

Dry Eye and Glaucoma



Dry eye is a common eye condition that affects thousands of New Zealanders every day. Like open-angle glaucoma, dry eye syndrome increases with ageing. Studies suggest that 40-50% of glaucoma patients have dry eye syndrome, with women more likely to have it. People with glaucoma are at higher risk of dry eye because some glaucoma treatments can worsen the ocular surface and increase the tendency to dry the eye.

How to recognise dry eye?

In some cases, dry eye can affect your quality of vision and sensitivity to light. The most frequently reported symptom of dry eye is grittiness and burning, or a sensation of sand or grit in the eyes. However, you might be surprised to learn that dry eyes can also cause excessive watering of the eyes (epiphora).

Why does this happen?

The thin layer of tears that covers your eye keeping it lubricated, nourished and protected, has three main components. Mucins are at the base to help anchor the tear film in place, a watery layer with nutrients is in the middle (aqueous), and oil is on the surface to reduce evaporation. The various glands that produce these components

must all be functioning to create a healthy tear film of adequate quantity and quality. Unfortunately, when this tear film becomes dysfunctional in dry eye, it causes the eye's surface to become inflamed and result in chronically red, irritated eyes.

This is either because of a lack of tear quantity (volume) or quality.

A tear film lacking in quantity is described as 'aqueous deficient'; a truly 'dry' dry eye.

One lacking in quality is described as 'evaporative' since it evaporates too rapidly, causing irritation and consequent eye-watering.

Evaporative dry eye is approximately 9 times more common than aqueous deficiency.

Why has my tear film become dysfunctional? Continued on page 2

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Dry eye and glaucoma continued

Usually, a combination of factors is responsible for the development of a dry eye.

Age and Gender: Dry eye becomes more common with increasing age and is particularly prevalent in post-menopausal women.

Environment: Air-conditioned, centrally heated, smoky, polluted, or windy settings, as well as computer use, can aggravate dry eye.

Contact lenses: Contact lenses disrupt the natural tear film, causing excessive tear film evaporation.

General health: Rheumatoid arthritis, diabetes, thyroid disease, and other immune disorders such as Sjögren's syndrome are more commonly associated with dry eye

Oral medications: Some medications for allergy, depression or acne, for example, can affect the tear film, as can eye surgery, especially laser surgery (e.g. LASIK).

Eye drops: Glaucoma is associated with a higher incidence of dry eye because of the medications used to treat the condition.

Many eye drops for glaucoma contain preservatives, which after repeated application over a long period, can damage the superficial cells of the ocular surface, making the eye's surface less wettable and resulting in the development of dry eye signs and symptoms.

What can you do about it?

Improve the environmental conditions: Minimise exposure to air-conditioning and central

heating. Enhance the moisture level around the eyes by using humidifiers and/or by wearing wrap-around style spectacles to create a humidified local environment protecting the eyes from airflow. Breathable foam inserts that form a seal around the eye are another option with wrap-around style glasses or sunglasses.

Minimise use of cosmetics:

Cosmetics and face creams worn on the face and around the eyes will migrate over the eyelid margins. This can disrupt the tear film, exacerbating dry eye, so try to minimise make-up use.

Optimise your workspace: Adopt a slight downward gaze at the computer screen where possible to minimise the exposed eye area. Lowering your screen prevents excessive drying of the eye surface. Improve your blinking by practising complete and regular blinking, at least once every 5 or 6 seconds. Blinking exercises can be done anywhere at any time, and there are even Phone Apps to help.

'Blink reminder' apps are designed for dry eye computer users and are free to download.(go to glaucoma.org.nz to find out more).

Adjust your diet: Increasing your dietary intake of essential fatty acids (EFA) to include Omega 3 can help reduce dry eye symptoms. Good natural sources of Omega 3 include oily fish (such as salmon) and linseed (or flaxseed) oil. In addition, Lacritec and Theratears have supplemental products specially formulated for dry eyes. AFT's eye drops NovaTears®+

Omega-3 also contains it.

It is also essential to stay hydrated, so drink plenty of water.

Perform eyelid hygiene: Tear film quality can be adversely affected by a lack of oils due to blocked glands (meibomian gland dysfunction). Thoroughly cleaning your eyelid margins, ideally with commercially prepared lid cleansers from your pharmacy, helps to remove any crusts (signs of blepharitis) from around the eyelashes. Warm compresses, using a wheat or flaxseed bag heated in the microwave, for example (Figure1), helps to unblock the glands by melting the oils.

After warming the eyelids for at least 5 minutes, it's helpful to roll your finger gently over your eyelids, towards the edges (i.e. rolling with a downwards motion for the top lid and upwards for the bottom lid) to help squeeze the oils out of the glands (Figure 2). It should be noted that these are chronic eyelid conditions and so need regular and long-term commitment if management is to be successful.

Eyelid position correction: The eye's surface can also become dry because tears are not evenly distributed across the ocular surface. This can occur because of lumps on the surface of the eye, such as pingueculums or pterygia. For tears to be spread evenly over the surface, the eyelids must be in good contact with the eye surface. Eyelids that are floppy, lax or turned out (ectropion) can also



Figure 1: Warm the eyelids for at least 5 mins with a wheat or flaxseed bag.

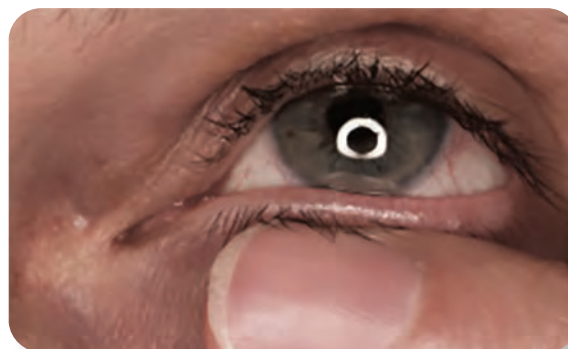


Figure 2: Gentle pressure outwards helps keep oils out of the gland.

worsen the tendency to dry eye. Whilst artificial tear drops can help, sometimes surgical correction of the eyelid position may be required.

Use artificial tears: Dry eye is most commonly managed with artificial tear supplements (drops). Most drops are best suited to aqueous deficient dry eye by boosting tear volume, but some newer products target the tear film oil layer to cater to those with evaporative dry eye. The products range in viscosity (runniness). Most gloopy eye drops provide the most prolonged relief from symptoms but cause transient blurring on application. This is well

tolerated by patients with moderate to severe dry eye, but those with milder symptoms might prefer a less viscous (a runnier) eye drop. Whilst generally safe to use, the main precaution is to restrict excessive use of preparations with preservatives as preservatives can cause ocular surface toxicity and worsen dry eye symptoms.

Consider punctal plugs: Punctal plugs are another solution for a dry eye that doesn't produce enough tears (aqueous deficient). Punctal plugging is when tiny silicone plugs are inserted in the drainage channels (puncta) on the inner

corners of your eyelid margins, stopping the tears (or tear supplements) from draining out, keeping the eye's surface wetter for longer.

Review medication side effects:

If prescribed medications cause significant dry eye, your general practitioner (GP) or specialist might consider alternatives. However, often the benefits of your medication outweigh the side effect of dry eye, and other means of managing the dry eye must be sought.

What does the future hold for dry eye: In May, Prof Jennifer Craig's presentation at the GNZ Professional Education Symposium 2021 focused on the intersection of dry eye and glaucoma. Professor Craig outlined why preserving the ocular surface is key in glaucoma patients, particularly for those with existing dry eye disease and in younger patients who may have decades of therapy ahead of them. New Zealand's lack of subsidised preservative free glaucoma medications was highlighted as a future challenge for ophthalmologists treating glaucoma. It is GNZ's hope that we will see a wider variety of eye drops available for New Zealand glaucoma patients in the not-too-distant future.

There are many reasons to be hopeful as researchers, leading pharmaceutical companies, and clinicians are continuously developing new products or improving existing ones. Your eye specialist will be able to advise you, on an individual basis, about the suitability of new products for you, as they become available locally.

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The Gordon Sanderson Scholars 2020/2019

Recent recipients of the Gordon Sanderson Scholarship, Aqeeda Singh and Blair Lowry discuss their research projects and the important role data plays in the future of glaucoma research and treatment.



Aqeeda Singh

2020 Scholarship recipient Aqeeda Singh outlines her project working with the Dunedin Study. This world-renowned longitudinal study has been following the lives of 1037 babies born in the greater Dunedin metropolitan area between 1 April 1972 and 31 March 1973 to understand the factors leading to significant health and social

outcomes in life. These participants have been interviewed, examined and investigated in numerous ways throughout their lives, with a very high follow-up rate of 94% at 45 years of age.

Now in middle age, the participants were recently subject to a broad range of tests relating to their eyes, including intraocular pressure (IOP), retinal photography, optical coherence tomography (OCT) and visual field testing; this study has examined their eye-related data to measure the prevalence of glaucoma in this cohort.

This is one of the first population-based studies of glaucoma prevalence using modern OCT scans to determine a diagnosis of early glaucoma (when visual fields are normal) and has implications for public health planning in New Zealand.

Two independent ophthalmologists examine the optic disc photographs, masked to other data, and then

suspicious discs were re-examined with more information such as IOP, corneal thickness, OCT and visual fields.

The prevalence amongst 45-year-olds was 0.79%, in line with other populations. At this age, the prevalence was low, so it was impossible to ascertain risk factors or undertake subgroup analyses such as risk by ethnicity. Interestingly, those with raised IOP had a low rate of glaucoma; nearly all of the cases we found had normal IOP. None of the glaucoma cases had significant visual field loss, so these are mild early cases.

Aqeeda said, "We are interested in modelling how the use of OCT at a population level detects many low-risk glaucoma suspect cases, generating a new burden for monitoring to prevent rare cases of vision loss". She is grateful to Prof Helen Danesh-Meyer for the support and direction with this project, as well as the Gordon Sanderson Scholarship committee and Glaucoma NZ.



Blair Lowry

Blair Lowry was a Gordon Sanderson Scholarship recipient in 2019, where he conducted research investigating the epidemiology of acute angle-closure glaucoma presentations over 2018 and 2019 at Greenlane Clinical Centre. Glaucoma is one of the leading causes of preventable blindness in New Zealand, which can be treated with effective treatment such as intraocular pressure lowering medications or surgical treatment to correct the rise in intraocular pressure. Acute angle closure glaucoma is an uncommon presentation, but it is a medical emergency that requires rapid identification and management to prevent blindness when it occurs.

My research focused on identifying important demographic parameters that might identify which groups are more susceptible to developing an acute angle closure crisis, and ideally, those at-risk populations. Through our sample recruitment, we included sixty-five cases of AACG from Greenlane Clinical Centre over the two years. We looked at the demographic parameters between these cases: age, sex, ethnicity, family history of glaucoma, and we will be looking at follow-up data of these individuals.

In our sample population, AACG mostly affected those greater than 40 years of age, with the mean age of AACG episodes being 63 years of



Glaucoma NZ Professional Symposium Summary

Glaucoma NZ Symposium 21: Dr Divya Perumal, Dr Hannah Kersten, Prof Jennifer Craig, Dr Sam Kain, Prof Helen Danesh-Meyer, Dr Keli Matheos.

Blair Lowry continued

age. Women had higher representation in this population than men (60% vs 40% respectively) – which is consistent with the literature. Caucasian individuals made up a significant proportion (47%) of cases, with Asian individuals being the 2nd most predominant ethnicity affected by AACG - due to anatomical predisposition to narrower angles – with 30% of the cases. Pasifika was represented quite significantly in this data, with 14% of Pacific people being affected by AACG, whilst Pasifika people represent 8.1% of the NZ population based on the 2018 census. 9% of the AACG cases were Māori individuals, which is lower than the reported 16.5% of Māori within the NZ population. A link between family history of glaucoma and AACG could not be established within this population as data is limited by the underreporting of family history of the cases. However, the literature identifies that there is a link between positive family history of glaucoma and AACG.

Blair would like to thank his supervisors Dr Hussain Patel and Prof Helen Danesh-Meyer for their support and direction with this project, as well as the Gordon Sanderson Scholarship committee and Glaucoma NZ for helping contribute to more ophthalmological research and a greater understanding of acute angle closure glaucoma in New Zealand. “It’s been an honour working with them all and getting a taste of what ophthalmology entails”.

It was a case of third time lucky for the 2021 Glaucoma New Zealand Symposium: After two postponements over the past year, the Symposium successfully went ahead at Alexandra Park, Auckland, on the 16th of May. Given that in-person educational events have been few and far between recently, it was a fantastic opportunity for colleagues, students and sponsors to network, and to enjoy some excellent glaucoma education from a diverse range of speakers. This year, GNZ also provided the opportunity for delegates to attend remotely (with 26 people live-streaming the event, in addition to the 100+ attendees in the room), which prompted positive feedback.

Professor Graham Lee, presenting via (almost completely glitch-free) Zoom from Brisbane, was this year’s keynote speaker. Professor Lee gave a whistle stop tour of the red eye in glaucoma. As he is fellowship trained in both cornea and glaucoma, he is uniquely qualified to cover this topic. The presentation covered a wide variety of causes and cases of red eye in glaucoma, ranging from sight-threatening emergencies (acute angle closure) to mild iatrogenic red eye (this part of the talk also highlighted the wider range of preservative-free medications available for glaucoma treatment in Australia compared with New Zealand). Prof Lee raised the importance of asking about a patient’s nutrition, and covered an interesting case of reduced conjunctival healing in a patient suffering from alcoholism.

The day concluded with the the Gordon Sanderson Scholarship presentation. The scholarship, named for Associate Professor Gordon Sanderson, one of the founding trustees of GNZ and a titan of eye care in New Zealand, has been awarded every year since Gordon passed away in 2017.

Overall, it was a fantastic day of glaucoma education. Special thanks goes to Pippa and Tersia at GNZ for organising the event, all of the excellent speakers, and to the sponsors, AFT Pharmaceuticals, Allergan, Designs for Vision, Device Technologies, Glaukos, Humanware, Medix21, OIC and Tristel for their generous support of the event.

If you want to know more please go to glaucoma.org.nz to read our in depth report and learn more from the eye health professionals who presented at the symposium.

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Ref 1 Jones, Lyndon, et al. "TFOS DEWS II management and therapy report." The ocular surface 15.3 (2017):575-628.

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Content in 'Eyelights' is intended to help readers understand glaucoma. Every effort is made to ensure the accuracy of this information. This information is not a substitute for the advice and recommendations of health professionals. Always consult a health professional prior to any decision regarding your eyes or other health concerns.

