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Corneal Disease Report

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- Clinical Pearls in Corneal Foreign Body Removal, p. 32
- Fine-Tune Your Corneal Disease Diagnostic Skills, p. 42
- Improving the Gold Standard for CXL, p. 50
- Piecing Together the HSVK Puzzle, p. 56

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Pivotal study designs: Two Phase 3, randomized, multicenter, parallel-group studies, APOLLO and LUNAR, evaluating noninferiority of once-daily VYZULTA vs twice-daily timolol maleate 0.5% in patients with open-angle glaucoma or ocular hypertension. Primary endpoint was IOP measured at 9 assessment time points in study eye. APOLLO (VYZULTA, n=284; timolol, n=133) and LUNAR (VYZULTA, n=278; timolol, n=136).

INDICATION
VYZULTA® (latanoprostene bunod ophthalmic solution), 0.024% is indicated for the reduction of intraocular pressure (IOP) in patients with open-angle glaucoma or ocular hypertension.

IMPORTANT SAFETY INFORMATION
- Increased pigmentation of the iris and periorbital tissue (eyelid) can occur. Iris pigmentation is likely to be permanent
- Gradual changes to eyelashes, including increased length, increased thickness, and number of eyelashes, may occur. These changes are usually reversible upon treatment discontinuation
- Use with caution in patients with a history of intraocular inflammation (iritis/uveitis). VYZULTA should generally not be used in patients with active intraocular inflammation
- Macular edema, including cystoid macular edema, has been reported during treatment with prostaglandin analogs. Use with caution in aphakic patients, in pseudophakic patients with a torn posterior lens capsule, or in patients with known risk factors for macular edema
- There have been reports of bacterial keratitis associated with the use of multiple-dose containers of topical ophthalmic products that were inadvertently contaminated by patients
- Contact lenses should be removed prior to the administration of VYZULTA and may be reinserted 15 minutes after administration
- Most common ocular adverse reactions with incidence ≥2% are conjunctival hyperemia (6%), eye irritation (4%), eye pain (3%), and instillation site pain (2%)

For more information, please see Brief Summary of full Prescribing Information on adjacent page.


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INDICATIONS AND USAGE
Initial U.S. Approval: 2017

ophthalmic use.
Iris color change may not be noticeable for several months to years. Typically, the brown pigmentation
pigmentation are not known.
of increased pigmentation, including permanent changes. The long-term effects of increased
periorbital tissue and eyelash changes are likely to be reversible in most patients. Patients
of VYZULTA, pigmentation of the iris is likely to be permanent, while pigmentation of the
melanocytes rather than to an increase in the number of melanocytes. After discontinuation
is administered. The pigmentation change is due to increased melanin content in the
pigmented tissues. The most frequently reported changes with prostaglandin analogs
VYZULTA® (latanoprostene bunod ophthalmic solution), 0.024% may cause changes to
5.1 Pigmentation
VYZULTA® (latanoprostene bunod ophthalmic solution), 0.024% may cause changes to
pigmented tissues. The most frequently reported changes with prostaglandin analogs have been increased pigmentation of the iris and periorbital tissue (eyelid).

Pigmentation is expected to increase as long as latanoprostene bunod ophthalmic solution is administered. The pigmentation change is due to increased melanin content in the melanocytes rather than to an increase in the number of melanocytes. After discontinuation of VYZULTA, pigmentation of the iris is likely to be permanent, while pigmentation of the periorbital tissue and eyelash changes are likely to be reversible in most patients. Patients who receive prostaglandin analogs, including VYZULTA, should be informed of the possibility of increased pigmentation, including permanent changes. The long-term effects of increased pigmentation are not known.

Iris color change may not be noticeable for several months to years. Typically, the brown pigmentation around the pupil spreads concentrically towards the periphery of the iris and the entire iris or parts of the iris become more brownish. Neither new nor freckles of the iris appear to be affected by treatment. While treatment with VYZULTA® (latanoprostene bunod ophthalmic solution), 0.024% can be continued in patients who develop noticeable increased iris pigmentation, these patients should be examined regularly [see Patient Counseling Information (17) in full Prescribing Information].

5.2 Eyelash Changes
VYZULTA may gradually change eyelashes and vellus hair in the treated eye. These changes include increased length, thickness, and the number of lashes or hairs. Eyelash changes are usually reversible upon discontinuation of treatment.

5.3 Intraocular Inflammation
VYZULTA should be used with caution in patients with a history of intraocular inflammation (iritis) and should not be used in patients with active intraocular inflammation as it may exacerbate this condition.

5.4 Macular Edema
Macular edema, including cystoid macular edema, has been reported during treatment with prostaglandin analogs. VYZULTA should be used with caution in aphakic patients, in pseudophakic patients with a torn posterior lens capsule, or in patients with known risk factors for macular edema.

5.5 Bacterial Keratitis
There have been reports of bacterial keratitis associated with the use of multiple-dose containers of topical ophthalmic products. These containers had been inadvertently contaminated by patients who, in most cases, had a concurrent corneal disease or a disruption of the ocular epithelial surface.

5.6 Use with Contact Lens
Contact lenses should be removed prior to the administration of VYZULTA because this product contains benzalkonium chloride. Lenses may be reinserted 15 minutes after administration.

6 ADVERSE REACTIONS
The following adverse reactions are described in the Warnings and Precautions section: pigmentation (5.1), eyelash changes (5.2), intraocular inflammation (5.3), macular edema (5.4), bacterial keratitis (5.5), use with contact lens (5.6).

6.1 Clinical Trials Experience
Because clinical trials are conducted under widely varying conditions, adverse reaction rates observed in the clinical trials of a drug cannot be directly compared to rates in the clinical trials of another drug and may not reflect the rates observed in practice.
VYZULTA was evaluated in 811 patients in 2 controlled clinical trials of up to 12 months duration. The most common ocular adverse reactions observed in patients treated with latanoprostene bunod were: conjunctival hyperemia (6%), eye irritation (4%), eye pain (3%), and instillation site pain (2%). Approximately 0.6% of patients discontinued therapy due to ocular adverse reactions including conjunctival hyperemia, conjunctival irritation, eye irritation, eye pain, conjunctival edema, vision blurred, punctate keratitis and foreign body sensation.

8 USE IN SPECIFIC POPULATIONS
8.1 Pregnancy
Risk Summary
There are no available human data for the use of VYZULTA during pregnancy to inform any associated risks.

Latanoprostene bunod has caused miscarriages, abortion, and fetal harm in rabbits. Latanoprostene bunod was shown to be abortifacient and teratogenic when administered intravenously (iv) to pregnant rabbits at exposures ≥ 0.28 times the clinical dose. Doses ≥ 20 μg/kg/day (23 times the clinical dose) produced 100% embryofetal lethality. Structural abnormalities observed in rabbit fetuses included anomalies of the great vessels and aortic arch vessels, domed head, sternobrachial and vertebral skeletal anomalies, limb hyperextension and malrotation, abdominal distension and edema. Latanoprostene bunod was not teratogenic in the rat when administered iv at 150 mcg/kg/day (67 times the clinical dose) [see Data].
The background risk of major birth defects and miscarriage for the indicated population is unknown. However, the background risk in the U.S. general population of major birth defects is 2 to 4%, and of miscarriage is 15 to 20%, of clinically recognized pregnancies.

Data

Animal Data
Embryofetal studies were conducted in pregnant rabbits administered latanoprostene bunod daily by intravenous injection on gestation days 7 through 19, to target the period of organogenesis. The doses administered ranged from 0.24 to 80 mcg/kg/day. Animal data were obtained at doses ≥ 0.24 mcg/kg/day latanoprostene bunod (0.28 times the clinical dose, on a body surface area basis, assuming 100% absorption). Embryofetal lethality (resorption) was increased in latanoprostene bunod treatment groups, as evidenced by increases in early resorptions at doses ≥ 0.24 mcg/kg/day and late resorptions at doses ≥ 6 mcg/kg/day (approximately 7 times the clinical dose). No fetuses survived in any rabbit pregnancy at doses of 20 mcg/kg/day (23 times the clinical dose) or greater. Latanoprostene bunod produced structural abnormalities at doses ≥ 0.24 mcg/kg/day (0.28 times the clinical dose). Malformations included anomalies of sternum, coarctation of the aorta with pulmonary trunk dilation, retroperitoneal subclavian artery with absent brachiocephalic artery, domed head, forepaw hyperextension and hindlimb malrotation, abdominal distension/edema, and missing/fused caudal vertebrae.

An embryofetal study was conducted in pregnant rats administered latanoprostene bunod daily by intravenous injection on gestation days 7 through 17, to target the period of organogenesis. The doses administered ranged from 150 to 1500 mcg/kg/day. Maternal toxicity was produced at 1500 mcg/kg/day (670 times the clinical dose, on a body surface area basis, assuming 100% absorption), as evidenced by reduced maternal weight gain. Embryofetal lethality (resorption and fetal death) and structural anomalies were produced at doses ≥ 300 mcg/kg/day (174 times the clinical dose). Malformations included anomalies of the sternum, domed head, forepaw hyperextension and hindlimb malrotation, ventral anomalies and delayed ossification of distal limb bones. A no observed adverse effect level (NOAEL) was established at 150 mcg/kg/day (87 times the clinical dose) in this study.

8.2 Lactation
Risk Summary
There are no data on the presence of VYZULTA in human milk, the effects on the breastfed infant, or the effects on milk production. The development and health benefits of breastfeeding should be considered, along with the mother’s clinical need for VYZULTA, and any potential adverse effects on the breastfed infant from VYZULTA.

8.4 Pediatric Use
Use in pediatric patients aged 16 years and younger is not recommended because of potential safety concerns related to increased pigmentation following long-term chronic use.

8.5 Geriatric Use
No overall clinical differences in safety or effectiveness have been observed between elderly and other adult patients.

13 NONCLINICAL TOXICOLOGY
13.1 Carcinogenesis, Mutagenesis, Impairment of Fertility
Latanoprostene bunod was not mutagenic in bacteria and did not induce micronuclei formation in the in vivo rat bone marrow micronucleus assay. Chromosomal aberrations were observed in vitro with human lymphocytes in the absence of metabolic activation.

Latanoprostene bunod has not been tested for carcinogenic activity in long-term animal studies. Latanoprost acid is a main metabolite of latanoprostene bunod. Exposure of rats and mice to latanoprost acid, resulting from oral dosing with latanoprost in lifetime rodent bioassays, was not carcinogenic.

Fertility studies have not been conducted with latanoprostene bunod. The potential to impact fertility can be partially characterized by exposure to latanoprost acid, a common metabolite of both latanoprostene bunod and latanoprost. Latanoprost acid has not been found to have any effect on male or female fertility in animal studies.

13.2 Animal Toxicology and/or Pharmacology
A 9-month toxicity study administered topical ocular doses of latanoprostene bunod to one eye of cynomolgus monkeys: control (vehicle only), one drop of 0.024% bid, one drop of 0.04% bid and two drops of 0.04% per dose, bid. The systemic exposures are equivalent to 4.2-fold, 7.9-fold, and 13.5-fold the clinical dose, respectively, on a body surface area basis (assuming 100% absorption). Microscopic evaluation of the lungs after 9 months observed pleural/subpleural chronic fibrosis/inflammation in the 0.04% dose male groups, with increasing incidence and severity compared to controls. Lung toxicity was not observed at the 0.024% dose.


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Mask-associated Dry Eye Affects 70% of Healthcare Providers

Given its high prevalence in light of COVID-19, efforts to address this condition are necessary.

The pandemic has had a deleterious impact on daily life in a variety of ways, including the ongoing use of masks and other personal protective equipment—particularly for individuals working in healthcare. As a result, there has been an increased interest in understanding the negative effects of regular mask usage. In a recent prospective, cross-sectional study, researchers sought to better understand the prevalence of self-reported mask-associated dry eye in healthcare professionals and the possible risk factors for this condition.

The study authors created a web-based, self-administered questionnaire that included 12 questions about mask-associated dry eye and its risk factors. It was sent to 437 healthcare professionals; of these, 333 were included in the study. The data collected included demographic and clinical characteristics. Participants who reported at least one symptom of dry eye were deemed to potentially have mask-associated dry eye and asked to undergo a clinical evaluation.

Findings revealed that the prevalence of self-reported mask-associated dry eye among individuals working in healthcare was 70%. The remaining 30% said they did not feel any discomfort while wearing a mask. Additionally, 19.2% noted that they already had at least one symptom of dry eye before they started using masks. However, 90.6% of these individuals also reported that their complaints associated with dry eye were exacerbated after they began wearing masks during the pandemic.

Overall, 81.9% of the study participants with mask-associated dry eye symptoms said their work performance was affected. Of the participants who had self-reported mask-associated dry eye and agreed to a clinical examination (n=195), 30.7% had aqueous-type dryness with staining on the ocular surface with fluorescein. In terms of potential risk factors, the researchers reported that having at least one of the symptoms of dry eye while not wearing a mask and older age were both associated with the condition.

This research suggests, according to the authors, a high prevalence of self-reported mask-associated dry eye among medical professionals, which was substantiated by clinical findings in a number of participants. These results highlight the importance of addressing this issue in eyecare practices.

“It is likely that COVID-19 will be in our lives for years to come, which would only make the problems that the healthcare professionals face, such as the mask-associated dry eye in question, worse if not addressed,” they concluded. “Hence, any symptoms of healthcare professionals to this effect should not be ignored during the period of pandemic in particular. Implementation of measures to reduce the symptoms related to dry eye will likely positively affect the work performance of healthcare professionals.”


IN BRIEF

Many Seniors Miss Out on Assistive Apps for Low Vision. Though dozens of downloadable apps are available as visual assistive aids for low vision patients, authors of a recent article determined the population most in need of these tools—adults over age 55—aren’t taking advantage. Explaining the value of these aids to low vision patients may help increase rates of usage.

One review of low vision exam records from the UCLA Vision Rehabilitation Center found that 90% of low vision patients have a smartphone, yet only 6% use visual assistive apps. To help get some answers as to why these apps are underutilized by patients over age 55, researchers are currently conducting a randomized clinical trial called Community Access through Remote EyeSight (CARE) on several free visual assistive apps, including SuperVision+2, Seeing AI3 and Aira.

“The primary reason for not using these three visual assistive apps was a lack of awareness of such apps, reported by 63% of the first 50 participants in the CARE trial, whose mean age was 73,” they noted in the study. According to the study, “The second most commonly reported reason for not using these apps was because they didn’t know how and had not received training, indicated by roughly one in five participants (21%).” These findings suggest this patient population has a lack of awareness about visual assistive apps that lends to their underutilization. The team recommended providing specific information to visually impaired patients about the mobile accessibility apps available to them.

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1To limit burning when using contact lenses, remove contacts, apply drops, then insert contacts.

Retinal Thinning May Occur After FLACS

Other research found that patients with prediabetes may also suffer loss of the RNFL.

Few studies have been conducted reporting the effect of femto-second laser-assisted cataract surgery (FLACS) on the posterior segment, and they have shown differing results. Between the surgery growing in popularity and the safety concerns it brings for the optic nerve, authors recently aimed to compare changes in the peripapillary retinal nerve fiber layer (pRNFL) after FLACS and conventional phacoemulsification cataract surgery.

A total of 261 patients with age-related cataract scheduled for FLACS (222 eyes) or conventional phaco (39 eyes) were included.1 Average and quadrant pRNFL thicknesses were measured using OCT before surgery and one, three and six months post-op.

The rate of clinically significant thinning six months post-op was higher in the FLACS group (17.5%) than in the phaco group (5.1%), but there were no significant differences between the groups for any of the quadrants or for average pRNFL. According to the researchers, this may be due to the insufficiency of the statistical power to reveal differences between the groups due to the small phaco group size.

FLACS eyes demonstrated a stable decrease of average pRNFL thickness throughout and a gradual decrease in pRNFL thickness up to six months in all quadrants except the temporal quadrant where the decrease stabilized after three months. “This indicates that pRNFL thinning completes first in the temporal quadrant,” the authors wrote. “The consistent decrease in pRNFL thickness in all quadrants and in the average pRNFL of FLACS eyes suggests that pRNFL thinning may occur after surgery.”

Although no relationship was found between vacuum time and decline in pRNFL thickness, the authors aren’t ruling out the possibility that an increase in IOP during docking of the femto laser may be the primary factor influencing pRNFL thinning.

Patient pRNFL status may be another factor. “Preoperative pRNFL thickness was associated with temporal quadrant thinning, corresponding to the location where pRNFL thinning was completed,” they wrote. “This suggests that eyes with thinner pRNFL can be prone to structural damage in the temporal quadrant after FLACS, which can have important implications regarding the use of the surgery in glaucoma patients, with generally thinner pRNFL measurements.”

Researchers in another study revealed a second population at risk of retinal thinning after examining the retinas of patients with elevated blood sugar.2 They observed that many showed thinning of the RNFL despite the absence of obvious vascular damage. While sometimes overlooked, diabetic retinopathy (DR) is not exclusive to patients with diabetes; rather, DR can cause early neuroretinal changes in prediabetes patients before developing into overt retinopathy.

Fifty patients with diagnosed prediabetes and 50 healthy controls were included in the study. The researchers measured RNFL using spectral-domain OCT. Not only did prediabetic patients have thinning in the temporal quadrant, but most showed loss in all four. The difference between the RNFL thicknesses of both groups was statistically significant. The average RNFL thicknesses measured in each group are shown in Table 1.

“The prevalence of prediabetes varies between 19.8% and 34.6% and is more common than diabetes,” the authors wrote. “In about two out of three prediabetic cases, the metabolic disorder progresses to obvious diabetes in later life. However, microvascular and macrovascular complications due to diabetes may occur in prediabetes even before diabetes develops,” they noted.

As an optometrist, you have the opportunity to recognize these early signs of disease in your patients before progression or adverse outcomes occur. Be vigilant when performing retinal exams and RNFL thickness measurements on all your patients, not only those with diabetes.

Table 1. Average RNFL Thicknesses in Prediabetics vs. Controls

<table>
<thead>
<tr>
<th></th>
<th>Mean RNFL Thickness</th>
<th>Inferior Quadrant</th>
<th>Superior Quadrant</th>
<th>Nasal Quadrant</th>
<th>Temporal Quadrant</th>
</tr>
</thead>
<tbody>
<tr>
<td>Prediabetes Group</td>
<td>94.7μm</td>
<td>120μm</td>
<td>112.3μm</td>
<td>71μm</td>
<td>65.3μm</td>
</tr>
<tr>
<td>Control Group</td>
<td>98.9μm</td>
<td>128μm</td>
<td>116.3μm</td>
<td>77μm</td>
<td>71.2μm</td>
</tr>
</tbody>
</table>

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Strabismus Increases Risk of Mental Illness

A second study also showed that those with the condition reported a lower quality of life compared with visually normal children.

Children with strabismus often have decreased visual function, including difficulties with schoolwork and sports. They have also reported problems with self-image and teasing for eye misalignment, glasses and/or patching. Researchers recently investigated whether patients with strabismus may have a higher risk of developing mental illness, reporting that this demographic has higher odds of having anxiety, schizophrenia, bipolar and depression compared with children without eye disease. There was no strong association with substance use.

The study used data from over 12 million patients in a longitudinal commercial insurance claims database 18 years and younger at the time of their strabismus diagnosis (50.8% boys, mean age: eight years old). Adjusted odds ratios for the association of mental illnesses with strabismus were 2.01 for anxiety, 1.83 for schizophrenia, 1.64 for bipolar, 1.61 for depression and 0.99 for substance use.

The researchers noted that there was a moderate association between each strabismus type (esotropia, exotropia and hypertropia) with anxiety, schizophrenia, bipolar and depression; odds ratios ranged from 1.23 for the association between esotropia and bipolar to 2.70 for the association between exotropia and anxiety. However, they still found it unclear whether the type of strabismus has an association with the magnitude of mental illness risk.

“These results should alert ophthalmologists and optometrists to counsel children and their caregivers regarding the risk for mental illness,” the authors concluded in their paper. “They should consider incorporating a screening tool for mental health problems for patients with strabismus and referral of pediatric patients with strabismus for mental health evaluation.”

Not only may strabismus be associated with mood disorders, but another study showed that it also has an impact on overall life satisfaction, even after surgery.

To better understand how the condition impacts a child’s quality of life, as well as their family’s, researchers evaluated the functional vision and eye-related quality of life in children with strabismus and their parents using a recently developed and validated questionnaire called the Pediatric Eye Questionnaire (PedEyeQ). The answers revealed that strabismus has an important impact on affected children and their families. Scores were significantly lower in children with strabismus compared with healthy children. Interestingly, children with prior successful corrective strabismus surgery also had worse PedEyeQ scores.

The PedEyeQ was applied to non-operated children with strabismus (n=18), operated children with strabismus (n=24) and visually normal children (n=21). This instrument is composed of three components. The Child PedEyeQ component has two versions (for ages 5-11 or 12-17) and is completed by the children. The Proxy and Parent PedEyeQ components have three versions (ages 0-4, 5-11 or 12-17) and are answered by the parent or caregiver.

All PedEyeQ domain scores were significantly lower in children with strabismus compared with visually normal children, except the functional vision domain in the child version. Despite previous evidence of a positive effect (Continued on p. 11)
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INDICATION
Tyrvaya™ (varenicline solution) Nasal Spray is indicated for the treatment of the signs and symptoms of dry eye disease.

IMPORTANT SAFETY INFORMATION
Adverse Reactions
The most common adverse reaction reported in 82% of patients was sneezing. Events that were reported in 5-16% of patients were cough, throat irritation, and instillation-site (nose) irritation.


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REVIEW OF OPTOMETRY
APRIL 15, 2022
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IN BRIEF

Retinal Thickness Fluctuations During Anti-VEGF Associated With Poor Outcomes.
Researchers recently found that fluctuations in retinal thickness may be associated with ranibizumab treatment response in patients with neovascular age-related macular degeneration (nAMD). A total of 1,097 patients with nAMD received monthly or as-needed (PRN) ranibizumab (0.5mg or 2.0mg) for 24 months. Fluctuation scores were grouped into four categories and used to assess central foveal thickness variability, with a change of 50μm considered insignificant.

Patients with lower fluctuations scores had the greatest vision gains at month 24, with changes from baseline of 9.0 to 10.8 and 8.7 to 10.0 letters in the monthly and PRN arms, respectively. Corresponding changes for high fluctuation patients were 6.7 and 6.5 letters, respectively. There were no differences between groups for association between fluctuations in subretinal fluid height and BCVA gains. There were inverse correlations between magnitude of fluctuations in neurosensory and inner retina thickness and BCVA gains for the high fluctuation group vs. the others. Patients with more anatomical stability showed rapid, robust BCVA gains, while those with variability had worse responses.

“Fluctuations in retinal thickness during anti-VEGF treatment may be associated with treatment response,” the study authors concluded. “Patients with the greatest fluctuation scores may still be able to attain vision gains, although less than those among patients with the lowest fluctuation scores. Monitoring fluctuations in retinal thickness during treatment may help prognosticate the response.”


Diabetes Patients More Likely to Have Accommodative, Binocular Disorders. It’s been shown that diabetes mellitus (DM) may impair ocular accommodation and binocular vision performance. To investigate, researchers studied 30 patients with DM (hemoglobin A1c above 6.5%) and 30 controls (ages 18-40).

The study showed that aspects of accommodative and binocular vision performance are strongly affected by diabetes and there’s a correlation between accommodative/binocular disorders and severity of disease. A significant percentage of young individuals with DM have severe vision-related symptoms—26.6% compared with the control group at 6.6%.

The near point of convergence was more remote in the diabetic group. Mean accommodation amplitude and vergence facility and the median monocular accommodative facility were significantly lower in the diabetic group.

The authors suggested exams of young diabetic patients should not just focus on pathological anterior or posterior segment changes but also on accommodative and binocular functions.

Mental Illness and Eye Disease in Children, Teens

Anxiety, schizophrenia, bipolar and depression were potential considerations.

Multiple studies have found depression and anxiety to be associated with eye disease, but few have focused on other mental illnesses such as anxiety, schizophrenia, bipolar disorder and depression. In a recent study, researchers sought to evaluate the association between five eye diseases—glaucoma, cataract, congenital optic nerve disease, congenital retinal disease and blindness/low vision—and mental illness in children and teens.

Nearly 12 million children 19 years and younger at the time of eye diagnosis were included. Of the patients with at least one of the five eye diseases, 30.5% had glaucoma, 9.5% had cataract, 21.4% had congenital optic nerve disease, 26.9% had congenital retinal disease and 25.9% had blindness or low vision.

These diseases were associated with anxiety, schizophrenia, bipolar disorder and depression. The chance of having these psychiatric diagnoses was higher among children with at least one of the five eye diseases than among children without eye disease, with odds ratios ranging from 1.26 to 1.54.

Inverse associations were found with substance use, with the exception of non-significant associations between glaucoma and bipolar and between cataracts and substance use.

“It is possible that since our study grouped substance use disorder together, we were not able to distinguish which substances were involved; for example, it is possible that eye disease can increase the risk of certain substances but not others,” they explained in their report.

“Nearly 12 million children 19 years and younger at the time of eye diagnosis were included. Of the patients with at least one of the five eye diseases, 30.5% had glaucoma, 9.5% had cataract, 21.4% had congenital optic nerve disease, 26.9% had congenital retinal disease and 25.9% had blindness or low vision.”

The researchers noted in their paper, published in Clinical Ophthalmology, a wide range of visual symptoms, it is unlikely that this feature of disease underlies the association with serious eye disease observed here.  

Strabismus Increases Mental Illness, Lowers QOL

(Continued from p. 8)  

of strabismus surgery, kids with successful outcomes had worse PedEyeQ scores in many domains compared with visually normal children.

In the child component, the frustration/worry domain score was significantly lower in kids with strabismus. In the proxy component, frustration/worry and eyecare domain scores were both significantly lower than functional vision, bothered by eyes/vision and social domain scores. In the parent component, worry about child’s eye condition domain score was significantly lower than all other parents’ scores, while the impact on parent and family domain score was significantly higher than all others.

The researchers said they believe educational programs and psychosocial rehabilitation interventions that aim to improve the quality of life, participation in society and psychosocial functioning should be implemented in children with strabismus and their families.

“It is also possible that eye disease can increase the use of substances but not necessarily increase the risk of substance use disorder, as there is a threshold of substance use that is deemed pathologic.”

The authors noted that these studies reinforce the need to investigate the role eye disease plays in mental illness in children and teens; however, they have been limited in their definitions of mental illness beyond depression and anxiety, and in the type of eye diseases they studied.

“They have been limited in their definitions of mental illness beyond depression and anxiety, and in the type of eye diseases they studied.”


More than 77% of patients with primary angle-closure glaucoma (PACG) are Asian. The visual outcomes of this form of glaucoma can be devastating, and compared with open-angle glaucoma, the condition is associated with a three-fold increased risk of severe bilateral vision impairment. Its prevalence is also increasing globally, and particularly in Asia, it is emerging as a serious public health concern. Less is known about the incidence and progression of primary angle-closure disease (PACD), which precedes PACG.

To gain a better understanding of the incidence and risk factors of PACD in a multi-ethnic Asian population, researchers analyzed data over six years from the Singapore Epidemiology of Eye Diseases study, a population-based cohort study that included participants aged 40 or older living in Singapore. All completed ophthalmic examinations at baseline and six-year follow-up visits. Of 6,762 participants with valid data from both examinations, those included in the analysis were the 5,298 determined to be at risk for PACG and the 5,060 at risk for PACD.

The six-year age-adjusted PACD incidence in the cohort was 3.5%. For PACG, primary angle-closure and primary angle-closure suspect, it was 0.29%, 0.46% and 2.54%, respectively. In addition, the team found that 9.38% of primary angle-closure suspects progressed to primary angle-closure or PACG over the six-year period.

Several baseline parameters were identified as risk factors for PACD development, including increasing age per decade (odds ratio: 1.35), higher intraocular pressure (odds ratio: 1.04) and shallower anterior chamber depth (odds ratio: 1.11). On the other hand, late posterior subcapsular cataract was associated with a decreased likelihood of developing PACD (odds ratio: 0.60).

Another important finding: as the rate of cataract surgeries increased, the number of patients who developed PACD decreased. “Studies have shown that cataract removal is associated with a deeper anterior chamber depth and wider anterior chamber angles, which subsequently reduces the risk of PACD,” the researchers wrote. As an example, they pointed to the former Liwan Eye study, where “the highest PACD incidence observed could be largely explained by the much lower rate of cataract surgery reported in the study.”

Although age was shown to be a significant risk factor for PACD in this study, it, along with gender, didn’t have a significant effect on progression across disease stages.

The researchers concluded, “These findings may aid clinicians in deciding the frequency of monitoring or urgency of treatment and patient counseling.” In addition, they suggested the data could help “policy administrators ensure adequate resources are allocated to screening and intervention initiatives to address angle-closure glaucoma-related vision impairment.”

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Low Vitamin A May Cause Childhood Chalazion

By contrast, vitamin D had no correlation with the condition.

Evidence shows that vitamin deficiency is a key risk factor for children with chalazion. In a recent case-control study, researchers demonstrated that low serum vitamin A was significantly associated with this condition, suggesting vitamin A deficiency is a possible cause of chalazion among these patients.

The study authors enrolled 180 participants—90 chalazion patients and 90 healthy controls. Serum from collected blood samples was used to test levels of vitamins A and D (for the latter, 25(OH)D specifically was measured). Serum level of <0.7 μmol/L and <50 nmol/L were defined as a deficiency of vitamin A and D, respectively.

Baseline characteristics, including age and BMI, were similar between the two groups. The study authors reported that the average serum vitamin A levels in chalazion patients were significantly lower than in the control group. They observed no significant difference in the serum D levels of the chalazion and healthy cohorts.

The association between serum level of vitamin A and chalazion occurrence was also analyzed. Data showed that the percentage of deficiency among chalazion patients (52.2%) was considerably higher than their healthy counterparts (28.6%), which researchers note suggests a negative relationship between serum vitamin A level and chalazion instance.

"Due to lower storage of vitamin A in liver, children face a bigger risk of vitamin A deficiency than adults," the researchers wrote. "Vitamin A deficiency may cause Bitot’s spots, conjunctival and corneal desiccation, multiple yellow-white peripheral focal retinal pigment epithelium defects and finally lead to serious clinical manifestations such as dry eye disease, and night blindness."

They added, "Vitamin A plays a crucial role on inflammatory response. Lower levels of serum vitamin A lead to inflammation-related diseases including chalazion."

No significant difference in vitamin D deficiency was observed between the chalazion and control groups (58.9% vs 56.7%).

“We found that serum vitamin A levels were significantly lower in patients with chalazion than in control subjects, while the serum levels of 25(OH)D were not significantly different between the two groups,” the study authors concluded. “Furthermore, we found a negative relationship between the serum vitamin A level and the morbidity of chalazion in children. These data demonstrate that vitamin A deficiency is a potential risk factor for chalazion in children.”

While the link between chalazion and vitamin A deficiency in children has been established, the researchers noted a need for further investigation of the molecular mechanisms.


IN BRIEF

- **Case Report: Severe Dietary Restriction Leads to Fungal Keratitis.** Though rare in advantaged countries, vitamin A deficiency may still occur as a result of malnutrition or extreme dieting, as was the issue in a recent case report published in the British Medical Journal. An Australian woman in her 60s presented with bilateral corneal perforations and decreased visual acuity. Culturing revealed a yeast colony comprised of Candida albicans in her left cornea.

  Her diagnosis was corneal perforations secondary to bilateral fungal keratitis and hypovitaminosis A. The researchers wrote in their paper that her vitamin deficiency was likely due to a diet of nothing but bananas and yogurt.

  She underwent tectonic corneal grafts for the perforations and received retinyl palmitate as well as oral and topical voriconazole. She declined epithelial defect management, despite being informed of the risk of recurrent microbial keratitis, and did not change her diet. Her final vision was hand movement OS and 6/30 OD.

  The ocular sequelae of vitamin A deficiency include nyctalopia, corneal ulcer and scarring. **Ocular features may include Bitot’s spots (elevated, “foamy” conjunctival plaques) and conjunctival and corneal xerosis;** however, the authors noted in their paper that, like this case, those classically described features may not be seen on examination. Nevertheless, they wrote, “their absence shouldn’t preclude clinical suspicion of vitamin A deficiency.”

  "Early and specific questioning to explore the causative condition is imperative in routine clinical practice to avoid omitting the history of hypovitaminosis A,” they continued. “This case highlights the importance of considering a multifactorial cause in corneal perforations” and diligently obtaining a thorough case history. "Consider vitamin A deficiency in cases of corneal perforation or impending perforations, especially in the context of an unusual or nutrient-deficient diet,” the report concluded.

While treatment with VYZULTA® (latanoprostene bunod ophthalmic solution), 0.024% can be continued of increased pigmentation, including permanent changes. The long-term effects of increased who receive prostaglandin analogs, including VYZULTA, should be informed of the possibility of increased pigmentation, including permanent changes. The long-term effects of increased pigmentation are not known. Iris color change may not be noticeable for several months to years. Typically, the brown pigmentation around the pupal spread concentrically towards the periphery of the iris and the entire iris or parts of the iris become more brownish. Neither nevi nor freckles of the iris appear to be affected by treatment. While treatment with VYZULTA® (latanoprostene bunod ophthalmic solution), 0.024% can be continued in patients who develop noticeably increased iris pigmentation, these patients should be examined regularly (see Patient Counseling Information (17) in full Prescribing Information).

5.2 Eyelash Changes
VYZULTA may gradually change eyelashes and vellus hair in the treated eye. These changes include increased length, thickness, and the number of lashes or hairs. Eyelash changes are usually reversible upon discontinuation of treatment.

5.3 Intraocular Inflammation
VYZULTA should be used with caution in patients with a history of intraocular inflammation (iritis/uveitis) and should generally not be used in patients with active intraocular inflammation as it may exacerbate this condition.

5.4 Macular Edema
Macular edema, including cystoid macular edema, has been reported during treatment with prostaglandin analogs. VYZULTA should be used with caution in aphakic patients, in pseudophakic patients with a torn posterior lens capsule, or in patients with known risk factors for macular edema.

5.5 Bacterial Keratitis
There have been reports of bacterial keratitis associated with the use of multiple-dose containers of topical ophthalmic products. These containers had been inadvertently contaminated by patients who, in most cases, had a concurrent corneal disease or a disruption of the ocular epithelial surface.

5.6 Use with Contact Lens
Contact lenses should be removed prior to the administration of VYZULTA because this product contains benzalkonium chloride. Lenses may be reinserted 15 minutes after administration.

6 ADVERSE REACTIONS
The following adverse reactions are described in the Warnings and Precautions section: pigmentation (5.1), eyelash changes (5.2), intraocular inflammation (5.3), macular edema (5.4), bacterial keratitis (5.5), use with contact lens (5.6).

6.1 Clinical Trials Experience
Because clinical trials are conducted under widely varying conditions, adverse reaction rates observed in the clinical trials of a drug cannot be directly compared to rates in the clinical trials of another drug and may not reflect the rates observed in practice.

VYZULTA was evaluated in 811 patients in 2 controlled clinical trials of up to 12 months duration. The most common ocular adverse reactions observed in patients treated with latanoprostene bunod were: conjunctival hyperemia (6%), eye irritation (4%), eye pain (3%), and instillation site pain (2%). Approximately 0.6% of patients discontinued therapy due to ocular adverse reactions including ocular hyperemia, conjunctival irritation, eye irritation, eye pain, conjunctival edema, vision blurred, punctate keratitis and foreign body sensation.

8 USE IN SPECIFIC POPULATIONS
8.1 Pregnancy
There are no available human data for the use of VYZULTA during pregnancy to inform any associated risks.

Latanoprostene bunod has caused miscarriages, abortion, and fetal harm in rabbits. Latanoprostene bunod was shown to be abortifacient and teratogenic when administered intravenously (iv) to pregnant rabbits at exposures ≥ 20 μg/kg/day (23 times the clinical dose) produced 100% embryofetal lethality. Structural abnormalities observed in rabbit fetuses included anomalies of the great vessels and aortic arch vessels, domed head, sternebral and vertebral skeletal anomalies, limb malrotation, abdominal distension and edema. Latanoprostene bunod was not teratogenic in the rat when administered iv at 150 mcg/kg/day (67 times the clinical dose) (see Data). The background risk of major birth defects and miscarriage for the indicated population is unknown. However, the background risk in the U.S. general population of major birth defects is 2 to 4%, and of miscarriage is 15 to 20%, of clinically recognized pregnancies.

Data
Animal Data
Embryofetal studies were conducted in pregnant rabbits administered latanoprostene bunod daily by intravenous injection on gestation days 7 through 17, to target the period of organogenesis. The doses administered ranged from 0.24 to 80 mcg/kg/day. Abortion occurred at doses ≥ 0.24 mcg/kg/day latanoprostene bunod (0.28 times the clinical dose, on a body surface area basis, assuming 100% absorption). Embryofetal lethality (resorption) was increased in latanoprostene bunod treatment groups, as evidenced by increases in early resorptions at doses ≥ 0.24 mcg/kg/day and late resorptions at doses ≥ 6 mcg/kg/day (approximately 7 times the clinical dose). No fetuses survived in any rabbit pregnancy at doses of 20 mcg/kg/day (23 times the clinical dose) or greater. Latanoprostene bunod produced structural abnormalities at doses ≥ 0.24 mcg/kg/day (0.23 times the clinical dose). Malformations included anomalies of sternum, coarctation of the aorta with pulmonary trunk dilation, retroesophageal subclavian artery with absent brachiocephalic artery, domed head, forepaw hyperextension and hindlimb malrotation, abdominal distension/edema, and missing/fused caudal vertebrae.

An embryofetal study was conducted in pregnant rats administered latanoprostene bunod daily by intravenous injection on gestation days 7 through 17, to target the period of organogenesis. The doses administered ranged from 150 to 1500 mcg/kg/day. Maternal toxicity was produced at 1500 mcg/kg/day (670 times the clinical dose, on a body surface area basis, assuming 100% absorption), as evidenced by reduced maternal weight gain. Embryofetal lethality (resorption and fetal death) and structural anomalies were produced at doses ≥ 300 mcg/kg/day (174 times the clinical dose). Malformations included anomalies of the sternum, domed head, forepaw hyperextension and hindlimb malrotation, vertebral anomalies and delayed calcification of distal limb bones. A no observed adverse effect level (NOAEL) was established at 150 mcg/kg/day (87 times the clinical dose) in this study.

8.2 Lactation
Risk Summary
There are no data on the presence of VYZULTA in human milk, the effects on the breastfed infant, or the effects on milk production. The development and health benefits of breastfeeding should be considered, along with the mother’s clinical need for VYZULTA, and any potential adverse effects on the breastfed infant from VYZULTA.

8.4 Pediatric Use
Use in pediatric patients aged 16 years and younger is not recommended because of potential safety concerns related to increased pigmentation following long-term chronic use.

8.5 Geriatric Use
No overall clinical differences in safety or effectiveness have been observed between elderly and other adult patients.

13 NONCLINICAL TOXICOLOGY
13.1 Carcinogenesis, Mutagenesis, Impairment of Fertility
Latanoprostene bunod was not mutagenic in bacteria and did not induce micronucleus formation in the in vivo rat bone marrow micronucleus assay. Chromosomal aberrations were observed in vitro with human lymphocytes in the absence of metabolic activation.

Latanoprostene bunod has not been tested for carcinogenic activity in long-term animal studies. Latanoprost acid is a main metabolite of latanoprostene bunod. Exposure of rats and mice to latanoprost acid, resulting from oral dosing with latanoprost in lifetime rodent bioassays, was not carcinogenic.

Fertility studies have not been conducted with latanoprostene bunod. The potential to impact fertility can be partially characterized by exposure to latanoprost acid, a common metabolite of both latanoprostene bunod and latanoprost. Latanoprost acid has not been found to have any effect on male or female fertility in animal studies.

13.2 Animal Toxicology and/or Pharmacology
A 9-month toxicity study administered topical ocular doses of latanoprostene bunod to one eye of cynomolgus monkeys: control (vehicle only), one drop of 0.024% bid, one drop of 0.04% bid and two drops of 0.04% per dose, bid. The systemic exposures are equivalent to 4.2-fold, 7.9-fold, and 13.5-fold the clinical dose, respectively, on a body surface area basis (assuming 100% absorption). Microscopic evaluation of the lungs after 9 months observed pleural/subpleural chronic fibrosis/inflammation in the 0.04% dose male groups, with increasing incidence and severity compared to controls. Lung toxicity was not observed at the 0.024% dose.


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Pivotal study designs: Two Phase 3, randomized, multicenter, parallel-group studies, APOLLO and LUNAR, evaluating noninferiority of once-daily VYZULTA vs twice-daily timolol maleate 0.5% in patients with open-angle glaucoma or ocular hypertension. Primary endpoint was IOP measured at 9 assessment time points in study eye. APOLLO (VYZULTA, n=284; timolol, n=133) and LUNAR (VYZULTA, n=278; timolol, n=136).

INDICATION

VYZULTA® (latanoprostene bunod ophthalmic solution), 0.024% is indicated for the reduction of intraocular pressure (IOP) in patients with open-angle glaucoma or ocular hypertension.

IMPORTANT SAFETY INFORMATION

- Increased pigmentation of the iris and periorbital tissue (eyelid) can occur. Iris pigmentation is likely to be permanent

- Gradual changes to eyelashes, including increased length, increased thickness, and number of eyelashes, may occur. These changes are usually reversible upon treatment discontinuation

- Use with caution in patients with a history of intraocular inflammation (iritis/uveitis). VYZULTA should generally not be used in patients with active intraocular inflammation

- Macular edema, including cystoid macular edema, has been reported during treatment with prostaglandin analogs. Use with caution in aphakic patients, in pseudophakic patients with a torn posterior lens capsule, or in patients with known risk factors for macular edema

- There have been reports of bacterial keratitis associated with the use of multiple-dose containers of topical ophthalmic products that were inadvertently contaminated by patients

- Contact lenses should be removed prior to the administration of VYZULTA and may be reinserted 15 minutes after administration

- Most common ocular adverse reactions with incidence ≥2% are conjunctival hyperemia (6%), eye irritation (4%), eye pain (3%), and instillation site pain (2%)

For more information, please see Brief Summary of full Prescribing Information on adjacent page.


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This month’s cover photo was taken by optometrist Irving Martínez Navé, a reader from León, Guanajuato, Mexico. You can follow him on Instagram at @martinez_photography for more great clinical images from his patient base.
Do you want to have your clinical photos showcased in Review of Optometry? Write to us at editor@reviewofoptometry.com.
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Optometry’s Role in the Patient Journey

Gloria Chiu, OD, FAAO, FSLS
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USC Roski Eye Institute, USC Keck School of Medicine
Los Angeles

Keratoconus (KC) is a degenerative condition with onset in early adolescence. It is characterized by gradual thinning of the corneal stroma, causing a cone-shaped protrusion and worsening vision. As doctors of optometry, our top priority with these patients should be to manage their disease—and only secondarily to correct their vision.

A referral for corneal collagen cross-linking, which has been shown to halt progression in 92%-100% of cases, may be able to preserve vision. As with any surgical procedure, there is the potential for complications and cross-linking may not be right for everyone. After treatment, patients will still need regular optometric care. Follow-up care is similar to that with any surgical procedure, with 10%-20% of cases requiring a penetrating keratoplasty (PKP). When patients reach the advanced stages of keratoconus, it becomes a debilitating disease that affects every aspect of their lives. Worsening KC severity is associated with significant declines in reading, mobility, and emotional well-being quality of life (QoL) scores. The impact on QoL can be even greater than that of retinal diseases and can be felt even when one eye still has good vision so it is important that patients get help as early as possible.

In the U.S., when cross-linking is performed with the iLink™ platform (Glaukos), the only FDA-approved cross-linking system, it is generally covered by insurance for 96% of those with commercial insurance. In a recent simulation model, treatment with iLink™ was found to be highly cost effective, resulting in a 26% reduction in PKPs and patients spending 28 fewer years in the advanced stages of KC. Young patients who can be treated early while their vision is still good have the most to gain.

That’s where optometrists’ role becomes so critical. Our awareness of early progressive KC signs and risk factors can be nothing short of life changing for that young myope in our chair. There is no need to wait until a patient has lost vision or has slit lamp signs (e.g., thinning or striae) to refer for a more in-depth KC evaluation. It is standard of care to intervene with cross-linking upon detection of progression.

Advanced tomography/topography provides the most sensitive and accurate diagnostic information. However, there are a number of signs and symptoms that should heighten suspicion of KC and prompt further testing, either in the practice or by referral. These include myopic shift, rapidly changing astigmatism, vision that won’t correct to 20/20 (with no other known reason), distorted mires on examination, and scissoring or an irregular retinoscopy reflex. Patients with a history of eye rubbing, connective tissue disease, Down syndrome, or family history of KC are also at higher risk.

By promptly referring these patients for further testing and, if warranted, iLink™ cross-linking treatment, optometrists are uniquely positioned to protect and preserve patients’ vision over their entire lifetime.

KEY TAKEAWAYS

- Cross-linking with the only FDA-approved iLink™ System can stop or slow progressive keratoconus.
- Early diagnosis and treatment are essential to preserve as much vision as possible.
- Optometrists are uniquely positioned to change lives and protect vision by identifying at-risk patients in the mild stages of the disease.

With Cross-Linking

26% fewer PKPs  28 fewer years in late-stage KC
Debating the Value of Genetic Testing

Early accounts of AvaGen’s role in clinical practice stimulate discussion and highlight the need for greater dialog.

Editor’s note: The January/February edition of Review of Cornea & Contact Lenses included a feature on genetic testing in keratoconus that included anecdotal impressions from several optometrists, as well as commentary from representatives of Avellino, makers of the AvaGen test. In one section of the discussion, Brian Chou, OD, reported receiving differing results on repeat testing of the same patient.

Here, representatives of Avellino offer elaboration and Dr. Chou responds to the new information.

Genetic Testing for Keratoconus Will Continue to Evolve, but Patients Can Benefit Now
By Mile Brujic, OD, and Sarvari Panchumarthi, PhD

We want to thank the editorial team for their article, “KCN Genetic Testing: Where Does It Fit In?”

One area optometrists should be devoting an increasing amount of attention and interest to is genetic testing. As with many new technological advancements in eye care, an increasing understanding of how these tests work and how to integrate them in the practice of optometry will be of the utmost importance today and in the next several years.

AvaGen by Avellino helps detect the genetic risk for keratoconus and the presence of corneal dystrophies. A rather new type of test to enter the field, there are three main aspects to it that are critical to help understand its clinical utility.

First is understanding the polygenic risk score (PRS) for keratoconus. The AvaGen PRS scale ranges from 0 to 100 and is divided into low-, medium- and high-risk segments. The risk score for a patient broadly falls into one of those three segments/categories. As an example, a polygenic risk score of 9 and a score of 22 are both in the low genetic risk category.

The PRS is based on the number and types of multitudes of genetic variants found in each tested individual. As long as the PRS remains in the same risk category, the minor differences in the score number are not indicative of significant differences in disease risk. Via next-generation sequencing (the technology used in the AvaGen genetic test), the analytical sensitivity and specificity of the test is 100%.

With respect to Dr. Chou’s testing of AvaGen using the same sample patient’s DNA with three different names, dates of birth and ethnicities, we would like to point out that the difference between a PRS of 9 and 22 (found in the first and second samples of the patient) is not clinically meaningful. Both scores place the patient’s genetic risk in the same category: low-risk. The genes identified in the risk of keratoconus are always detected, and if a specific variant used in the PRS calculation is ‘dropped out’ due to low quality of read depth, the PRS value would be expected to differ slightly.

Avellino does quality control by running each patient’s sample in triplicates in its test process for each run. In the third testing of the same patient, which was received in mid-October 2021, the PRS yielded a higher value (61), as the company released an improved version of the PRS analysis in this same month.

Second, understanding how a polygenic risk score is developed for complex genetic diseases is key. Complex genetic diseases occur due to the cumulative effect of several single-nucleotide variants (SNVs) of very small effect size. Unlike monogenic diseases (where the presence of a single pathogenic variant yields a more definitive predisposition for the disease), polygenic diseases are more complex in nature. There is always an interplay of genes and environment in the manifestation of a polygenic disease. This makes the prediction of disease risk in polygenic diseases more elusive.

As genomics knowledge improves for a certain disease, the polygenic risk score’s predictive value can be refined and improved over time. For AvaGen,
INDICATIONS
DEXTENZA is a corticosteroid indicated for:

• The treatment of ocular inflammation and pain following ophthalmic surgery.
• The treatment of ocular itching associated with allergic conjunctivitis.

IMPORTANT SAFETY INFORMATION
CONTRAINDICATIONS
DEXTENZA is contraindicated in patients with active corneal, conjunctival or canalicular infections, including epithelial herpes simplex keratitis (dendritic keratitis), vaccinia, varicella; mycobacterial infections; fungal diseases of the eye, and dacryocystitis.

WARNINGS AND PRECAUTIONS
Intraocular Pressure Increase - Prolonged use of corticosteroids may result in glaucoma with damage to the optic nerve, defects in visual acuity and fields of vision. Steroids should be used with caution in the presence of glaucoma. Intraocular pressure should be monitored during treatment.

Bacterial Infections - Corticosteroids may suppress the host response and thus increase the hazard for secondary ocular infections. In acute purulent conditions, steroids may mask infection and enhance existing infection.

Viral Infections - Use of ocular steroids may prolong the course and may exacerbate the severity of many viral infections of the eye (including herpes simplex).

ADVERSE REACTIONS
Ocular Inflammation and Pain Following Ophthalmic Surgery
The most common ocular adverse reactions that occurred in patients treated with DEXTENZA were: anterior chamber inflammation including iritis and iridocyclitis (10%), intraocular pressure increased (6%), visual acuity reduced (2%), cystoid macular edema (1%), corneal edema (1%), eye pain (1%), and conjunctival hyperemia (1%). The most common non-ocular adverse reaction was headache (1%).

Other Potential Corticosteroid Complications - The initial prescription and renewal of the medication order of DEXTENZA should be made by a physician only after examination of the patient with the aid of magnification, such as slit lamp biomicroscopy, and, where appropriate, fluorescein staining. If signs and symptoms fail to improve after 2 days, the patient should be re-evaluated.

Fungal Infections - Fungus invasion must be considered in any persistent corneal ulceration where a steroid has been used or is in use. Fungal culture should be taken when appropriate.

Delayed Healing - Use of steroids after cataract surgery may delay healing and increase the incidence of bleb formation.

References:

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ABOUT RICK

Rick Bay served as the publisher of the Review Group for more than 20 years. To those who worked for him, he was a leader whose essence was based in a fierce and boundless loyalty. To those in the industry and the professions he served, he will be remembered for his unique array of skills and for his dedication to exceeding the expectations of his customers, making many of them fast friends.

The Rick Bay Foundation for Excellence in Eyecare Education

(The Rick Bay Foundation for Excellence in Eyecare Education is a nonprofit, tax-exempt organization under section 501(c)(3) of the Internal Revenue Code.)
the PRS was updated in October 2021 with refinements in its calculation due to identification of a fewer number of more prevalent variants (common variants), making the test more refined in the information it generates. Ethnicity differences were also accounted for in variant selection in the new version; therefore, ethnicity details are no longer required to estimate individual polygenic risk score, as they were in the calculation.

Last, with complex genetic diseases, correlation is not causation. A high PRS for a given disease may never see the disease manifest and vice-versa. This is due to the interplay between genetic risk factors and environment/lifestyle. Likewise, someone with a low PRS may experience a given disease, as their lifestyle and other environmental factors may trigger the disease due to non-genetic causes.

This is where using our current clinical skill set to detect conditions such as keratoconus—including refractive error analysis, slit lamp examination, keratometry, topography, tomography, as well as the knowledge and skill of the eyecare practitioner (ECP)—play a crucial role in patient care. A genetic test should never be used in isolation or as the stand-alone diagnostic method for monogenic or polygenic diseases.

A resource for ECPs is the availability of genetic counselors, who can further help interpret results of a patient’s genetic test, including PRS, and provide additional guidance, such as helping determine if it is appropriate to test family members. Avellino provides this service for both ECPs and patients.

Genetic testing is a valuable data source for the medical community, with the eyecare field being no exception. While we may be in the early stages of this journey, patients can benefit now from our collective expertise and knowledge of the human genome.

Mile Brujic, OD, is a practicing doctor at Premier Vision Group in Bowling Green, OH. He provides consulting services to Avellino.

Sarcvari Panchumarthi, PhD, is a molecular geneticist with more than 20 years of research experience in various subdisciplines of genetics, in particular the genetics concerning cancer, cardiovascular, neurological and ophthalmological disorders.

Progress Toward Answers is Welcome—But Questions Remain
By Brian Chou, OD

I offer my thanks to Drs. Brujic and Panchumarthi for their interest in my report. Their explanation about the unusual AvaGen results I received provides insight and also raises new questions.

Still left unanswered: what is the explanation for the same keratoconus subject’s identified risk genes getting reported as different each of the three times AvaGen was run? The revised polygenic scoring implemented in October 2021 may explain the subject’s higher polygenic risk score of 61 vs. the prior PRS values of 9 and 22. However, the KCN-associated gene profiles themselves changed from test to test. Was this also a function of the update to the PRS algorithm? In other words, were the same genetic variants detected in all three tests but reported differently to reflect updates to the PRS algorithm?

I wonder, too, if other clinicians using AvaGen were notified of the scoring change. I learned of this by reading the companion sidebar Q&A to my own report. Due to the potentially high stakes of medical testing, future modifications to the PRS algorithm should include contemporaneous proactive notice to ECPs, not notice after the fact.

I agree that the test-retest PRS change from 9 to 22 would not have altered clinical decision-making, as both values were within AvaGen’s “low risk” category; even so, how would a clinician interpret a test-retest PRS change of the same increment, if for instance the PRS shifted from 22 to 35, thereby moving into the “moderate” risk category?

Although high analytical validity is requisite and expected for CLIA certification, it does not inform the ECP, nor patient, about the more important clinical validity and utility.

I look forward to Avellino and others elucidating these areas. A pure genetic test for keratoconus that is clinically valid and useful is welcome, even if epigenetics plays a notable role.

Dr. Chou practices at ReVision Optometry, a referral clinic for keratoconus and scleral lenses in San Diego. He is a past recipient of the National Keratoconus Foundation’s Top Doctor award.

Is the Joy in Cataract Surgery Fading?
By Don Stover, OD

Cataract surgery 40 years ago meant two months post-op of branded Pred Forte 1% QID, no NSAID and a week of antibiotics. There were limited cases of cystoid macular edema (CME) because it was an all-surgical procedure with no phacoemulsification. High astigmatism (above three diopters) was common, but patients—typically with 20/20+ corrected acuity—did not complain and strangely did not wear their glasses as much as would be predicted. Their color vision was exceptional for the rest of their lives.

Then phaco started and best-corrected vision was reduced to 20/25 or 20/30—explained wrongly as corneal edema, but it really was macular damage. Color vision was commonly the same before surgery as afterwards.
Even though the resultant refractive error was almost devoid of astigmatism, patients were wearing their glasses more, not less. Simultaneously, there were improvements in phaco and pharmaceuticals, such as the addition of NSAIDs, stronger steroids and better injectable drugs for dilation during surgery. Meds that worked just as well once a day became available. Good acuity and improved color vision were now possible again.

But then the economics of cataract surgery changed. Reimbursement was cut from $1800 to the present-day rate of roughly $550. The post-op treatment course went from two months to no more than three weeks. For 30 years, attempts were made to use intracameral injectable medication to eliminate or at least minimize drop use after surgery. Recently reintroduced, this no-drop cataract surgery still poses some problems, such as visual or anatomic concerns at the one-year follow-up visit, which suggest an insufficient post-op regimen.

An article was published stating that post-op cataract surgery meds could be taken five minutes apart and still would work as well, despite known differences between generic and branded products, such as lower dosages and different corneal penetration rates. Unfortunately, most patients are told now to take post-op drops five minutes apart.

Also, recent changes in the availability of steroid medications have taken place. Inveltys, a higher concentration, better-penetrating form of loteprednol, seems to have overtaken the one-day post-op. If a surgeon counts the day of surgery and the one-day follow-up back and forth. We’ve had patients stay overnight in the town where they had surgery and their acuity is always a couple of lines better in the first few days after surgery.

Also, use of 1% cyclogyl right after surgery resulted in a dilated but 20/20-acuity at the one day post-op. Failure to use cyclogyl for the drive home yielded vision of 20/25 or 20/30 almost always at the one-day post-op. If a surgeon counts seconds of phaco time, they should also count how many miles that patient has to drive in to get surgery and follow-up.

In 2020, the cataract surgeon I used for 14 years retired; it was a sad day. In my small optometry practice, sometimes two patients a day were referred there for cataract surgery.

I think “optometry-referred” cataract surgery patients are different. They often have to travel, and thus need the extra cyclogyl drop and should not travel far again the day after surgery. This is the reason for letting the optometrist see patients the next day instead of the surgeon, but this is hard for most surgeons to appreciate. These patients are most likely more prone to subclinical chronic CME and thus stronger and longer post-op medications in my opinion are recommended.

There is a financial component to “optometry-referred” patients for the patient, the surgeon and the OD. Branded drugs are very costly to the patient. Thirty years ago, the pharmaceutical companies gave branded post-op meds to patients if their brand of intraocular implant was used. Now, there are no gift bags anymore and samples are hard to come by. Branded meds can cost over $600 and the surgeon may be out of pocket for some injections used during surgery because Medicare won’t allow a supply fee. Certain generic NSAIDs, when relied upon at the expense of better anti-inflammatories, may even cause CME.

Then there is the cost of seeing cataract patients for two months. I generally see them for three visits if I see the patient on the first day post-op. I generally don’t make money on post-op visits. Wouldn’t it be nice if cataract surgery was one fee and the post-op care was another fee? This might support better post-op care.

I must say cataract surgery is quite remarkable. Most surgeons do a very good job; if anything, there continue to be improvements in the surgery itself.

All I know is that I miss what cataract surgery was just a few years ago. The patients from those days still come in and can’t believe how good their vision is.
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Upstairs, Downstairs

The medical/optical dichotomy of optometry was on display once again at Vision Expo East.

In New York earlier this month for Vision Expo East, one morning I shared a cab over to the Javits Center with a coworker from the division of our company that produces 20/20 and Vision Monday, both fixtures of the retailing world of eye care.

The cabbie asked us what was happening at the Javits. “It’s an optical show,” my colleague said, not incorrectly. But his framing (no pun intended, I swear) of the conference around eye wear products surprised me. Had I been riding solo that day, I likely would have said, “It’s a meeting for eye doctors.”

And there in a nutshell lies the somewhat schizophrenic identity of optometry.

We all bring our own priorities to our day-to-day experiences, and I’ll be the first to admit that mine are on the medical aspects of eye care. But of course optical products are a huge driver of practice success and, furthermore, probably still the one aspect of eye care the public mostly associates with optometry. Still, it can be odd to see these elements sometimes manifest not merely as two halves of one whole but as entirely different events.

Walking the show floor with a newer editorial colleague on the Review team, in town for her first Vision Expo East, I wryly pointed out that, as usual, the stuff we came to see was “in the basement.” For as long as I can remember (my first VEE was in 1992), the layout of the conference puts retail exhibitors in the spacious upstairs showroom and the med/pharma events in the windowless bottom hall. A fan of British costume dramas, I notice this upstairs, downstairs feel of the place every year.

That show, like the more recent Downton Abbey, chronicled the lives of a wealthy family and their staff, each with their own living quarters. Now, don’t get me wrong: I’m not saying the medically minded attendees are treated like lesser participants. But VEE does often feel like two separate lives being lived in the same house. That’s likely inevitable, given the high attendance by dispensary staff there.

Opticians and dispensing professionals comprise a huge contingent of the attendees, as it should be. They’ll always want and need to know all about the latest trends in frames and lenses. Vision Expo, like its precursor OptiFair, has always been one of the premier outlets for the retail and dispensing side of eye care. A meeting with deep roots in the multibillion-dollar optical industry owes nothing to the medical side of optometry; if they wanted to hold a purely retail event, it would be justifiable and probably highly successful. But it wouldn’t wholly reflect what optometry is today.

The organizers are to be commended for investing in the optometric CE side in recent years. The addition of education co-chairs Mark Dunbar and Ben Gaddie several years ago made medical topics a much bigger priority, and these two exceptional ODs once again put together an outstanding program this year. We shared dozens of excellent clinical insights from the optometric program through live coverage in our Twitter feed and other social media outlets during the meeting.

I learned a lot at VEE this year and it’ll be put to good use in these pages. Next year, I’d like to spend a little more time upstairs.
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Cornea Cornerstone

How to properly manage this important part of optometric care.

While we manage all aspects of primary eye care, what sets us apart as a profession is managing cornea and external disease. Fortunately, the cornea requires little extra equipment beyond our slit lamp to visualize pathology effectively. Consider the opportunities ranging from dry eye disease, epithelial basement membrane dystrophy/ recurrent corneal erosion and Fuchs’ dystrophy to keratoconus, abrasions, foreign bodies and numerous forms of keratitis—just to name a few!

Dry Eye Disease

It’s essential to differentiate the type of dry eye first; otherwise, your treatment strategy isn’t likely to work. Expression of the meibomian glands can be done in two to five seconds with an expression paddle. Abnormal expression will indicate an evaporative dry eye. In contrast, normal meibomian gland expression with significant corneal staining (using NaFI dye) and a thin meniscus will indicate an aqueous-deficient dry eye. Keep in mind that the location of the stain is critical—inferior stain will indicate incomplete overnight lid closure or potentially lagophthalmos.

NaFI dye will also aid in the visualization of a foreign body, papillae, conjunctival staining, conjunctivochalasis, limbal stem cell deficiency, persistent epithelial corneal defects, abrasions and corneal ulcers.

Corneal Foreign Body Removal

When managing corneal foreign bodies, begin with visual acuity testing followed by a slit lamp examination and determine the location, depth and possible material you are dealing with. Rule out an infiltrative process in the cornea indicative of a secondary infection. Observe the anterior chamber for cell and flare. Next, place NaFI dye in the eye to help find the foreign body and always evert the upper eyelid.

While we manage all aspects of primary eye care, what sets us apart as a profession is managing cornea and external disease.

Determining the depth is critical and using a slit beam will help you see how far into the cornea it has penetrated. If there is no risk of intraocular foreign bodies, intraocular pressure assessment will confirm and you can begin working on removal. Consider using a 30-gauge needle—which is beveled—and keep the bevel toward the cornea to lift the foreign bodies (sterile jeweler’s forceps can also be used). This is especially helpful for a metallic foreign body. Be sure to remove significant residual rust in the cornea with an Alger brush. An antibiotic drop should be instilled and prescribed until the epithelium is healed. Corticosteroids may need to be considered after re-epithelialization if significant inflammation and/or an anterior chamber reaction is noted.

The Evolution of Crosslinking

There are two potential future treatments. The first involves a scleral lens with a built-in transducer that emits UV light. The scleral lens bowl is filled with riboflavin solution and placed on the cornea, and the patient wears the lens, which is hooked up to a machine that administers the light treatments.

The second, soon to enter Phase III FDA clinical studies, involves lysi oxidase, a substance that is deficient in patients with keratoconus. Phase IIb trials with drops containing this ingredient show a decrease of 1.8D in K values, which is technically a greater response than currently approved corneal crosslinking options.

Herpes Zoster Ophthalmics (HZO)

Optometry is seeing more of these cases and we have to be prepared to manage these patients. Although the cornea is heavily involved, with signs ranging from pseudodendrites to eventual neurotrophic keratitis and lipid keratopathy, the most common sign of HZO is iritis. Since this is a systemic disease with ocular manifestations, the primary therapy is oral antivirals. I prefer valacyclovir 1000mg three times a day for 10 to 14 days.

Additionally, treat the ocular inflammation early with potent topical steroids to prevent corneal scarring and quiet the uveitis. Tapering can occur over six weeks and you may need to maintain a prophylactic dose of oral valacyclovir of 1000mg once a day. Also consider ophthalmic gel Zirgan (Bausch + Lomb) five times per day for a week followed by three times a day for a week when the cornea is involved. This aggressive management will help spare vision loss.

The cornea is a comfortable place for optometry. Keep your skills up and gain confidence as you work on a structure that is readily visible and will make a difference for you and your patients in this cornerstone of optometric care.
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Is there such a thing as truth? I was brought up by Betty and Earl to believe there is. Oh, yes, my parents, coincidently also named Betty and Earl, thought so too, but I’m talking about Betty and Earl down at the drive-in. Betty and Earl knew that if someone came to the drive-in, they would probably want a couple of hot dogs and some popcorn, and you could hang your hat on that, my friends.

So, having been taught that there is truth to certain things, I am concerned that optometry and optometrists might be living in a haze rather than clearly seeing the truths in our profession that we can count on no matter what. Here are some truths in optometry:

1. People want to have good vision. You cannot hurt somebody if you help them see better. I know we can all give examples of folks who come in at 20/50 and say, “I’m just here for a checkup. My eyes are fine.” What about them, you may ask? I think we can make them see 20/200, but wearing a clean shirt is not a bad move. Don’t leave home without it. Oh, and bathe. You know who you are. Yes, I am talking about you.

2. Each exam has an endpoint. Of course, sometimes the endpoint is that we just cannot help the patient. Not everyone can choose which is better, number one or number two. For some folks, the endpoint is the wilderness, not the mountaintop. Let them continue to wander visually until they decide where to make camp. Personally, I love when a patient comes in with a bag of glasses from 10 different eye doctors, none of whom are any good at optometry. My goal shifts to not being the dumbest doctor in the bag. If you make the patient be the boss of their own eyeballs, they will never rank you lower than the fifth dumbest eye doctor in the bag. Now there’s an endpoint for you.

3. Consulting online reviews is no way to run a practice. Patients who gripe online are not your best practice consultants. People who harass you online also harass everybody they ever meet online. Do your best to help them receive the care they need from the ophthalmologist who disrespects optometry the most. Karma.

4. People care about your appearance. I am living proof that you don’t have to be pretty to be successful in optometry, but wearing a clean shirt is not a bad move. Don’t leave home without it.

5. It’s okay to drive a cool car. I know doctors who own a tricked-out, luxurious car but have an old beater they drive to work because they are afraid the patient will think they are just there to make money. (1) There is nothing wrong with making money. Your patients make money too, right? (2) If somebody is messing with my eye, I want them to drive a nice vehicle. That means they see lots of patients. And, taking it even further, if somebody is doing my heart bypass, I want them to be driving a gold-plated Bentley with a personalized license plate that reads, “BSTHRTDR.”

6. You cannot save your stuff from their choices. I spent my early years trying to make my staffers better and happier. This did not work. But when I changed my approach to make them better employees, they automatically became better and happier or they hit the road, which also, I think, helped them be better and happier while simultaneously making me better and happier.

7. You must accept you aren’t perfect. If you do, then you are actually perfect after all. Yes, you will blunder through more often than you will ever know. Welcome to humankind. Wanna be perfect? Ask your spouse how to do it.

8. You need to get paid. I know, 50% of your townspeople are covered by a crappy vision insurance plan. Still, assuming you would like to keep the lights on, then, obviously, you need to get paid. It’s okay.

9. People, you have to stop and spend time just breathing. Close your eyes. Be here right now. Oh, I should have mentioned, pull your car off the highway first. My bad.

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About Dr. Vickers

Dr. Vickers received his optometry degree from the Pennsylvania College of Optometry in 1979 and was clinical director at Vision Associates in St. Albans, WV, for 36 years. He is now in private practice in Dallas, where he continues to practice full-scope optometry. He has no financial interests to disclose.
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Q A patient presents to me with a significantly swollen left upper lid. What are the next steps?

A While there are plenty of possible causes of swollen eyelids, the first that comes to mind is preseptal cellulitis. “While this is the most probable, we must always keep the differential with the highest chance of morbidity or mortality in mind,” says Mark Miriello, OD, of The Eye Institute in Philadelphia. “With swollen eyelids, that is orbital cellulitis.”

Other differentials to consider involve severe anterior segment inflammation such as viral conjunctivitis, corneal abrasion and even uveitis. Taking a careful symptom inventory and thorough slit lamp examination can help rule these differentials out.

Trace the Source

The orbital septum is a shield of fibrous tissue that acts as a barrier between the eyelids and the orbit itself. Preseptal cellulitis, then, is a bacterial infection of the skin—effectively isolated to the eyelids.1,2

Patients present with swollen, red lids that may be tender to the touch and febrile. Since the inflammation is confined to the eyelid, they present with unaffected vision, ocular motility and pupils and without proptosis.1,3

The source of the infection is often a result of other infectious eyelid lesions. Sinusitis, particularly of the ethmoid sinus, is a common culprit. Other routes of infection include superficial facial traumas and surgeries, foreign bodies and insect bites.1

A patient with preseptal cellulitis who responded well to oral antibiotics.

When taking history, determine if the patient is at risk for any of these etiologies. “Ask about precipitating styes, traumas and sinus infections,” Dr. Miriello says. “If the patient can localize the part of the lids where their symptoms first presented, pay special attention there during slit lamp examination.”

Preseptal cellulitis treatment involves oral antimicrobial therapy. With gram-positive cocci, such as Staphylococcus and Streptococcus species, being the most common causative organisms, select antibiotics accordingly.1 Keflex (cephalexin, Advancis Pharmaceutical) is a great choice and is commonly dosed 500mg PO BID for adults.3 This dosage is well tolerated and can be safely taken up to QID.4

Dr. Miriello prefers the more frequent dosing so he never has to wonder if a persistent case would have responded better. The treatment course can range from seven to 10 days, depending on the severity of the initial presentation.5

Check Motility

Orbital cellulitis occurs when these infections gain access to the orbit itself. While these patients may look preseptal, there are several symptoms and signs that readily set this more serious orbital infection apart. Look for some degree of ophthalmoplegia and pain with eye movement due to inflammatory infiltration of the extraocular muscles, Dr. Miriello says. Exophthalmometry is also recommended, since ipsilateral proptosis signals likely orbital involvement.1,3,5

As the disease progresses and the retrobulbar optic nerve becomes involved, visual acuity, color vision and pupillary reactions can become affected. Untreated orbital cellulitis may also result in orbital abscesses, cavernous sinus thrombosis or even intracranial extension of the infection, which can be deadly.1,3,5

Orbital cases may stem from preseptal ones if timely treatment is not initiated. Sinusitis is overwhelmingly the most common cause. Others include the spread of dental and middle ear infections, infected adnexal injection sites and traumas or surgeries that allow access to the orbit.1,3,5

“These patients require intravenous antibiotics and hospitalization, so prompt referral is crucial when you suspect orbital involvement,” Dr. Miriello says. Multidisciplinary management can greatly improve outcomes.1,3,5

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US-LAS-220057 02/22
CLINICAL PEARLS IN CORNEAL FOREIGN BODY REMOVAL

How to properly identify, treat and manage patients with this ocular trauma.

BY BRIDGET HENDRICKS, OD
WORCESTER, MA

It’s a typical afternoon in your practice and a 35-year-old male presents with the following complaint: “I was installing shelves in my home a few days ago and I think I got something in my eye.” He mentions that he could see something dark gray on his eye when he looked in the mirror. He tried to rinse it out, but it wouldn’t budge—now the eye is red and painful.

Ocular foreign bodies are a common form of ocular trauma, accounting for approximately 40% of all such traumas and leading to about 2% of all emergency room visits.1 Patients’ symptoms can vary widely depending on the nature and timing of the foreign body injury.

Regardless of your practice mode, most of us have encountered situations similar to this. But what is the best way to proceed? In this article, we will review how to evaluate, treat and manage patients with an ocular foreign body, as well as provide helpful clinical pearls for successful removal.

#1: TAKE A THOROUGH PATIENT HISTORY

A proper detailed history of the patient will help guide your examination, determine the type of prophylactic care needed as well as provide documentation for any workers’ compensation or other legal issues that may be associated with the incident. The case history should be problem-focused with an emphasis on what, when, where and how.

• Ask what the foreign body is. Knowing the kind of material involved can help give you an idea of the risk for infection and guide your decision-making on the type of postoperative treatment that may be indicated. Organic or vegetative materials are associated with a higher rate of infection. Metallic materials are likely to result in a rust ring. Inert materials (i.e., glass, ceramic or high-grade plastic) tend to be non-reactive and carry less risk of an inflammatory response.3,4

• Ask when the injury occurred. The longer the foreign body has been in place, the higher the risk for infection, inflammation and rust formation (with metallic ferrous material). Salt present in the tears can react with the iron in metallic foreign bodies to create a corneal rust ring around the foreign body. This reaction

Fig. 1. Preoperative slit lamp biomicroscopy of the patient’s left eye shows the embedded corneal foreign body.

Dr. Hendricks is an associate professor of optometry at Massachusetts College of Pharmacy and Health Sciences (MCPHS) University School of Optometry, chief of services for the Glaucoma Clinic (The Eye and Vision Center at MCPHS) and practices at the Eye and Vision Center at MCPHS and Edward M. Kennedy Community Health Center. She completed her residency in ocular disease in the VA Boston Healthcare System and served as faculty at the New England College of Optometry before joining MCPHS University in 2017. She has no financial disclosures.
Fig. 2. Preoperative imaging of the patient’s left eye. AS-OCT shows hyper-reflectivity and shadowing due to the corneal foreign body (left). The corneal foreign body did not completely penetrate the cornea (right).

#2: DON’T CUT CORNERS OR RUSH EXAMS

Attention to detail during foreign body removal can make the difference between successful outcomes and sight-threatening complications. Immediately following the case history, be sure to obtain and document BCVAs before performing any procedures or instilling drops. This will determine the level of suspicion for penetrating injuries, as these injuries are associated with more significantly decreased vision. Baseline visual acuities are also important for medicolegal reasons.

For example, if the patient has amblyopia or another pre-existing condition causing reduced acuity, it is important to have documentation that the acuity was reduced prior to the procedure that you performed. This could protect you from allegations that your procedure is responsible for the patient’s reduced acuity. An exception to this rule is if the patient is experiencing intense pain with blepharospasm, rendering them unable to open their eyes. In this case, a drop of topical anesthetic such as proparacaine or tetracaine may be instilled prior to visual acuity measurement.

Other entrance tests such as pupil and extraocular motility testing should be performed, as abnormalities in pupil shape (e.g., a peaked iris) and limited mobility of the eye can be a sign of penetrating injury.

A thorough slit lamp exam should be performed, first without fluorescein dye. Examine all structures of both eyes looking for injection, chemosis, anterior chamber cells and flare and presence of foreign bodies. Be sure to check the conjunctiva for any signs of perforation—this will appear as an area of injection and chemosis surrounding an entrance point. Always perform lid eversion to check for foreign bodies trapped under the eyelid.

After your initial scan of the ocular structures, add fluorescein dye and examine the eye using a Cobalt blue filter. Look for signs of epithelial defects, corneal abrasions or lacerations. Keep in mind that the presence of vertically oriented linear abrasions could result from foreign bodies embedded under the eyelid.

Always perform the Seidel test to help rule out globe perforation. A positive test will appear as a “dark waterfall” of aqueous leakage within the fluorescein, indicating that ocular penetration has occurred. Other signs consistent with a penetrating injury include shallow anterior chamber, hyphema, defects of the iris or pupil, a break in Descemet’s membrane and lens opacities.

As a word of caution, small foreign bodies characterized by high heat or high velocity can penetrate the globe and result in a self-sealing entry point. Such wounds may not result in a positive Seidel sign or shallow anterior chamber, but a tunnel through the cornea may be visible. In addition, there may be associated anterior uveitis. If there are any signs or suspicion of penetrating injury, apply a rigid shield (e.g., Fox shield or inverted Styrofoam cup) and refer the patient immediately to the emergency department or ophthalmic surgeon for specialty evaluation and management.

Once the eye is stabilized, further investigation for retained foreign...
bodies is indicated with tests such as dilated fundus examination, gonioscopy and imaging studies. Potential imaging methods include computed tomography scans, B-Scan ultrasonography, ultrasound biomicroscopy, plain film x-rays and magnetic resonance imaging. The latter is contraindicated if suspected foreign body is metallic because it can cause migration of the object, further damaging the ocular tissues.

After you’ve ruled out penetrating injury and located the foreign body, determine its exact location, size and depth. Keep in mind there can be more than one foreign body present, or one in the fellow eye that the patient is unaware of. An optic section should be used to determine the depth; however, in some cases—especially if there is corneal edema—it is difficult to visualize.

Anterior segment OCT (AS-OCT) can help determine the depth in unclear cases; it will typically show a hyper-reflective lesion representing the foreign body material. The foreign body will also cause shadowing of the corneal layers corresponding to its location. Thus, the high resolution of AS-OCT provides detailed information that can be used to determine the most appropriate removal technique (Figures 1 and 2).

Understanding exact depth and location is also helpful in predicting prognosis. Objects that have penetrated the stroma will produce a scar, which can result in decreased vision if the object is within or near the visual axis. In addition to detailed characteristics of the foreign body, AS-OCT can provide information on the status of the surrounding tissue, the integrity of Descemet’s membrane (i.e., risk for impending perforation) and even identify any missed lesions that were not previously visible on slit lamp examination.

Next comes determining whether or not the foreign body actually needs to be removed. If it is not penetrating the globe, it’s best to go ahead and remove it. Inert objects such as sand and glass are well-tolerated in the cornea without resulting in a tissue reaction. If an inert object cannot be removed without significant risk and is not in the visual axis, it can be left in place with close monitoring of the patient. In contrast, organic substances (vegetative materials) and metals are poorly tolerated by the cornea, resulting in edema, scarring, inflammation, neovascularization and stromal necrosis. These substances must be promptly removed.

#3: CHOOSE THE BEST TOOL FOR THE JOB
If the foreign body is superficial, it can sometimes be rinsed away via ocular irrigation with sterile saline or wiped away with a moist cotton-tipped applicator. If the foreign body is embedded, more invasive tools are necessary: golf spud, jewelers forceps, magnetic probe and small-gauge needles (Figure 3). Rust rings may be removed with either a small-gauge needle or an ophthalmic burr/Algerbrush.

Here are some rules of thumb to take into consideration when selecting which instrument to use:
• Make sure instruments are sterile.
• Use the least invasive technique that will allow for safe and effective removal.
• Attempt to remove superficial objects with a moistened sterile
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*These statements have not been evaluated by the Food and Drug Administration. These products are not intended to diagnose, treat, cure, or prevent any disease.
#4: ENSURE PATIENT CONSENT
Prior to foreign body removal, explain the procedure, risks, benefits and alternatives to the patient and obtain written consent (or document verbal consent in your chart).

Ensure patient comfort and cooperation by properly anesthetizing the ocular surface with topical anesthetic drops such as proparacaine 0.5% or tetracaine 0.5%. Instill one or two drops of anesthetic into the lower cul-de-sac of the eye and wait at least 30 seconds before starting the procedure. Instillation of topical anesthetic into the fellow eye will aid in reducing blink reflex and increasing patient cooperation.

For best stabilization and visualization, a slit lamp should be used with the patient’s forehead securely placed against the forehead rest. Identify a fixation target that will ensure proper and steady alignment of the eye and reduce risk of eye movement during the procedure. Instruct the patient to fixate on the target using the opposite eye. Occasionally, the use of an eyelid speculum is required for patients who are less cooperative or have a strong blink reflex.

#5: OPTIMIZE STABILITY DURING REMOVAL
Stability is key when performing foreign body removal. Stabilize your arm using the slit lamp table or an arm rest. Hold the instrument like a pencil between your thumb and forefinger. Rest your fourth and fifth fingers holding the instrument on the patient’s check, bridge of nose or upright bar of the slit lamp. Align your instrument by sighting outside of the slit lamp, positioning it in front of the foreign body at an angle tangential to the ocular surface. It’s imperative that you always approach the eye tangentially in order to prevent perforating the eye or further embedding the foreign body into the eye. Once properly aligned, look through the oculars as you perform the remainder of the procedure. Secure the upper eyelid with your free hand and use the instrument to remove the foreign body.

When using a cotton-tipped applicator, foreign bodies can be removed by gently tapping the moistened tip to the object and lifting it off of the tissue. If the foreign body is conjunctival, you can use a gentle swiping motion if necessary. Avoid any swiping motions on the cornea as this could cause an abrasion.

When using forceps, gently clasp the object and lift it off of the tissue, and avoid pinching the surface of the eye.

When using a needle, position it with the bevel facing outward, toward you. With small strokes, use it to loosen the edges of the foreign body. Then, with the tip of the needle positioned just under the edge of the foreign body, use a subtle flicking motion to release it from the corneal surface. Make sure to maintain a tangential angle to the globe throughout this procedure.

When using a spud, the instrument should be handled in a similar manner as the needle, using the tip of the spud to tease out the edges of the foreign body and then using a flicking motion to release it from the surface of the cornea (Figure 4).

#6: DOUBLE CHECK, IRRIGATE AND DOCUMENT
Once you have successfully removed the foreign body, irrigate the eye with sterile saline solution and re-examine with the slit lamp. Document the size, location and depth of the resultant epithelial defect.

#7: REMOVE ANY RUST AT INITIAL VISIT
Metallic (ferrous) ocular foreign bodies begin to oxidize, forming rust at the injury site within just a few hours of becoming embedded in the cornea. The resulting rust usually forms a ring around the metallic foreign
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body (Figure 5). If left there, this rust ring can cause an immune response, resulting in extended healing time, inflammation, tissue damage and stromal necrosis. Therefore, it is best to remove as much of the rust as possible, as soon as possible.

An Algerbrush—a small, low-speed, battery-operated drill that is fitted with a tiny dental burr—is commonly used for removal. To alleviate patient apprehension during this technique, mention that the Algerbrush makes a soft whirring noise. The device has a built-in clutch mechanism that will stop the instrument when a certain degree of resistance is encountered. Therefore, when an Algerbrush contacts Bowman’s membrane, it will automatically shut off. This safety measure assumes proper use of the instrument, using a tangential approach, and that the foreign body itself is not penetrating Bowman’s membrane. In cases where the foreign body is penetrating Bowman’s, there is no longer enough resistance to initiate the stop.

When using the Algerbrush, approach the cornea tangentially and apply it to the rust ring with very little pressure; allow the spinning motion of the burr on the cornea to do the work for you. Lightly contact the rotating burr with all areas of the rust ring. There should be minimal hand movement, with the exception of moving the instrument to a different part of the rust ring. Occasionally, it is necessary to switch the Algerbrush to your opposite hand in order to take advantage of the now reversed rotation of the burr to remove any resistant rust.

Alternatively, you may use a spud or needle to remove the rust ring, always keeping the instrument tangential to the cornea. Use small motions to scrape away the epithelium that contains the rust.

Ideally, you should remove as much rust as possible during the initial visit in avoid an inflammatory response (Figure 6). However, very deep stromal rust may be left in place if you feel it would be unsafe to attempt to remove it at that visit. Keep in mind: aggressive and widespread use of the burr within the stroma can lead to increased scarring. Thus, in cases of deep stromal rust, it is best to remove only the superficial rust and leave the rest for removal at a follow-up visit.

As the cornea heals, the residual rust will move more anteriorly, allowing for easier removal with less risk of long-term scarring. In some cases, it takes several visits to remove all of the rust. At the conclusion of the procedure, rinse the eye with sterile saline, re-examine the eye with the slit lamp and document the extent of the corneal defect. You may instill a drop of broad-spectrum topical antibiotic for prophylactic coverage.

Fig. 5. Metallic foreign body with rust ring.

Fig. 6. Corneal epithelial defect immediately following foreign body removal with an Algerbrush.
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STOP FLARES FAST

• >50% reduction in symptoms of blepharitis/blepharoconjunctivitis in 1 week of dosing. No IOP spikes reported during first week of treatment.

• Greater bactericidal activity—more effective at killing MRSA than TobraDex* (>99.9% kill rate vs 0%)²

• Delivers 12.5x higher tobramycin concentration in ocular tissue compared to TobraDex²

Indications and Usage
For steroid responsive inflammatory ocular conditions of the palpebral and bulbar conjunctiva, cornea, and anterior segment of the globe and chronic anterior uveitis, corneal injury from chemical, radiation or thermal burns, or penetration of foreign bodies for which a corticosteroid is indicated and where the risk of superficial bacterial ocular infection is high or where there is an expectation that potentially dangerous numbers of bacteria will be present in the eye.

Important Safety Information
CONTRAINDICATIONS:
Most viral disease of the cornea and conjunctiva including epithelial herpes simplex keratitis (dendritic keratitis), vaccinia, and varicella, and also in mycobacterial infection of the eye and fungal disease of ocular structures. Hypersensitivity to any components of the medication.

WARNINGS & PRECAUTIONS:
• IOP increase – Prolonged use may result in glaucoma with damage to the optic nerve, defects in visual acuity and fields of vision. IOP should be monitored.

• Aminoglycoside sensitivity – Sensitivity to topicaly applied aminoglycosides may occur.

• Cataracts – Posterior subcapsular cataract formation may occur.

• Delayed healing – May delay healing and increase the incidence of bleb formation. Perforations of the cornea or sclera have occurred. Slit lamp biomicroscopy, and fluorescein staining should be conducted.

• Bacterial infections – May suppress host response and increase secondary ocular infections. In acute purulent conditions, steroids may mask infection or enhance existing infection. If signs and symptoms fail to improve after 2 days, the patient should be re-evaluated.

• Viral infections – Use with history of herpes simplex requires great caution. The course and severity of many viral infections of the eye (including herpes simplex) may be exacerbated.

• Fungal infections – Fungal infections of the cornea may occur and should be considered in any persistent corneal ulceration.

• Use with systemic aminoglycosides – Total serum concentration of tobramycin should be monitored.

ADVERSE REACTIONS:
The most frequent adverse reactions (<4%) to topical ocular tobramycin are hypersensitivity and localized ocular toxicity, including eye pain, eyelid pruritus, eyelid edema, and conjunctival hyperemia.

The reactions due to the steroid component are increased intraocular pressure with possible development of glaucoma, and infrequent optic nerve disorder; subcap-sular cataract; and impaired healing.

The development of secondary infection has occurred. Fungal infections of the cornea may occur. Secondary bacterial ocular infection following suppression of host responses also occurs.

Non-ocular adverse events (0.5% to 1%) included headache and increased blood pressure.

Please see Brief Summary of full Prescribing Information on the adjacent page.

INDICATIONS AND USAGE

TOBRADEX ST is a topical antibiotic and corticosteroid combination for steroid-responsive inflammatory ocular conditions for which a corticosteroid is indicated and where superficial bacterial ocular infection or a risk of bacterial ocular infection exists.

Ocular steroids are indicated in inflammatory conditions of the palpebral and bulbar conjunctiva, cornea and anterior segment of the globe where the inherent risk of steroid use in certain infective conjunctivitides is accepted to obtain a diminution in edema and inflammation. They are also indicated in chronic anterior uveitis and corneal injury from chemical, radiation or thermal burns, or penetration of foreign bodies.

The use of a combination drug with an anti-infective component is indicated where the risk of superficial ocular infection is high or where there is an expectation that potentially dangerous numbers of bacteria will be present in the eye.

DOSAGE AND ADMINISTRATION

Recommended Dosing: Instill one drop into the conjunctival sac(s) every four to six hours. During the initial 24 to 48 hours, dosage may be increased to one drop every 2 hours. Frequency should be decreased gradually as warranted by improvement in clinical signs. Care should be taken not to discontinue therapy prematurely.

CONTRAINDICATIONS

Nonbacterial Etiology: TOBRADEX ST is contraindicated in most viral diseases of the cornea and conjunctiva including epithelial herpes simplex keratitis (dendritic keratitis), vaccinia, and varicella, and also in mycobacterial infection of the eye and fungal diseases of ocular structures.

Hypersensitivity: Hypersensitivity to any component of the medication.

WARNINGS AND PRECAUTIONS

IOP increase: Prolonged use of corticosteroids may result in glaucoma with damage to the optic nerve, defects in visual acuity and fields of vision. IOP should be monitored.

Aminoglycoside sensitivity: Sensitivity to topically applied aminoglycosides may occur.

Cataracts: May result in posterior subcapsular cataract formation.

Delayed healing: May delay healing and increase the incidence of bleb formation after cataract surgery. In those diseases causing thinning of the cornea or sclera, perforations have been known to occur with the use of topical steroids.

Bacterial infections: May suppress the host response and thus increase the hazard of secondary ocular infections. In acute purulent conditions, steroids may mask infection or enhance existing infection. If signs and symptoms fail to improve after 2 days, the patient should be re-evaluated.

Viral infections: Treatment in patients with a history of herpes simplex requires great caution. Use of ocular steroids may prolong the course and may exacerbate the severity of many viral infections of the eye (including herpes simplex).

Fungal infections: Fungal infections of the cornea are particularly prone to develop with long-term use. Fungal invasion must be considered in any persistent corneal ulceration.

Use with systemic aminoglycosides: Use with systemic aminoglycoside antibiotics requires monitoring for total serum concentration of tobramycin.

ADVERSE REACTIONS

The most frequent adverse reactions to topical ocular tobramycin (TOBREX®) are hypersensitivity and localized ocular toxicity, including eye pain, eyelids pruritis, eyelid edema, and conjunctival hyperemia. These reactions occur in less than 4% of patients. Similar reactions may occur with the topical use of other aminoglycoside antibiotics.

Non-ocular adverse events occurring at an incidence of 0.5% to 1% included headache and increased blood pressure.

The reactions due to the steroid component are: increased intraocular pressure (IOP) with possible development of glaucoma, and infrequent optic nerve disorder; subcapsular cataract; and impaired healing.

Secondary Infection: The development of secondary infection has occurred. Fungal infections of the cornea are particularly prone to develop with long-term use. Fungal invasion must be considered in any persistent corneal ulceration. Secondary bacterial ocular infection following suppression of host responses also occurs.

USE IN SPECIFIC POPULATIONS

Pregnancy and Nursing Mothers

There are no adequate and well controlled studies in pregnant women. TOBRADEX® ST ophthalmic suspension should be used during pregnancy only if the potential benefit justifies the potential risk to the fetus. Caution should be exercised when TOBRADEX® ST is administered to a nursing woman.

Pediatric Use: Safety and effectiveness in pediatric patients below the age of 2 years have not been established.

Geriatric Use: No overall differences in safety or effectiveness have been observed between elderly and younger patients.

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#8: TREAT WITH ANTIBIOTICS
Postoperative care involves prevention of infection, control of pain and inflammation and monitoring for complications. As a rule of thumb, the patient should be treated with a broad-spectrum antibiotic, such as a fluoroquinolone, four times per day for one week or until the defect has re-epithelialized. In cases of large corneal defects or conjunctival defects, antibiotic ointment may be preferred due to longer coverage as well as cushioning for comfort. A cycloplegic, such as cyclopentolate or homatropine, can be prescribed twice per day if there is significant pain or inflammation.

Topical ophthalmic NSAIDs can also be considered for treatment of pain in foreign body patients; however, these medications are not ideal in such cases, as they can lead to compromised corneal healing. Over-the-counter oral analgesics such as acetaminophen and ibuprofen are rarely indicated but can be considered if there is considerable pain.

In cases of excessive inflammation, topical corticosteroids can be used but only after re-epithelialization of the defect. Steroids should also be considered if the foreign body is deep/stromal and central to help reduce the amount of scarring. However, steroids should not be used with vegetative/organic/soil-based foreign bodies due to increased risk of fungal infection. Amniotic membranes are another treatment option for deep central foreign bodies.

For larger corneal defects, treatment with a bandage contact lens is particularly effective for reducing pain and minimizing tissue disruption that is associated with the movement of the eyelid over the defect. However, this would be contraindicated in cases in which the foreign body was organic or vegetative due to risk of fungal keratitis. If a bandage contact lens is used, it is important to see the patient in 24 hours.

At the follow-up visit, the contact lens should be removed and the cornea examined. Continue bandage contact lens treatment until the epithelial defect has healed. Monitor closely, as bandage contact lenses can create an environment more susceptible to infection.

If the foreign body material is organic or vegetative and the patient is at high risk for fungal keratitis, treatment with topical and/or oral antifungal medication can be considered.

#9: MONITOR OUTCOMES
Patients should be seen for follow-up in 24 hours. Typically, there is significant healing and reduction in pain within the first 24 to 72 hours (Figure 7). Monitor for complications such as non-healing infection, edema, iritis, recurrent corneal erosions and visually significant scarring. If any complications are found, treat appropriately and refer to a specialist when indicated.

#10: PREVENT FUTURE INJURIES
One of the most important aspects of postoperative care is providing patient education on the importance of wearing well-fitted safety goggles while performing high-risk activities. Make sure your patient has appropriate safety eyewear, and if not, help them obtain it, as well as explain and demonstrate proper use.

Ocular foreign body removal is a rewarding and skilled service that you can provide to your surrounding community. By following these clinical pearls, you can remove them with confidence, case and without undue complications, giving your patients much-needed relief.

FOR A STRUCTURE ONLY HALF A MILLIMETER THICK, THERE ARE A WIDE VARIETY OF PROBLEMS THAT CAN ArISE IN THE CORNEA. MOST OF THESE PATHOLOGIES CAN BE GROUPED INTO ONE OF SIX CATEGORIES: INJURY, INFLAMMATION, INFECTION, DYSPLASIA, DEGENERATION OR DEPOSITION.

KMK Optometry has helped 98% of optometry students nationwide prepare for their board exams since 2010, and this month we’re going to fine-tune your cornea differential diagnosis skills in particular. Below are some of our favorite cases with photos from our Instagram page that represent a few of the pathology groups mentioned above for you to test your knowledge. Have fun and good luck! Answers to all questions appear at the conclusion of this article on page 49.

**CASE ONE**

A 55-year-old woman was referred from the emergency room with gradually worsening vision over the previous week in her left eye with increasing light sensitivity and a headache. Her history included cataracts, LASIK surgery in 2006 and herpes zoster ophthalmicus in 2019. She noted she had experienced
recent stress due to being furloughed. Her vision was 20/50 in the left eye. Her exam findings were notable for 1+ anterior chamber cell and the corneal findings seen in the photo.

**Corneal Findings**

1. **What is your tentative diagnosis?**
   a. Corneal hydrops from post-LASIK ectasia.
   b. Recurrent zoster keratitis.
   c. Salzmann’s nodular degeneration.
   d. Diffuse lamellar keratitis.

2. **Which treatment is most appropriate?**
   a. Corneal crosslinking.
   b. Superficial keratectomy.
   c. Oral valacyclovir and topical prednisolone.
   d. Flap lift with irrigation.
   e. Topical gancyclovir.

**Discussion**

Interstitial keratitis is an infectious condition that affects the stromal layer of the cornea and most commonly arises from the herpes simplex virus or the varicella zoster virus, sometimes triggered by stress or fatigue. The treatment is similar for both viruses, involving oral antivirals to inhibit viral replication and topical steroid eye drops to reduce the inflammatory response to the viral antigens. Recurrence can be common as seen with this patient, as well as visually debilitating; herpetic keratitis is one of the leading causes of corneal scarring.

Fortunately, this patient improved from 20/50 to 20/20 vision on oral Valtrex (valacyclovir, GlaxoSmithKline) 1g TID and prednisolone acetate six times a day followed by a six-month taper, and continues to take a QD prophylactic dose of each ongoing.

**CASE TWO**

Referred from a local optometrist for “corneal thinning,” this patient thought her eyes were perfectly healthy other than having a higher than average amount of astigmatism. She had no pain, redness or tearing, and best-corrected visual acuity (BCVA) was 20/20 OD and 20/25 OS with against-the-rule 1.5D of cylinder OD and 3.5D of cylinder OS. On slit lamp exam, there was paralimbal corneal thinning with neovascularization and lipid deposits bilaterally, most prominent superiorly and inferiorly, and the epithelium was intact.

**Corneal Findings**

3. **What is your tentative diagnosis?**
   a. Pellucid marginal degeneration.
   b. Peripheral ulcerative keratitis.
   c. Terrien’s marginal degeneration.
   d. Furrow degeneration.

4. **Which treatment is most appropriate?**
   a. Corneal crosslinking.
   b. Bandage contact lens, topical moxifloxacin and oral vitamin C.
   c. Culture.
   d. Observation.
   e. Amniotic membrane graft.

**Discussion**

Given the lack of inflammation in this case, the presence of peripheral thinning greatest superiorly and inferiorly accompanied by lipid and neovascularization, the diagnosis of Terrien’s marginal degeneration was made. This condition is slowly progressive, causing high levels of against-the-rule astigmatism—usually bilaterally—resulting in normal to mildly reduced visual acuity depending. Vision is usually well-corrected with glasses or scleral contact lenses, and serious complications like spontaneous perforation are quite rare, usually associated with ocular trauma.

This patient was content with her glasses, deferred a scleral lens consult and was referred back to her OD for yearly monitoring.
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**Based on a survey of presbyopes who stopped wearing contact lenses after symptoms of presbyopia emerged.

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**CASE THREE**

This female patient had a history of severe trauma 20 years prior, necessitating vitrectomy, scleral buckle, scleral-fixated IOL and a trabeculectomy, which had scarred down. She was currently taking four different glaucoma drops and had no pain, but vision had gradually declining over the last year along with slight progression of her arcuate (non-central) visual field defects. BCVA was 20/100 with an IOP of 25. There was a diffuse whorling keratitis that picked up NaFl stain, no stromal edema and no inflammation in the anterior chamber.

**Corneal Findings**

5. What is your tentative diagnosis?
   a. Microcystic corneal edema.
   b. Rhopressa-associated verticillata.
   c. Anterior basement membrane dystrophy.
   d. Limbal stem cell deficiency.

6. Which treatment is most appropriate?
   a. Tube shunt and discontinue Rhopressa (netarsudil, Aerie).
   b. Phototherapeutic keratectomy with Prokera (Bio-Tissue).
   c. Descemet’s membrane endothelial keratoplasty (DMEK) and Xen implant (Allergan).
   d. Micropulse cyclophotocoagulation and platelet-rich plasma eye drops.

**Discussion**

This patient is in a difficult predicament, with uncontrolled IOP on maximal medical therapy and limbal stem cell deficiency (LSCD) from glaucoma drop toxicity. The swirling pattern is reminiscent of verticillata, but that does not stain with NaFl and rarely affects visual acuity. This LSCD pattern arises from the centrifugal nature of epithelial replication from the palisades of Vogt at the limbus as the cells migrate centrally. For IOP reduction, it was decided to perform micropulse cyclophotocoagulation to avoid incisional surgery that may further damage the cornea. She was also switched to preservative-free Tim-Brim-Dorz Qam and Tim-Brim-Dorz-Lat eye drops Qhs (Imprimis). Platelet-rich plasma eye drops were formulated at 40% for maximum efficacy and used six times a day.

Fortunately, vision recovered to 20/50 after two months of treatment and IOP improved to the upper teens.

**CASE FOUR**

This 61-year-old male had not had an eye exam in over 10 years and reported that his vision was not as good as it used to be. His eyes were occasionally dry, but otherwise comfortable. He worked as a roofer and reported smoking a pack of cigarettes per day. He had no history of any ocular surgery or injury. His exam was remarkable for 2+ nuclear cataracts on exam and visual acuity was 20/30. The left cornea had an irregular swath of hazy epithelium without vascularization or ulceration, as documented in the photo.

**Corneal Findings**

7. What is your tentative diagnosis?
   a. Ocular surface squamous neoplasia (OSSN).
   b. Atypical pterygium.
   c. Salzmann’s nodular degeneration.
   d. Limbal dermoid.

8. Which treatment is most appropriate?
   a. Topical chemotherapy eye drops.
   b. Surgical resection with amniotic membrane graft and mitomycin-C.
   c. Superficial keratectomy with amniotic membrane graft.
   d. Diamond burr polishing plus bandage contact lens.

**Discussion**

Sun exposure is a strong risk factor for developing OSSN and this patient was no different. The photograph shows hazy epithelium in an irregular pattern unlike a pterygium or Salzmann’s degeneration. In recent years, topical chemotherapeutic agents such as interferon alpha-2b, mitomycin-C and 5-fluorouracil have gained popularity in their ability to clear OSSN from the conjunctiva and cornea without surgery, and they show superiority in clearing microscopic disease that can be missed with traditional excisional biopsy.
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Fortunately, this patient regained a perfectly clear cornea after several rounds of topical interferon drops and is now doing well without signs of recurrence.

CASE FIVE

A 49-year-old female reported a history of recurrent redness in her left eye the last three years, as well as light sensitivity and mild blur as of this spring. She had a history of chalazion removal two years prior, did not wear contact lenses and was already taking ciprofloxacin eye drops from her PCP, which were not helping.

Slit lamp exam showed significant telangiectatic blepharitis, a corneal infiltrate with a leash of superficial neovascularization from the nasal limbus and a smaller-sized epithelial defect overlying the infiltrate. She denied recent travel out of the country and new sexual partners.

Corneal Findings

9. What is your tentative diagnosis?
   a. Interstitial keratitis.
   b. Phlyctenular keratoconjunctivitis.
   c. Vernal keratoconjunctivitis.
   d. Recurrent corneal erosion.

10. Which treatment is most appropriate?
   a. Laboratory workup and penicillin.
   b. Oral cetirizine, topical olopatadine and loteprednol.
   c. Debridement, diamond burr polish and bandage contact lens.
   d. Oral doxycycline and tobramycin-dexamethasone eye drops.

Discussion

The recurrent nature of the condition, concurrent rosacea blepharitis and classic corneal appearance led to a diagnosis of phlyctenular keratoconjunctivitis. The ciprofloxacin from the PCP was not helpful, as this condition is primarily due to a hypersensitivity reaction, which is inflammatory in nature. Staphylococcal and tuberculosis antigens are the most common causes, and there is often concomitant ocular rosacea noted as well. The recurrent inflammation creates a nodule that can progress centrally across the cornea as what is termed a “marching phlyctenule,” trailed by a leash of superficial neovascularization.

Fortunately, this patient responded very well to treatment with 50% improvement in symptoms in 48 hours and is currently quiescent on prophylactic therapy of cyclosporine 0.05% eye drops and hypochlorous acid 0.02% lid spray.

Case Six

A 59-year-old female came to our clinic reporting recurrent sharp eye pain in the morning in the right eye over the past month. She had penetrating keratoplasty (PKP) in the left eye that was doing well and had not seen an eye doctor for quite some time. Her vision was 20/50 in the right eye and 20/25 in the left eye. She mentioned decreased vision along with worsening glare at night over the last six months in the right eye. There was no inflammation in either cornea, no arcus, no sutures re-
remaining in the transplant and the slit lamp exam revealed the findings in the photograph in the right eye only.

**Corneal Findings**

11. *What is your tentative diagnosis?*
   a. Schnyder’s dystrophy with recurrent corneal erosions.
   b. Infectious crystalline keratopathy.
   c. Anterior basement membrane dystrophy.
   d. Lattice dystrophy with recurrent corneal erosions.

12. *Which treatment is most appropriate?*
   a. Superficial keratectomy and amniotic membrane.
   b. Descemet’s membrane endothelial keratoplasty.
   c. Phototherapeutic keratectomy (PTK).
   d. Culture and fortified antibiotics.

**Discussion**

Corneal dystrophies can be difficult to differentiate, but this appearance of glass-like branching lines in the anterior stroma is classic for lattice corneal dystrophy, the most common of the stromal dystrophies. These patients suffer from recurrent corneal erosions, which can be managed medically to start, but often end up undergoing PTK to reduce symptoms and improve vision. The opacities tend to coalesce with age and can impact visual acuity as the central cornea becomes more involved.

Historically, PKP was the surgery of choice to restore vision, but now deep anterior lamellar keratoplasty and PTK are excellent options as well and have gained popularity, given their lower risk profiles. The amyloid deposits can recur both after PTK and in corneal grafts after five to 10 years.

This patient opted for PTK and was educated on the likelihood of family members being affected with an autosomal-dominant inheritance pattern.

**THANKS FOR PLAYING!**

We hope you not only fine-tuned your corneal diagnostic skills but had fun learning about these cases, which we believe can help you assess patients with similar conditions that may make their way to your office.

Check out our Instagram page, @kmkoptometrypro, to see more interesting cases and continue testing your knowledge, and let us know your thoughts!

<table>
<thead>
<tr>
<th>Case 6. Notice the glass-like branching lines in the anterior stroma in this patient.</th>
<th>Answers</th>
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<tbody>
<tr>
<td>1. b</td>
<td>7. a</td>
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<tr>
<td>2. c</td>
<td>8. a</td>
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<td>5. d</td>
<td>11. d</td>
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<tr>
<td>6. d</td>
<td>12. c</td>
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Improving the Gold Standard for CXL

Here’s a look at what’s currently available, what’s in the works and what’s on the horizon.

BY LILY ARENDT, OD
SAN ANTONIO, TX

In 2016, we gained the ability to treat our keratoconus patients suffering from progressive vision loss through the FDA approval of epi-off corneal crosslinking. Historically, keratoconus patients were given glasses or contact lenses to help optimize their vision in the hopes that they would not progress. Thankfully, we can now slow and even halt keratoconus progression, reducing the number of patients who need corneal transplants.

As we continue to perform conventional epi-off corneal crosslinking, outside the United States crosslinking protocols are evolving and researchers are proposing new protocols that aim to offer a faster, more comfortable and effective way to help our patients. This article will look at various crosslinking methods, including accelerated, pulsed light, transepithelial and others that seek to improve on the gold standard, and discuss what these advancements and innovations could mean for us and our patients in 2022 and in the future.

Conventional Epi-off Crosslinking

The current gold standard for corneal crosslinking in patients with progressive keratoconus and other corneal ectasia is the epi-off Dresden protocol created in 2003. This process involves the removal of 8mm to 9mm of epithelium from the central cornea, followed by the application of a riboflavin solution every two minutes for 30 minutes. After the cornea is saturated, it is irradiated using 370nm UVA at an intensity of 3mW/cm² while riboflavin is applied every two minutes for another 30 minutes. A minimum corneal thickness of 400μm is required before UV irradiation or intraoperatively. This is the only FDA-approved crosslinking protocol in the United States, and it has proven to be effective at strengthening the cornea and halting progression in patients with keratoconus.

While effective, the Dresden protocol has its disadvantages. Removal of the epithelium has a higher risk of pain and increased discomfort. It is greatest in the first three days postoperatively and often requires the use...

About the author

Dr. Arendt is an ocular disease and refractive surgery resident at Parkhurst Nuvision in San Antonio, TX. She is a member of the Refractive Surgery Alliance, American Optometric Association and Texas Optometric Association. She has no financial interests to disclose.
of topical or oral pain medications. There is a chance that the patient will develop reduced visual acuity, but per the data submitted to the FDA, only 1% to 2% of patients had corneal opacity or scarring at the 12-month follow-up.

Typical postoperative protocol includes application of a bandage contact lens immediately after treatment and frequent use of topical antibiotics during the first three to five days as the epithelium heals. Upon removal of the bandage contact lens, the patient is often switched to topical steroids to decrease inflammation and reduce the risk of persistent corneal haze. This is different than the postoperative haze typically seen after crosslinking between months one through six, which is not visually significant and does not require steroids for resolution. Adding topical steroids before the corneal epithelium has fully healed has been linked to an increased risk of microbial keratitis.

**Epi-off Accelerated Crosslinking**

Conventional crosslinking lasts a minimum of one hour and is one of the longest procedures performed in ophthalmology clinics today. To make the process more efficient and less time-consuming, researchers and clinicians have been trying out new accelerated protocols with higher UVA irradiation employed for a shorter amount of time. This is based on the Busen-Roscoe law of reciprocity: “a response is determined by the product of the intensity and the duration of the stimulus.” This means that for an accelerated protocol to achieve similar results as the Dresden protocol, it would need to maintain the same or have a cumulative energy of 5.4J/cm².

In a 2021 study, the seven-year data comparing epi-off accelerated crosslinking and conventional epi-off crosslinking in progressive keratoconus was shared. The accelerated protocol used UVA irradiation of 9mW/cm² for a duration of 10 minutes instead of the typical 30 minutes. There was no significant difference between protocols.

Conversely, another study included multiple accelerated protocols, including three minutes at 30mW/cm², five minutes at 18mW/cm² and 10 minutes at 9mW/cm², and found that each accelerated protocol was less effective at flattening the cornea when compared with conventional crosslinking. While there is some promise of achieving successful crosslinking in shorter duration, more research is needed to find the ideal energy and duration.

**The Importance of Oxygen**

Not only is the energy and duration important but we now know that oxygen is also key in the crosslinking process. For corneal crosslinking to occur, UVA must interact with both riboflavin and oxygen in the stroma to create oxygen free-radicals that kick-start the formation of covalent bonds between collagen molecules.

To illustrate the importance of oxygen in crosslinking, *ex vivo* epi-off crosslinking was performed on porcine corneas in both a regular oxygen environment (21%) and a low-oxygen environment (<0.1%). The corneas that were crosslinked in the normal oxygen environment showed improved biomechanical stability, while the corneas in the low-oxygen environment had results similar to the non-treated controls. This ultimately raised concerns over the effectiveness of transepithelial crosslinking (epi-on) where the corneal epithelium limits the amount of oxygen diffusion into the stroma.

In an effort to make epi-off crosslinking even more effective, it was determined that the level of oxygen in the stroma depletes within seconds after exposure to UVA irradiation and takes three to four minutes to replenish to normal levels. From this came a crosslinking method called pulsed light accelerated crosslinking, in which UVA irradiation is applied on and off repeatedly to allow for oxygen to diffuse back into the stroma. When compared with conventional crosslinking, this method alone was not as effective because it did not penetrate as far down into the cornea as conventional crosslinking but may prove to be a key component in future crosslinking protocols.

To determine how much of the cornea is crosslinked, researchers measure what is called the stromal demarcation line, which represents a transition zone between crosslinked and non-crosslinked stromal tissue. It has been found to occur in histopathological studies as early as 24 hours and has been seen as early as two weeks on slit lamp biomicroscopy. The presence and depth of a stromal demarcation line has been considered...
This technique was performed in a study that produced unsuccessful results, showing no change in corrected distance visual acuity or Kmax postoperatively. This was attributed to the fact that contact lenses act as a barrier by preventing most UVA from reaching the stroma and interfering with oxygen diffusion.

A new protocol was developed in 2021 to crosslink ultrathin corneas. Referred to as the sub400 protocol, it uses a previously published algorithm to perform customized crosslinking based on each eye’s corneal thickness after epithelial removal. Each eye receives the same UVA irradiation (3mW/cm²), but the duration varies in order to crosslink without harming the endothelium. Study results showed successful crosslinking in corneas ranging in thickness from 214µm to 398µm with keratoconus progression halted in 90% of eyes.

While many researchers have been trying to effectively crosslink patients with thinner corneas, others are focused on creating a quicker, safer and more comfortable procedure for patients all while leaving the epithelium in place. This new wave in crosslinking is called transepithelial crosslinking, or epi-on crosslinking.

Epi-On Crosslinking

Transepithelial crosslinking leaves the epithelium intact and has significantly less postoperative pain, offers a quicker visual recovery (with vision often returning to preoperative levels within two or three days) and allows patients to resume contact lens wear in one week. There is also a decreased risk of postoperative infection and persistent corneal haze infrequently seen with conventional crosslinking. With the quick postoperative recovery associated with transepithelial crosslinking, it raises the possibility of performing bilateral crosslinking on the same day.

While the benefits may seem to speak for themselves, an intact epithelium is quite literally a barrier to a successful crosslinking procedure. Riboflavin is a hydrophilic, high molecular weight molecule that cannot penetrate through an intact epithelium on its own. In a study evaluating the difference in biomechanical effect for both epi-on and epi-off crosslinking in rabbits, it was concluded that epi-on crosslinking produced only one fifth of the biomechanical rigidity in the cornea found after conventional crosslinking. When this is translated to human corneas, it would likely equate to a 64% increase in rigidity with transepithelial vs. a 320% increase with conventional crosslinking. The authors attributed this disparity to insufficient riboflavin diffusion through an intact epithelium.

Certain chemicals can be used to enhance riboflavin permeability and often include benzalkonium chloride, gentamicin and ethylenediamine tetra-acetic acid (EDTA). Transepithelial crosslinking was performed on 26 eyes of patients between the ages of 11 and 26 with a riboflavin solution containing two permeability enhancers, trometamol and EDTA. Results showed that both uncorrected and corrected distance acuity improved in the first three to six months before gradually returning to baseline over 24 months. A similar trend was found with Kmax values. Due to the ineffectiveness of treatment at the 12-month follow-up, the team ultimately retreated 50% of the patients younger than 18 with epi-off crosslinking. It was concluded that transepithelial crosslinking was likely not a good option for halting progression in patients younger than 26 who have more aggressive forms of keratoconus.

Iontophoresis

Another way that researchers have tried improving riboflavin permeabil-
When it comes to ocular surface inflammation associated with dry eye disease, FLAREX® IS A PROVEN WINNER.

The power of Pred Forte* (prednisolone acetate ophthalmic suspension, USP) 1% with the safety of FML*1 (fluorometholone ophthalmic suspension, USP) 0.1%.

97% of ocular surface inflammation was resolved or improved with FLAREX vs 89% with Pred Forte.1

Head-to-head with FML:

FLAREX was significantly more effective in the resolution of external non-infectious inflammatory conditions of the eye (P=0.03)1

Head-to-head with Pred Forte:

FLAREX had comparable, non-inferiority efficacy in the treatment of external non-infectious inflammatory conditions of the eye1

In the FDA pivotal clinical evaluation:

No reported adverse events in any treatment group when evaluated versus Pred Forte and FML1

DISPENSE AS WRITTEN. THERE IS NO GENERIC EQUIVALENT OF FLAREX. BE SURE TO PRESCRIBE IT BY NAME.

INDICATIONS AND USAGE
FLAREX® (fluorometholone acetate ophthalmic suspension) is indicated for use in the treatment of steroid-responsive inflammatory conditions of the patebral and bulbar conjunctiva, cornea, and anterior segment of the eye.

IMPORTANT SAFETY INFORMATION
ADVERSE REACTIONS: Glaucoma with optic nerve damage, visual acuity and field defects, cataract formation, secondary ocular infection following suppression of host response, and perforation of the globe may occur.

Please see the Full Prescribing Information on the next page.


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FLAREX NDC NUMBER: 71776-100-05
FLAREX® (fluorometholone acetate ophthalmic suspension) 0.1%

Brief Summary

INDICATIONS AND USAGE
FLAREX (fluorometholone acetate ophthalmic suspension) is indicated for use in the treatment of steroid-responsive inflammatory conditions of the palpebral and bulbar conjunctiva, cornea, and anterior segment of the eye.

DOSE AND ADMINISTRATION
Shake Well Before Using. One to two drops instilled into the conjunctival sac(s) four times daily. During the initial 24 to 48 hours, the dosage may be safely increased to two drops every two hours. If no improvement after two weeks, consult physician. Care should be taken not to discontinue therapy prematurely.

CONTRAINDICATIONS
Contraindicated in acute superficial herpes simplex keratitis, vaccinia, varicella, and most other viral diseases of the cornea and conjunctiva; mycobacterial infection of the eye; fungal diseases; acute purulent untreated infections, which like other diseases caused by microorganisms, may be masked or enhanced by the presence of the steroid; and in those persons who have known hypersensitivity to any component of this preparation.

WARNINGS AND PRECAUTIONS
Topical Ophthalmic Use Only
For topical ophthalmic use only. Not for injection.

Intraocular Pressure Increase
Prolonged use may result in glaucoma, damage to the optic nerve, and defects in visual acuity and visual field. It is advisable that the intraocular pressure be checked frequently.

Cataracts
Use of corticosteroids may result in cataract formation.

Delayed Healing
Topical ophthalmic corticosteroids may slow corneal wound healing. In those diseases causing thinning of the cornea or sclera, perforation has been known to occur with chronic use of topical steroids.

Viral Infections
Use in the treatment of herpes simplex infection requires great caution.

Bacterial Infections
Use of corticosteroids may suppress the host response and thus aid in the establishment of secondary ocular infections. Acute purulent infections of the eye may be masked or exacerbated by the presence of steroid medication.

Fungal Infections
Fungal infections of the cornea are particularly prone to develop coincidentally with long-term local steroid application. Fungus invasion must be considered in any persistent corneal ulceration where a steroid has been used or is in use.

Contamination
Do not touch dropper tip to any surface, as this may contaminate the suspension.

Contact Lens Wear
Contact lenses should be removed during instillation of FLAREX but may be reinserted after 15 minutes.

Temporarily Blurred Vision
Vision may be temporarily blurred following dosing with FLAREX. Care should be exercised in operating machinery or driving a motor vehicle.

ADVERSE REACTIONS

Clinical Trials Experience
Glaucome with optic nerve damage, visual acuity and field defects, cataract formation, secondary ocular infection following suppression of host response, and perforation of the globe may occur.

Postmarketing Experience
The following reaction has been identified during postmarketing use of FLAREX in clinical practice. Because reactions are reported voluntarily from a population of unknown size, estimates of frequency cannot be made. The reaction, which has been chosen for inclusion due to either its seriousness, frequency of reporting, possible causal connection to FLAREX, or a combination of these factors, includes dysgeusia.

USE IN SPECIFIC POPULATIONS

Pregnancy
Fluorometholone has been shown to be embryocidal and teratogenic in rabbits when administered at low multiples of the human ocular dose. Fluorometholone was applied ocularly to rabbits daily on days 6-18 of gestation, and dose-related fetal loss and fetal abnormalities including cleft palate, deformed rib cage, anomalous limbs, and neural abnormalities, such as encephalocoele, craniorachischisis, and spina bifida, were observed. There are no adequate and well-controlled studies of fluorometholone in pregnant women, and it is not known whether fluorometholone can cause fetal harm when administered to a pregnant woman. Fluorometholone should be used during pregnancy only if the potential benefit justifies the potential risk to the fetus.

Lactation
Systemically administered corticosteroids appear in human milk and could suppress growth, interfere with endogenous corticosteroid production, or cause other untoward effects. It is not known whether topical administration of corticosteroids could result in sufficient systemic absorption to produce detectable quantities in human milk. Because many drugs are excreted in human milk, caution should be exercised when FLAREX is administered to a nursing woman.

Pediatric Use
Safety and effectiveness in pediatric patients have not been established.

Geriatric Use
No overall differences in safety or effectiveness have been observed between elderly and younger patients.

NONCLINICAL TOXICOLOGY

Carcinogenesis, Mutagenesis, Impairment of Fertility
No studies have been conducted in animals or in humans to evaluate the possibility of these effects with fluorometholone.

PATIENT COUNSELING INFORMATION

Risk of Contamination
Do not touch dropper tip to any surface, as this may contaminate the suspension.

Use with Contact Lenses
The preservative in FLAREX, benzalkonium chloride, may be absorbed by soft contact lenses. Contact lenses should be removed during instillation of FLAREX but may be reinserted 15 minutes after instillation.

Temporarily Blurred Vision
Patients should be advised that their vision may be temporarily blurred following dosing with FLAREX. Care should be exercised in operating machinery or driving a motor vehicle.

Rx Only
Distributed by: Santen Pharmaceutical Co., Ltd.
Fort Worth, TX 76102
Corneal crosslinking (CXL) is a procedure that uses ultraviolet A (UVA) to strengthen the cornea by chemically crosslinking the collagen fibers. This process helps to stabilize the cornea and prevent further progression of keratoconus, a condition in which the cornea thinning occurs, causing vision problems.

**Procedure Details**

The C-Eye by Emagine AG, a UVA illumination device, was designed to be mounted onto the slit lamp, allowing patients to receive crosslinking while seated in an exam room. For epi-off crosslinking, the epithelium is removed at the slit lamp using an ethanol-based technique. After the patient is relocated to a reclining chair where corneal thickness is measured while riboflavin is instilled, they are then moved back to the slit lamp for UVA irradiation treatment lasting for 10 minutes. The creator of this device says that it will “increase accessibility to treatment globally and reduce overall costs related to the procedure” by moving it from the operating room to the exam room. Technology like this may be the next step toward optometrists performing crosslinking. At the start of 2022, there are currently three states whose scope of practice allows ODs to perform the procedure: Virginia, Louisiana, and Oklahoma.

**What’s Next?**

Glauskos, the creator of the first-generation iLink therapy currently FDA-approved for epi-off corneal crosslinking, is currently vying for FDA approval of an epi-on crosslinking protocol. In February 2021, Glauskos shared positive results from its pivotal Phase III trial for iLink epi-on therapy. The study included 279 eyes randomized in a 2:1 ratio to receive transepithelial crosslinking or placebo-controlled treatment. The study demonstrated a statistically significant improvement in Kmax at six months from baseline (-1.00 D), meeting the primary efficacy endpoint. The company also shared that the treatment was well-tolerated, with no patients leaving the study due to adverse events. The next step for Glauskos includes submitting an NDA this year, and the company is aiming for FDA approval in 2023.

**Takeaways**

There are many protocols that have been created to try and improve on the gold standard of crosslinking, the epi-off Dresden protocol, which is currently the only FDA-approved crosslinking method in the United States. Researchers from all over the world are looking for ways to make the procedure faster, more effective, more comfortable and even more accessible to patients. Until we find a better option, we can rest assured knowing that we do currently have a way to halt progression in our patients with progressive keratoconus and improve the quality of their lives in invaluable ways.

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Throughout our careers as eye-care professionals, we encounter many concerning conditions with the potential for devastating visual outcomes. One in particular is the often-frustrating herpes simplex virus keratitis (HSVK). Due to its varied presentation and recurrence rate, delays in proper diagnosis and treatment often lead to corneal scarring and poor visual outcomes. Thus, it has become increasingly important to recognize the signs and symptoms of this condition and promptly initiate appropriate management. By increasing our understanding of HSVK, we can improve our patients’ overall comfort and their visual prognosis.

Understanding the Virus
To truly recognize why HSVK can be so visually devastating, we must first understand the pathophysiology that makes this pathogen so prevalent.

Epithelial disease presents with dendritic lesions that are characteristic of the condition.

The herpes simplex virus (HSV) is a linear, double-stranded DNA virus that belongs to the Herpesviridae family. There are over 100 known herpes viruses within this family, including HSV-1, HSV-2, varicella-zoster virus, cytomegalovirus and the Epstein-Barr virus.

HSV is unique in its ability to reproduce quickly in a variety of tissues and establish latency, which can be reactivated at a later time. This has resulted in HSVK becoming the leading cause of infectious corneal blindness among developed nations. In fact, over 10 million people worldwide may...
have herpetic eye disease, with the seroprevalence being over 50% in the United States alone. HSV-1 is the most common serotype associated with ocular infections. Ocular involvement rarely stems from HSV-2, a disease typically sexually transmitted through the transferral of infected secretions. There are two main stages of infection in the pathophysiology of HSV-1: primary infection and reactivation (or recurrence). Primary infection occurs through direct contact with mucous membranes of the face, lips or eyes due to trigeminal nerve innervation of all three structures. These primary infections commonly cause contagious cold sores or fever blisters in and around the mouth. They rarely involve the cornea. Signs and symptoms of corneal manifestations associated with this phase of the infection are often subclinical and mild.

Once a primary infection occurs in the mucous membrane of the eye, the virus then travels down to the trigeminal ganglion, where it becomes latent. Any event that stresses the immune system, whether it be physiological (e.g., fever) or otherwise (e.g., environmental stressors), can then result in reactivation of the latent virus. After the virus is reactivated it travels along the ophthalmic branch of the trigeminal ganglion to the cornea, resulting in either superficial epithelial replication, which presents as dendritic keratitis, or an immune-mediated response (stromal or endothelial keratitis).

Making the Correct Diagnosis
HSV can have various clinical manifestations, making it challenging for the clinician to determine a proper diagnosis. It is essential to first obtain a thorough case history to gather any relevant information that can assist in the diagnosis. Important questions to ask include recent use of topical or oral corticosteroids, ocular trauma, ocular surgeries, psychological stress, history of HSVK, recent illnesses, fever, extraordinary ultraviolet light exposure, immunosuppression, history of sexually transmitted diseases and hormonal changes. Physical stimuli, emotional triggers and corticosteroid use can weaken the immune system, allowing the virus to reactivate and causing ocular HSV.

Use of vital dye staining is extremely helpful in distinguishing dendrites and making an accurate diagnosis.
Next, a gross examination of the facial adnexa can reveal vesicles on the forehead, eyelids or nose that may respect the vertical midline, which would narrow down the diagnosis to herpes zoster ophthalmicus, not HSVK. The latter typically presents with an acute onset of symptoms of unilateral ocular pain, redness, photophobia, tearing, itching, foreign body sensation, irritation and blurred vision.1 HSVK can usually be diagnosed with a slit lamp biomicroscopy exam alone and without any laboratory testing. Corneal sensitivity evaluation is also important and is easily performed using a cotton wisp by first testing the unaffected cornea to determine any differences in corneal sensation compared with the non-affected eye. Decreased corneal sensation is a hallmark for herpetic keratitis, unlike other microbial infections where there is an increase in sensation.8

Corneal sensory loss is a common sequela from both herpes simplex keratitis and herpes zoster ophthalmicus, with more severity in the latter.9 The degree of corneal sensory loss is directly proportional to the number of recurrences.9 During the slit lamp examination, dyes such as fluorescein, rose bengal and lissamine green are essential in assisting the clinical diagnosis of HSVK. Corneal dendrites are easily seen with fluorescein dye, since the body of the dendrite will illuminate as there is epithelial breakdown and decaying cells present. On the other hand, rose bengal and lissamine green will stain the dendrite’s viral infected cells at the margins of the swollen terminal bulbs.6,7,10 These dyes also assist the clinician in correctly distinguishing a corneal dendrite from a pseudo-dendrite caused by herpes zoster ophthalmicus. Pseudodendrites have raised centers filled with swollen cells without terminal bulbs, ulceration or branching.11,12 They also stain poorly with vital dyes, producing a negative staining with fluorescein and rose bengal.6,12

A slit lamp examination is typically sufficient in diagnosing HSVK; however, laboratory and diagnostic tests are available for atypical cases where a diagnosis of HSVK cannot easily be made. The main diagnostic tests for HSVK include culturing, direct fluorescent antibody (DFA) and polymerase chain reaction (PCR). Other, less commonly used diagnostic tests include cytology, enzyme-linked immunosorbent assay and serology.13 Culturing HSV-1 is considered the

<table>
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<th>Table 1. Topical Antiviral Therapy Dosing13</th>
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<tr>
<td><strong>Antiviral</strong></td>
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<tr>
<td>Trifluridine ophthalmic solution 1% (Viroptic)</td>
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<tr>
<td>Ganciclovir gel 0.15% (Zirgan)</td>
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<tr>
<td>Acyclovir ointment 3% (Avacyl)</td>
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Even patients who only demonstrate small dendrites such as this one can present with significant pain and discomfort.
gold standard. Culturing a virus is a misnomer as a virus cannot grow on a culture plate of agar but can infect other cells on the plate, thus proliferating the virus. Only cases of HSV epithelial keratitis can be cultured, since a virus cannot be cultured from the stroma or endothelium.6,13 Culturing tends to have high specificity and low sensitivity and typically takes up to 10 days to obtain the results.13

DFA and PCR have proven to be reliable alternatives in determining the presence of HSV-1. DFA detects any HSV-1 antigens and provides rapid results with lower specificity than PCR, which detects viral DNA and is shown to be as sensitive and specific as a cell culture.13 A limitation of PCR is its inability to differentiate between pathological levels of HSV and normal shedding of HSV in the tear film.13

Serology is not commonly used, since a majority of people have already had prior exposure to HSV; thus, it has low specificity.8,13 Serology is more useful in younger patients where primary infections are more common.13

**Clinical Presentations**

HSV can present with different corneal manifestations depending on what layer of the cornea the virus raids: the epithelium, stroma and/or endothelium.1 Herpes simplex epithelial keratitis is the most common subtype of HSVK and is responsible for 50% to 80% of all ocular herpes infections.14

Within 12 to 24 hours, infected epithelial cells form punctate vesicles. These vesicles, the beginning stages of dendrites, are formed by the swollen cell nuclei filled with replicating DNA viral load.5,15 As epithelial cells swell, apoptosis and shed the virus, adjacent cells become infiltrated, ultimately causing a corneal dendrite.

These dendritic lesions present with granular epithelium, branching linear pattern, terminal bulbs and raised gray edges.5,16 The body of the dendrite stains with fluorescein dye while the borders of the terminal bulbs stain with rose bengal or lissamine green. The dendritic lesion may progress and form a geographic ulcer in 25% of cases.5 HSV epithelial keratitis is classically the most painful type of HSVK as epithelial cells are invaded by the herpes virus, causing epithelial cell death. Common symptoms include pain, tearing, photophobia, foreign body sensation, conjunctival injection and decreased vision.15 HSV stromal keratitis can be described as diffuse, linear or disciform.5,16

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**Table 2. Oral Antiviral Therapy Dosing**

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<tr>
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<tbody>
<tr>
<td>Acyclovir</td>
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<tr>
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<td>800mg five times daily</td>
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<tr>
<td>Acyclovir</td>
<td>400mg two times daily</td>
</tr>
<tr>
<td>Valacyclovir</td>
<td>500mg one time daily</td>
</tr>
<tr>
<td>Famciclovir</td>
<td>250mg two times daily</td>
</tr>
</tbody>
</table>

* Prophylactic dose of oral antiviral with a therapeutic dose of topical corticosteroid.
** Therapeutic dose of oral antiviral with a limited dose of topical corticosteroid. Oral antiviral is reduced to prophylactic dose after seven to 10 days and maintained as long as topical corticosteroids are used.
*** Therapeutic dose of antiviral with a therapeutic dose of topical corticosteroid. Oral antiviral is reduced to prophylactic dose after seven to 10 days and maintained as long as topical corticosteroids are used.
with ulceration is commonly misdiagnosed as bacterial keratitis due to its clinical presentation of an epithelial defect with a stromal infiltrate. Corneal neovascularization can originate from the periphery and encroach toward the inflammatory site of the stroma. If left untreated, HSV stromal keratitis can develop corneal thinning, ulceration and conclusive perforation. Other complications include the formation of a corneal abscess, keratic precipitates, anterior uveitis and elevated intraocular pressure (IOP). Common symptoms for both HSV stromal keratitis with or without ulceration include photophobia, blurred vision, redness, photophobia and tearing. HSV stromal keratitis without ulceration is the most common form of HSV stromal keratitis. This subtype is sometimes referred to as “immune-mediated” or “non-necrotizing” and can be recurrent. It is caused by an inflammatory response against retained HSV antigens from a previous HSV episode inducing an antibody-complement cascade in the stroma. This stromal inflammatory response can occur in the latent or active stages of the infection. Herpes simplex stromal keratitis without epithelial ulceration presents as a stromal infiltrate without any epithelial disruption. The stromal inflammation may be focal, multifocal, diffuse and/or associated with anterior uveitis. If chronic, it can cause neovascularization, corneal scarring and corneal thinning.

HSV endothelial keratitis is another form of an immune-mediated response where the herpes virus spreads to the endothelium, causing endothelial cell dysfunction. Unlike HSV stromal keratitis, about half of these cases have no prior history of HSV epithelial keratitis and can occur independently from other forms of HSVK. HSV endothelial keratitis is commonly referred to as disciform keratitis; contrastingly, disciform is not completely accurate as HSV endothelial keratitis can also be considered linear or diffuse. The orientation of the associated keratic precipitates built up on the posterior wall of the endothelium and the extent of the stromal edema determines its classification. Disciform keratitis is the most common presentation of HSV endothelial keratitis, with disc-shaped stromal edema with underlying semicircle-shaped keratic precipitates.

Like the names suggest, linear endothelial HSVK has a vertical lined pattern of keratic precipitates and diffuse endothelial HSVK has keratic precipitates diffusely scattered across the entirety of the cornea. Clinical features of HSV endothelial keratitis include stromal edema caused by stromal deturgescence, overlying epithelial edema, inflammation of the endothelium with underlying keratic precipitates without significant anterior uveitis and elevated IOP.

Common symptoms are pain, redness, tearing, blurred vision and photophobia. Recurrent endothelial keratitis can cause endothelial cell loss, chronic corneal edema, scarring, corneal structural damage and irregular astigmatism.

Differential Diagnosis
One of the most difficult aspects of diagnosing and managing patients with HSVK is its wide variety of clinical manifestations. Due to its prevalence, consider HSVK as a possible diagnosis in any patient who presents with a unilateral red eye, especially with staining on the cornea. Familiarizing yourself with the differentials that may exhibit similar characteristics to HSVK will help to prevent delay in diagnosis and initiation of treatment.

Due to the fact that HSVK can impact multiple layers of the cornea, the differentials may vary based on which layer of tissue is impacted. For example, dendritic epithelial keratitis and linear or diffuse endothelial keratitis may have different differentials than...
those of stromal or endothelial keratitis, although in some cases these may overlap. HSV epithelial keratitis differentials should include other types of infectious and noninfectious keratitis that may cause dendrite-like or geographic-like lesions on the cornea. This list may be extensive and should include varicella zoster viral keratitis, *Acanthamoeba* keratitis, microbial keratitis without stromal involvement (*i.e.*, bacterial/fungal), *Chlamydia* epithelial keratitis, Epstein-Barr epithelial keratitis, recurrent corneal erosions, exposure keratopathy, Thygeson’s superficial punctate keratitis and epithelial basement membrane dystrophy (EBMD).13

Many of these conditions, such as *Acanthamoeba* keratitis and microbial keratitis, may be difficult to distinguish due to dendrite-like corneal presentations. Many times, these conditions can be differentiated based on a thorough clinical history. *Acanthamoeba* and bacterial keratitis are commonly associated with contact lens wear and water exposure. Fungal keratitis is likely associated with corneal insult or injury due to vegetation. HSVK can occur without contact lens use, water exposure or vegetative trauma. Other conditions such as Thygeson’s and EBMD differ from HSVK in that they most commonly present bilaterally, whereas bilateral HSVK is very rare.

Apart from HSV epithelial keratitis, HSV stromal and endothelial keratitis have their own list of differentials to consider. These separate conditions can also present with interstitial keratitis and keratouveitis that mimic the presentation of HSV stromal and endothelial disease.13 These differentials may include microbial keratitis with stromal involvement (any type), syphilis, Cogan’s syndrome, measles keratitis, mumps keratitis, Lyme disease, Posner-Schlossman syndrome, cytomegalovirus endothelial keratitis and corneal graft rejection.15

In many cases of HSVK differentials, possible diagnoses may be eliminated based on a thorough case history and clinical discussion with the patient. However, in cases where the puzzle cannot be solved based on verbal communication alone, use of corneal staining or, in some cases, laboratory testing may be needed to diagnose HSVK with confidence.

**Treatment and Management**

Complicating the proper diagnosis and treatment of HSV are the multiple corneal layers affected. The indicated medications and their dosages vary depending on the subtype of HSVK present. Treatment for HSVK includes oral antivirals, topical antivirals and topical corticosteroids; however, not all available treatment options are appropriate for each HSVK subtype. Thus, it is important to correctly diagnose the subtype of HSVK to formulate an appropriate treatment plan.

There are three topical antivirals approved by the FDA as treatment for HSVK: trifluridine ophthalmic solution 1%, ganciclovir gel 0.15% and acyclovir ointment 3% (*Table 1*).13 Acyclovir ointment 3% is currently not available to be prescribed as it is still on the discontinued drug product list for reasons other than safety and effectiveness.18 Trifluridine ophthalmic solution 1% is known to be more toxic to the ocular surface than ganciclovir gel 0.15% since it is nonselective against DNA synthesis of both normal and viral-infected cells.5 The cellular and ocular surface toxicity caused by trifluridine ophthalmic solution 1% can result in epithelial keratitis and delayed reepithelization. In contrast, ganciclovir gel 0.15% does not target the DNA of healthy cells, thus causing less ocular surface toxicity.

Oral antiviral agents are considered off-label for the treatment of HSVK. Currently, the available oral antivirals are acyclovir, valacyclovir and famciclovir (*Table 2*).8,13 These oral antiviral agents have excellent corneal and anterior chamber penetrance making
them effective treatment options for HSVK. However, due to the possibility of renal injury caused by oral antiviral agents, there needs to be special consideration when prescribing them for patients with kidney disease as there can be issues with the metabolism and excretion of the medication.\(^9\)

**HSV epithelial keratitis.** Oral antiviral agents are commonly used as treatment for HSVK epithelial keratitis without the use of topical corticosteroids, which are contraindicated as they can prolong epithelialization and cause the dendrite to progress into a geographic ulcer.\(^16\)

The benefits of treating with oral antivirals instead of topical antivirals are that they are more affordable, have reduced dosing schedules and are not toxic to the cornea. Both topical and oral antivirals have independently been shown to be equally effective in treating HSV epithelial keratitis; however, there has not been sufficient evidence that treating HSV keratitis with both oral and topical antivirals hastens the healing process.\(^5,16\)

The HEDS study showed no added benefit in concurrently treating HSVK with topical and oral antivirals.\(^13\) For HSV epithelial keratitis, oral acyclovir is dosed 400mg five times a day for seven to 10 days, valacyclovir is dosed 500mg two times a day for seven to 10 days and famciclovir is dosed 250mg two times a day for seven to 10 days.\(^6,13\)

If treating with topical antivirals, trifluridine ophthalmic solution 1% is dosed one drop nine times a day for seven days and then reduced to five times a day if the ulcer has healed after day seven. Treatment with trifluridine ophthalmic solution 1% should not exceed 21 days due to the associated risk of ocular toxicity. Ganciclovir gel 0.15% is dosed one drop five times a day until the ulcer resolves and then reduced to three times a day for seven days.\(^13\) For geographic ulcers, the dosages for oral antivirals are doubled for a treatment period of 15 to 21 days.

**HSV stromal keratitis without ulceration.** HSV stromal keratitis is considered an immune-mediated response to viral antigens in the stroma. Thus, a therapeutic dose of topical corticosteroids is needed to quell the inflammatory response alongside a dose of oral antivirals. It is important that oral antivirals are used while the patient is being treated with topical corticosteroids. In HSV stromal keratitis without ulceration, oral acyclovir is dosed 400mg twice daily, valacyclovir is dosed 500mg once daily and famciclovir is dosed 250mg twice daily. Topical corticosteroids can be dosed six to eight times daily and should be used for at least 10 weeks and tapered slowly as the cornea improves. Developing a treatment plan for HSVK is more of an art than a science, as the treatment is based on the patient’s medical and ocular history, clinical presentation and rate of healing. Also, considering HSV stromal and endothelial keratitis are unpredictable in nature with the risk of permanent vision loss, refer these patients to a board-certified corneal specialist to help with comanagement.

**HSV stromal keratitis with ulceration.** When ulceration is present, the treatment plan needs to change. A limited dose of topical corticosteroids can be used along with a therapeutic dose of an oral antiviral. Oral acyclovir is dosed 800mg three to five times a day for seven to 10 days, valacyclovir is dosed 1g three times a day for seven to 10 days and famciclovir is dosed 500mg two times a day for seven to 10 days. The oral antiviral should then be reduced to a prophylactic dose and maintained throughout the duration of treatment with the topical corticosteroid. There is a lack of data that supports any specific recommended length of treatment for

**Optometric Study Center**

**HSV KERATITIS**

HSV stromal keratitis with or without ulceration can be focal, multifocal or diffuse. This patient has HSV stromal keratitis with epithelial involvement.
this subtype of HSVK; therefore, clinical experience and presentation greatly aid in developing the treatment plan. Again, patients with HSV stromal keratitis should be further managed by a board-certified corneal specialist. Oral antivirals are preferred in treating any form of HSV stromal keratitis or endothelial keratitis since trifluridine ophthalmic solution 1% and ganciclovir gel 0.15% do not have adequate corneal stroma penetration.

**HSV endothelial keratitis.** This is relatively uncommon, and there are only a few studies that provide recommended treatment plans. Treatment with topical corticosteroids is needed, since it is an inflammatory-mediated response to the virus in the corneal endothelium. HSV endothelial keratitis responds extremely well to topical corticosteroids and, in comparison to HSV stromal keratitis, HSV endothelial keratitis heals at a remarkably faster rate.13 Therapeutic doses of both topical corticosteroids and oral antiviral agents are needed for this subtype of HSVK. Topical corticosteroids can be dosed one drop six to eight times a day and tapered slowly. Oral acyclovir is dosed 400mg three to five times daily, valacyclovir is dosed 500mg two times daily and famciclovir is dosed 250mg two times daily. The therapeutic dose of oral antiviral should be reduced to seven to 10 days and maintained throughout the entire course of the topical corticosteroid taper schedule. Ganciclovir gel 0.15% and trifluridine ophthalmic solution 1% are not recommended in treating HSV endothelial keratitis due to their poor corneal penetrance. Given the challenging nature of this condition, comanagement with a specialist can be useful for both the optometrist and their patient.

**Life After HSVK**

A previous history of HSV stromal keratitis and recurrent HSVK increases the risk of future HSVK recurrence.13 An oral antiviral maintenance dose can be introduced for patients who are at a higher risk to prevent recurrence of HSVK. The HEDS II study found a 45% reduction in recurrent episodes of ocular HSVK when on prophylactic treatment.19 The maintenance dose for acyclovir 400mg is two times daily for one year, valacyclovir 500mg is one time daily for one year and famciclovir is two times daily for one year. This long-term maintenance dose can be extended past one year for special cases that present a higher risk of recurrence, such as in immunocompromised patients. As for prophylaxis, a patient with a history of HSVK may also be on a short-term maintenance dose if they plan to undergo any ocular surgeries such as cataract surgery. Lastly, an oral antiviral maintenance dose is needed if a patient were to be prescribed a corticosteroid at any time. HSV stromal keratitis and endothelial keratitis, whether initial or recurrent episodes, are the greatest threat to permanent corneal structural damage and vision loss. These inflammatory reactions can cause corneal scarring, neovascularization, corneal thinning, decreased corneal sensation and fibrosis. Recurrent episodes place the patient at a higher risk of developing permanent corneal structural abnormalities. Approximately one million people worldwide are affected by permanent visual impairment due to ocular HSV.13 There are also a reported 1,000 penetrating keratoplasties annually in the United States due to visually significant corneal scarring from ocular HSV.

It is imperative to be vigilant in recognizing early clinical signs and symptoms of HSVK to help prevent severe cases from causing visually significant corneal damage. It is also important to educate patients and discuss possible visual outcomes and options. Comanaging with a corneal specialist is crucial in reducing your liability as a provider.

Depending on the course of the condition, the patient may benefit from an amniotic membrane for improved epithelial healing or a specialty contact lens for improved vision. A patient with significant corneal scarring and decreased vision may experience an improvement in vision from specialty lenses such as scleral lenses. HSVK is a common and often visually devastating condition. Doctors of optometry can better manage these patients through a detailed understanding of the clinical presentation, differentials and treatments. This not only increases confidence in managing the condition but also improves outcomes and overall visual prognosis.

1. HSV-1 is which classification of virus?
   a. Linear, single-stranded DNA virus.
   b. Linear, double-stranded DNA virus.
   c. Linear, triple-stranded DNA virus.
   d. None of the above.

2. Which is a member of the Herpesviridae family?
   a. HSV-2.
   b. Epstein-Barr virus.
   c. Cytomegalovirus.
   d. All of the above.

3. HSV-1 reactivation occurs along which branch of the trigeminal ganglion resulting in corneal involvement?
   a. Mandibular.
   b. Maxillary.
   c. Ophthalmic.
   d. None of the above.

4. Which type of HSVK is thought to be due to an immune-mediated response?
   a. Epithelial keratitis.
   b. Stromal keratitis.
   c. Endothelial keratitis.
   d. Both b and c.

5. Which is considered to be a differential diagnosis of HSV epithelial keratitis?
   a. Acanthamoeba keratitis.
   b. Cogan’s syndrome.
   c. Ruberosis irides.
   d. Arcus senilis.

6. Which is considered to be a differential diagnosis of HSV stromal keratitis?
   a. Cogan’s syndrome.
   b. Syphilis.
   c. Lyme disease.
   d. All of the above.

7. Which is not one of the three main diagnostic tests for HSVK?
   a. Culture.
   b. Serology.
   c. DFA.
   d. PCR.

8. Which is the most common subtype of HSVK?
   a. HSV endothelial keratitis.
   b. HSV stromal keratitis with ulceration.
   c. HSV epithelial keratitis.
   d. Disciform keratitis.

9. Which can trigger a recurrence of HSVK?
   a. Immunosuppression.
   b. Psychological stress.
   c. Ocular surgeries.
   d. All of the above.

10. Which is a form of HSV endothelial keratitis?
    a. Disciform.
    b. Diffuse.
    c. Linear.
    d. All of the above.

11. Which is commonly misdiagnosed as a bacterial infection?
    a. HSV epithelial keratitis.
    b. Disciform keratitis.
    c. HSV stromal keratitis without ulceration.
    d. HSV stromal keratitis with ulceration.

12. Which subtype of HSVK can manifest independently from other forms of HSVK?
    a. HSV stromal keratitis with ulceration.
    b. HSV endothelial keratitis.
    c. Necrotizing stromal keratitis.
    d. HSV stromal keratitis without ulceration.

13. Which vital dye stains the body of a dendrite?
    a. Lissamine green.
    b. Rose bengal.
    c. Fluorescein.
    d. Giemsa stain.

14. Which is not an FDA-approved topical antiviral medication?
    a. Trifluridine solution 1%.
    b. Vidarabine ointment 3%.
    c. Ganciclovir gel 0.15%.
    d. Acyclovir ointment 3%.

15. Which topical antiviral is dosed one drop nine times a day for seven days and then reduced to five times a day after seven days if the ulcer is healed?
    a. Trifluridine ophthalmic solution 1%.
    b. Vidarabine ointment 3%.
    c. Ganciclovir gel 0.15%.
    d. Acyclovir ointment 3%.

16. What is the maximum number of days a patient can be on trifluridine ophthalmic solution 1% treatment?
    a. 15 days.
    b. 21 days.
    c. 30 days.
    d. 10 days.

17. Topical antiviral medications are not recommended for which subtype of HSVK?
    a. HSV stromal keratitis.
    b. HSV epithelial keratitis.
    c. HSV endothelial keratitis.
    d. Both a and c.

18. Which is the correct initial dose for an oral antiviral agent in treating HSVK stromal keratitis with ulceration?
    a. Acyclovir 800mg three to five times daily.
    b. Valacyclovir 1g three times daily.
    c. Famciclovir 500mg two times daily.
    d. All of the above.

19. Which is the recommended initial treatment for HSVK stromal keratitis without ulceration?
    a. Prophylactic dose of an oral antiviral agent.
    b. Therapeutic dose of topical corticosteroid.
    c. Limited dose of topical corticosteroid.
    d. Both a and b.

20. Which reacts extremely well to topical corticosteroids and heals at a faster rate compared with other subtypes of HSVK?
    a. HSV epithelial keratitis.
    b. HSV endothelial keratitis.
    c. HSV stromal keratitis without ulceration.
    d. HSV stromal keratitis with ulceration.
Examination Answer Sheet

Piecing Together the HSVK Puzzle
Valid for credit through April 15, 2025

Online: This exam can be taken online at revieweducationgroup.com. Upon passing the exam, you can view your results immediately and download a real-time CE certificate. You can also view your test history at any time from the website.

Directions: Select one answer for each question in the exam and completely darken the appropriate circle. A minimum score of 70% is required to earn credit.

Answers to CE exam: Post-activity evaluation questions:
1. A A A A A
2. A A A A A
3. A A A A A
4. A A A A A
5. A A A A A
6. A A A A A
7. A A A A A
8. A A A A A
9. A A A A A
10. A A A A A
11. A A A A A
12. A A A A A
13. A A A A A
14. A A A A A
15. A A A A A
16. A A A A A
17. A A A A A
18. A A A A A
19. A A A A A
20. A A A A A
21. Understand the presentations of herpes simplex virus keratitis. Rate how well the activity supported your achievement of these learning objectives. 1=Poor, 2=Fair, 3=Neutral, 4=Good, 5=Excellent
22. Distinguish between HSVK and differential diagnoses.
23. Diagnose patients with herpes simplex virus keratitis.
24. Medically manage herpes simplex virus keratitis cases.
25. Based upon your participation in this activity, do you intend to change your practice behavior? (Choose only one of the following options.)
26. Thinking about how your participation in this activity will influence your patient care, how many of your patients are likely to benefit? (please use a number): 
27. If you plan to change your practice behavior, what type of changes do you plan to implement? (Check all that apply.)
28. How confident are you that you will be able to make your intended changes?
29. Which of the following do you anticipate will be the primary barrier to implementing these changes?
30. Additional comments on this course: ____________________________________________________________

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City
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Rate the quality of the material provided:
1=Strongly disagree, 2=Somewhat disagree, 3=Neutral, 4=Somewhat agree, 5=Strongly agree
31. The content was evidence-based. 
32. The content was balanced and free of bias. 
33. The presentation was clear and effective.

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APRIL 15, 2022 | REVIEW OF OPTOMETRY 33
Brush Up on Your Low Vision Skills

You can make rehabilitation a priority in your practice with fewer hassles than you may think.

The term “low vision” refers to vision loss that is uncorrectable with current medical and surgical interventions or using traditional spectacle and contact lenses. While the Centers for Medicare and Medicaid Services recognize distinct categories of low vision based primarily on best-corrected visual acuity (VA), a much broader range of patients may also be defined as having “low vision” based on compromised visual fields, impaired contrast sensitivity, or other deficits in visual function.

The practice modality known as vision rehabilitation involves development of an individual rehabilitation plan (IRP) to help care for the patient with low vision. According to the American Optometric Association, an IRP may include “prescription glasses or contact lenses, optical and electronic magnification devices, assistive technology, glare control with therapeutic filters, contrast enhancement, eccentric viewing, visual field enhancement, non-optical options and referral for additional services with other professionals.”

A 2017 analysis estimated that 7.08 million people in the United States were living with uncorrectable VA loss. The same study indicated that 1.08 million of these individuals demonstrated VA loss that qualified them to meet the federal definition of “legal blindness.” When considering the other deficits in visual function that could convey “low vision” or “legal blindness” status, these numbers grow larger yet. Projections estimate the number of legally blind individuals will double by the year 2050, due in large part to the prevalence of age-related eye disease and the general aging of the US population.

Our optometric training and scope of practice makes us the ideal providers of vision rehabilitation care. We are well-equipped to combine our understanding of optics with our knowledge of ophthalmic diseases and their functional implications to effectively serve the visually impaired patients in our practices. However, for those of us outside of academic and not-for-profit practice settings, it may be difficult to tackle IRP development and provide comprehensive vision rehabilitation care. For this reason, it is important to discuss the ways in which all practicing optometrists can readily incorporate some foundational vision rehabilitation components into their practice to better serve this growing patient population.

Levels of Vision Rehabilitation

In 2010, the Association of Schools and Colleges of Optometry (ASCO) convened a working group of optometric vision rehabilitation educators from across the country to define and delineate tiered competencies for vision rehabilitation care among graduates from schools and colleges of optometry.

The result of this group’s work is summarized in Table 1, which clearly lays out 20 entry level competencies for vision rehabilitation care that can be expected of any practicing optometrist regardless of their practice setting. It is valuable to review each of the points laid out in Table 1 and elaborate on the ways in which these can be applied in most optometric practices.

About the author

Dr. Robinson is an assistant professor in the department of ophthalmology and visual sciences and director of Low Vision Rehabilitation at the Vanderbilt Eye Institute. He is a Fellow of the American Academy of Optometry and a Clinical Diplomate in the Academy’s Low Vision Section. He has no financial disclosures.
Patient History Competencies
The first of the 20 ASCO competencies shown in Table 1 has to do with the foundational knowledge of epidemiology and vision impairment we all obtain through our strong optometric education and ongoing continuing education efforts. Our understanding of, and ability to communicate, these basic principles are rightfully at the top of the ASCO list of entry level competencies.

Competency number two requires that the optometrist obtain a thorough case history. A comprehensive understanding of a visually impaired patient’s rehabilitative needs requires a case history that goes beyond that of a routine eye exam and dives into specific functional difficulties caused by one’s vision loss, goals and expectations for the vision rehabilitation process.

While the exact set of intake questions used may vary from clinic to clinic, the National Eye Institute’s Visual Functioning Questionnaire-25 (VFQ-25) provides a solid bank of foundational questions from which to adapt an intake questionnaire for visually impaired patients. Listening when patients voice functional concerns relative to their vision loss, along with asking the appropriate questions to elicit functional concerns when needed, are foundational to the development of an effective IRP.

The next four ASCO competencies describe the ability to recognize various patient-specific factors that may influence the vision rehabilitation process. These include familial, psychological, and social factors as well as physical and neurologic comorbidities that may impact an IRP. Recognition of these factors may take place during the case history discussion or at any time throughout the vision rehabilitation evaluation.

If appreciation for any of these factors seems to fall outside of what we all took from our optometric education, Faye’s Clinical Low Vision (2nd Ed.) remains a proven and useful resource.

<table>
<thead>
<tr>
<th>TABLE 1. ASCO ENTRY LEVEL LOW VISION REHABILITATION COMPETENCIES7</th>
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<tbody>
<tr>
<td>1. Be able to apply epidemiologic aspects of visual impairment, appropriate terminology and classifications of visual impairment in order to communicate with patients, the public and other health care providers.</td>
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<tr>
<td>2. In addition to performing a standard case history, be able to ask basic questions about symptoms, functional difficulties and rehabilitation goals to anticipate the level of care that patients with visual impairment may require.</td>
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<tr>
<td>3. Be able to recognize functional implications, hereditary factors and prognoses of common causes of visual impairment and explain them in language understandable to patients, families and other care providers.</td>
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<tr>
<td>4. Be able to recognize psychological factors (e.g., depression, grief, motivation) that may affect adjustment to vision loss and the potential for rehabilitation.</td>
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<tr>
<td>5. Be able to recognize pertinent social factors (e.g., social support system, education level, vocation, physical environment) and how they may influence the rehabilitation plan and process.</td>
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<tr>
<td>6. Be able to recognize significant physical and neurological comorbidities (e.g., Parkinson’s disease, stroke dementia) that influence low vision rehabilitation and modify evaluation strategies and rehabilitation.</td>
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<td>7. Be able to perform visual acuity testing at both distance and near on patients with visual impairment using appropriate charts with proper documentation (e.g., working distance eccentric viewing illumination).</td>
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<td>8. Be able to perform trial lens refraction and modify refractive techniques for the patient with visual impairment (e.g., bracketing handheld Jackson cross cylinder).</td>
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<td>9. Be able to recognize common symptoms of contrast sensitivity loss, screen for loss, recommend basic modifications (e.g., filter, lens lighting and environmental options) and refer for comprehensive low vision rehabilitation when indicated.</td>
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<tr>
<td>10. Be able to detect scotomas of the central visual field, understand their impact on visual acuity and visual function and educate patients about their implications for activities of daily living.</td>
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<td>11. Understand basic optical principles of low vision rehabilitation devices and be able to predict magnification levels needed to achieve patient goals.</td>
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<tr>
<td>12. Be able to prescribe basic optical and non-optical low vision rehabilitation devices, provide training in their use and refer for comprehensive low vision rehabilitation when indicated.</td>
</tr>
<tr>
<td>13. Be able to recognize availability of and indications for use of adaptive technology (e.g., video magnification, software) and refer for comprehensive low vision rehabilitation when indicated.</td>
</tr>
<tr>
<td>14. Be cognizant of rehabilitation strategies for visual field deficits (e.g., sighted guide technique, orientation and mobility, visual field enhancement devices and equipment, scanning training) and refer for comprehensive low vision rehabilitation when indicated.</td>
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<tr>
<td>15. Develop an understanding of the special considerations for examining children, the elderly and the multiply handicapped and educate about referral options and potential for rehabilitation.</td>
</tr>
<tr>
<td>16. Understand relevant vision standards for driving, provide necessary assessment and documentation and refer for comprehensive low vision rehabilitation, driver education/training and medical evaluation when indicated.</td>
</tr>
<tr>
<td>17. Be aware of the criteria for legal blindness determination and be able to educate patients on the basic social and legal ramifications of legal blindness certification.</td>
</tr>
<tr>
<td>18. Understand that the needs of patients with visual impairment may require professional collaboration and be able to coordinate care with available rehabilitative, educational and social service resources.</td>
</tr>
<tr>
<td>19. Identify governmental, private and consumer organizations that offer support and information to individuals with visual impairment (e.g., NEI, Veterans Administration, state rehabilitation agencies, foundations for the blind, consumer advocacy groups and support groups).</td>
</tr>
<tr>
<td>20. Be familiar with third-party reimbursement for low vision rehabilitation services and materials.</td>
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Assessment Competencies

Competency number seven begins the actual vision examination process, detailing a thorough measurement of VA at distance and near. Depending on the practice setting and some patient-specific factors, various VA charts may be used. For distance VA measurement, a stand-based Early Treatment Diabetic Retinopathy Study (ETDRS; Precision Vision) chart remains the gold standard (Figure 1).

This chart is mobile and conveniently allows for adjustment of the patient-to-target distance in cases when a patient is unable to read the largest optotypes from the standard 4m viewing distance. The backlit version also eliminates the need to provide proper illumination of the surface of the chart. A useful alternative distance VA chart is the William Feinbloom Distance Test Chart for the Partially Sighted (Designs for Vision; Figure 2). In employing this option, it is important to use and document appropriate illumination of the target.

For measuring near VA, various options exist that use single letters/numbers, words or continuous text. One of the more commonly used continuous text options is the MNRead Reading Test (Lighthouse Low Vision Products; Figure 3). Regardless of the chart being used, documenting distance and near VA is completed in standard form of “testing distance/optotype size” in a common (metric or imperial) measurement, always noting eccentric view (if employed) and lighting used (where pertinent).

A careful trial frame refraction, described in competency number eight, dictates everything that comes thereafter in a vision rehabilitation evaluation. This type of refraction is not all that different from the expert phoropter refinements being performed on our patients on a daily basis. The main difference lies in the use of a trial frame, loose lenses and a handheld Jackson cross cylinder (JCC) lens, which simplify the refraction of a patient who uses an eccentric view or atypical head posture to reach a null point and dampen nystagmus.

Further, the trial frame and loose lenses allow for larger and more efficient jumps in lens power when a patient’s acuity dictates a larger “just noticeable difference” in lens power be demonstrated. Otherwise, the trial frame refraction uses the same order of spherocylindrical lens power refinement steps and bracketing technique as a refraction in the phoropter. The goal of the trial frame refraction is the same as any other refraction we perform: to optimize a patient’s visual clarity through correction of ametropia.

Competency number nine pertains to contrast sensitivity (CS), a measure of visual function that is often reduced in the visually impaired patient and is likely not a primary focus in most routine eye care settings. Formal clinical testing of contrast sensitivity is simple and may be completed with any number of validated clinical tests. The Pelli-Robson CS Chart and Mars Letter Contrast Sensitivity Test are two common options for efficiently testing CS.

The practicing optometrist should be comfortable recognizing and screening for contrast sensitivity limitations, potentially using one of the aforementioned tests, and should also be able to recommend simple modifications (filters and environmental lighting) to aid in these patients’ function. If not, they should be prepared to make a referral to a comprehensive vision rehabilitation program.

Another important visual function consideration is the recognition of, and modifications to account for, scotomas. Many patients needing vision rehabilitation care have central visual deficits from commonly encountered conditions such as age-related macular degeneration, various inherited macular dystrophies, retinal vascular occlusions and some optic nerve disorders. Presence of scotomas may be detected through knowledge of a patient’s diagnosis, responses during the intake interview, or formal tests such as microperimetry or the California Central Field Test.

Competency number 10 states that the optometrist should be able to detect scotomas and educate the patient on their presence and strategies, such as eccentric viewing, for working around them. Again, if they are unable to do so, referral to a comprehensive vision rehabilitation program should be initiated.

Treatment Competencies

A comprehensive understanding of optics is foundational to our optometric education and to competency number 11. Being able to predict magnification needs based on a patient’s visual function and goals and understanding the angular magnification properties of various devices used in vision rehabilitation are considered entry-level competencies.

For example, using a ratio of the target print size to the currently legible print size helps to determine the dioptic power needed to reach the target print size. This can...
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predict the appropriate handheld or spectacle-based tools for a given patient and guide the optometrist’s handling of the device demonstration, prescription and training process.

In general, one should be comfortable prescribing near reading add powers up to around six diopters and accounting for the corresponding convergence demand through the appropriate amount of base in prism. Otherwise, referral for comprehensive vision rehabilitation is indicated.

Basic vision rehabilitation device prescription and training are the focus of point number 12. The optometrist should be equipped to demonstrate low-powered optical magnification devices and non-optical devices such as lighting options and filters. If they choose not to perform these demonstrations, referral to a comprehensive vision rehabilitation program is the standard.

Recognition of pertinent assistive technology options for a given patient is an important part of developing an IRP. Standalone desktop and portable video-based magnifiers, as well as the mobile phone and tablet devices already owned by many of our patients, are examples of helpful assistive technology options that can benefit our visually impaired patients. Competency number 13 notes that it is adequate to recognize the need for, and refer to, appropriate assistive technology training programs when indicated.

When a patient’s visual impairment entails visual field compromise, the optometrist should be comfortable recognizing its impact on patient function and addressing this appropriately. As outlined in competency number 14, this may include prescription of devices that enhance visual field awareness (e.g., prisms, reverse telescopes) or referral for appropriate training such as sighted guide and formal orientation and mobility programs.

Just as in primary optometric care, the practitioner is likely to encounter children and people with multiple disabilities among their visually impaired patient base. Competency number 15 notes that the optometrist should be comfortable employing specialized examination techniques and making appropriate referral for services and training in these cases, when indicated.

Competencies number 16 and 17 stress the importance of knowing legal blindness and licensure laws to better serve visually impaired patients and to provide documentation or referral for additional training or services as needed. A refresher on legal blindness criteria can be found on the Social Security Administration’s website, while state-by-state vision testing standards for licensure are kept up to date by Prevent Blindness on their website.

Competency points 18 and 19 pertain to the multidisciplinary nature of comprehensive vision rehabilitation care. The optometrist should know when to include referral to other rehabilitation professionals and social or psychological support services in a patient’s IRP. Further, connecting a patient with the appropriate governmental or community support networks can be an important part in helping to meet that patient’s individual needs.

The American Printing House for the Blind, the National Federation of the Blind and the American Foundation for the Blind all maintain helpful online directories of available vision rehabilitation resources in one’s state or region.
Enable your patients to enjoy reading again

A genus of herbaceous flowering plants

Many flowers have a symmetry. When the petals are bisected through the center, we have any point and symmetrical halves are produced. The flower is said to be isosceles or regular, e.g., ...one or trilobatum. This is an example of radial symmetry.
Finally, competency point number 20 pertains to one’s understanding of vision rehabilitation billing, coding and reimbursement structures. These factors likely play a role in limiting the number of private practice optometrists providing entry level vision rehabilitation services, but it doesn’t have to be that way. The practicing optometrist should understand that vision rehabilitation billing is typically based on time spent and uses the Evaluation & Management (E&M, 99xxx) codes.

The time spent includes any face-to-face time with the patient, as well as time spent on the same day preparing for the case and documenting findings or communicating with other professionals regarding the case. It does not include the refraction or other diagnostic testing that are being billed as separate services.

Table 2 lays out time requirements for billing each E&M code for both new and established outpatients. Any additional time spent beyond the Level 5 allotment in each column can be billed in additional 15-minute increments using code 99417. The ability to bill visits based on time can help to offset the inherent reduction in patient volumes when providing vision rehabilitation care in a private practice setting.

As for the prescription of vision rehabilitation devices, it should be noted that most are not covered as durable medical equipment by Medicare and other medical insurance carriers. While we are hopeful that this will change in the coming years, it is important for the vision rehabilitation practitioner to be able to tap into other coverage options for these devices. Options may include state blind or vocational rehabilitation services, private grant opportunities, flexible spending accounts or other local funding resources.

Clinical Takeaways
Caring for visually impaired patients falls squarely in optometry’s wheelhouse. Our comprehensive understanding of both ophthalmic disease and optics puts us all in a position to help our visually impaired patients to maximize their quality of life.

As the prevalence of vision impairment continues to grow, provision of vision rehabilitation services will fall to optometrists in all practice settings.1 While not all of us are equipped to provide comprehensive vision rehabilitation services, the core competencies defined by the ASCO provide a road map to starting the vision rehabilitation process for patients who need it.2

As was noted in the preceding discussion of all 20 ASCO competencies, there will come a point in many vision rehabilitation cases where referral for more comprehensive services is indicated. This may be dictated by the need for near add powers above a certain threshold, prescription a biopic telescope, eccentric viewing training, or referral for vocational rehabilitation, among many other factors.

When the needs of the patient are simpler, private practice optometrists can use the ASCO competencies to enhance their care for this growing, and grateful, patient population.

In summary, it’s important to view vision rehab as a spectrum of interventions—many of which are accessible to primary care optometrists—rather than a binary choice that forces ODs to be “all-in” or “all-out.”

### TABLE 2. OUTPATIENT TIME-BASED E&M CODING LEVELS FOR NEW AND ESTABLISHED PATIENTS

<table>
<thead>
<tr>
<th>Level</th>
<th>New (Minutes)</th>
<th>Established (Minutes)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Level 1</td>
<td>99201 (10)</td>
<td>99211 (N/A)</td>
</tr>
<tr>
<td>Level 2</td>
<td>99202 (15)</td>
<td>99212 (10)</td>
</tr>
<tr>
<td>Level 3</td>
<td>99203 (30)</td>
<td>99213 (20)</td>
</tr>
<tr>
<td>Level 4</td>
<td>99204 (45)</td>
<td>99212 (30)</td>
</tr>
<tr>
<td>Level 5</td>
<td>99205 (60)</td>
<td>99215 (40)</td>
</tr>
</tbody>
</table>

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TO CUT OR NOT TO CUT?
Weighing Strabismus Surgery Referral

BY BRENDA MONTECALVO, OD
BEAVERCREEK, OH

There is no rush to immediately recommend surgery for strabismus. Collaborative efforts between the surgeon and optometrist allow for better outcomes. Developing all visual skills benefits the patient with strabismus, providing sensory fusion. The critical stage for sensory fusion is less limiting with new approaches used in vision therapy (VT). When amblyopia exists, VT should be the first-line approach in order to gain the best possible outcome for misalignment.

The Consequences of Strabismus
Patients living with constant strabismus face a lifetime of difficulties. They are more likely to be diagnosed with psychological disorders, often have low self-esteem, are less likely to find a partner, marry or have a family. They even have more difficulty being a competitive job applicant, compared with those without strabismus. Patients in their third decade of life with uncorrected strabismus are more likely to face long-term psychological disorders. One study showed that individuals with constant exotropia were three-times more likely to be diagnosed with a psychological disorder than controls. Patients with uncorrected strabismus also had higher rates of homicidal and suicidal ideation.

Strabismus puts people at a considerable disadvantage due to the noticeable nature of having a turned eye, resulting in negative social stigma. In addition, there are many professions that depend on stereopsis. Achieving the best possible outcomes for these patients is critical for them to avoid negative psychological, socioeconomic and employment consequences. New evidence shines light on understanding outcomes of surgical intervention.

As optometrists, we meet patients going through life with these challenges because of their strabismus. We are obligated to at least make all possible treatments known and available to those interested in pursuing better vision.

Creating the very best possible outcome involves a collaborative effort between the optometrist and the strabismic surgeon. Good communication and respect for each other’s skills allows for comprehensive care. First, take time to visit each other’s practices. Second, educate the staff on how each profession contributes to successful outcomes. Third, schedule regular meetings between the two teams to discuss specific cases.

Understanding when to refer for surgery and combining the skills of a good optometric vision therapist will give your patient the best opportunity for cosmetically straight eyes with some level of stereopsis. To facilitate this for the strabismus patient, follow these steps:

1. Measure best-correlated visual acuity (VA) of each eye.
2. Evaluate binocularity to determine deviation and if any immediate spectacle compensation will improve the binocular stasis.
3. Evaluate the ocular motor and accommodative systems.
4. Determine if there is any appreciation of diplopia under any condition.
5. Identify level of suppression and fusion.
6. If there is some level of visual motor function that can be improved by VT, eight sessions should be completed. Then re-evaluate the system for possible improvements, such as less suppression, better cosmetic alignment, improved oculomotor function and better accommodative flexibility.

7. If improvements are identified, continue with eight more sessions and re-evaluate.

8. If there is a plateau in improvement and the deviation is still obvious, referral for surgery should be suggested.

9. After surgery, re-evaluation of visual motor skills should be conducted and post-VT activities should be instituted to maintain gains made through surgery.

Procedure Risks
Short- and long-term complications from surgical intervention reinforce why collaboration is important. Post-op complications of strabismus surgery include conjunctivitis, scleritis, sub-Tenon’s abscess, orbital cellulitis, endophthalmitis, hypopyon, vitreous haze and scleral perforation.

Intraoperative surgical site complications can include scleral perforation, lost muscle, slipped muscle and oculocardiac reflex, while post-op surgical site complications can include postoperative infection, allergic reaction, foreign body granuloma, conjunctival inclusion cyst, conjunctival scarring, fat adherence, dellen, anterior segment ischemia, eyelid retraction, ptosis and possible change in refraction. Post-op strabismus complications can involve diplopia, hyperphoria, anti-elevation syndrome and iatrogenic Brown syndrome.

There are several risk factors to be aware of regarding strabismus surgery. Unsatisfactory eye alignment is more likely after surgery in patients with poor fusion potential or with more complicated types of strabismus. Patients with dense amblyopia or structural problems in one or both eyes have limited potential for binocular vision and do not employ fusional mechanisms to improve or maintain eye alignment.

Similarly, patients with neurodevelopmental anomalies have been shown to have higher rates of undercorrection and overcorrection after strabismus surgery. Also, patients with more unusual and severe forms of strabismus, such as cranial nerve III palsies, are more difficult to align satisfactorily with surgery.

Surgical Timing Without Amblyopia
The outcomes of infantile strabismus surgery are based on two main factors: ocular alignment and stereopsis. Stereopsis is measured several different ways. The optimal standard is full random dot stereopsis. However, there are lesser degrees of stereo that do qualify as successful.

We are often misled into believing that strabismus surgical outcomes are high. We should note that they are considerably lower than one would hope for. Pediatric strabismus surgery has low success rates and high reoperation rates because of difficult alignment measurements and the nature of different strabismus types.

This is true when you have an inexperienced surgeon such as a first-year resident. Multiple surgeries per case are also common. The more surgeries one has, the greater chance for exotropia, vertical deviations, no stereopsis, constant diplopia or constant uncorrectable strabismus.

A successful outcome can be based on a variety of measurements. If cosmetic alignment is the criteria, the success rate is higher. Full stereopsis criteria have a much lower success rate. Socio-psychological effects are negative for those with poor eye alignment, so all efforts should be made for the optometrist and ophthalmologist to collaborate in order to achieve the best outcome possible.

The definition of success in strabismus surgery varies among different studies. Most studies set their success criteria as 8pd to 10pd deviations from orthophoria at either the three-month, six-month or last follow-up.

Recently, an evidence-based study concluded that performing surgery later in life in patients with infantile esotropia increased the motor success rate of surgery. In addition, orthophoria is achieved with fewer surgical operations. Another study showed that more favorable long-term outcomes and less vertical deviations occurred when fewer surgeries were performed. The Cochrane study also concluded that there is no clear consensus on the optimal timing of surgery.

Clinically, a sandwich approach—VT before and after surgery—has demonstrated that the highest outcomes are possible for these patients. This might consist of vision therapy prior to strabismus surgery to optimize sensory readiness for motor fusion and/or post-surgical therapy to stabilize or safeguard binocular vision.

VT Skills
Oculomotor, localization and accommodative skills build a foundation prior to
sensory fusion. Management of these foundational skills prior to working on sensory fusion allows the optometrist providing pre- and post-VT to gain better outcomes. Clinically, my patients often gain cosmetic alignment simply by developing these skills, especially opposite to the direction of deviation. For example, with esotropia, working pursuits, saccades, localization and accommodative flexibility in temporal secondary gaze improves the ability to begin to appreciate fusion in primary gaze. For exotropia, these skills are emphasized in nasal secondary gaze. Poor eye movements reduce the success of gaining sensory fusion.

An overall limiting factor for patient outcomes is misaligned oculocentric localization. The patient’s awareness of where objects are judged to be located can be displaced in the direction of deviation. This needs to be addressed for both surgery and VT cases. Oculocentric localization should match the visual axis of each eye.

After the above skills are developed, in esotropia, depending on the type, VT is used to improve sensory fusion (the patient should have good second-degree fusion prior to surgery if possible) and relative motor fusion ranges if possible. In exotropia with some sensory fusion, VT is used to improve motor fusion recoveries and ranges as much as possible prior to surgery.

Much care is needed when working with anomalous correspondence. The safest approach is to overemphasize the skills of oculomotor, localization and accommodation. Use caution when attempting to gain normal correspondence with VT prior to surgery. Attempts to break suppression should only be done for short periods (five minutes or less).

VT post-op depends on the patient’s alignment and fusional status after surgery. Generally, VT focuses on improving all visual skills including eye movements, localization, accommodation, sensory and motor fusion, again around the new angle. The ability to recover fusion is more beneficial than simply building ranges.

One study on resolution without surgery showed that esotropia with onset in early infancy frequently resolves in patients first examined at less than 20 weeks of age when the deviation is less than 40pd and is intermittent or variable. Cases with a constant deviation of greater than or equal to 40pd presenting after 10 weeks of age have a low likelihood of spontaneous resolution. Child development of fusion occurs at about three months; thus, misalignment is normal prior to this age.

A large percentage of patients with either esotropia or exotropia gain sensory fusion with a visual motor-based VT program. The few that do not are then referred for surgical intervention. This gives the patient the best possible outcome.

**Surgical Timing with Amblyopia**

There is conflicting evidence whether or not to treat the strabismic amblyope before or after treating the eye deviation. One study suggested the patient obtain the best VA possible prior to surgery. Recent studies suggest that correcting eye alignment earlier will give the amblyopic eye the opportunity to develop better VA and binocular vision. Surgery is generally reserved for children with anisotropic amblyopia who do not respond to standard treatment, or children with serious vision impairment who are unable to wear glasses for developmental, sensory or other reasons. As a result, only a small percentage of children are suitable candidates for this surgery.

The Cochrane study stated that the optimum timing of when to perform strabismus surgery in children with amblyopia is unknown. It noted that treatment for amblyopia involves methods of exercising the amblyopic eye by encouraging its increased use. The researchers referenced several methods used to reduce the vision of the non-amblyopic eye temporarily, including penalization with cycloplegic or dilating eye drops, optical blurring of vision with high-plus lenses or physical occlusion of the normal eye. They acknowledged the newer binocular training approaches for amblyopia treatment available. These usually involve using a digital device that deprives the non-amblyopic eye in a biocular viewing task.
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Today, innovative technology, such as this virtual reality headset, allows us to treat amblyopia, strabismus, convergence disorders and other binocular vision problems.

To date, very few studies been conducted that represent VT. Treatment often involves a multitude of activities other than simply sitting and playing video games. These would include bi-ocular pursuits, saccades, accommodative tasks, visual spatial activities, anti-suppression decision-making procedures and visually guided movement activities. It is likely that when assessing all VT tools, successful outcomes would be much higher than those in controlled studies.

Binocular training should aim to treat amblyopia by restoring the underlying issue of reduced or absent binocularity.13

**New Approaches to Amblyopia Treatment**

Based on new research, the very best intervention for patients with amblyopia is to first develop a visual system in the amblyopic eye that can fixate, pursue and saccade with equal accuracy as compared with the nonamblyopic eye. The second step is to create binocularity for long term maintenance of acuity gains. Recent research has shown that by developing better systems, such as eye movements, accommodation and binocularity, the amblyopic eye improves without patching.13-15 This new concept has positive implications for patients, especially children who would prefer not to look different from their peers. This is a more positive psychological approach for these patients. Also, the lasting effect of VA gains is much greater using these methods than by full-time patching.

Classic teaching dictates that amblyopia must be corrected to the maximum extent possible before realignment surgery is undertaken.11 Based on my clinical experience, VT is extremely effective for amblyopia when it involves changing how the brain processes information for the amblyopic eye. This style of vision therapy uses a wide variety of eye movements and accommodative flexibility activities prior to attempting anti-suppression and fusion activities. The amblyopic eye also has opportunities to participate in a variety of perceptual activities. Typically, you will begin to see improved VA within about six to eight weeks.

After the eye movements and accommodative abilities are as equal as possible between the two eyes, incorporate the anti-suppression, fusion and convergence and divergence activities. A patient with amblyopia who develops good binocularity more readily maintains gains made in VA improvement. This is accomplished without having to continue with blur patch or deprivation activities.

Bottom line: there is no rush to jump into the deep end and recommend surgery as a first-line treatment for strabismus, and more specifically for exotropia. Wait until you have exhausted all the tools and techniques we have at our disposal.

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Out for Blood

This dry eye treatment may not be first on your list, but it’s worth taking into consideration.

Q List the blood product options for dry eye patients. What are their advantages and disadvantages? What contraindications and risks do they carry?

A “Dry eye disease is an increasingly common ophthalmic condition that negatively impacts the quality of life of millions worldwide,” says Andrew Fischer, OD, of Professional Eyecare Associates in Indiana. As our understanding of this disease continues to expand, so does the availability of our treatment options, he adds.

Blood-based dry eye treatments have increased in popularity and aided many dry eye patients for whom “traditional” treatment methods may have been unable to provide adequate relief. Blood-based products can be drawn from the patient themselves (autologous) or from a donor (allogeneic) and can be prepared in various treatment forms, including serum eye drops and platelet-derived eye drops. Blood-based drops are indicated for moderate to severe dry eye, persistent epithelial defects, graft-vs.-host disease, recurrent corneal erosions, neurotrophic keratitis and limbal stem cell deficiency, to name a few.

In short, blood-derived dry eye treatments help accelerate corneal epithelial cell healing by increasing the growth factor available to the ocular surface. While there is an abundance of clinical research supporting the use of blood-based eye drops for dry eye, this treatment is not FDA-approved; therefore, there is not a universally accepted standardized protocol preparation.

Serum Drops

This is perhaps the most common blood-based product used in dry eye therapy, says Dr. Fischer. To prepare serum eye drops, a sample of blood is obtained, allowed to clot at room temperature and centrifuged to separate the blood cell components from the serum. The serum is then collected and diluted with a balanced salt solution to prespecified concentrations, most commonly 20% and 50% serum by volume, but higher concentrations can be prepared.

These drops have potential limitations, as blood serum can contain pro-inflammatory cytokines, especially in patients with autoimmune diseases such as Sjögren’s syndrome or graft-vs.-host disease. For these patients, autologous serum may be counterproductive and lead to further degradation of the ocular surface.¹

Platelet-derived Drops

Options in this category include platelet-rich plasma (PRP), platelet-rich growth factor (PRGF) and platelet lysate (PL). Unlike serum eye drops, these preparations contain blood platelets, which inherently have high levels of growth hormones, promoting wound healing and corneal epithelial regeneration.²

To produce platelet-based drops, an anticoagulant is added to a blood sample, which is then centrifuged and separated into platelet-poor plasma, PRP and red and white blood cells. The PRP is extracted and transferred into dropper bottles for use. Unlike serum eye drops, PRP drops aren’t diluted. PRGF can be produced from PRP by adding calcium chloride, which significantly increases growth factor concentration. PL is derived from PRF, diluted to 30% by volume and frozen then thawed to increase growth factor concentration due to platelet thermolysis.

Takeaways

Contraindications to autologous blood products are largely relative, as the product is ultimately derived from the patient’s own body; however, blood infections, anemia and clotting disorders can be considered contraindications. In these cases, allogenic samples are an option.

Blood-derived dry eye treatments are especially useful in patients who are sensitive to preservatives or request a more holistic approach to dry eye treatment. While the TFOS DEWS II considers blood-based therapy a third-line treatment, it may be worth taking into consideration earlier in the process, especially in patients who could benefit from the robust healing properties of natural growth factors. ■


About Dr. Shovlin

Dr. Shovlin, a senior optometrist at Northeastern Eye Institute in Scranton, PA, is a fellow and past president of the American Academy of Optometry and a clinical editor of Review of Optometry and Review of Cornea & Contact Lenses. He consults for Kala, Aerie, AbbVie, Novartis, Hubble and Bausch + Lomb and is on the medical advisory panel for Lentechs.
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ABOUT RICK
Rick Bay served as the publisher of The Review Group for more than 20 years. To those who worked for him, he was a leader whose essence was based in a fierce and boundless loyalty. To those in the industry and the professions he served, he will be remembered for his unique array of skills and for his dedication to exceeding the expectations of his customers, making many of them fast friends.

(The Rick Bay Foundation for Excellence in Eyecare Education is a nonprofit, tax-exempt organization under section 501(c)(3) of the Internal Revenue Code.)
Herpes Hurts

Temporal pain often portends this condition, so learn how to best manage it.

A 79-year-old man presented with a three-day history of headache and pain and tenderness on the right side of his head. He also had mild ocular discomfort OD and related that it felt similar to when he develops styes; he used artificial tears to no improvement. He denied weight loss, appetite loss and jaw claudication. He was otherwise feeling well.

Beware a Vesicular Rash

His uncorrected visual acuity was 20/40 OD and 20/30 OS, commensurate with his cataracts. Intraocular pressure (IOP) was 13mm Hg OD and 10mm Hg OS. The remainder of his exam was normal. The only abnormality was a small, isolated vesicular lesion on his right lower eyelid. He did also have diffuse redness to his right forehead scalp, and he reported pain when his right temporal head was palpated.

Due to the head and scalp pain, two alternate diagnoses were considered; namely, herpes zoster dermatitis as the primary consideration and temporal arteritis as a secondary possibility. He was referred to obtain an erythrocyte sedimentation rate, C-reactive protein, platelet count and hemoglobin level, as well as prescribed valacyclovir 1,000mg TID PO to the regimen.

He returned emergently the next day with rapidly worsening vesicular eruptions across the right side of his face. There was increased ocular discomfort. His acuity and IOP were unchanged, but he now had a blistering rash on his forehead extending to his scalp, upper and lower eyelid edema, several subepithelial infiltrates and keratic precipitates on his right cornea and a mild degree of flare in the anterior chamber.

His tests for temporal arteritis were normal. His diagnosis now clearly was herpes zoster dermatitis and ophthalmicus. He was maintained on valacyclovir and prescribed prednisolone acetate 1% QID OS. He was immediately referred to his primary care physician who then added a Medrol Dosepak (oral methylprednisolone, Pfizer) and gabapentin 300mg TID PO to the regimen.

Burning Red

Herpes zoster ophthalmicus (HZO) may vary from dermatologic involvement alone to ocular manifestations such as lid retraction, keratitis, scleritis, uveitis, glaucoma, retinitis (acute retinal necrosis and progressive outer retinal necrosis), optic neuritis and panophthalmitis. When ophthalmic manifestations arise, the condition is termed HZO; it occurs in 7% of all zoster patients. Anecdotally, cases involving the eye have been observed to produce adnexal pain over months, without iritis, uveitis or vesicular breakout, making the chief complaint a mystery until the skin manifestations appear. The lesions of herpes zoster generally complete resolve within one to three weeks.
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REVIEW OF OPTOMETRY

The News Feed
Published March 7, 2022 • By Staff
DRY EYES, NEWS

Smoking, High BMI Associated with Disruption
Interferometry identified these populations at-risk

Researchers recently found that smokers and individuals with a high BMI showed significantly lower fluid layer grades and tear meniscus height scores compared with controls. They noted that their assessment of tear film parameters supports the findings of previous studies that implicate smoking and obesity as risk factors for dry eye.
Varicella zoster virus (VZV) causes chickenpox and herpes zoster represents a separate clinical condition caused by the same virus.1–12 The VZV is a member of the herpesvirus family. Varicella is spread by direct contact with active skin lesions or airborne via droplets and is highly contagious. The virus has an affinity for the upper respiratory tract and typically enters the human system through the conjunctiva and/or nasal or oral mucosa, producing the characteristic pox appearance.1–12 When the host’s immunity fails, the dormant virus leaves the confines of the dorsal root ganglion, where it lies dormant to produce chickenpox upon first infection or shingles (the zoster presentation) upon recurrence.1–12

Ninety percent of the population develops serological infection by adolescence, with nearly 100% of the population having some evidence of antibodies to the disease by age 60.1 Only 20% of patients suffer reactivation after initial infection.2 Any condition that decreases immune status, such as human immunodeficiency virus infection, chemotherapy, malignancy and long-term oral corticosteroid or other immunosuppressant use, increases the risk of herpes zoster activation. A second reactivation is even more rare and occurs mostly in immunosuppressed patients such as organ transplant recipients or those who suffer from AIDS or neoplasm.1

Treatment

Herpes zoster is managed using orally administered antiviral medications such as acyclovir (800mg PO five times a day), famciclovir (500mg PO TID) and valacyclovir (500mg to 1,000mg BID-TID for one week).1.10,12,13 The antiviral medications are most effective when initiated within 72 hours of the onset of the rash, especially for reducing the degree of post-herpetic neuralgia.12

Orally administered corticosteroids can reduce pain and potentially the onset of postherpetic neuralgia.10,12 Since the disease is self-limiting, most care is palliative, including the use of astringents, such as calamine lotion or aluminum acetate solution to minimize weeping and soothe the affected area, and topical anti-irritants and ointments to prevent secondary infection. Patients with postherpetic neuralgia may require narcotics for pain control.12 Tricyclic antidepressants and anticonvulsants in low dosages are potential options for unremitting pain.12 Capsaicin cream (based on the chemical in chili peppers that makes them hot), lidocaine patches and injectable nerve blockers can be used in the worst cases.12

Antiviral agents can play role in preventing varicella zoster disease in immunocompetent and immunosuppressed patients.18 As with herpes zoster, acyclovir has been documented as effective in preventing disease reactivation, but the proper dose, duration and circumstance of its uses are controversial.19

Since VZV-specific cell-mediated immunity and cell-mediated immunity in general declines with age, it should be expected that the incidence of herpes zoster virus (HZV) increases with age. This may explain the increased incidence of HZV in immunocompromised individuals or those who are undergoing immunosuppressive therapy.2,26 Therefore, any patient younger than 50 who presents with signs of an acute zoster infection must be referred to rule out an immunocompromised state.2

It is now well-accepted that herpes zoster should be combated prophylactically through vaccination. Shingrix (GlaxoSmithKline), a recombinant zoster vaccine, reduces the incidence of herpes zoster by more than 90% and is preferred to the live, attenuated herpes zoster (shingles) and postherpetic neuralgia. Am Fam Physician. 2018;98(9):437-48.15

For the patient reported here, upon his last follow-up eight weeks after his initial presentation, his ocular inflammation resolved on topical steroids and had been discontinued, as had the valacyclovir. He reported a great improvement in all findings, though he was still on gabapentin for postherpetic neuralgia.
EAST COAST OPTOMETRIC GLAUCOMA SYMPOSIUM

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OVERVIEW
The East Coast Optometric Glaucoma Symposium (ECOGS) meeting will be held as an in-person event. This two-day biannual symposium is designed to provide optometrists with exposure to current thinking on evolving standards of care, state-of-the-art technology and breaking research that will guide current and future glaucoma care in the optometric setting. Incorporating cases, clinical pearls, and discussion sessions, the program will maximize the opportunity for participant/faculty engagement.

The OGS conferences are long-running and trusted programs for optometrists managing patients with glaucoma. Each East Coast symposium focuses on glaucoma diagnosis and management, with the West Coast symposium focusing on therapies and innovations for comprehensive glaucoma coverage.

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A Tale of Two Patients
One had more structural damage, the other had more progression. Here’s how to proceed in both cases of glaucoma.

Two Caucasian females, both in their late 70s, presented as new patients in order to establish care. Both had similar histories insofar as having been diagnosed with glaucoma for several years and managed with medications and surgery. One patient had SLTs performed bilaterally, whereas the other had bilateral SLTs and a trabeculectomy in one eye. Upon presentation, both were clearly in the advanced stage of their glaucoma. And then their stories diverged somewhat.

They were initially seen by me approximately 12 months apart, with the latter having been a patient of mine for only nine months or so. Where I practice has a very large influx of patients who are moving to the area for retirement. Inheriting someone else’s patient is part and parcel of having a glaucoma practice, but at the end of the day, successfully managing these patients is a win-win for everyone: the patient, myself and the practice.

Management Challenges
While the majority (about 75%) of glaucoma patients fall into the mild to moderate category, some will certainly fall into the advanced category (about 15%), and the remaining will fall into the refractory category (about 10%), meaning that despite appropriate therapy, they continue to progress or have difficulty maintaining stability. Optometry is well positioned to manage the 75% of glaucoma cases that fall into the mild to moderate stage and often can do so without the assistance of our non-optometric ophthalmic colleagues. Even the 15% of patients who have advanced disease can be managed by optometry and surgical intervention. But oftentimes, in cases of advanced disease, there are some refractory patients who need further care by fellowship-trained glaucoma surgeons.

Our challenge in managing patients with advanced disease lies in making sure they remain stable. In essence, this is our goal for all glaucoma patients. Certainly, in mild to moderate disease states, there is a bit of room for some progression throughout the patient’s lifetime without significant effects on quality of life, but in patients with advanced disease, there is little room for progression. Once progression is even subtly determined, a change in therapy is warranted, often involving a surgical strategy.

Determining progression in glaucoma involves both structural and functional testing. Studies seem to indicate that patients with minimal to no visual field loss are best identified as having progressed by changes seen in their OCT findings, whereas individuals with more advanced disease seem to demonstrate progression more readily with visual field changes.1 This is certainly partly attributable to the floor effect seen when there is significant loss of retinal ganglion cells, but these rules are not steadfast in all cases. At the end of the day, both structural and functional testing is required for all glaucoma patients, independent of disease staging.

One of the inherent difficulties in visual field studies is the subjectivity of the testing, not to mention the simple truth that many patients do not enjoy sitting for a visual field test. Reliability indices can clearly give an indication as to the usefulness of the visual field in determining whether or not there is progression. Perhaps newer visual field platforms, such as virtual reality headset-based field testing, may make the testing experience more enjoyable for the patient and more reliable as well. We’ve introduced virtual reality visual field testing in my office, and to my surprise, the large majority of patients who have used the device have actually commented that they enjoy that modality.

Case Details
Both patients’ BMO-MRW printouts showed significant erosion of the neuroretinal rims in just about all sectors of the optic nerves, especially temporally. One of the places to look for structural progression of glaucoma (in addition to the circumpapillary...
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*Included in registration.
RNFL) is the neuroretinal rim. OCT of these patients should include scans of the RNFL, BMO and macular ganglion cell layer, all the while looking for subtle signs of progression.

Interestingly, the patient with slightly more advanced structural damage, the most recent of the two, has remained stable throughout the time I have seen her, but the other has progressed, which is especially evident in the ganglion cell layer analysis.

She has been a challenge from the first visit, presenting with an extensive history of red irritated eyes, sensitivity to several glaucoma medications, a failed trabeculectomy in one eye and IOPs in the mid-teens OU. She was on a steroid drop TID to control for inflammation and three separate glaucoma medications. Frankly, her eyes were unacceptably inflamed.

My initial goal of her treatment was threefold: (1) to decrease her medication burden, (2) to reduce her chronic inflammation and (3) to obtain better IOP control. After reviewing her previous records and adjusting a variety of her medications, slowly but surely, and with very close monitoring, we were able to achieve a reasonably good outcome of all three goals. This took, however, the better part of six months to accomplish. All the while, her fields and OCT findings remained stable.

Unfortunately, her OCTs began to show subtle change recently, and without any realistic medical options left, she was referred to the most appropriate glaucoma surgeon for further intervention. While her disease was not quite as advanced as the other patient, she is the one who has been experiencing consistent progression while in my care.

Will this patient be seeing the glaucoma surgeon exclusively from here on out? No, the glaucoma surgeon will render what surgical intervention they believe would best suit the patient, but she will ultimately return to my care for continued management. It’s important to find the right glaucoma surgeon, one who provides the best care for the patient and has mutual respect for all members of their care team. Great glaucoma surgeons do great things in the operating room and recognize that great optometrists do great things in the office. And sometimes, that great glaucoma surgeon is not in your home town.  

OVERVIEW
The Intrepid Eye Society, in tandem with Review Education Group and MedscapeLIVE!, look forward to bringing you "Intrepid Presents" in an in-person format in June 2022.

The Intrepid Eye Society is a group of emerging thought-leaders in optometry looking to promote excellence and growth in our field through innovation and implementation. We'll discuss future medical therapeutics, diagnostics, practice development, research and development, and collaborative care.

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Visually Handicapped

This patient’s medical history and genetics played a key role in finding the correct diagnosis.

A 35-year-old mentally handicapped Hispanic male presented with his mother for a second opinion on possible treatment options to help him see better. He reported a gradual painless, progressive loss of vision in both eyes that started at about nine years old. His mother reported that his vision was so poor by age 14 that he was only able to detect light.

His medical history is significant for mental delay. He also has hearing loss and polycystic kidney disease. As a child, he had strabismus surgery on the right eye to improve alignment and also had surgery to remove extra toes.

On examination, he had only light perception vision in each eye. Though his extraocular alignment appeared normal, he did have roving eye movements. His pupils were equal, round and reactive to light; there was no afferent pupillary defect. Tensions by applanation tonometry measured 12mm Hg in each eye. The anterior segment was unremarkable.

Dilated fundus exam was significant for vitreous syneresis and a posterior vitreous detachment in each eye. Other changes can be seen in Figures 1, 2, 3 and 4.

Take the Retina Quiz
1. What is the likely diagnosis for this patient?
   a. Fundus albipunctatus.
   b. Retinitis pigmentosa.
   c. Stargardt’s macular disease.
   d. Gyrate atrophy.

2. What is the likely etiology?
   a. Bardet-Biedl syndrome.
   b. Defect in ABC4 gene.
   c. Usher’s syndrome.
   d. Non-specific hereditary retinal degeneration.

3. What other systemic findings would you expect to see in this patient?
   a. Tall, thin stature.
   b. Cardiac abnormalities.
   c. Testicular hypogonadism.
   d. Normal development.

4. What treatment options should be considered for this patient?
   a. Stem cell implantation.
   b. Genetic therapy.
   c. Argus retinal implant.
   d. All of the above.

For answers, see page 98.

Diagnosis
Sadly, our patient has advanced retinitis pigmentosa (RP). There is extensive bone pigment spicules throughout the arcades and extending into the posterior pole. The vessels are severely attenuated and there is some pallor of the optic nerve, but not as much as you would expect considering how advanced his disease is.

Figs. 1 & 2. Here is a widefield view of the right and left eye of our patient. How do you explain the clinical findings?

About Dr. Dunbar
Dr. Dunbar is the director of optometric services and optometry residency supervisor at the Bascom Palmer Eye Institute at the University of Miami. He is a founding member of the Optometric Glaucoma Society and the Optometric Retina Society. Dr. Dunbar is a consultant for Carl Zeiss Meditec, Allergan, Regeneron and Genentech.
RP is a genetically heterogeneous group of disorders with varying modes of inheritance from autosomal dominant to X-linked recessive. Patients will develop bilateral, progressive vision loss, typically between the ages of nine and 19 with no sex or race predilection. Symptoms include night blindness (most common), loss of peripheral vision, reduced color vision and blurred vision. Signs include the triad of perivascular bone-spicule pigmentation in the mid-periphery, waxy pallor of the optic disc and vessel attenuation. Other findings commonly seen in RP include cystoid macular edema, diffuse RPE atrophy, optic disc drusen (10%), epiretinal membrane, vitreous condensation, keratoconus and posterior subcapsular cataracts (35% to 51% of cases).

Discussion
An estimated 100,000 people in the United States have RP and more than 80 mutated genes are known to cause this condition. In fact, RP may represent many different genetic disorders leading to a final common pathway that is clinically recognized as RP. The vast majority of cases affect only the eyes; however, there are several systemic conditions that are associated with RP, including Usher’s syndrome and Bardet-Biedl syndrome (BBS), among others. It is evident based on our patient’s medical history and clinical findings that he has BBS.

BBS is a genetic condition that impacts multiple body systems. It is classically defined by six features: obesity (specifically, fat deposition along the abdomen), intellectual impairments, kidney problems, genital and hormonal problems (hypogonadism), extra toes and digits (polydactylism) and visual impairment in the form of a rod-cone dystrophy, which occurs in 90% of patients with BBS. This condition develops as the result of mutations in more than 21 different genes and is usually autosomal recessive. Of the 21 genes, the most common are BBS1 (28% of cases), BBS10 (10% of cases), BBS2, BBS12 and ARL.

Interestingly, there is no clear link between the different mutations identified and disease severity, but some trends have emerged. Patients with mutations in the BBS1 gene seem to have milder ophthalmologic involvement, whereas patients with mutations in the BBS2, BBS3 and BBS4 genes experience classic deterioration in their vision. Our patient had all the classic features of BBS including, unfortunately, severe RP.

There is no specific cure for RP. For blind individuals with this condition, the Argus II Retinal Prosthesis System is considered with the hopes of providing useful vision to those that are severely impacted. The device combines a miniature eye implant with a patient-worn camera and a processor; it stimulates the visual cortex to sense shimmering, light or dark patterns or spots. The patient must learn how to interpret these signals as shapes and objects. The vision generated by Argus II is very different from traditional sight and the patient learns to interpret the new “language” of sight.

Unfortunately, our patient was not a candidate for the Argus II because of his roving eye movements and mental delay. In these types of situations, we make sure visually handicapped patients are referred to the Miami Lighthouse for the Blind, but our patient was already well-connected. In fact, he had been using a mobile cane for close to 20 years. He travels by himself to places such as the Lighthouse, the gym and the mall while using public transportation, and he is fluent in Braille.

Ring Me Up

The IC-8 IOL creates a permanent pinhole effect, improving the prospects for a monovision approach.

BY JAKE WYSIADLOWSKI, OD
AUSTIN, TX

As surgical technology evolves, the number of options we have to improve post-cataract vision continues to grow. FDA approval seems imminent for the IC-8 intraocular lens (IOL) from AcuFocus, which stands out among the rest by combining small-aperture optics with a monofocal lens. This lens adds a small mini-ring in its center that extends depth of focus. The lens is designed to mitigate the visual effects of unfocused peripheral light.

Range of Benefits
The 1.36mm aperture of the ring only allows focused rays of light onto the retina, while the peripheral, defocused rays are blocked by the 3.23mm mask, a descendant of the now-discontinued Kamra corneal inlay. It is a one-piece hydrophobic, acrylic IOL with a 6mm optic zone, 12.5mm length and powers that range from +10D to +30D in 0.50D steps.

A toric design has not been released; however, the IOL can work for up to 1.50D of astigmatism due to the pinhole effect. The lens has been reported to provide up to 3D of extended depth of focus with a monovision offset, reduce effects of lower and higher-order aberrations and allows plus or minus 1D of deviation from target refraction.

Implanting the Lens
In most cases, the IC-8 lens is used in combination with a low amount of monovision, typically -0.75D to -1D. The lens is implanted in the non-dominant eye and a monofocal IOL with an aim of plano is placed in the dominant eye. Assuring the lens is well-centered with the visual axis to allow the optics to work properly is important.

Also, refracting these patients can yield interesting results due to the extended depth of focus. For this reason, a midpoint refraction or a red/green balance refraction is recommended. For the midpoint refraction, begin with a manifest refraction, then determine the maximum plus and maximum minus lenses to blur, and then determine the midpoint. For the red/green balance refraction, you want the patient to report the letters on the green and red side being equal in clarity.

Surgeons have reported good near and intermediate vision with better distance vision compared to traditional monovision. There is also less ametropia due a lower amount of offset for the IC-8 IOL eye.

Who Can Benefit from IC-8?
Patients with irregular corneas from keratoconus or previous refractive surgeries such as radial keratotomy could benefit from this lens because only central light rays reach the retina. However, due to the reduced amount of light entering the light with this modality, it is important to be cautious if considering implanting bilaterally. The lens may also be beneficial in those reporting light sensitivity or problems with glare.

Patients with central corneal scarring, large pupils, macular pathology, severe glaucoma, AMD, vitreous opacities or even those with high need for near vision should not be considered candidates for this IOL. Patients with pupils larger than the mask can notice dysphotopsias. Reduced contrast sensitivity can occur, as well as a minor reduction in visual field. Patients may report difficulties in low-light conditions due to the IOL design.

Postoperative management is fairly similar to other IOLs with a few extra considerations, including thorough patient education on this type of lens modality, which will provide a smoother postoperative period.

From patients desiring presbyopia correction to those looking for greater functional range of vision, this unique IOL should provide solutions for appropriately selected and properly educated patients.

Surgical Minute

For a video of the procedure, read this article online at www.reviewofoptometry.com.

About the Author

Dr. Wysiadlowski is an ocular disease resident at Dell Laser Consultants in Austin, TX.

About Drs. Cunningham and Whitley

Dr. Cunningham is the director of optometry at Dell Laser Consultants in Austin, TX. He has no financial interests to disclose. Dr. Whitley is the director of professional relations and residency program supervisor at Virginia Eye Consultants in Norfolk, VA. He is a consultant for Alcon.
Dear Colleagues,

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Three Sides to Every Survey

This patient history tool is simple to implement yet significant for outcomes.

(1) Dr. Taub) was interviewing for a job early in my career, and I was asked a question that I occasionally now use in my resident and faculty interviews. The answer I gave was instantaneous and, perhaps in hindsight, a little smug, but ultimately, I got the job. The question from the interviewer was, “If you were stranded on a deserted island and could have one piece of equipment to help you give an eye exam, what would it be?” Of course, you might be thinking of things like a slit lamp, phoropter or retinoscope. Those are all answers I have heard as the questioner. My answer: “Me!” Keep in mind that this was 20 years ago when I was a snot-nosed punk, but even today, I would still give the same answer.

My reasoning was, and remains, that through a good history with proper questioning and sleuthing, almost every condition can be diagnosed without picking up a single piece of equipment. This is not just hubris; according to Nobel Peace Prize laureate Bernard Lown, medical history provides enough information in about 75% of patient encounters to make the diagnosis even before the physical examination and additional testing is completed.1 One study revealed that the initial (upon reading the referral letter and taking the history) and final diagnoses matched in 83% of new patients.2 Another found a success rate of 76%.3 Confidence in the correct diagnosis in this investigation increased from 7.1 on a scale of one to 10 after the patient history to 8.2 after the physical examination and 9.3 after the laboratory testing.3

The Trio

History-taking, for most optometrists, starts when the examination begins and hopefully lasts throughout the entire process. However, it can also start before the patient even steps into the room through symptom surveys. These assist by guiding the exam and helping patients make connections.

This checklist focuses on quality-of-life for those suffering from vision impairment. To download a full-sized copy, please visit www.reviewofoptometry.com.
between their symptoms and their vision. There are quality-of-life surveys for dry eye, computer vision syndrome and almost every other visual condition in the book. Since this column concentrates on prescribing and binocular vision topics, we will narrow our focus a bit and introduce you to several surveys that you might consider adding to your paperwork when you see a pediatric patient on your schedule.

When the landmark Convergence Insufficiency Treatment Trial found that in-office vision therapy is the superior treatment for convergence insufficiency compared with home therapy and pencil push-ups, the primary measure used to show improvement was the Convergence Insufficiency Symptom Survey (CISS).4,5 A 15-question Likert-scale survey, the CISS was designed to quantify the severity of symptoms associated with convergence insufficiency, showing good validity and reliability.6,7

Questions include, “Do you have double vision when reading or doing close work?” and, “Do you have headaches when reading or doing close work?” Each question is rated on a scale of zero to four based on severity, with zero indicating “never” and four indicating “always.” A score of 16 or higher was found to differentiate children with convergence insufficiency from those with normal binocular vision. Even though one study found that children with oculomotor dys-function, but not those with accommodative insufficiency, had higher CISS scores than those with normal binocular function, the survey remains invaluable in identifying children who suffer from all types of binocular vision issues.8

The Quality-of-Life checklist developed by the College of Optometrists in Vision Development (COVD-QOL) was originally a 30-question Likert scale questionnaire but was later shortened to 19 questions. It documents improvement when treating patients with vision therapy, correlates with academic performance and has good test-retest reliability.9-11 We use this checklist as part of all exams that we perform for children older than six. A parent fills it out if the child is younger, but older children can complete it independently. Like the CISS, it uses a zero to four scale, with a four indicating that the symptom “always” occurs. The COVD-QOL overlaps many questions on the CISS but also includes questions relating to forgetfulness, poor memory and saying “I can’t” before even trying a task. A score of 20 or higher indicates the need for a complete binocular vision evaluation.

The largest area of growth that we have seen in our practice involves patients suffering from a brain injury. These injuries can take place shortly before the exam, or they may have happened years to decades earlier. The patient may not realize that their symptoms are linked to the injury, so the Brain Injury Vision Symptom Survey (BIVSS) aims to help make that connection and allow the doctor to grade the severity of the visual symptoms. The 28-question, four-point Likert scale survey assesses eight areas of vision, including visual clarity, double vision and photosensitivity. It has been shown to have good validity and test-retest reliability.12,13 A score of 31 is a red flag and an indication that further evaluation must be considered.

Even though the BIVSS was designed for traumatic brain injuries, we use it for all acquired injuries, as well

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**CONVERGENCE INSUFFICIENCY SYMPTOM SURVEY (CISS)**

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<thead>
<tr>
<th>Question</th>
<th>Never</th>
<th>Infrequently</th>
<th>Sometimes</th>
<th>Fairly Often</th>
<th>Always</th>
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<tbody>
<tr>
<td>1. Do your eyes feel tired when reading or doing close work?</td>
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<td>2. Do your eyes feel uncomfortable when reading or doing close work?</td>
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<td>3. Do you have headaches when reading or doing close work?</td>
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<td>4. Do you feel sleepy when reading or doing close work?</td>
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<td>5. Do you lose concentration when reading or doing close work?</td>
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<td>6. Do you have trouble remembering what you have read?</td>
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<td>7. Do you have double vision when reading or doing close work?</td>
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<td>8. Do you see the words move, jump, swim or appear to float on the page when reading or doing close work?</td>
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<td>9. Do you feel like you read slowly?</td>
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<tr>
<td>10. Do your eyes ever hurt when reading or doing close work?</td>
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<td>11. Do your eyes ever feel sore when reading or doing close work?</td>
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<td>12. Do you feel a “pulling” feeling around your eyes when reading or doing close work?</td>
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<td>13. Do you notice the words blurring or coming in and out of focus when reading or doing close work?</td>
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<td>14. Do you lose your place while reading or doing close work?</td>
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<td>15. Do you have to re-read the same line of words when reading?</td>
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To obtain score: Total the number of “X”s in each column. Multiply by the column value

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<thead>
<tr>
<th>Column</th>
<th>Value</th>
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Sum

**SCORE:**

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This form helps pinpoint convergency insufficiency impact. To download, please visit [www.reviewofoptometry.com](http://www.reviewofoptometry.com) and look for this article’s page.
as for patients suffering from neurological conditions like MS and ALS. We also use it to determine potential treatment benefits from vision therapy, tints and occlusion.

**Takeaways**

There is no one data point on which an entire diagnosis can be based. The same can be said for symptom surveys. They are yet another tool at your disposal to enhance and create a complete patient history that can, in turn, be used to guide your exam and the testing choices that you make, allowing for more effective and efficient diagnosis and treatment.

Since these surveys represent only one data point, there are times when the results match the objective data and times when they do not. If the survey score is high but the objective data is normal or relatively normal, look closely at who filled out the survey. We have seen circumstances in which the parent fills out the survey quite differently than their child does, and vice-versa. As there are three sides to every story, perhaps there are also three sides to every survey. Take the time to consider each to help paint a fuller picture.

If the survey score is low but the objective data indicates a problem, that doesn’t necessarily mean that the survey is wrong; keep in mind that the patient may not be symptomatic yet or ever. The absence of symptoms may in fact be a symptom. The entirety of the examination must be considered altogether in the decision-making process.

The amount of output needed to start using surveys is minimal, but the impact on your clinical exams and patient outcomes can be enormous. Incorporating these tools into your care process in the clinic should be a no-brainer.

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A 75-year-old African-American female presented to the office acutely with a chief complaint of ocular pain and redness OS of four days’ duration. Her ocular history was remarkable for uneventful phaco-emulsification surgery in that eye, completed two weeks prior. Systemic history was positive for hypertension, for which she was properly medicated and compliant. The patient denied allergies of any kind.

Clinical Findings
Her best-corrected entering visual acuities were 20/20 OD and 20/400 OS, with no improvement using pinhole at distance or near. Her external examination was normal. There was excessive lacrimation with conjunctival injection OS. There was no evidence of afferent pupillary defect. The pertinent biomicroscopic examination of the anterior segment OS is demonstrated in the photographs. Goldmann applanation tonometry measured 15mm Hg OD and 65mm Hg OS.

For More Information
Additional studies might have included gonioscopy, B-scan ultrasonography, anterior segment and angle optical coherence tomography and ultrasound biomicroscopy (obtained in this case and shown below).

Additionally, history questions should be asked regarding inadvertent trauma, which may have produced accidental hypotony with or without a cyclodialysis cleft. More questions might include use of any medication from the sulfonamide group (e.g., oral acetazolamide, topiramate), which may cause an idiiosyncratic uveal effusion with forward movement of both the choroid and ciliary body.

Your Diagnosis
What would be your diagnosis in this case? What is the patient’s likely prognosis? To find out, please read the online version of this article at www.reviewofoptometry.com.

Biomicroscopy and ultrasound biomicroscopy yielded the above findings. What do they—and the case history—point toward?

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Retina Quiz Answers (from page 90)—Q1: b, Q2: a, Q3: c, Q4: c

NEXT MONTH IN THE MAG
In May, we present our annual dry eye report. Articles will include:
- How to Ease into Next-Level Dry Eye Care
- Understanding Patient Lifestyle Influences on Dry Eye
- How to Get Patients Pumped Up for Dry Eye Therapy

Also in this issue:
- How Dry Eye Screening Questionnaires Can Build Your Practice
- How to Work up Charles Bonnet Syndrome and Similar Conditions
- Scope Expansion: How to Add Incisions and Injections to Your Practice
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