (2.3—9.7) 1.4 (0.7—2.8) 1.1 (0.5—2.8) 1.0 (0.5—2.9) 1.8 (0.0—8.9) 5.7 (2.6—2.1) 1.9 (0.0—4.9) 1.0 (0.6—3.9) 0.9 (0.4—4.9) 1.1 (1.0—2.9) 1.1 (1.0—2.9)	0.25 -0.000 0.4 0.5 0.9 0.1 -0.000 0.28 0.2 0.2 0.2 0.2 0.2 0.2 0.2 0.2 0.2 0.2
*Compared with subjects who did not nepo Adjusted for age. *Adjusted for age with case vision *Supricion that gluccom well case vision *Supricion that discount medication will	lon.			

Newman-Casey P, Robin A, Lee P, et al. Original article: The Most Common Bansiers to Glaucoma Medication Adherence. A Cross-Sectional Survey. Ophthalmology. July 1 2015;122:1308-1316.

What can we do to overcome barriers to adherence?



What can we do to overcome barriers to adherence?

the 3. Strategies for Improving Adher

identify poor adherence Look for markers of nonadherence: missed appointments ("no-shows"), lack of response to medication, missed refile.

frontational Emphasize the value of the regimen and the effect of

Elicit patient's feelings about his or her ability to follow the regimen, and if necessary, design supports to promote adherence

Provide simple, clear instructions and simplify the reg imen as much as possible

Encourage the use of a medication-taking system Listen to the patient, and customize the regimen in

accordance with the patient's wishes

Obtain the help from family members, friends, and community services when needed

priate
Consider more "forgiving" medications when ad
ence appears unfilely!
Medications with lone half-lives

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

Table 1. Summary of Strategies to Address Adherence Issues

- Be proactive. Assume most putients are nonadherent.
 Use tailored, patient-centered approaches.
 Learn which interventions work best in your practice, with y
- Use a combination of methods.
- Specific

 Actively address at-risk patients.
- Simplify and optimize treatment regimens when possible polypharmacy, side effects).
- Neduce drugs costs when possitie.
 Undentand the patient's health beliefs abou
 Use patient education.
- Use verbal and written delivery.
 Adapt information to those with poor vision or low
 Make use of office staff.
- Review drug administration at each visit.
 Suggest that parients keep a medication diary, which will be
- Use telephone or mail reminders when possible.
 Suggest the patient incorporate drops into daily activities.
 Involve a helpful construct family member to main with
- Involve a helpful caregiver/family member to ass applying drops or reminding to take drops.
 Be supportive. Use open communication.
- Be supportive. Use open communication.
 Ask-tell-ask dialog.
 Motivational interviewing.
- Stages of readiness for change.

 Be resoured to not as conclusionated and



Buders D. A.Clinician's Guide to the Assessment and Management of Nonadherence in Glaucoma. Ophthalmology, January 1

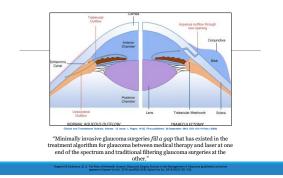




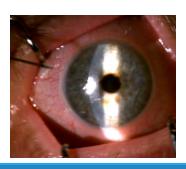
Oscan K, Sman B, Murat U. Review on the Eyedrop Self-instillation Techniques and Factory Affecting These Techniques in Glaucom Patients, Scientifica, Vol 2016 (2016).













Glaucoma and Dry Eyes "Frenemies for Life"

- To increase awareness of both chronic, progressive, and unyielding conditions.
- To increase awareness of the strong association between both chronic, progressive, and unyielding conditions.

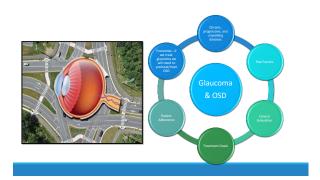
 They love to be together.

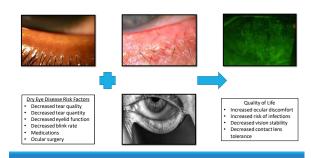
 Inseparable and Inevitable
- To increase understanding of the similarities of both chronic, progressive, and unyielding conditions.

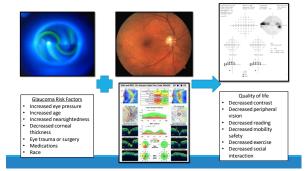
 They have a lot in common.
- To increase understanding of the collateral damage between both chronic, progressive, and unyielding conditions when together.

 They are worse when together.

 It does not happen overnight.







The Science and Art of Managing Glaucoma...and Dry Eyes



Better providers...Better care

"Positive Deviance (PD) is based on the observation that in every community there are certain individuals or groups whose uncommon behaviors and strategies enable them to find better solutions to problems than their peers, while having access to the same resources and facing similar or worse challenges."



