Ocular Surface Preservation for the Glaucoma Patient.

Prevent Defense is the Best Offense.

Jacob Lang O.D., F.A.A.O.

Drjakelang@gmail.com

Board Certified, American Board of Optometry Intrepid Eye Society Board Member, Membership Chair Lead Optometrist & Optometric Residency Coordinator Associated Eye Care Stillwater, MN



Disclosures

 Lang- Aerie, Allergan, Avellino, Horizon, Novartis, Ocular Therapeutix, Orasis, Scope Sun Pharma, AOS

Ocular Surface Preservation

What are we looking at? What are we up against?

• DED is one of the most common ocular disease

- Approximately 30 million Americans (aprox. 3 million with glaucoma)
- Incidence of OSD increases with AGE as well as many other factors
 - DM
 - Migraines
 - Surgery
 - Associated with Anxiety, Depression, and Mental Health Conditions
 - Eye Drops (Blessing or a Curse?) (Gentamicin)
 - Environmental factors including screen time

What are we looking at? What are we up against?

Glaucoma Incidence

- Affects 60 Million Worldwide
- Prevalence increases with AGE
 - 8% of people older than 80

Shares many risk factors with DED

- Age
- DM
- Migraines
- PXF?
- Inflammation?

What are we looking at? What are we up against?

Co-Incidence of OSD and Glaucoma

 The prevalence of OSD is greatly increased in glaucoma patients, reaching 50% of patients in some studies.

Rate of SPK in Glaucoma patients

• As high as 54% of glaucoma patients have SPK

TBUT and Schirmer scores

• Abnormal in >60% of Glaucoma patients

Inflammation and the OSD/Glaucoma Flip-Flop

OSD has longstanding "roots" in inflammation and inflammatory pathways But what about Glaucoma? Is Glaucoma an **inflammatory condition**?

- There's a growing body of evidence suggests that neuro-inflammation and immune response are part of the sequence of pathological events leading to the optic neuropathy of glaucoma.
- Azithromycin
 - Strokes, MGD
- Minocycline
 - Strokes, MGD, Corneal Thinning
- Immunomodulation

Inflammation and the OSD/Glaucoma Flip-Flop

OSD has longstanding "roots" in inflammation and inflammatory pathways But what about Glaucoma? Is Glaucoma an inflammatory condition?

We Might be treating **BOTH** conditions when targeting inflammation!!!

Ocular Surface Preservation

Glaucoma Therapies and the Ocular Surface

- What are we going to talk about here???
- Insta @SeeOneTeachOne

What's the "**BAK**" Deal?

(Cue the BAK Rant, but remember everything is a balance...)

What is **BAK**?

- Benzalkonium chloride
- Quaternary Ammonium Compound
- Bacteriocidal, Bacteriostatic, and surfactant properties
- BAK destroys cell membranes to kill pathogens
- Indiscriminate killer
- BAK exposure is an independent risk factor for DED and OSD
- Has a **detergent** like effect which destabilizes the **lipid** layer
- Especially prevalent in generic formulations

What's the "**BAK**" Deal?

What is **BAK**, does it cause inflammation?

- BAK has pro-apoptotic effects and causes oxidative stress similar to those found in the TM of glaucoma patients with accelerated aging, trabecular cell death and extracellular matrix accumulation.
- BAK is a lipophilic molecule that accumulates deeply.
- Measurable levels of BAK can be found in the conjunctiva up to 1 week after administration of a single drop.

What's the "**BAK**" Deal?

What is **BAK**?

- BAK has been retrieved using the mass spectrometry imaging technique in deep structures such as the trabecular meshwork or the lens in rabbit eyes treated with BAK for several months, which was also confirmed in human tissues.
- At higher concentrations, sub-conjunctival injection of BAK is capable of causing trabecular cell death, inflammatory infiltration, and increased IOP.
- This explains why IOP improves when switching over to PF meds.

Medical Glaucoma Therapies

Prostaglandins

- Known Inflammatory Mediator
- Stimulate the expression of matrix metalloproteinases (MMPs) which hydrolyze excessive ECM, opening up extracellular spaces and decrease fluid resistance flowing through these spaces. In addition, they induce relaxation of the TM and ciliary muscle, which reduces tension and increases the outflow pathways.
- Yet it's our first line of treatment due to it's effectiveness

Medical Glaucoma Therapies

Alpha Agonists

- Brimonidine
 - α-2 adrenergic receptor agonist that decreases aqueous humor secretion and enhances aqueous humor resorption by the uveoscleral channels.
 - Allergenic and/or pro-inflammatory properties
 - Induces a Granulomatous Uveitis (and increased IOP) in some patients
 - Case reports of INCREASED IOP with use (inflammatory response?)

Other Medical Therapies

Lumify???



Medical Therapies - Medicamentosa-We are our own worst enemy

Medicamentosa

- Allergic manifestations of therapy
 - Hyperemia, chemosis, edema, follicular reaction, etc.
- Prostaglandins 1.5%
- CAIs **3-4%**
- Beta-Blockers 11-13%
- Alpha-Agonists **11.5%**
 - (Combigan?)

A Patient's Story



Medical Therapies - Medicamentosa-

Medicamentosa

- Pseudopemphigoid
 - Cicatrizing conjunctivitis that mimics true pemphigoid
 - 28.3% of all pseudopemphagoid cases were due to topical glaucoma medications



Medical Therapies

TABLE 2. COMMONLY PRESCRIBED GLAUCOMA MEDICATIONS WITH THEIR CORRESPONDING PRESERVATIVE^a

Medication	Preservative
Xalatan	BAK 0.02%
Lumigan	BAK 0.02%
Azopt	BAK 0.01%
Timoptic	BAK 0.01%
Trusopt	BAK 0.0075%
Cosopt	BAK 0.0075%
Combigan	BAK 0.005%
Travatan Z	SofZia
Alphagan P	Purite
Zioptan	None
Cosopt PF	None
Timoptic in Ocudose	None



Preservative-Free Alternatives Options for decreasing ocular toxicity in patients with

glaucoma. By Arksdiy Yadgarov, MD, and Reens A. Garg, MD

VEWPDF

🚺 | 📴 | E-MAIL | PRINT | BOOKMARK

Rhopressa 0.015% BAK Simbrinza 0.003% BAK Rocklatan 0.02% BAK

Vyzulta 0.02% BAK

IOP-Lowering medications	BAK concentration %	
Xalatan	0.02	
Travatan	0.015	
Betoptic S	0.01	
Azopt	0.01	
Timoptic	0.01	
Simbrinza	0.003	
Brimonidine	0.005	
Lumigan	0.005	
Betagan	0.005	
Combigan	0.005	
Cosopt	0.0075	
Trusopt	0.0075	
Www. Tear Oamelarity in a Oleveense		



Tear Osmolarity in a Glaucoma Practice

The role of point-of-care testing in dry eye disease and glaucoma management. By Leslie E. O'Dell. OD

VIEW PDF



Instill 1 drop in the affected eye(s) once dail

since it has been shown that more frequent administration of prostaglandin analogs may decrease the IOP-lowering effect.

Reduction of IOP starts approximately 2 hours after the first administration with maximum effect reached after 12 hours.

 May be used concomitantly with other topical ophthalmic drug products to lower IOP. If more than 1 topical ophthalmic drug is being used, the drugs should be administered at least 5 minutes apart.
 VYZULTA (latanoprostene bunod (0.024%) ophthalmic

🧏 💿 🚥

Prostaglandin Analogs Indications/Usage:

Indicated for the reduction of intraocular pressure (IOP) in patients with open-angle glaucoma or ocular hypertension.

Typical Dosing:

Elevated intraocular pressure (IOP) in patients with open-angle glaucoma or ocular hypertension, recommended:

instill 1 drop in the conjunctival sac of the affected eye(s) once daily in the evening.
Do not administer more than once daily since it has

been shown that more frequent administration of prostaglandin analogs may lessen the IOP-lowering effect. • If used concomitantly with other topical ophthalmic

drug products to lower IOP, administer each drug product at least 5 minutes apart.

How Supplied: 2.5 mL bottles 5 mL bottles

ZIOPTAN (tafluprost (0.0015%) ophthalmic solution)

Prostaglandin Analogs

Indications/Usage: Indicated for reducing elevated intraocular pressure (IOP) in patients with open-angle glaucoma or ocular hypertension.

. . . .

Typical Dosing: ► Elevated intraocular pressure (IOP) in patients with open-angle glaucoma or ocular hypertension, recommended:

- Instill 1 drop in the conjunctival sac of the affected eye(s) once daily in the evening.
- The dose should not exceed once daily since it has been shown that more frequent administration of prostaglandin analogs may lessen the IOP-lowering effect.
- Reduction of the IOP starts approximately 2 4 hours after the first administration with the maximum effect reached after 12 hours.
- May be used concomitantly with other topical ophthalmic drug products to lower IOP. If more than 1 topical ophthalmic product is being used, each one should be administered at least 5 minutes apart.
- should be administered at least 5 minutes apart. • The solution from 1 individual unit is to be used immediately after opening for administration to one or both eyes. Since sterility cannot be maintained after the individual unit is opened, the remaining contents should be discarded immediately after

administration.

0.3 mL in 10 single-use containers per pouch

Preservatives:

None listed in package insert.

Instill 1 drop in the affected eye(s) 3 times daily.
 May be used concomitantly with other topical ophthalmic drug products to lower IOP. If more than

optimalmic drug products to lower IOP, it more than 1 topical ophthalmic drug is being used, the drugs should be administered at least 5 minutes apart. How Supplied:

10 mL bottles
Preservatives:

benzalkonium chloride

?

XALATAN (latanoprost (0.005%) ophthalmic solution)

Indications/Usage: Indicated for the reduction of elevated intraocular pressure (IOP) in patients with open-angle glaucoma or ocular hypertension.

- Typical Dosing: ► Elevated intraocular pressure (IOP) in patients with open-angle glaucoma or ocular hypertension, recommended:
- Instill 1 drop in the affected eye(s) 1 time daily in the evening. If 1 dose is missed, treatment should

ontinue with the next dose as normal.
The dosage should not exceed once daily; the combined use of 2 or more prostaglandins, or

- prostaglandin analogs including XALATAN is not recommended. It has been shown that administration of these prostaglandin drug products more than once daily may decrease the 10P-lowering effect or cause paradoxical elevations in 10P.
- Reduction of the IOP starts approximately 3 4 hours after administration and the maximum effect is reached after 8 – 12 hours.
- May be used concomitantly with other topical ophthalmic drug products to lower IOP. If more than 1 topical ophthalmic drug is being used, the drugs should be administered at least 5 minutes apart.
 Contact leases should be removed note to the
- administration of XALATAN, and may be reinserted 15 minutes after administration.

How Supplied: • 2.5 mL bottles Preservatives: benzalkonium chloride

Storage:

Protect from light.

- Store unopened bottle(s) under refrigeration at 36° - 46°F (2° - 8°C).
- During shipment to the patient, the bottle may be maintained at temperatures up to 104°F (40°C) for a period not exceeding 8 days.
- Once a bottle is opened for use, it may be stored at room temperature up to 77°F (25°C) for 6 weeks.

Cost: Compare Prices

Assistance: Payment Assistance

Show More

dosage may be changed to 1 drop of 0.5% solution in the affected evels)

 Since in some patients the pressure-lowering response to Preservative-free TIMOPTIC in OCUDDSE may require a few weeks to stabilize, evaluation should include a determination of IOP after approximately 4 weeks of treatment with Preservative-free TIMOPTIC in OCUDDSE.

- If the IOP is maintained at satisfactory levels, the dosage schedule may be changed to 1 drop once a day in the affected eye(s).
- Because of diurnal variations in IOP, satisfactory response to the once-a-day dose is best dotormined by measuring the IOP at different.

XELPROS (latanoprost (0.005%) ophthalmic emulsion)

Prostaglandin Analogs

Indications/Usage: Indicated for the reduction of elevated intraocular pressure (IOP) in patients with open-angle glaucoma or ocular hypertension.

Typical Dosing:

► Elevated intraocular pressure (IOP) in patients with open-angle glaucoma or ocular hypertension, recommended:

- Instill 1 drop in the affected eye(s) once daily in the evening.
- If 1 dose is missed, treatment should continue with the next dose as normal.
- The dosage should not exceed once daily; the combined use of 2 or more prostaglandins, or prostaglandin analogs including XELPROS is not recommended. It has been shown that administration of these prostaglandin drug products more than once daily may decrease the IOP-lowering effect or cause paradoxical elevations in IOP.
- Reduction of the IOP starts approximately 3 4 hours after administration and the maximum effect is reached after 8 – 12 hours.
- May be used concomitantly with other topical ophthalmic drug products to lower IOP. If more than 1 topical ophthalmic drug is being used, the drugs should be administered at least 5 minutes apart.
- Contact lenses should be removed prior to the administration of XELPROS, and may be reinserted 15 minutes after administration.

How Supplied: • 2.5 mL bottles

Preservatives:

potassium sorbate

- Storage: Protect from light.
- Store at 36° 77°F (2°C 25°C).
- During shipment to the patient, the bottle may be maintained at temperatures up to 104°F (40°C) for a
- Period not exceeding 8 days.
 After opening, XELPROS can be used until the
- expiration date stamped on bottle and then discarded.
 - Cost: Compare Prices
 - Assistance: Payment Assistance

Assistance: Payment Assistance

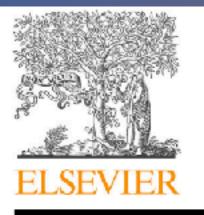
36 items found

Ophthalmic Drugs OTC Tears Lid Hygiene Our Team About

Ocular Surface VS. Optic Nerve Testing

- What do you do when you see a patient with Glaucomatous Optic Neuropathy???
- What do you **do** when you see a patient on Plaquenil?
- How do you track the effectiveness of your therapy?
- How do you know there's been a change without a baseline?
- Do you tell refractive surgery patients that it may worsen their OSD?

Ocular Surface VS. Optic Nerve Testing



Contents lists available at ScienceDirect

The Ocular Surface

journal homepage: www.theocularsurface.com

Original Research

Exploring topical anti-glaucoma medication effects on the ocular surface in the context of the current understanding of dry eye

Aaron B.C. Wong, Michael T.M. Wang, Kevin Liu, Zak J. Prime, Helen V. Danesh-Meyer, Jennifer P. Craig^{*}

Department of Ophthalmology, New Zealand National Eye Centre, The University of Auckland, New Zealand

Ocular Surface and Medical Glaucoma Tx

- Of note, decreased tear film stability and elevated tear osmolarity, both global indices of dry eye disease, were observed in eyes treated with topical anti-glaucoma medications.
- This was associated with increased levels of bulbar conjunctival hyperemia and eyelid margin changes.
- These findings suggest that inflammatory mechanisms may play a role in the propensity of dry eye development in patients receiving long term topical anti-glaucoma medications.

Baseline DED/OSD Testing

- Osmolarity
- Inflammation
- Staining
 - Fluorescein,
 - Lissamine green
- Tear Prism
- Tear Production
 - Schirmer

- TBUT
- Symptom Surveys
 - SPEED
 - DEQ-5
- Meibography & Gland Analysis/Expression
- Lid Anatomy, Skin Conditions, Demodex

OPEN ACCESS



July 2018 Volume 59, Issue 9 ARVO Annual Meeting Abstract | July 2018 Thirty-minute ocular application of benzalkonium chloride (BAK) leads to a disappearance of neuronal spikes associated with a marked fragmentation of tight junctions and corneal nerves in rodents: implications for dry eye disease.

Evguenia Ivakhnitskaia; Kamila Mizerska; Valentina Dallacasagrande; Victor H Guaiquil; Mark Rosenblatt; Harumitsu Hirata

+ Author Affiliations & Notes

Investigative Ophthalmology & Visual Science July 2018, Vol.59, 3284. doi:

DEWS II Definition; "Dry eye is a multifactorial disease of the ocular surface characterized by a loss of homeostasis of the tear film, and accompanied by ocular symptoms, in which tear film instability and hyperosmolarity, ocular surface inflammation and damage, and <u>neurosensory</u> <u>abnormalities</u> play etiological roles."

Conclusions : Our results revealed that short exposure to ophthalmic solutions with low BAK concentrations used in routine eye examinations has profound consequences on corneal nerve and epithelial integrity; hence, caution is warranted with such uses. The long-term consequences of this short exposure on corneal nerves and tight junctions is currently under investigation. Overall, these studies may lead to a better understanding of the pathogenesis of dry eye disease.

This is an abstract that was submitted for the 2018 ARVO Annual Meeting, held in Honolulu, Hawaii, April 29 - May 3, 2018.



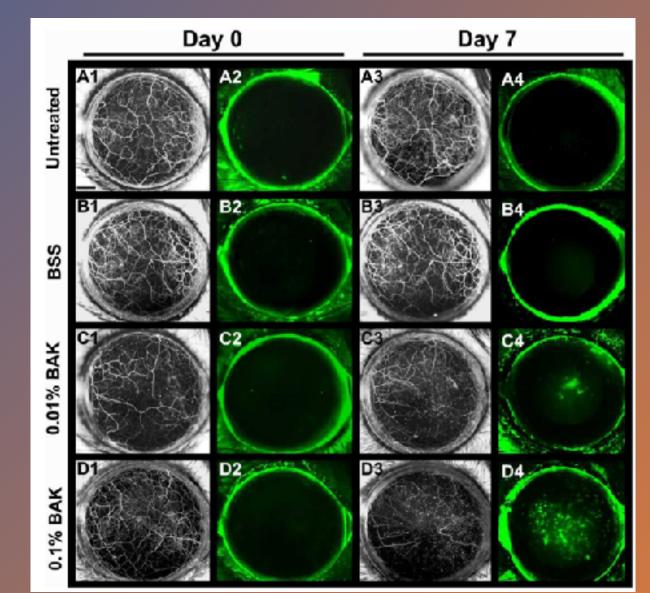
Invest Ophthalmol Vis Sci. 2012 Apr; 53(4): 1792–1802. Published online 2012 Apr 18. doi: <u>10.1167/iovs.11-8775</u> PMCID: PMC3995561 PMID: 22410563

Corneal Neurotoxicity Due to Topical Benzalkonium Chloride

Joy Sarkar, Shweta Chaudhary, Abed Namavari, Okan Ozturk, Jin-Hong Chang, Lisette Yco, Snehal Sonawane, Vishakha Khanolkar, Joelle Hallak, and Sandeep Jain

Conclusion.

Topical application of BAK to the eye causes corneal neurotoxicity, inflammation, and reduced aqueous tear production.



Influence of Treating Ocular Surface Disease on Intraocular Pressure in Glaucoma Patients Intolerant to Their Topical Treatments: A Report of 10 Cases

> Pierre Dubrulle, MD,*† Antoine Labbé, MD, PhD,*†‡§ Emmanuelle Brasnu, MD, PhD,* Hong Liang, MD, PhD,*†‡ Pascale Hamard, MD, PhD,* Lyes Meziani, MD,* and Christophe Baudouin, MD, PhD*†‡§

- Case Series (10) patients referred for filtering glaucoma surgery.
- The main treatments were
 - change of topical anti-glaucoma medications to preservative-free equivalents
 - <u>removal</u> of allergenic treatments or those identified as causing side effects
 - switch to another therapeutic class with the same efficacy but with a better safety profile
- Treatment of OSD especially MGD.

- After a minimum follow-up of 6 months, we observed improved ocular surface in all patients, associated with an intraocular pressure (IOP) decrease or stabilization even if some anti-glaucoma medications were removed.
- The mean IOP significantly decreased from 23.75±9.98mmHg to 15.15±4.75mmHg (-36.2%; P=0.0001).
- The mean number of IOP-lowering medications was 3.7±1.06 at presentation and 2.8±0.63 after treatment (P=0.01).
- The Oxford staining score also decreased from a mean 1.7±0.67 to 0.4±0.51 (-76.5%; P<0.001).

- For 2 patients, IOP was not sufficiently reduced after treatment and they finally underwent filtering surgery.
- You need surgery? **PROVE IT!**
- Who wants to do surgery on a sick ocular surface?

How do I get my prescription to Eagle Pharmacy?

• There are two ways to have your prescription sent to Eagle Pharmacy:

CHOOSE HOW TO GET YOUR PRESCRIPTION TO EAGLE PHARMACY	
Doctor's Office Sends the Rx	You Send the Rx
The fastest and best way to get your prescription to Eagle Pharmacy is to ask your eye doctor's office staff to call, fax, or ePrescribe your prescription directly to Eagle Pharmacy, just like they do with other pharmacies.	If your eye doctor gave you a paper prescription, you can mail the original paper prescription directly to Eagle Pharmacy along with your completed enrollment form.
Your doctor's office may need the contact information for Eagle Pharmacy so be sure to provide them the information below.	
Call Eagle Pharmacy at 844-813-3864 Mon to Fri 9AM – 7PM EST	Mail to Akorn EyeRx Direct c/o Eagle Pharmacy PO Box 90937 Lakeland, FL 33804 Click here to download an enrollment form
Fax Eagle Pharmacy at 855-618-4610	
ePrescribe to Eagle Pharmacy, Lakeland FL 33810	

NPI: 1487905840 NCPDP: 5711975

Decrease Inflammation

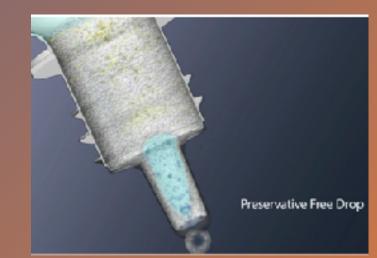
- Decrease BAK (other preservatives) Load
- Anti-Inflamatory Medications
 - Lifitegrast
 - Cyclosporine
 - NSAIDS
 - Acuvail is **PF**
 - Steroids
 - Nanoparticles

Decrease Inflammation

• Decrease BAK (other preservatives) Load

NANO DROPPER

- ~10 μL vs. 40 μL
- Eliminate BAK (other preservatives) Load
 - TearClear





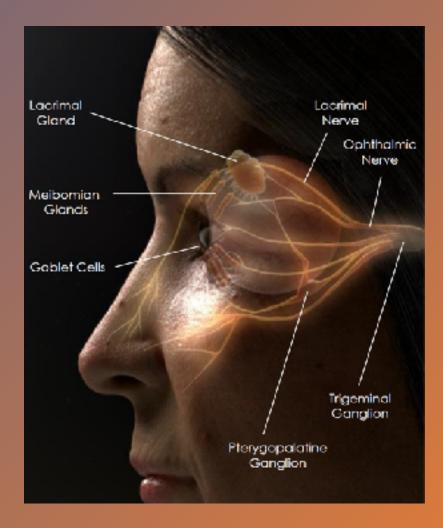
Decrease Inflammation

- Tear Stimulation
 - TrueTear (discontinued June 2020), iTear100
- "Fill" the Swamp? (Plugs)
- Treat the Lids
 - Lipiflow, ILux, TearCare
 - Lid Hygiene
 - IPL
- Don't forget about
 - Contact Lens Care & Solutions
 - Allergic Conjunctivitis



Decrease Inflammation

- Tear Stimulation
 - OC-01 Nasal Spray



Intense Pulse Light (IPL) and Inflammation

- Proposed MoA
 - Lowers cytokine levels (IL-17A, IL-6, TNF-α)

> Sci Rep. 2019 May 21;9(1):7648. doi: 10.1038/s41598-019-44000-0.

- Meibum Expressibility Improvement as a Therapeutic Target of Intense Pulsed Light Treatment in
- A Meibomian Gland Dysfunction and Its Association
- **C** with Tear Inflammatory Cytokines

Moonjung Choi¹, Soo Jung Han², Yong Woo Ji^{2³}, Young Joon Choi^{2⁴}, Ikhyun Jun², Mutlaq Hamad Alotaibi^{2⁵}, Byung Yi Ko¹, Eung Kweon Kim^{2⁶}, Tae-Im Kim², Sang Min Nam⁷, Kyoung Yul Seo⁸

Decrease Inflammation

- Medication Vacation (AKA Ocular Surface <u>Quarantine</u>)
 - Oral CAIs as a transitional therapy

What about **Punctal Plugs**?

Effect of a punctal plug on ocular surface disease in patients using topical prostaglandin analogues: a randomized controlled trial

Justin C Sherwin MPhil FRANZCO,^{1,2,3} ^(D) Gokulan Ratnarajan MD FRCOphth,⁴ Babar Elahi FRCOphth,⁵ Anna Bilkiewicz-Pawelec MD¹ and John F Salmon MD FRCOphth¹

¹Oxford Eye Hospital, John Radcliffe Hospital, Oxford, ⁴The Queen Victoria Hospital, East Grinstead, ⁵Dudley Hospital NHS Trust, Dudley, UK; and ²Ophthalmology, University of Melbourne Department of Surgery, and ³Centre for Eye Research Australia, Royal Victorian Eye and Ear Hospital, Melbourne, Australia

What about **Punctal Plugs**?

- Conclusions and Relevance: Punctal plug insertion improves subjective and objective measures of OSD and results in a reduced IOP in patients with symptomatic ocular surface disease using prostaglandin analogue monotherapy.
- TBUT increased
- Oxford Staining Score decreased
- Tear Osmolarity decreased

What about **Surgery**?

- SLT and MIGS
 - Should SLT be the **first option** for POAG with concomitant OSD???
- At Cataract Extraction
 - PHACO Study
- Filtering procedures (Trabs and Tubes)
 - What is the first/most important intra-operative indicator of success

Should we LiGHT them up?

- SLT first line in Europe
- Delay topical treatments x 3 years

ARTICLES | VOLUME 393, ISSUE 10180, P1505-1516, APRIL 13, 2019

Selective laser trabeculoplasty versus eye drops for first-line treatment of ocular hypertension and glaucoma (LiGHT): a multicentre randomised controlled trial

Gus Gazzard, FRCOphth \times \times \times evgenia Konstantakopoulou, PhD \times Prof David Garway-Heath, MD \times Anurag Garg, FRCOphth \times Victoria Vickerstaff, MSc \times Rachael Hunter, MSc \times et al. Show all authors \times Show footnotes

Open Access • Published: March 09, 2019 • DOI: https://doi.org/10.1016/S0140-6736(18)32213-X •

What about MIGS?

Options?

- TM bypass
 - iStent original vs iStent Inject
 - Hydrus
- Trabeculotomy
 - Trabectome
 - Kahook dual blade
 - OMNI (viscodilation + trabeculotomy)

Ophthalmol Ther (2020) 9:941-953 https://doi.org/10.1007/s40123-020-00290-6



ORIGINAL RESEARCH

Prospective Interventional Cohort Study of Ocular Surface Disease Changes in Eyes After Trabecular Micro-Bypass Stent(s) Implantation (iStent or iStent *inject*) with Phacoemulsification

Justin A. Schweitzer 💿 · Whitney H. Hauser · Mitch Ibach · Brandon Baartman · Subba R. Gollamudi · Andrew W. Crothers ·

John E. Linn · John P. Berdahl

Received: July 5, 2020/Published online: August 13, 2020 © The Author(s) 2020

Ophthalmol Ther (2020) 9:941–953 hnps://doi.org/10.1007/s40123-020-00290-6

Prospective Interventional Cohort Study of Ocular Surface Disease Changes in Eves After Trabecular

Table 2 Preoperative and month 3 ocular surface parameters, intraocular pressure, and number of medications eyes undergoing istent + phacoemulsification $(n = 45)^a$

	Conjunctival hyperemia (Efron Scale)	Corneal/conjunc- tival staining (Oxford Schema)	FTBUT (no. seconds)	Ocular Surface Disease Index (OSDI)	Intraocular pressure (mmHg)	Number of medications
Preoperative	1.4 ± 0.7	1.4 ± 1.0	4.4 ± 2.4	39.8 ± 21.9	17.5 ± 4.2	1.5 ± 0.9
Month 3	1.2 ± 0.6	0.4 ± 0.5	6.5 ± 2.5	16.4 ± 14.8	14.5 ± 3.3	0.6 ± 0.8
<i>p</i> value vs. preoperatively (paired t-test)	0.128	< 0.0001	< 0.0001	< 0.0001	< 0.001	< 0.0001
% Change vs. preoperatively	14% reduction	71% reduction	48% increase (improvement)	56% reduction	17% reduction	60% reduction

^a Analysis does not include two eyes receiving iStent *inject*

Ophthalmol Ther (2020) 9:941-953 https://doi.org/10.1007/s40123-020-00290-6 Check for updates

ORIGINAL RESEARCH

Prospective Interventional Cohort Study of Ocular Surface Disease Changes in Eyes After Trabecular Micro-Bypass Stent(s) Implantation (iStent or iStent *inject*) with Phacoemulsification

Justin A. Schweitzer 💿 · Whitney H. Hauser · Mitch Ibach · Brandon Baartman · Subba R. Gollamudi · Andrew W. Crothers · John E. Linn · John P. Berdahl

Received: July 5, 2020 / Published online: August 13, 2020 © The Author(s) 2020

Implantation of trabecular micro- bypass stent(s) (iStent or iStent inject) with cataract surgery produced significant improvements in ocular surface health, alongside significant reductions in IOP and medications.

10 mcg Bimatoprost Implant (Durysta™)

- Intracameral injection
- Biodegradable implant
- Last 4-6 months
- Consistent IOP lowering effects without the topical side effects of preservatives



2 Weeks

9 Months

12 Months

10 mcg Bimatoprost Implant (Durysta™)

•30% IOP ↓ over 12 weeks
•80% of patients on no additional drops over 9 month study

•Long term--40% had 30% reduction at 1 year, 28% at 2 years.

Medeiros FA, et al Ophthalmology June 2020



2 Weeks

9 Months

12 Months

Other Devices for Drug Delivery?

- Plugs
 - Dextensa derivative?
- Subconjunctival?
- Fornix?
 - Lacrisert

THANK YOU!

Jacob Lang OD, FAAO, Dipl. ABO

- drjakelang@gmail.com
- Instagram @SeeOneTeachOne



Patia Jensen

Live. Work. Create

<u>pmjensen@outlook.com</u> https://www.linkedin.com/in/patiajensen/

- 1. Wong ABC, Wang MTM, Liu K, Prime ZJ, Danesh-Meyer H V., Craig JP. Exploring topical antiglaucoma medication effects on the ocular surface in the context of the current understanding of dry eye. *Ocul Surf.* 2018;16(3):289-293. doi:10.1016/j.jtos.2018.03.002.
- Dubrulle P, Labbé A. Influence of Treating Ocular Surface Disease on Intraocular Pressure in Glaucoma Patients Intolerant to Their Topical Treatments : A Report of 10 Cases. 2018;27(12):1105-1111. doi:10.1097/IJG.000000000001041.
- 3. Sherwin JC, Ratnarajan G, Elahi B, Bilkiewicz-Pawelec A, Salmon JF. Effect of a punctal plug on ocular surface disease in patients using topical prostaglandin analogues: a randomized controlled trial. *Clin Exp Ophthalmol*. 2018;46(8):888-894. doi:10.1111/ceo.13311.
- 4. Jin SW, Min JS. Clinical evaluation of the effect of diquafosol ophthalmic solution in glaucoma patients with dry eye syndrome. *Jpn J Ophthalmol*. 2016;60(3):150-155. doi:10.1007/s10384-016-0430-8.
- Stringham J, Ashkenazy N, Galor A, Wellik SR. Barriers To Glaucoma Medication Compliance Among Veterans: Dry Eye Symptoms and Anxiety Disorders HHS Public Access. *Eye Contact Lens*. 2018;44(1):50-54. doi:10.1097/ICL.00000000000000301.

- 6. Jalkh AE, El Hajj Moussa WG, Schakal AR, et al. Comparison of Efficacy and Ocular Surface Disease Index Score between Bimatoprost, Latanoprost, Travoprost, and Tafluprost in Glaucoma Patients. *J Ophthalmol*. 2018;2018:1-7. doi:10.1155/2018/1319628.
- 7. rodriguez-garcia alejandro, loya-garcia D, hernandez-Quintela everardo, navas alejandro. Clinical Ophthalmology risk factors for ocular surface damage in Mexican patients with dry eye disease: a population-based study. 2019. doi:10.2147/OPTh.s190803.
- 8. Banitt M, Jung H. Ocular Surface Disease in the Glaucoma Patient. *Int Ophthalmol Clin.* 2018;58(3):23-33. doi:10.1097/iio.0000000000232.
- 9. Ji H, Zhu Y, Zhang Y, et al. The Effect of Dry Eye Disease on Scar Formation in Rabbit Glaucoma Filtration Surgery. *Int J Mol Sci Artic*. doi:10.3390/ijms18061150.
- 10. alejandro Rodriguez-garcia, Loya-garcia D, everardo Hernandez-Quintela, alejandro Navas. Clinical Ophthalmology risk factors for ocular surface damage in Mexican patients with dry eye disease: a population-based study. 2019. doi:10.2147/OPTh.s190803.

- 11. Williams PA, Marsh-Armstrong N, Howell GR, et al. Neuroinflammation in glaucoma: A new opportunity [1] The Lasker/IRRF Initiative on Astrocytes and Glaucomatous Neurodegeneration Participants † HHS Public Access. *Exp Eye Res.* 2017;157:20-27. doi:10.1016/j.exer.2017.02.014.
- 12. Hamrah P, Alipour F, Jiang S, Sohn J-H, Foulks GN. Optimizing evaluation of Lissamine Green parameters for ocular surface staining. *Eye*. 2011;25:1429-1434. doi:10.1038/eye.2011.184.
- 13. Gumus K, Schuetzle KL, Pflugfelder SC. Randomized Controlled Crossover Trial Comparing the Impact of Sham or Intranasal Tear Neurostimulation on Conjunctival Goblet Cell Degranulation. 2017. doi:10.1016/j.ajo.2017.03.002.
- 14. Stringham J, Ashkenazy N, Galor A, Wellik SR. Barriers to Glaucoma Medication Compliance Among Veterans. *Eye Contact Lens Sci Clin Pract*. 2017;44(1):1. doi:10.1097/ICL.00000000000000301.
- 15. Alvaro Gomes JP, Azar DT, Baudouin C, et al. TFOS DEWS II iatrogenic report. *Ocul Surf*. 2017;15:511-538. doi:10.1016/j.jtos.2017.05.004.

- 16. Craig JP, Nelson JD, Azar DT, et al. TFOS DEWS II Report Executive Summary. *Ocul Surf*. 2017;15(4):802-812. doi:10.1016/j.jtos.2017.08.003.
- 17. Honjo M, Tanihara H, Kameda T, Kawaji T, Yoshimura N, Araie M. Potential role of rho-associated protein kinase inhibitor Y-27632 in glaucoma filtration surgery. *Investig Ophthalmol Vis Sci*. 2007;48(12):5549-5557. doi:10.1167/iovs.07-0878.
- 18. Birnbaum FA, Hamrah P, Jacobs DS, Song BJ. Acquired corneal neuropathy and photoallodynia associated with malposition of an Ex-PRESS shunt. *J Glaucoma*. 2017;26(1):e19-e21. doi:10.1097/ IJG.000000000000000502.
- Gazzard G, Konstantakopoulou E, Garway-Heath D, et al; LiGHT Trial Study Group. Selective laser trabeculoplasty versus eye drops for the first-line treatment of ocular hypertension and glaucoma (LiGHT): a multicentre randomised controlled trial [published online ahead of print March 8, 2019]. Lancet. doi: 10.1016/S0140-6736(18)32213-X.