

On behalf of Vision Expo, we sincerely thank you for being with us this year.

Vision Expo Has Gone Green!

We have eliminated all paper session evaluation forms. Please be sure to complete your electronic session evaluations online when you login to request your CE Letter for each course you attended! Your feedback is important to us as our Education Planning Committee considers content and speakers for future meetings to provide you with the best education possible.



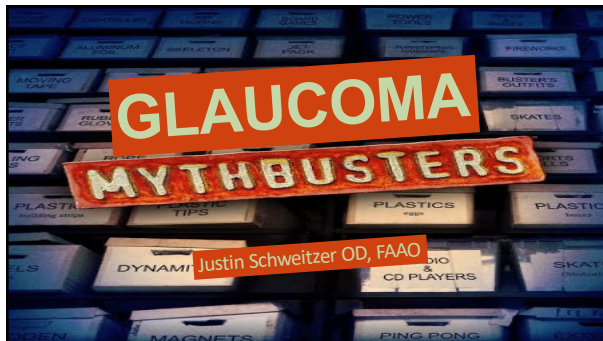
1

Financial Disclosure – Justin Schweitzer, OD, FAAO

- Alton - C/L
- Aldeyra - C
- Allergan - C/L
- Bausch + Lomb - C/L
- Bruder - C
- Sight Sciences - C/L
- Dompe - C/L
- Zeiss - C/L
- Visus - C
- Science Based Health - C
- Tarius - C/L
- Santen - C
- Sun - C/L
- Reichert - C
- Glaukos - C/L
- MedPrint - C
- LVC - C/L
- Avellino - C
- heart bio - C
- Ocuphire - C
- Viatrix - C
- Thea - C
- Heru - C
- Eyenovia - C

All relevant relationships have been mitigated

2



3



Myth 1 ?

“Controlled Glaucoma”
Is
A
Stable
Visual Field
And
Stable
OCT?

4

Redefining “Controlled Glaucoma”



IOP/VF/ON + Quality of life
Fluctuating vision
Headache, fatigue
Can't remember the bottle color

“Not Controlled”

Slide Courtesy of Paul Singh MD

5

The Burden of Caring for and Treating Glaucoma
The Patient Perspective




Quality-of-Life Consequences of Glaucoma Treatment Burden

1. Use of Time
2. Role and Activity Limitations
3. Financial Difficulties
4. Emotional Stress
5. Side Effects
6. Strain on Family and Friends

Stagg BC, Granger A, Gaetterman TC, Hess R, Lee PP. The Burden of Caring for and Treating Glaucoma: The Patient Perspective. Ophthalmol Glaucoma. 2022 Jan-Feb;5(1):32-39. doi: 10.1016/j.ogla.2021.04.011. Epub 2021 May 10. PMID: 33984555.

6

Impact of Multiple Glaucoma Medications on Dry Eye Disease

Number of Drops	Incidence of DED among 61 glaucoma patients ¹	Incidence of DED among 19,665 glaucoma patients ²
1 	11%	51%
2 	39%	55%
3+ 	40%	60%

1. Fakhour RD et al. Cornea. 2010;29:1418-1421. 2. Shi C et al. Graefes Arch Clin Exp Ophthalmol. 2008;46:1593-1601. 3. Leung EW et al. J Glaucoma. 2008;17:350-355.

Slide Courtesy of Paul Singh MD

7


Effects on Meibomian Glands

Effect on lids/meibomian glands
 Study on glaucoma patients 18mo stable treatment with different medications.
 Reduced number of meibomian glands
 Reduced numbers of acinae and increased dysfunction in patients
 Patients on multiple medications with preservatives = increased dysfunction and reduced number of acinae

	OSR score	BSR	IT	Control staining	Meibin score	Meibin score
Control	4.5±0.8*	12.6±1.1*	18.8±0.9*	9.7±1.4*	0.15±0.05*	1.36±0.31*
Group 1	16.8±1.9***	7.2±0.9**	9.2±0.7**	1.9±0.3	0.48±0.12	2.05±0.51
Group 2	16.3±1.9***	7.1±0.9**	9.1±0.7**	1.9±0.3	0.48±0.12	1.95±0.48
Preserved Group	16.3±1.9***	8.1±0.9***	9.6±0.7***	1.9±0.3	0.53±0.16	2.15±0.55
Group 1	16.8±1.9***	4.1±0.5**	7.1±0.5*	2.3±0.1	0.75±0.18	2.25±0.55
Group 2	16.8±1.9***	3.9±0.5**	6.9±0.5*	2.3±0.1	0.75±0.18	2.25±0.55

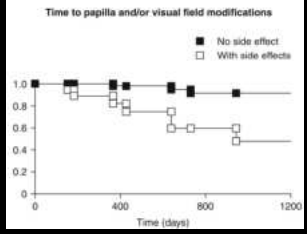
*p<0.05 vs group 1, 2 and 3. **p<0.05 vs group 1 and 2. ***p<0.05 vs group 1, preserved group and vs group 2 and 3. BSR: mean±SD (SD) (range) (n=10) (preservative free IT), (n=10) (control).

Agarwal L, et al. Br J Ophthalmol 2013.



8

Side Effects Lead to Progression



* Davis, Philippe, et al. Medical outcomes of glaucoma therapy from a nationwide representative survey

9

Solutions?

Preservative-Free Formulations
SLT
Glaucoma Drug Delivery
Minimally Invasive Glaucoma Surgery

10

Preservative-Free Formulations

N=349, Significant improvement in both signs and symptoms of OSD with switch to PF meds

Table 4 Frequency of symptoms and signs at visits 1 and 2 in PF group

	Visit 1 (preserved)		Visit 2 (preservative free)		p-Value
	N ^a	(%)	N ^a	(%)	
Patient symptoms					
Discomfort upon instillation	186/245	77.4%	40/243	17.7%	<0.001
Patients presenting with at least one symptom between instillations	283/343	82.2%	123/244	50.8%	<0.001
Objective signs found at the clinical examination (patients presenting with at least one)					
Pupillary sign (Dyspupils)	123/242	51.7%	30/246	14.3%	<0.001
Conjunctival sign	232/238	98.9%	76/238	31.9%	<0.001
Superficial punctate keratitis	43/238	18.4%	18/237	7.6%	<0.001

^aNumber of patients for which the variable had been recorded

Rosta, P.J., P. Rouiquin, and C. Baudouin. Prevalence of ocular symptoms and signs with preserved and preservative free glaucoma medication

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Preservative-Free Solutions

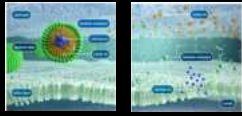
PF-Latanoprost

	Phase 3 (US) Trial (n=325)		Phase 3 (Europe) Trial (n=353)	
	PF-Latanoprost	Xalatan	PF-Latanoprost	Xalatan
Mean baseline IOP ± SD (mmHg)	18.8 ± 2.9	19.2 ± 3.1	24.1 ± 1.8	24.0 ± 1.7
Mean IOP reduction from baseline (mmHg) (range)	2.7 (2.2 - 3.0)	3.4 (2.9 - 3.8)	8.6 (8.3 - 8.8)	8.9 (8.8 - 9.0)


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BAK-Free Latanoprost

- Following instillation, micelles mix with the tear film
- As the micelles migrate toward the ocular surface, they break apart, releasing latanoprost



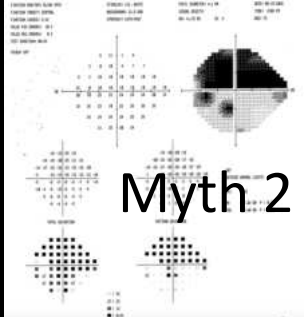
Preservative-Free



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Myth 2 ?

Surgical Intervention Should Be Delayed For As Long As Possible?

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Visual Field Damage and QOL^{1,2}

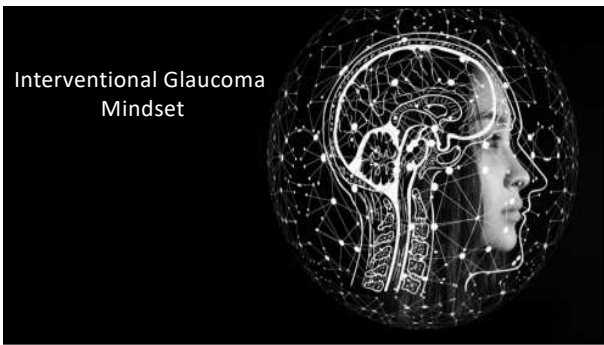


"At worse levels of VF damage, glaucoma patients demonstrate shorter, more fragmented bouts of physical activity throughout the day and lower activity levels during typical waking hours, reflecting low physiologic functioning"



1. Jari H, E. Javerik A, Schack, Aleksandra M, Pollock, David A, Wang J, Wang, Shella K, West, David S, Friedman, Laura N, Giffy, Tianping U, Pradeep Y, Ramulu. Patterns of Daily Physical Activity across the Spectrum of Visual Field Damage in Glaucoma Patients. *Ophthalmology*, Volume 128, Issue 1, 2021, Pages 70-77, DOI:10.1016/j.ophtha.2020.06.013.
 2. Ratti, R, Casanova, L, Riva, F, et al. Visual field loss and vision-related quality of life in the Italian Primary Open Angle Glaucoma Study. *Sci Rep* 8, 619 (2018). <https://doi.org/10.1038/s41598-017-19113-z>

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Safety Profile of MIGS

	ibent	ibent Hyflex	Hyflex	KDB	Treectome	GATT	TRAB360	VISCOS60	OMNI System	ASIC
IOP spikes	1.8-22.2	1.08-18.8	1.9-8.48	1.0-18.3	2.06-28.8	0-18.7	1.2	0.9-1.1	3.7	0-22.2
Hypotonia	1.85-11.4	0-0	1.02-0.45	0-34.9	4.72-05	0.07-38	50.6	1-13.1	3.7	1.0-20
Corneal oedema	2.1-6.87	0-10	0-3.23	1.0-15.5	0	0	0.2	0	4.9	0
Blat flooding	N/A	N/A	N/A	N/A	N/A	N/A	N/A	N/A	N/A	N/A
Device obstruction	0-13.2	0-6.2	0-4.2	N/A	N/A	N/A	N/A	N/A	N/A	N/A
ICH	1.9-2.27	0	0	0	1.47	0	0	1	0	0

1. Wood, K, Gedde, S. Safety profile of minimally invasive glaucoma surgery. *Curr Opin Ophthalmol*. 2021 Mar 13(2):160-168. doi: 10.1097/ICU.0000000000000791. PMID: 3313724.
 2. Rowson, A.C., Hogarty, D.T., Maher, D., Liu, L. Minimally Invasive Glaucoma Surgery: Safety of Individual Devices. *J. Clin. Med.* 2022, 11, 6833. <https://doi.org/10.3390/jcm11226833>

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Schlemm's Canal/TM Procedures

	Stents	SC Dilation	TM Cutting
Fibrosis Risk	(-)	(+)/(-)	(+)(+)
Hyphema	(-)	(+)/(-)	(+)(+)
PAS Risk	(-)	(-)	(+)
IOP Lowering	(+)	(+)	(+)(+)
Data	(+)(+)(+)	(+)/(-)	(+)(+)

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Trabecular Microbypass Stent (Stent Inject W) Schlemm Canal Microstent (Hydrus)



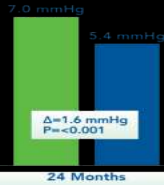
20

Trabecular MicroBypass X 2 PIVOTAL TRIAL

PRIMARY ENDPOINT:
≥ 20% Reduction in Unmedicated DIOP



SECONDARY ENDPOINT:
Mean Unmedicated DIOP Reduction



Shenolikov TM, Sridharan SR, Luboff DM, Stone MC, Quill TE, Kozlov EA, Golezovskaya G, Hertzberg BM, Katz SE. Short-Term Study Group. Prospective, Randomized, Controlled Pivotal Trial of an Ab Interno Implanted Trabecular Micro-Bypass in Primary Open-Angle Glaucoma and Cornea: Two-Year Results. Ophthalmology. 2019 Jun;126(6):1812-1821. doi: 10.1016/j.ophtha.2019.01.006. Epub 2019 Mar 14. PMID: 30880358.

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HORIZON Trial – 5 Year

	Stent + Cataract (n=369)	Cataract Only (n=187)
Change in diurnal IOP (mean)	-8.3 mm HG (+/-3.8)	-6.5 mm HG (+/-4.0)
60 months medication free	66%	46%
60 months mean IOP (mm Hg)	16.6 (+/-3.2)	17.6 (+/-3.6)
1 preoperative med	52.6%	54%
2 to 4 preoperative med	47.4%	46%

Arnold M, De Francesco T, Rhee S, McCabe C, Powers R, Gussard S, Sarrafian TW, Singh K. HORIZON investigators. Long-term outcomes from the HORIZON randomized trial for a minimally-invasive procedure in combination cataract and glaucoma surgery. *Ophthalmology*. 2022 Feb 23;131(2):422-432.

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Trabecular MicroBypass X 2 PIVOTAL TRIAL¹

Conclusions: The overall safety profile of the treatment group was favorable and similar to that in the control group throughout the 2-year follow-up.

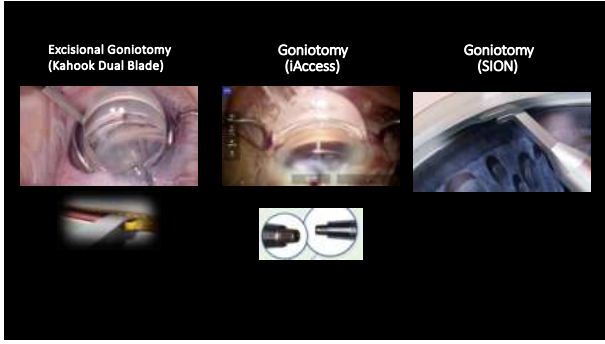
HORIZON Trial – 5 Year²

Conclusions: The addition of a Schlemm's canal microstent in conjunction with CS was safe, resulted in lowered IOP and medication use, and reduced the need for postoperative incisional glaucoma filtration surgery compared with CS after 5 years. Long-term presence of the implant did not affect the corneal endothelium adversely.

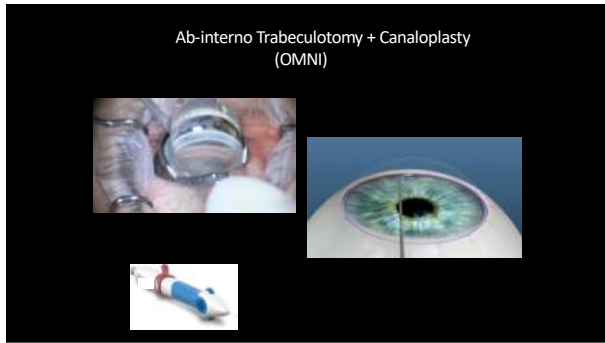
1. Sarrafian TW, Sarkissian SB Jr, Lubans DM, Schar MZ, Doh Y, Rama EA, Gnanapavan R, Warrick DM, Koo LS. Trabecular MicroBypass X2. Prospective, Randomized, Controlled Pivotal Trial of an Ab Interno Implanted Trabecular MicroBypass System Using Schlemm's Canal. *Ophthalmology*. 2019 Jun;126(6):1213-1224. Epub 2018 Mar 22. PMID: 29582255

2. Arnold M, De Francesco T, Rhee S, McCabe C, Powers R, Gussard S, Sarrafian TW, Singh K. HORIZON investigators. Long-term outcomes from the HORIZON randomized trial for a Schlemm's canal microstent in combination cataract and glaucoma surgery. *Ophthalmology*. 2022 Feb 23;131(2):422-432.

23



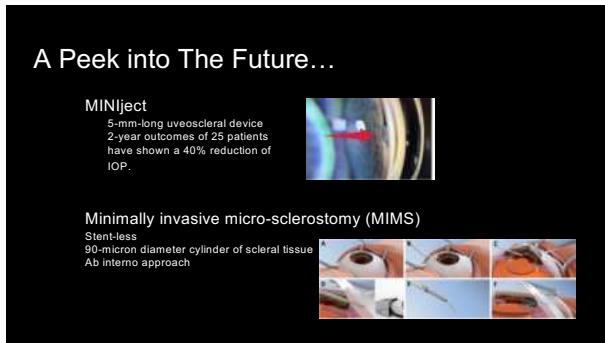
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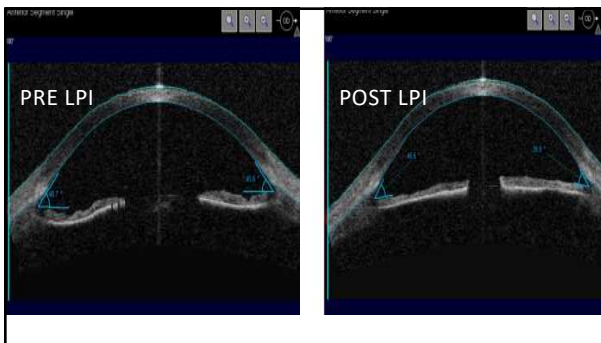
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Myth 3 ?

A close-up photograph of a clear ice cube with a silver pen nib resting on its surface, set against a black background.

Anterior Segment OCT Can Replace Gonioscopy When Analyzing The Angle?

29



31

Randomised Controlled Trial | Lancet. 2019 Apr 20;393(10181):1609-1616.
doi: 10.1016/S0140-6736(18)32607-2. Epub 2019 Mar 14.

Laser peripheral iridotomy for the prevention of angle closure: a single-centre, randomised controlled trial

Mingqiang He ¹, Yuchen Jiang ², Shengsong Huang ³, Dilyi S Chang ⁴, Beatriz Munoz ⁵,
Tui Aung ⁶, Paul J Foster ⁶, David S Friedman ⁶

889 subjects (Treated eye=LPI Untreated eye = control) – followed for 6 years

Criteria for Primary Angle Closure Suspect

1. No PAS present in any quadrant
2. No visible TM 6 clock hours or greater

Criteria considered as not preventative

1. IOP > or equal to 24 (measured twice)
2. PAS covering 1 or greater clock hours
3. Acute Angle Closure

4.19/1,000 – LPI arm
7.97/1000 – control arm
47% reduction

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Randomised Controlled Trial | Lancet. 2019 Apr 20;393(10181):1609-1616.
doi: 10.1016/S0140-6736(18)32607-2. Epub 2019 Mar 14.

Laser peripheral iridotomy for the prevention of angle closure: a single-centre, randomised controlled trial

Mingqiang He ¹, Yuchen Jiang ², Shengsong Huang ³, Dilyi S Chang ⁴, Beatriz Munoz ⁵,
Tui Aung ⁶, Paul J Foster ⁶, David S Friedman ⁶

Interpretation: Incidence of angle-closure disease was very low among individuals classified as primary angle closure suspects identified through community-based screening. Laser peripheral iridotomy had a modest, albeit significant, prophylactic effect. In view of the low incidence rate of outcomes that have no immediate threat to vision, the benefit of prophylactic laser peripheral iridotomy is limited; therefore, widespread prophylactic laser peripheral iridotomy for primary angle-closure suspects is not recommended.

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Diagnostic accuracy of AS-OCT vs gonioscopy for detecting angle closure: a systematic review and meta-analysis

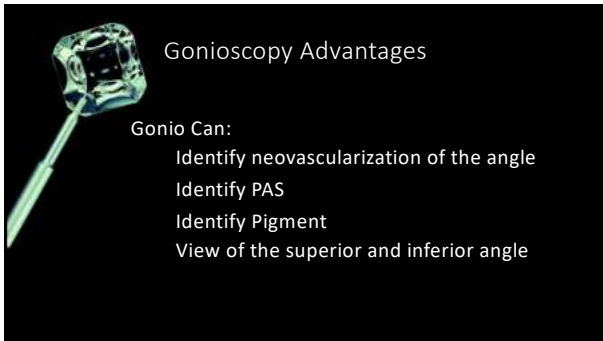
Thomas Desmond^{1,2} · Vincent Tran² · Monish Maharaj^{1,4} · Nicole Carnit^{1,2,5,6} · Andrew White^{1,2,3}

Received: 12 January 2021 / Revised: 13 May 2021 / Accepted: 3 June 2021 / Published online: 5 July 2021
© The Author(s), under exclusive licence to Springer-Verlag GmbH Germany, part of Springer Nature 2021

- It is currently unclear how AS-OCT fits into clinical practice for detecting angle closure.
- AS-OCT is sensitive for detecting angle closure.
- AS-OCT may be a good screening tool for angle closure.
- AS-OCT has a high rate of false positives when measured against gonioscopy.
- AS-OCT is not yet able to replace gonioscopy.

Desmond T, Tran V, Maharaj M, Carnit N, White A. Diagnostic accuracy of AS-OCT vs gonioscopy for detecting angle closure: a systematic review and meta-analysis. *Graves Arch Clin Exp Ophthalmol*. 2022; Jan;260(1):1-23. doi: 10.1007/s00417-021-05271-4. Epub 2021 Jul 5. Erratum in: *Graves Arch Clin Exp Ophthalmol*. 2021 Sep 28; PMID: 34222969; PMCID: PMC8253367.

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<p>PROS</p> <ol style="list-style-type: none"> 1. Improved patient comfort. 2. Increased accessibility. 3. Real-time data and analytics. 4. Customized testing. 5. Patient engagement. 	<p>CONS</p> <ol style="list-style-type: none"> 1. <i>Not Well Studied in Comparison</i> 1. <i>Questionable underestimation in advanced disease.</i>
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Virtual Reality-Based and Conventional Visual Field Examination Comparison in Healthy and Glaucoma Patients
Jan Stapelberg^{1,2}, Saeed Saeed Kucur^{3,4}, Nina Huber⁵, Rami Nohri⁶, and Raphael Sattmann⁷

"VR System slightly underestimated VF defects in glaucoma patients"



Stapelberg J, Kucur SS, Huber N, Nohri R, Sattmann R. Virtual Reality-Based and Conventional Visual Field Examination Comparison in Healthy and Glaucoma Patients. *Transl Vis Sci Technol*. 2024 Oct 4;13(12):10. doi: 10.1167/tvst.10.12.10. PMID: 34814166. PMCID: PMC6989417.

Assessment of Remote Training, At-Home Testing, and Test-Retest Variability of a Novel Test for Clustered Virtual Reality Perimetry
Chia Zi, Kong AM, Turner M, Safiee M, Demotte BE, Backus BT, Saha JJ, Schuman JS, Deiner MS, et al.



Chia Zi, Kong AM, Turner M, Safiee M, Demotte BE, Backus BT, Saha JJ, Schuman JS, Deiner MS, et al. Assessment of Remote Training, At-Home Testing, and Test-Retest Variability of a Novel Test for Clustered Virtual Reality Perimetry. *Ophthalmol Glaucoma*. 2023 Aug 22;5(258):4199232023006. doi: 10.1016/j.ogta.2023.08.006.

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Validation of a Wearable Virtual Reality Perimeter for Glaucoma Staging, The NOVA Trial: Novel Virtual Reality Field Assessment
Oliver Brachler^{1,2}, Saeed Saeed Kucur^{3,4}, Thomas W. Sattmann⁵, Michael Zingales⁶, Howard Barabes^{7,8}, Nelson Reichle⁹, and Jason Staszko¹⁰

Statistically noninferior to HFA when staging glaucoma using Medicare definitions

Limitations: Monitoring advanced glaucoma



Bradley C, Ahmed IK, Samuelson TW, Chaglassian M, Barnebey H, Radcliffe N, Bacharach J. Validation of a Wearable Virtual Reality Perimeter for Glaucoma Staging, The NOVA Trial: Novel Virtual Reality Field Assessment. *Transl Vis Sci Technol*. 2024 Mar 1;13(2):10. doi: 10.1167/tvst.13.2.10. PMID: 38488433. PMCID: PMC10946691.

PMCID: PMC10946691
PMID: 38488433

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Subjective/Binocular Visual Field Testing

39% faster than SAP in clinical testing and functions in ambient light.¹

Equivalent to SAP with repeatability.¹

Random binocular testing



1. Comparison between New Perimetry Device (IMovifa®) and Humphrey Field Analyzer®
M Eslani, T Nishida, S Moghimi, JM Arias, C Vassile, V Mohammadzadeh, RN Weinreb;
Invest. Ophthalmol. Vis. Sci. 2022;63(7):1272 – A0412.

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Objective Visual Field Testing

FDA 510(K) Cleared
Tests OU simultaneously in 7 minutes
Measures the response of the pupils to a stimulus



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Advantages of objective perimetry

- Nothing to learn for the patient
- One *bilateral* test
- Less susceptible to refractive error and media opacity
- Easy to take - patients report they prefer OFA
- Learning effect - results can improve with experience
- Two monocular tests
- Susceptible to refractive error and media opacity
- More susceptible to anxiety, frustration, fatigue - "I just guess"

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Advantages of objective perimetry

- No patient response required
- Patients just need to look straight ahead and not fall asleep
- Dark room *not* required
- **Predictable Exam time**
 - ~7 minutes, for *both eyes* (30-2 & 24-2 together) OR
 - ~90 seconds, for *both eyes*
- If analysis improves can refresh reports
- Patients must click a button
- Reliant upon the patient's ability, dexterity, cooperation
- Dark room required
- **Variable exam time (24-2)**
 - 3 to >7 mins per eye (longer for some patients)
- No, SAP discards raw data

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Objective Perimetry

Diagnostic Power and Reproducibility of Objective Perimetry in Glaucoma

Abstract

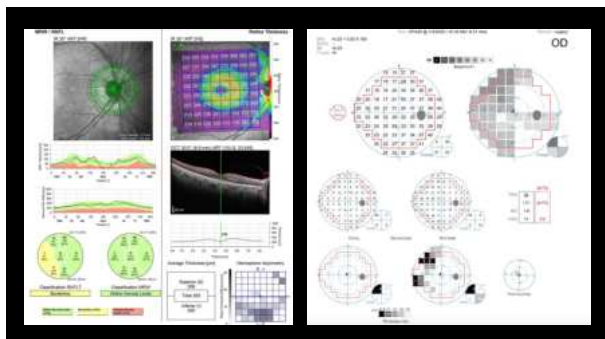
Purpose: To compare objective perimetry with 2 forms of standard automated perimetry (SAP) in glaucoma.

Methods: The study cohort consisted of 85 patients with glaucoma (PAG) and 84 normal control subjects. The PAG had both perimetric and computerized scans. Multifocal pupillary objective perimetry was performed with the objective field analyzer (OFA), which independently assesses the visual fields of both eyes concurrently. The OFA test assessed the central 30 degrees, and the OFA test assessed the central 10 degrees, both around 30-2 style reports. The OFA tests were repeated 2 weeks apart to assess test-retest variability (TRV). OFA was compared with Humphrey 24-2 and 30-2 threshold testing. Diagnostic power was quantified as the area under the receiver operating characteristic curve (AUROC). Test duration, mean fixation, and pattern standard deviation of the 4 tests were reported.

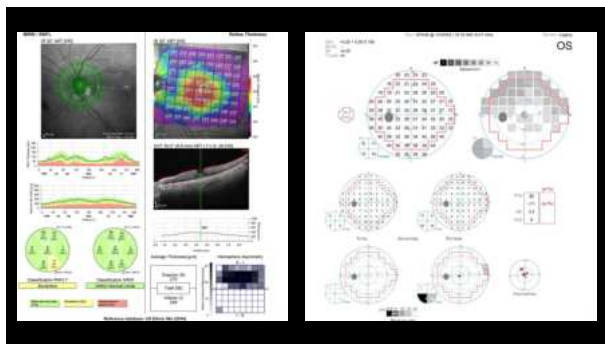
1. Ali, E.N., Maddess, T., James, A.C., Voicu, C., Lueck, C.J. Pupilary response to sparse multifocal stimuli in multiple sclerosis patients. *Mult. Scler.* 2014, 20, 854-861. [CrossRef]

2. Maddess, T., Ho, Y.L., Wong, S.S., Kolic, M., Goh, X.L., Carle, C.F., James, A.C. Multifocal pupillographic perimetry with white and colored stimuli. *J. Glaucoma* 2011, 20, 336-343. [CrossRef] [PubMed]

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Myth 5 ?

First Line Therapy Is Always A Drop?

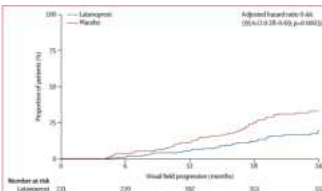
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Why are PGAs first line?

- Most efficacious
- Once daily dosing
- Minimal systemic SE's
- Uveoscleral outflow slows @night

25-33% ↓



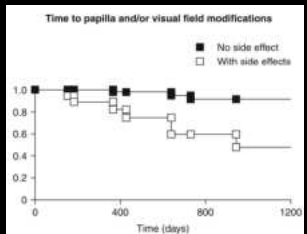
Number at risk	0	4	8	12	16	20	24
Latanoprost	213	209	187	163	133	105	85
Timolol	210	205	183	159	129	103	84

David G, David C, Luchsinger G, Amdurian F, Axtell N, Axtell-Brown A, ... Paper T (2015, April 6). Latanoprost for open-angle glaucoma (LORGES) randomised, double-masked, placebo-controlled trial. *PLoS One*. 10(4):e0121594.

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Treatment Challenges

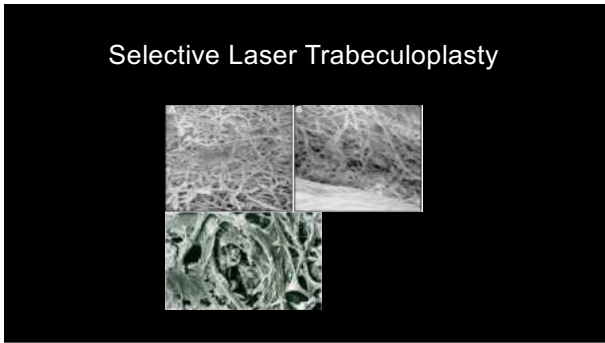
Time to papilla and/or visual field modifications



■ No side effect
□ With side effects

• Davis, Philippe, et al. Medical outcomes of glaucoma therapy from a nationwide representative survey.

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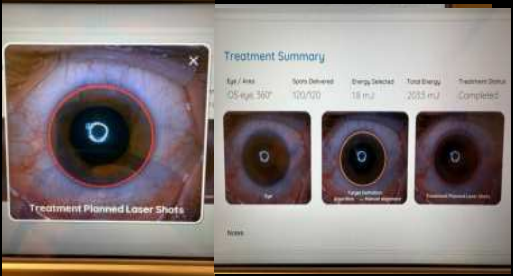
LIGHT trial: 6-year results of primary selective laser trabeculoplasty versus eye drops for the treatment of glaucoma and ocular hypertension

Gus Gazzard, Evgenia Konstantakopoulou, David Garway-Heath, Mariam Adeleke, Victoria Vickerstaff, Gareth Ambler, Rachael Hunter, Catey Bunce, Neil Nathwani, Keith Barton, on behalf of the LIGHT Trial Study Group

Primary Outcome - Quality of Life at 6 years
Secondary Outcome - clinical effectiveness and safety

Conclusions:
No significant difference in QOL
26.8% VS 19.6% progressed drops vs SLT
Trab required in 32 eyes in drops arm compared to 13 eyes in the SLT arm
69.8% of SLT Drop Free @ 6 Years

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Eye / Area	Spots Delivered	Energy Delivered	Total Energy	Treatment Status
OS eye, 360°	100/100	18 mJ	2033 mJ	Completed

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Patients Attitudes Towards Drug Delivery

- Triple Combination Eye Drop – 85%
- Microdose Eye Spray – 54%
- Drug-eluting Contact Lens – 31%
- Drug-eluting Periocular Ring Insert – 43%
- Injectable Subconjunctival Drug Insert- 32%
- Injectable Anterior Chamber Implant – 30%

attitude
is everything

Wang BB, Lin MM, Nguyen T, et al. Patient attitudes towards novel glaucoma drug delivery approaches. *Digit J Ophthalmol*. 2018; 24(3): 16-23

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Bimatoprost SR

(10-microgram bimatoprost sustained-release implant)

- Biodegradable bimatoprost sustained-release implant
- FDA-approved and indicated to reduce IOP in patients with open angle glaucoma or OHT
- Single intracameral administration
- Phase I/II/III Studies




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Travoprost intraocular implant

Resides in AC angle, anchored behind TM



- Length: 1.8 mm
- Diameter: 0.5 mm
- Titanium
- Non-ferrous

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36 Month Update

1. 70% and 68% of subjects in fast and slow-release were well-controlled on fewer or same medications as baseline.
2. Average IOP reductions were 8.3 mmHg and 8.5mmHg in the fast and slow-release arms.

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