



NOW FROM GLAUKOS

Using Photrex[®] Viscous (riboflavin 5'-phosphate in 20% dextran ophthalmic solution), Photrex[®] (riboflavin 5'-phosphate ophthalmic solution), and the KXL[®] system, the iLink[™] corneal cross-linking procedure from Glaukos is the only FDA-approved therapeutic treatment for patients with progressive keratoconus and corneal ectasia following refractive surgery.*¹



GET THERE IN TIME

iLink[™] is the only FDA-approved cross-linking procedure that slows or halts progressive keratoconus to help you preserve your patients' vision.

See back cover for Important Safety Information and visit livingwithkeratoconus.com for full prescribing information.

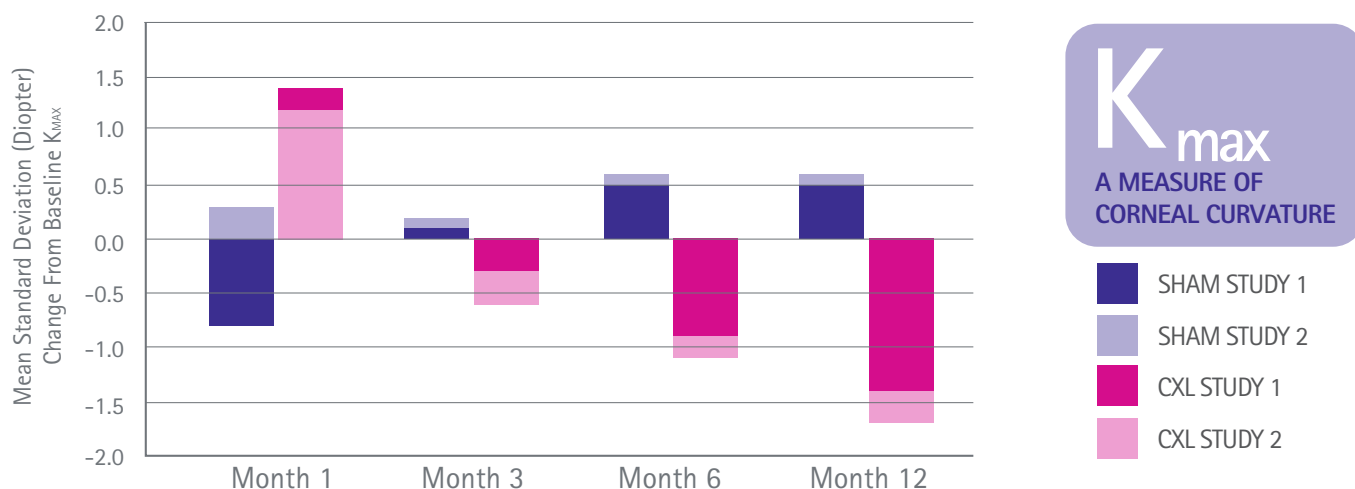
*Photrex[®] Viscous and Photrex[®] are manufactured for Avedro. The KXL System is manufactured by Avedro. Avedro is a wholly owned subsidiary of Glaukos Corporation.

GLAUKOS[®]

Transforming the standard of care for progressive keratoconus

In clinical studies, cross-linking-treated eyes showed increasing improvement in K_{\max} from month 3 through month 12, while in untreated, sham eyes, K_{\max} demonstrated steepening.¹

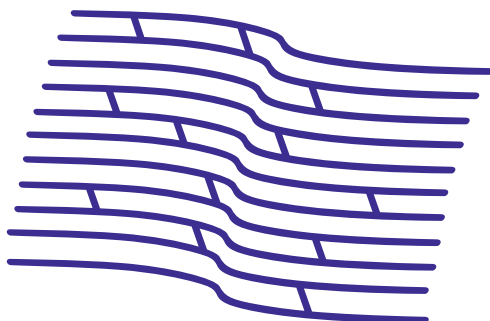
Change From Baseline (K_{\max})—Progressive Keratoconus Patients*¹



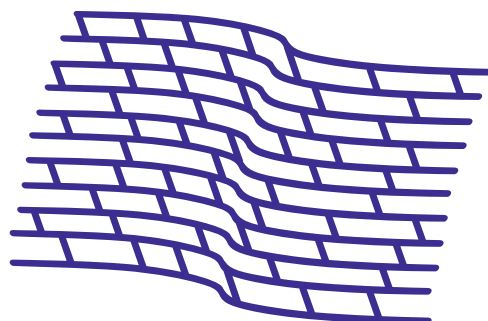
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CXL=corneal cross-linking.

Cross-linking using UVA and riboflavin²

- Creates new corneal collagen cross-links
- Results in a shortening and thickening of collagen fibrils
- Leads to stiffening of the cornea



LESS CROSS-LINKING (WEAKER)



MORE CROSS-LINKING (STRONGER)

Offers a proven safety profile¹

iLink™ keeps keratoconus from calling the shots

Left untreated, 1 in 5 patients with progressive keratoconus may require a corneal transplant. More than half of these patients could need multiple transplants within 20 years.^{3,4}

OVER 400,000

CROSS-LINKING PROCEDURES HAVE BEEN PERFORMED WORLDWIDE WITH OUR EQUIPMENT⁵

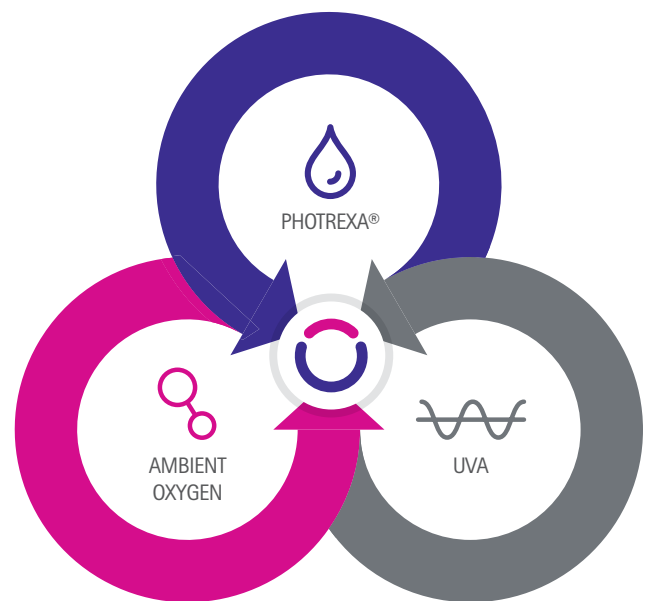
APPROVED IN 2016

IN THE UNITED STATES FOLLOWING 2 SUCCESSFUL PHASE III TRIALS¹

Understanding the science

iLink™ proprietary procedure⁶

- iLink™ combines ultraviolet light with ambient oxygen and Photrexa® bioactivated ophthalmic solution
- Absorption of UVA by Photrexa® generates radical riboflavin to form cross-links
- Precise, metered UV light delivery with the KXL system
- Backed by 20 years of science to achieve the right formulation



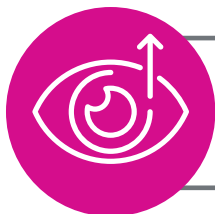
Both the American Academy of Ophthalmology and the Cornea Society recommend cross-linking for the treatment of progressive keratoconus.^{7,8}

Diagnose and treat progressive keratoconus

Keratoconus may be vastly underdiagnosed, underestimated, and underserved⁹



Left unchecked, progression can lead to a higher risk of vision loss, significantly reduced vision quality of life, and the potential need for a corneal transplant.^{10,11}



Prevalence was as high as **1 in 375** in a recent Netherlands study, which is **5 to 10** times higher than previously reported.¹²



Onset occurs from 12 years of age through 20 years of age for a vast majority of patients.¹³



Vision loss is associated with anxiety, depression, potential loss of independence, and a heavier burden on family members and/or social services.¹⁰

Offer your patients the benefits of iLink™

- Peace of mind
- The only therapeutic treatment option for progressive keratoconus
- May reduce or delay the need for more invasive procedures¹⁴
- One-time procedure widely covered by commercial medical insurance¹⁵

Take control with the iLink™ procedure

As a minimally invasive procedure, the iLink™ cross-linking procedure is the new standard of care for progressive keratoconus. With extensive clinical validations, iLink™ is proven safe and efficacious in numerous trials and peer-reviewed publications.^{1,7}



The proprietary iLink™ procedure incorporates:

The KXL System

- UVA irradiation: 30 minutes at 3 mW/cm²
- Laser alignment for patient positioning
- Wireless control for beam alignment in the X, Y, and Z axes
- Fully integrated stable delivery platform
- Touchscreen operation
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Photrexa® and Photrexa® Viscous

- Only FDA-approved bioactivated riboflavin ophthalmic solutions
- Manufactured using validated processes in a state-of-the-art drug manufacturing facility
- GMP-certified material
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Reimbursement support and tools give patients access to iLink™

OVER 95% COMMERCIAL INSURANCE COVERAGE



Financial feasibility tools available



National team of dedicated reimbursement experts



Billing, coding, preauthorization, claims, and appeals resources available



Flexible options for equipment acquisition



Uninsured and Medicaid available for qualified patients



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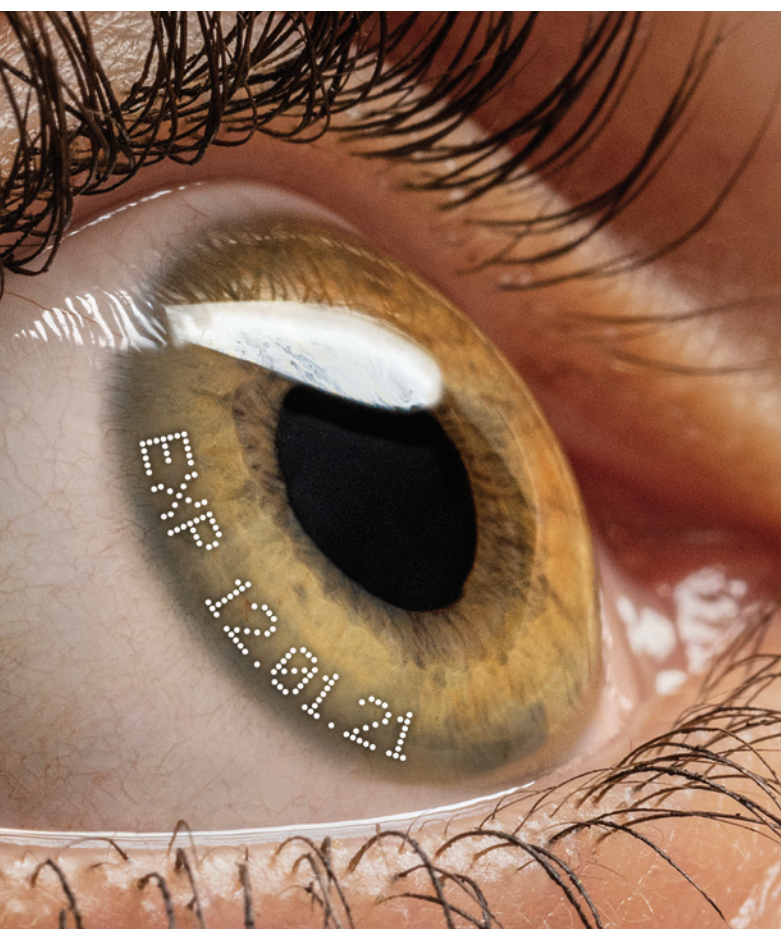
Slowing or halting disease progression with a single outpatient procedure may be a more cost-effective option for patients than ongoing management approaches.¹⁵



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Our focus is to develop and lead the global ophthalmic market with therapies for glaucoma and corneal and retinal diseases through continual innovation and investment.

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- Smallest medical device ever implanted in humans
- First and only FDA-approved procedure proven to slow or halt progression of keratoconus

iLink™ slows or halts progression to help you preserve vision

- Only FDA-approved cross-linking procedure
- Wide reimbursement coverage
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REFERENCES:

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IMPORTANT SAFETY INFORMATION

Ulcerative keratitis can occur. Patients should be monitored for resolution of epithelial defects.

The most common ocular adverse reaction was corneal opacity (haze). Other ocular side effects include punctate keratitis, corneal striae, dry eye, corneal epithelium defect, eye pain, light sensitivity, reduced visual acuity, and blurred vision.

These are not all of the side effects of the corneal collagen cross-linking treatment. For more information, go to www.livingwithkeratoconus.com to obtain the FDA-approved product labeling. You are encouraged to report all side effects to the FDA. Visit www.fda.gov/medwatch, or call 1-800-FDA-1088.

iLink™
CROSS-LINKING PROCEDURE

MA-01973A

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GLAUKOS®
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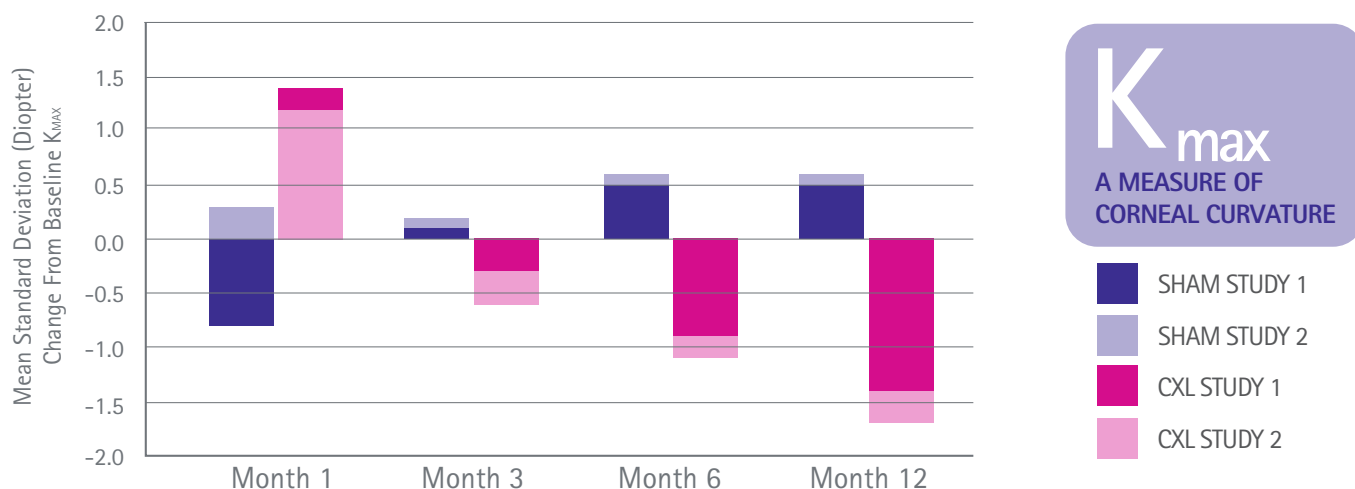
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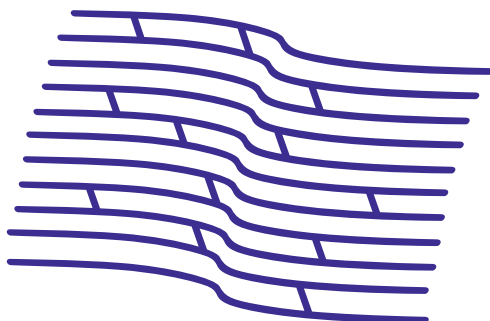
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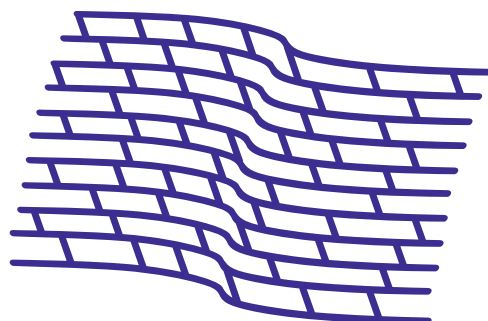
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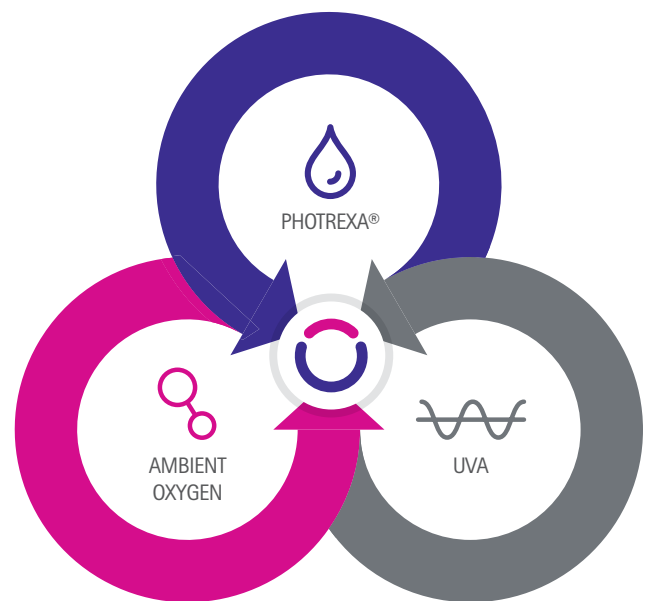
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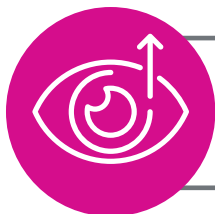
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