REVIEW

Corneal Collagen Crosslinking for Ectasia After Refractive Surgery: A Systematic Review and Meta-Analysis

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Correspondence: Dillan Cunha Amaral, Centro de Ciências da Saúde, Bloco K - Av. Carlos Chagas Filho, 373 - 2° andar, Sala 49 - Cidade Universitária, Rio de Janeiro, RJ, 21044-020, Brazil, Email dillanc.amaral@gmail.com

Introduction: Corneal ectasia leads to progressive irregular corneal curvature and reduced visual acuity.

Objective: To assess the safety and effectiveness of corneal collagen cross-linking (CXL) for managing corneal ectasia resulting from refractive laser surgery (RSL).

Methods: A systematic review and meta-analysis were realized according to PRISMA guidelines. We searched PubMed, EMBASE, Cochrane, and Web of Science databases for studies on CXL in patients with ectasia after RLS. The outcomes of interest included visual acuity, refractive outcomes, topographic parameters (Kmax, index surface variance (ISV), index of Vertical Asymmetry (IVA), keratoconus index (KI), central keratoconus index (CKI), index of height asymmetry (IHA), index of height decentration (IHD) and Rmin (minimum sagittal curvature)), central corneal thickness, endothelial cell count, and possible adverse events. Statistical analysis was performed using the R software (version 4.2.3, R Foundation for Statistical Computing, Vienna, Austria).

Results: 15 studies encompassing 421 patients (512 eyes) were included. The mean age was 32.03 ± 4.4 years. The pooled results showed a stable uncorrected visual acuity post-CXL, with a significant improvement in corrected distance visual acuity (SMD = 0.09; 95% CI: -0.07 to 0.26). The spherical equivalent decreased significantly (SMD = -0.09; 95% CI: -0.35, -0.02). The topographic parameter Kmax decreased significantly (SMD = 0.15; 95% CI:0.01 to 0.28); however, the other parameters, ISV, IVA, KI, CKI, IHA, IHD, and Rmin, did not change significantly. Central corneal thickness decreased significantly (SMD = 0.24; 95% CI:0.07 to 0.41), and the endothelial cell count remained stable The complications were rare.

Conclusion: CXL is a safe and effective technique for managing corneal ectasia after RLS.

Keywords: cross-linking, corneal ectasia, laser refractive surgery, meta-analysis, systematic review

Introduction

Corneal ectasia is a rare but potentially vision-threatening complication that may occur after refractive laser surgery (RSL), leading to deterioration of visual acuity and the need for additional treatment.^{1–3} Ectatic corneal diseases encompass conditions marked by gradual thinning and bulging of the corneal structure.⁴ This condition leads to irregular astigmatism (corneal distortion) and subsequent decrease in visual acuity.⁵ This may occur within weeks or even 45 months of the surgical procedure.⁶ Post-refractive ectasia was extremely rare in eyes without risk factors for PRK (0.020%), LASIK (0.090%), or SMILE (0.011%) per 100,000 eyes. The overall incidence was low even when including eyes with preoperative risk factors: 0.025% for PRK, 0.100% for LASIK, and 0.020% for SMILE per 100,000 eyes.⁷ A retrospective study of 30,000 LASIK cases revealed that the risk of postoperative corneal ectasia after LASIK is equal

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© 2024 Amaral et al. This work is published and licensed by Dove Medical Press Limited. The full terms of this license are available at https://www.dovepress.com/rems. work you hereby accept the Terms. Non-commercial uses of the work are permitted without any further permission from Dove Medical Press Limited, provided the work is properly attributed. For permission for commercial use of this work, is ese aparagraphs 4.2 and 5 of our Terms (http://www.dovepress.com/rems.ph). to or lower than the risk of developing keratoconus without LASIK.^{8,9} This suggests that corneal ectasia following RLS may occur by chance, even without such surgery. Furthermore, post-RLS ectasia causes have not been completely elucidated, but many risk factors have been described, such as personal/family history of keratoconus, younger age, abnormal corneal topography and tomography, high myopia, low residual stromal bed, excessive stromal ablation, high percentage of tissue altered, deep primary keratotomy leading to a thick corneal flap, forme fruste keratoconus, pellucid marginal degeneration, lamellar procedures, deeper ablations and low preoperative corneal thickness.^{10–18}

Current treatment options for visual rehabilitation include rigid contact lenses, penetrating or lamellar keratoplasty, intrastromal corneal ring segments, and corneal collagen cross-linking (CXL).^{19–22} This method increases the biomechanical stability and stiffness of the cornea in a photochemical process by adding additional polymer bands between the collagen fibers using ultraviolet (UV) light and riboflavin.^{23–25} As a consequence, additional covalent bonds are formed between collagen molecules, stabilizing the collagen structure.²⁵ To date, few studies have evaluated the use of CXL for treating corneal ectasia after refractive surgery, and the available results do not show good congruence. As it is a rare complication, the number of patients included in these studies was low. The last meta-analysis produced on the subject was in 2017,²⁶ and since then, there have been results from new high-impact clinical trials with more significant sample numbers. Hence, a new meta-analysis is essential to consolidate the findings from past and recent studies.

This systematic review and meta-analysis aimed to assess CXL's safety and efficacy in addressing corneal ectasia after RLS.

Methods

Search Strategy and Data Extraction

The protocol for this systematic review of the literature on CXL therapy in cases of corneal ectasia after RLS was registered in the PROSPERO International Prospective Register of Systematic Reviews (CRD42023452904). We followed the Preferred Reporting Items for Systematic Reviews and Meta-Analyses (PRISMA) guidelines for data extraction.²⁷ The terms ("Corneal Collagen Cross Linking" OR "Corneal Cross Linking" OR "Corneal" OR "Epi off Corneal Cross Linking" OR "Epi off CXL" OR "Epi on CXL" OR "Epi-off Corneal" OR "Cross-Linking" OR "Epi-off CXL" OR "Epi-on CXL" OR "Epithelium off Corneal Cross Linking" OR "Epithelium off CXL" OR "Epithelium on Corneal Cross Linking" OR "Epithelium on CXL" OR "Epithelium-off Corneal Cross-Linking" OR "Epithelium-off Corneal" OR "Cross-Linkings" OR "Epithelium-off CXL" OR "Epithelium-on Corneal Cross-Linking" OR "Epithelium-on CXL" OR "Transepithelial Corneal Cross Linking" OR "Transepithelial Corneal Cross-Linking" OR "Transepithelial CXL" OR "crosslinking" OR "cross-linking") AND ("post-LASIK keratectasia" OR "Postoperative Ectasia" OR "keratectasia" OR "corneal ectasia" OR "Post-LASIK ectasia") AND ("Keratorefractive Surgical Procedure" OR "Keratorefractive Surgical Procedures" OR "Refractive Surgeries" OR "Refractive Surgery" OR "Refractive Surgical Procedure") were used for the search. Search terms were used to query the PubMed, Embase, Cochrane, and Web of Science databases. The searches started on July 20, 2023, and ended on August 20, 2023. References from all the included studies, previous systematic reviews, and meta-analyses were manually searched for additional studies. Two authors (A.H. and L.C.) independently extracted data using predefined search criteria and quality assessment. The full articles of eligible publications were then scrutinized. Only trials that met the following criteria were included in the meta-analysis.

Eligibility Criteria

Inclusion in this meta-analysis was restricted to studies that met all the following eligibility criteria: (1) human study; (2) participants: patients with corneal ectasia after RLS, defined as progressive corneal curvature steepening, increasing myopia and astigmatism, and loss of visual acuity postoperative;^{18,28} (3) intervention: CXL; (4) at least one or more clinical outcomes representing intraoperative and/or postoperative outcome parameters must be assessed and published; and (5) design: randomized clinical trials (RCT) or non-randomized cohorts, and studies were included only if they reported any of the clinical outcomes of interest. We excluded studies with (1) animal studies or cadaver subjects and (2) studies that were not published in English.

Outcomes

Our study aimed to evaluate multiple outcomes, encompassing the endpoints of (1) visual acuity outcomes with corrected distance visual acuity (CDVA) and uncorrected distance visual acuity (UCVA); (2) refractive outcome with Spherical Equivalent (SE); (3) topographic and tomographic parameter outcomes with Kmax, Index surface variance (ISV), Index of Vertical Asymmetry (IVA), Keratoconus Index (KI), Center Keratoconus Index (CKI), Index of Height Asymmetry (IHA), Index of Height Decentration (IHD), and minimum sagittal curvature (Rmin); (4) endothelial cell count (EEC), central corneal thickness (CCT); and (5) serious complications.

Data Extraction

Two reviewers (L.C. and A.M.) independently evaluated the quality of the citations and extracted data. Any disagreements were resolved by another reviewer (D.A). The following information was extracted: name of first author, year of publication, trial location, research design, number of eyes, mean patient age, sex proportion, follow-up duration, and type of refractive surgery.

Adverse events and complications encompassed those possibly related to the procedure, such as epithelial ingrowth, infectious keratitis, flap margin elevation, retinal damage, persistent corneal edema, corneal scarring, and cataract formation in the patient population.

Statistical Analysis

This systematic review and meta-analysis were performed using the Cochrane Collaboration and Preferred Reporting Items for Systematic Reviews and Meta-Analysis (PRISMA) statement guidelines.²⁷ Relative risk (RR) with 95% confidence intervals was used to compare treatment effects for categorical outcomes. Continuous outcomes were compared using standardized mean differences (SMD). Outcomes were measured as the mean \pm standard deviation (SD). Statistical significance was set than 0.05. Cochran's Q-test and I² statistics were used to assess heterogeneity, and I² > 50% indicated substantial heterogeneity. We used a random-effects model for outcomes with significant heterogeneity. Statistical analysis was performed using the R software (version 4.2.3, R Foundation for Statistical Computing, Vienna, Austria).

Results

Study Selection and Baseline Characteristics

As detailed in Figure 1, we found 1324 articles, with 225 articles in PubMed, 493 in Embase, 606 in Web of Science, and 23 in Cochrane databases. Of these, 785 duplicates were excluded from the analysis. After removing duplicate records and ineligible studies, 31 studies remained and were thoroughly reviewed based on inclusion criteria. Eighteen articles were excluded based on these criteria. Finally, 15 studies were included in this review, including four RCTs^{29–32} and eleven non-randomized cohorts.^{33–43}

We analyzed 421 patients, including 512 eyes diagnosed with post-laser corneal ectasia. The sample sizes of these trials ranged from 11 to 91. These trials were performed in eight countries (three in the United States; two in China, Egypt, and Switzerland; and one in Australia, Germany, Lebanon, and Turkey). Eleven trials reported that their patients were followed-up for 12 months or more post-CXL. The mean age of the studies that provided this information was 43.05 ± 11.6 years old with range (20–67). The characteristics of other studies are listed in Table 1.

Visual Acuity and Refractive Outcomes

Visual acuity was recorded and analyzed as the logarithm of the minimum angle of resolution (logMAR) value. Compared to the preoperative baseline, there was no significant difference in UCVA after the CXL procedure (SMD = 0.09; 95% CI: -0.07, 0.26; Figure 2), and no statistically significant heterogeneity between studies was identified (P = 0.64, I² = 0%). However, compared with the preoperative values, the CDVA significantly improved post-CXL (SMD = 0.33; 95% CI:0.12 to 0.54; Figure 3), and heterogeneity was observed (P < 0.01, I² = 54%).

Compared to the preoperative baseline, SE significantly decreased after the CXL procedure (SMD = -0.09; 95% CI: -0.35 - -0.02; Figure 4), and no statistically significant heterogeneity between studies was identified (P = 0.82, I² = 0%).

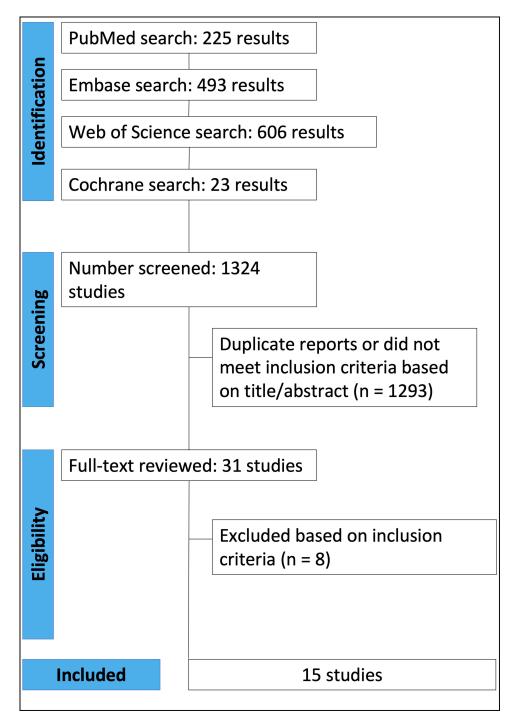


Figure I Flow diagram of study selection.

Topographic Results

Kmax significantly decreased after corneal CXL treatment (SMD = 0.15; 95% CI:0.01 0.28; Figure 5). No heterogeneity was observed (P = 0.94, $I^2 = 0\%$). However, ISV did not exhibit a significant difference post-CXL (SMD = 0.15; 95% CI: -0.24, 0.54; Figure 6), with no observed heterogeneity (P = 0.51, $I^2 = 0\%$). Similarly, the IVA did not show a significant difference post-CXL (SMD = 0.03; 95% CI: -0.35, 0.42; Figure 7), and there was no heterogeneity (P = 0.38, $I^2 = 0\%$). KI also demonstrated no significant difference post-CXL (SMD = -0.02; 95% CI: -0.40 0.37; Figure 8), and no heterogeneity was observed (P = 0.99, $I^2 = 0\%$). In contrast, CKI showed no significant difference post-CXL (SMD = -0.02; 95% CI: -0.40 0.37; Figure 8).

Author	Year	Country	Design	Eyes	Mean Age (Years)	Male/Female	Duration (Months)
Salgado ³³	2011	Germany	Prospective	20	38.4 (27–51)	9/6	12 month
Li G ³⁴	2012	China	Prospective	П	27.4 (20–36)	5/6	I2 month
Richoz ³⁵	2013	Switzerland	Retrospective	26	35 (23–46)	18/8	25 ± 13 month
Hallahan ²⁹	2014	United States	RCT	27	NA	17/10	3 month
Yildirim ³⁶	2014	Turkey	Retrospective	20	34 (25–45)	7/7	42 ± 7 month
Tong ³⁷	2017	Australia	Retrospective	14	NA	5/6	NA
Hersh ³⁰	2017	United States	RCT	91	NA	33/58	I2 month
Khairy ³¹	2019	Egypt	RCT	54	NA	NA	I2 month
Tian ³⁸	2021	China	Prospective	25	NA	19/6	NA
Chanbour ³⁹	2021	Lebanon	Retrospective	54	NA	26/28	36 month
Nasef ⁴⁰	2022	Egypt	Retrospective	21	NA	3/8	24 month
Margines ⁴¹	2023	United States	Prospective	82	42.8 (21–67)	NA	60 month
Rocha ³²	2014	Switzerland	RCT	14	NA	NA	3 month
Marino ⁴²	2015	Brazil	Prospective	40	33.8 (24–52)	15/9	24 month
Vinciguerra ⁴³	2010	Italy	Prospective	13	42 (30–59)	6/3	I2 month

Table I Baseline Characteristics of the Studies Included

Abbreviations: RCT, randomized clinical trial; NA, not applicable.

-0.73; 95% CI: -2.79 1.32; Figure 9), but heterogeneity was present (P < 0.01, I² = 93%). Rmin did not show significant changes post-CXL (SMD = -0.12; 95% CI: -0.51, 0.26; Figure 10), and there was no heterogeneity (P = 0.83, I² = 93%). The IHA did not differ significantly post-CXL (SMD = -0.21; 95% CI: -0.18, 0.59; Figure 11), and there was no heterogeneity (P = 0.65, I² = 0%). Lastly, IHD did not show significant differences post-CXL (SMD = 1.13; 95% CI: -1.01 to 3.27; Figure 12), but heterogeneity was observed (P < 0.01, I² = 92%).

Central Corneal Thickness and Endothelial Cell Count

The analysis of these data revealed that the pre- and post-CXL value differences in CCT were statistically significant (SMD = 0.24; 95% CI:0.07 to 0.41; Figure 13), with decreased values after treatment.

			e-CXL			st-CXL	5	Standar			I				Weight	Weigh
Study	Total	Mean	SD	Total	Mean	SD		Dif	ferend	ce		SMD	95	%-CI	(common)	(random
Yildirim 2014 (36)	20	0.78	0.6100	20	0.50	0.3100			++	-		0.57	[-0.07;	1.20]	7.0%	7.0%
Hersh 2017 (30)	91	1.41	1.4300	91	1.32	1.3760		-		_		0.06	[-0.23;	0.35]	33.0%	33.0%
Li G 2012 (34)	11	0.77	0.3200	11	0.70	0.7000			-			0.12	[-0.71;	0.96]	4.0%	4.0%
Margines 2023 (41)	82	0.75	0.3500	82	0.71	0.7100		-		_		0.07	[-0.24;	0.38]	29.8%	29.8%
Salgado 2011 (33)	20	0.53	0.3800	20	0.40	0.4000		_			_	0.33	[-0.30;	0.95]	7.2%	7.2%
Marino 2015 (42)	40	0.33	0.1800	40	0.37	0.1800			++-			-0.22	[-0.66;	0.22]	14.4%	14.4%
Vinciguerra 2010 (43)	13	1.08	0.4300	13	0.94	0.4600						0.30	[-0.47;	1.08]	4.7%	4.7%
Common effect model	277			277					4			0.09	[-0.07;	0.26]	100.0%	
Random effects model									\Rightarrow			0.09	[-0.07;	0.26]		100.0%
Heterogeneity: $I^2 = 0\%$, τ^2	= 0, p =	= 0.55					1			ſ	1					
							-1	-0.5	0	0.5	1					

Figure 2 UCVA forest plot.

Abbreviations: 1², heterogeneity; p. p-value; Pre-CXL, before crosslinking; Post-CXL, after crosslining; SMD, standardized mean difference.

Hersh 2017 (30) 91 0.96 1.4400 91 0.86 1.4000 Li G 2012 (34) 11 0.36 0.3000 11 0.23 0.2300 0.47 [-0.38; 1.32] 2.5% 4.4 Salgado 2011 (33) 20 0.19 0.2100 20 0.15 0.1400 0.22 [-0.40; 0.84] 4.7% 6.5 Richoz 2014 (35) 26 0.50 0.3000 14 0.20 0.400 0.84 [0.27; 1.41] 5.7% 7.2 Tong 2017 (37) 14 0.30 0.0600 14 0.20 0.0400 0.84 [0.27; 1.41] 5.7% 7.2 Chanbour 2021 (39) 28 0.11 0.1200 28 0.12 0.1200			Pre	e-CXL		Pos	st-CXL	Standardised Mean			Weight	Weight
Hersh 2017 (30) 91 0.96 1.4400 91 0.86 1.4000 0.07 [-0.22; 0.36] 21.7% 11.6 Li G 2012 (34) 11 0.36 0.3000 11 0.23 0.2300 0.47 [-0.38; 1.32] 2.5% 4.4 Salgado 2011 (33) 20 0.19 0.2100 20 0.15 0.1400 0.22 [-0.40; 0.84] 4.7% 6.5 Richoz 2014 (35) 26 0.50 0.3000 14 0.20 0.0400 - 0.84 [0.27; 1.41] 5.7% 7.2 Tong 2017 (37) 14 0.30 0.0600 14 0.20 0.0400 - 0.84 [0.27; 1.41] 5.7% 7.2 Chanbour 2021 (39) 28 0.11 0.1200 28 0.12 0.1200 - - 0.08 [-0.61; 0.44] 6.7% 7.8 Nasef 2022 (40) 21 0.22 0.2000 21 0.14 0.1100 - 0.38 [0.08; 0.69] 19.2% 11.3 Tian 2021 (38) 25 0.25 0.3100 25 0.15	Study	Total	Mean	SD	Total	Mean	SD	Difference	SMD	95%-CI	(common)	(random)
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Salgado 2011 (33) 20 0.19 0.2100 20 0.15 0.1400	Hersh 2017 (30)	91	0.96 1	1.4400	91	0.86	1.4000		0.07	[-0.22; 0.36]	21.7%	11.6%
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Nasef 2022 (40) 21 0.22 0.2000 21 0.14 0.1100 0.49 [-0.13; 1.10] 4.8% 6.6 Margines 2023 (41) 82 0.30 0.1600 82 0.23 0.2000 0.38 [0.08; 0.69] 19.2% 11.3 Tian 2021 (38) 25 0.25 0.3100 25 0.15 0.1700 0.39 [-0.17; 0.95] 5.8% 7.3 Rocha 2014 (32) 14 1.06 1.4200 14 1.00 1.4300 0.04 [-0.70; 0.78] 3.3% 5.3 Marino 2015 (42) 40 0.13 0.160 0.06 0.800 0.85 [0.04; 1.66] 2.8% 4.7 Common effect model 432 432 432 0.27 [0.14; 0.41] 100.0%	Chanbour 2021 (39)	28	0.11 (0.1200	28	0.12	0.1200		-0.08	[-0.61; 0.44]	6.7%	7.8%
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Rocha 2014 (32) 14 1.06 1.4200 14 1.00 1.4300 0.04 [-0.70; 0.78] 3.3% 5.3 Marino 2015 (42) 40 0.13 0.1000 40 0.15 0.1200 -0.18 [-0.62; 0.26] 9.5% 9.1 Vinciguerra 2010 (43) 13 0.16 0.1400 13 0.06 0.0800 -0.85 [0.04; 1.66] 2.8% 4.7 Common effect model 432 432 -0.27 [0.14; 0.41] 100.0%	Margines 2023 (41)	82	0.30 (0.1600	82	0.23	0.2000		0.38	[0.08; 0.69]	19.2%	11.3%
Marino 2015 (42) 40 0.13 0.1000 40 0.15 0.1200 -0.18 [-0.62; 0.26] 9.5% 9.1 Vinciguerra 2010 (43) 13 0.16 0.1400 13 0.06 0.0800 -0.85 [0.04; 1.66] 2.8% 4.7 Common effect model 432 432 -0.27 [0.14; 0.41] 100.0%	Tian 2021 (38)	25	0.25 (0.3100	25	0.15	0.1700		0.39	[-0.17; 0.95]	5.8%	7.3%
Vinciguerra 2010 (43) 13 0.16 0.1400 13 0.06 0.0800 0.85 [0.04; 1.66] 2.8% 4.7 Common effect model 432 432 0.27 [0.14; 0.41] 100.0%	Rocha 2014 (32)	14	1.06 1	1.4200	14	1.00	1.4300		0.04	[-0.70; 0.78]	3.3%	5.3%
Common effect model 432 432	Marino 2015 (42)	40	0.13 (0.1000	40	0.15	0.1200		-0.18	[-0.62; 0.26]	9.5%	9.1%
	Vinciguerra 2010 (43)	13	0.16 (0.1400	13	0.06	0.0800		0.85	[0.04; 1.66]	2.8%	4.7%
Dandam affaata madal 0.22 [0.12: 0.54] 100 0	Common effect model	432			432			\$	0.27	[0.14; 0.41]	100.0%	
Heterogeneity: $l^2 = 54\%$, $\tau^2 = 0.0788$, $p < 0.01$	Random effects model Hotorogonoity: $J^2 = 54\%$ r		188 n < 1	0.01					0.33	[0.12; 0.54]		100.0%

Figure 3 CDVA forest plot.

Abbreviations: 1², heterogeneity; p, p-value; Pre-CXL, before crosslinking; Post-CXL, after crosslinking; SMD, standardized mean difference.

		Pr	e-CXL		Pos	st-CXL	Sta	andardise	ed Mean				Weight	Weigh
Study	Total	Mean	SD	Total	Mean	SD		Differe	nce	SN	ID	95%-CI	(common)	(random
Khayri 2019 (31)	54	-3.75	1.2900	54	-3.12	1.2300				-0.	50 [-0.	88; -0.11]	17.9%	17.9%
Salgado 2011 (33)	20	-2.39	2.0300	20	-2.07	2.1800		-		-0.	15 [-0.	77; 0.47	6.8%	6.8%
Chanbour 2021 (39)	54	-2.64	3.7300	54	-2.49	4.0900				-0.	04 [-0.	42; 0.34]	18.5%	18.5%
Hallahan 2014 (29)	27	-3.37	4.2300	27	-3.02	4.3500	_			-0.	.0-] 80	61; 0.45]	9.2%	9.2%
Nasef 2022 (40)	21	-2.00	1.8000	21	-1.70	1.9000		-		-0.	16 [-0.	76; 0.45	7.2%	7.2%
Rocha 2014 (32)	14	-7.33	6.8400	14	-6.35	7.1800		-		-0.	14 [-0.	88; 0.61]	4.8%	4.8%
Vinciguerra 2010 (43)	13	-4.16	2.9000	13	-3.25	2.0500				-0.3	35 [-1.	13; 0.42]	4.4%	4.4%
Hersh 2017 (30)	91	-3.10	4.1000	91	-2.60	4.0000			-	-0.	12 [-0.	41; 0.17]	31.1%	31.1%
Common effect model	294			294				${\diamondsuit}$		-0.	19 [-0.	35; -0.02]	100.0%	-
Random effects model								\diamond		-0.	19 [-0.	35; -0.02]		100.0%
Heterogeneity: $I^2 = 0\%$, τ^2	= 0, p =	= 0.82							1					
							-1 -(.5 0	0.5	1				

Figure 4 SE forest plot.

Abbreviations: 1², heterogeneity; p, p-value; Pre-CXL, before crosslinking; Post-CXL, after crosslinking; SMD, standardized mean difference.

The difference in EEC was not statistically significant (SMD = 0.05; 95% CI: -0.15 to 0.26; Figure 14). Additionally, no statistically significant heterogeneity between studies was identified in the comparison of CCT (P = 0.58, I² = 0%) and EEC (P = 0.10, I² = 49%).

Adverse Events and Complications

The outcomes of severe complications are illustrated in Figure 15. A solitary instance of a serious complication emerged from one study, specifically a case involving epithelial ingrowth beneath the LASIK flap. Consequently, these findings indicate an aggregate occurrence rate of 0% (accompanied by a 95% confidence interval ranging from 0% to 1%). It is worth noting that there was no statistically significant variance among the studies (P = 1.00, $I^2 = 0$ %).

Discussion

Following the introduction of corneal collagen cross-linking in the treatment of keratoconus in 2003, many researchers have demonstrated the possibility of using CXL to slow or prevent corneal ectasia progression.^{23,44–49} Removal of the corneal epithelium remains essential with the current techniques and technology but raises the possibility of adverse

		Pi	e-CXL		Pos	st-CXL	Standardised Mean			Weight	Weigh
Study	Total	Mean	SD	Total	Mean	SD	Difference	SMD	95%-CI	(common)	(random
Yildirim 2014 (36)	20	46.00	4.4000	20	45.70	4.1000		0.07	[-0.55; 0.69]	4.8%	4.8%
Hersh 2017 (30)	91	55.40	6.9000	91	54.70	6.9000		0.10	[-0.19; 0.39]	21.9%	21.9%
Li G 2012 (34)	11	45.37	5.6400	11	43.23	5.3700		0.37	[-0.47; 1.22]	2.6%	2.6%
Salgado 2011 (33)	20	44.12	3.9700	20	44.43	4.0600		-0.08	[-0.70; 0.54]	4.8%	4.8%
Richoz 2014 (35)	26	52.80	5.0000	26	50.90	4.9000	- <u>+</u> ;	0.38	[-0.17; 0.93]	6.1%	6.1%
Tong 2017 (37)	14	52.90	2.1000	14	53.10	2.0000		-0.09	[-0.84; 0.65]	3.4%	3.4%
Chanbour 2021 (39)	28	50.99	6.3500	28	51.54	6.6100		-0.08	[-0.61; 0.44]	6.7%	6.7%
Hallahan 2014 (29)	27	55.54	6.1700	27	54.19	7.0200		0.20	[-0.33; 0.74]	6.5%	6.5%
Nasef 2022 (40)	21	48.60	4.6000	21	48.00	3.4000		0.15	[-0.46; 0.75]	5.0%	5.0%
Tian 2021 (38)	25	55.20	8.3300	25	54.19	8.0300		0.12	[-0.43; 0.68]	6.0%	6.0%
Margines 2023 (41)	82	46.50	4.8000	82	45.50	1.0900		0.29	[-0.02; 0.59]	19.5%	19.5%
Marino 2015 (42)	40	48.89	2.8500	40	48.91	2.9100		-0.01	[-0.45; 0.43]	9.6%	9.6%
Vinciguerra 2010 (43)	13	45.93	6.0300	13	42.49	4.8800		— 0.61	[-0.18; 1.40]	3.0%	3.0%
Common effect model	418			418			-	0.15	[0.01; 0.28]	100.0%	-
Random effects model							Ċ	0.15	[0.01; 0.28]		100.0%
Heterogeneity: $I^2 = 0\%$, τ^2	= 0, p =	= 0.94							_		
•							-1 -0.5 0 0.5 1				

Figure 5 Kmax forest plot.

Abbreviations: 1², heterogeneity: p, p-value; Pre-CXL, before crosslinking; Post-CXL, after crosslinking; SMD, standardized mean difference.

Study	Total	F Mean	Pre-CXL SD	Total		ost-CXL SD	Standardised Mean Difference	SMD	95%-CI	Weight (common)	
Tong 2017 (37) Tian 2021 (38) Vinciguerra 2010 (43)	25	88.16	15.7000 41.9200 33.2200	25	84.32	16.7000 41.5500 31.2200		0.09	[-0.25; 1.26] [-0.46; 0.65] [-0.88; 0.66]	26.2% 48.5% 25.2%	48.5%
Common effect model Random effects model Heterogeneity: $l^2 = 0\%$, τ^2	52 = 0, p =	= 0.51		52			-1 -0.5 0 0.5 1		[-0.24; 0.54] [-0.24; 0.54]	100.0% 	400.00/

Figure 6 ISV forest plot.

Abbreviations: I², heterogeneity; p, p-value; Pre-CXL, before crosslinking; Post-CXL, after crosslinking; SMD, standardized mean difference.

		F	Pre-CXL			st-CXL	Standardised Mean			Weight	Weigh
Study	Total	Mean	SD	Total	Mean	SD	Difference	SMD	95%-CI	(common)	(random
Tong 2017 (37)	14	1.10	0.2000	14	1.00	0.2000		0.49	[-0.27; 1.24]	26.3%	26.3%
Tian 2021 (38)	25	24.88	19.7800	25	26.85	20.4500		-0.10	[-0.65; 0.46]	48.5%	48.5%
Vinciguerra 2010 (43)	13	0.81	0.4500	13	0.90	0.4700		-0.19	[-0.96; 0.58]	25.1%	25.1%
Common effect model	52			52				0.03	[-0.35; 0.42]	100.0%	-
Random effects model								0.03	[-0.35; 0.42]		100.0%
Heterogeneity: $I^2 = 0\%$, τ^2	< 0.000	(1, p = 0)	0.38								
							-1 -0.5 0 0.5	1			

Figure 7 IVA forest plot.

Abbreviations: I², heterogeneity; p, p-value; Pre-CXL, before crosslinking; Post-CXL, after crosslinking; SMD, standardized mean difference.

events and complications such as infections and reduced corneal transparency.^{2,50} Therefore, this is an updated metaanalysis of epi-off CXL for corneal ectasia following refractive surgery.

In our systematic review and meta-analysis encompassing 15 studies with 421 patients, we assessed the safety and efficacy of CXL for treating corneal ectasia following RSL. Compared to a meta-analysis involving 118 patients, our analysis included additional outcome measures such as SE, IVA, CKI, IHA, ISV, and IHD. In contrast to the previous

	F 1	re-CXL		Pos	st-CXL	Standardised Mean			Weight	Weight
Γotal	Mean	SD	Total	Mean	SD	Difference	SMD	95%-CI	(common)	(random)
14	1.20	0.0700	14	1.20	0.0800		- 0.00	[-0.74; 0.74]	26.9%	26.9%
25	1.21	0.1600	25	1.21	0.1600		0.00	[-0.55; 0.55]	48.1%	48.1%
13	1.14	0.1300	13	1.15	0.1200 -		-0.08	[-0.85; 0.69]	25.0%	25.0%
52			52				-0.02	[-0.40; 0.37]	100.0%	
•	0.00						-0.02	[-0.40; 0.37]		100.0%
: 0, p =	: 0.99									
-	14 25 13 52	25 1.21 13 1.14	14 1.20 0.0700 25 1.21 0.1600 13 1.14 0.1300 52	14 1.20 0.0700 14 25 1.21 0.1600 25 13 1.14 0.1300 13 52 52 52	14 1.20 0.0700 14 1.20 25 1.21 0.1600 25 1.21 13 1.14 0.1300 13 1.15 52 52 52	14 1.20 0.0700 14 1.20 0.0800 25 1.21 0.1600 25 1.21 0.1600 13 1.14 0.1300 13 1.15 0.1200 - 52 52 52	14 1.20 0.0700 14 1.20 0.0800 25 1.21 0.1600 25 1.21 0.1600 13 1.14 0.1300 13 1.15 0.1200 52 52 1 1 1 0, p = 0.99 1 1 1 1	14 1.20 0.0700 14 1.20 0.0800 0.00 25 1.21 0.1600 25 1.21 0.1600 0.00 13 1.14 0.1300 13 1.15 0.1200 -0.08 52 52 -0.02 -0.02 -0.02 0, p = 0.99 -0.99 -0.02 -0.02	14 1.20 0.0700 14 1.20 0.0800 0.00 [-0.74; 0.74] 25 1.21 0.1600 25 1.21 0.1600 0.00 [-0.55; 0.55] 13 1.14 0.1300 13 1.15 0.1200 -0.08 [-0.85; 0.69] 52 52 -0.02 [-0.40; 0.37] -0.02 [-0.40; 0.37] 0, p = 0.99 -0.09 -0.02 [-0.40; 0.37] -0.02 [-0.40; 0.37]	14 1.20 0.0700 14 1.20 0.0800 0.00 [-0.74; 0.74] 26.9% 25 1.21 0.1600 25 1.21 0.1600 0.00 [-0.55; 0.55] 48.1% 13 1.14 0.1300 13 1.15 0.1200 -0.08 [-0.40; 0.37] 100.0% 52 52 -0.02 [-0.40; 0.37] 100.0%

Figure 8 KI forest plot.

Abbreviations: 1², heterogeneity: p, p-value; Pre-CXL, before crosslinking; Post-CXL, after crosslinking; SMD, standardized mean difference.

			e-CXL			st-CXL	Standardised Mean			Weight	•
Study	Total	Mean	SD	Total	Mean	SD	Difference	SMD	95%-CI	(common)	(random
Tong 2017 (37)	14	1.01	0.0100	14	1.04	0.0100		-2.91	[-4.02; -1.81]	14.4%	32.0%
Tian 2021 (38)	25	1.08	0.0700	25	1.07	0.0700		0.14	[-0.41; 0.70]	56.8%	34.4%
Vinciguerra 2010 (43)	13	1.01	0.0600	13	0.98	0.0700	+	0.45	[-0.33; 1.23]	28.8%	33.6%
Common effect model	52			52				-0.21	[-0.63; 0.21]	100.0%	-
Random effects model								-0.73	[-2.79; 1.32]		100.0%
Heterogeneity: I ² = 93%, τ	² = 3.12	224, p <	0.01								
						-	4 -2 0 2	4			

Figure 9 CKI forest plot.

Abbreviations: 1², heterogeneity; p, p-value; Pre-CXL, before crosslinking; Post-CXL, after crosslinking; SMD, standardized mean difference.

	0.2000	14	6.50	0 2000	1.4	0.00			-
		14	6.50						
6 24				0.2000		0.00	[-0.74; 0.74]	27.0%	27.0%
	0.9100	25	6.32	0.8700		-0.09	[-0.64: 0.47]	48.2%	48.2%
6.67	0.7700	13	6.88	0.4400 -		-0.32	[-1.10; 0.45]	24.7%	24.7%
		52				-0.12	[-0.51; 0.26]	100.0%	-
						-0.12	1-0.51: 0.261		100.0%
= 0.83						ר ד	[0.0., 0.20]		
- 0.00					1 05 0 05				
			52	52	52 = 0.83	52	52 -0.12 -0.12	52 = 0.83	52 = 0.83

Figure 10 Rmin forest plot.

Abbreviations: 1², heterogeneity; p. p-value; Pre-CXL, before crosslinking; Post-CXL, after crosslinking; SMD, standardized mean difference.

		F	Pre-CXL		Po	st-CXL	Standardised Me	ean		Weight	Weigh
Study	Total	Mean	SD	Total	Mean	SD	Difference	SMD	95%-CI	(common)	(random
Tong 2017 (37)	14	36.90	6.3000	14	33.60	8.5000		0.43	[-0.32; 1.18]	26.5%	26.5%
Tian 2021 (38)	25	0.85	0.5100	25	0.84	0.4800		- 0.02	[-0.53; 0.57]	48.6%	48.6%
Vinciguerra 2010 (43)	13	17.51	15.4700	13	12.82	11.4800		0.33	[-0.44; 1.11]	24.9%	24.9%
Common effect model	52			52				- 0.21	[-0.18; 0.59]	100.0%	
Random effects model							$\langle \rangle$	- 0.21	[-0.18; 0.59]		100.0%
Heterogeneity: $I^2 = 0\%$, τ^2	= 0, p =	= 0.65									
5,							-1 -0.5 0 0.	5 1			

Figure II IHA forest plot.

Abbreviations: I², heterogeneity; p, p-value; Pre-CXL, before crosslinking; Post-CXL, after crosslinking; SMD, standardized mean difference.

Study	Total	Pı Mean	re-CXL SD	Total	Po: Mean	st-CXL SD	Standardised Mean Difference	SMD	95%-CI	Weight (common)	•
Tong 2017 (37)	14	0.10	0.0200	14	0.01	0.0300	: <u> </u>	3.43	[2.21; 4.64]	12.0%	31.7%
Tian 2021 (38)	25	0.12	0.0800	25	0.11	0.0700		0.13	[-0.42; 0.69]	57.8%	34.5%
Vinciguerra 2010 (43)	13	0.06	0.0400	13	0.06	0.0300		0.00	[-0.77; 0.77]	30.1%	33.8%
Common effect model	52			52				0.49	[0.07; 0.91]	100.0%	
Random effects model								1.13	[-1.01; 3.27]		100.0%
Heterogeneity: $I^2 = 92\%$, τ	$^{2} = 3.37$	792, p <	: 0.01								
0,		.,					-4 -2 0 2 4				

Figure 12 IHD forest plot.

Abbreviations: 1², heterogeneity; p, p-value; Pre-CXL, before crosslinking; Post-CXL, after crosslinking; SMD, standardized mean difference.

			Pre-CXL		F	Post-CXL	Standardised Mean			Weight	Weigh
Study	Total	Mean	SD	Total	Mean	SD	Difference	SMD	95%-CI	(common)	(random
Hersch 2017	91	2538.00	525.0000	91	2610.00	438.0000		-0.15	[-0.44; 0.14]	51.1%	32.4%
Li G 2012 (34)	11	3096.22	288.6100	11	2930.11	249.0600		0.59	[-0.26; 1.45]	5.9%	11.2%
Tian 2021 (38)	25	2469.00	437.0000	25	2357.00	364.0000		0.27	[-0.28; 0.83]	13.9%	19.5%
Marino 2015 (42)	40	2997.00	119.0000	40	2997.00	122.0000	<u> </u>	0.00	[-0.44; 0.44]	22.5%	24.7%
Vinciguerra 2010 (43)	13	2555.00	470.0000	13	2120.00	517.0000	1	0.85	[0.04; 1.66]	6.6%	12.2%
Common effect model	180			180			\diamond	0.05	[-0.15; 0.26]	100.0%	-
Random effects model								0.18	[-0.16; 0.51]		100.0%
Heterogeneity: $I^2 = 49\%$, τ	2 = 0.06	674, p = 0.	.10								
							-1.5 -1 -0.5 0 0.5 1 1.5				

Figure 13 EEC forest plot.

Abbreviations: 1², heterogeneity; p, p-value; Pre-CXL, before crosslinking; Post-CXL, after crosslinking; SMD, standardized mean difference.

		Pre-C	XL	Post-CXL		Standardised Mean			Weight	Weight
Study	Total	Mean	SD Total	Mean	SD	Difference	SMD	95%-CI	(common)	(random
Yildirim 2014 (36)	20	477.00 27.00	00 20	465.00	23.0000	+ + =	0.47	[-0.16; 1.10]	7.0%	7.0%
Hallahan 2014 (29)	27	433.00 67.00	00 27	407.00	64.0000		0.39	[-0.15; 0.93]	9.6%	9.6%
Li G 2012 (34)	11	436.10 32.25	500 11	448.50	22.6100		-0.43	[-1.28; 0.42]	3.9%	3.9%
Tian 2021 (38)	25	414.92 40.96	600 25	415.50	41.8900		-0.01	[-0.57; 0.54]	9.0%	9.0%
Nasef 2022 (40)	21	457.60 48.40	00 21	442.60	56.9000		0.28	[-0.33; 0.89]	7.5%	7.5%
Margines 2023 (41)	82	451.00 55.00	00 82	436.00	25.0000		0.35	[0.04; 0.66]	29.2%	29.2%
Chanbour 2021 (39)	28	463.10 57.90	00 28	459.50	69.4000		0.06	[-0.47; 0.58]	10.1%	10.1%
Rocha 2014 (32)	14	418.00 45.00	000 14	381.00	45.0000		- 0.80	[0.02; 1.57]	4.6%	4.6%
Marino 2015 (42)	40	466.00 63.00	00 40	456.00	56.0000		0.17	[-0.27; 0.61]	14.4%	14.4%
Vinciguerra 2010 (43)	13	503.45 43.35	500 13	504.00	38.3000		-0.01	[-0.78; 0.76]	4.7%	4.7%
Common effect model	281		281			\$	0.24	[0.07; 0.41]	100.0%	-
Random effects model						Ś	0.24	[0.07; 0.41]		100.0%
Heterogeneity: $I^2 = 0\%$, τ^2	< 0.00	01, <i>p</i> = 0.58			Г					
- ,					-1.5	5 -1 -0.5 0 0.5 1	1.5			

Figure 14 CCT forest plot.

Abbreviations: I², heterogeneity; p, p-value; Pre-CXL, before crosslinking; Post-CXL, after crosslinking; SMD, standardized mean difference.

meta-analysis, our findings revealed improved results for Kmax and SE values and reduced in CCT post-CXL. Other outcomes analyzed in the previous study remained consistent.

The benefit of visual acuity improvement appears to be the leading indicator of CXL treatment. Although the UCVA remained stable, CDVA improved significantly. In this context, Hersch et al reported that CDVA improved by 1–9 letters in 32 (44%) eyes and by ten or more letters in 23 (32%) eyes after 12 months of follow-up. Potential explanations for the enhancement in CDVA, as opposed to UDVA, following CXL could be attributed to the effect of CXL on improving corneal anterior surface regularity and potentially facilitating partial repositioning of the optical zone. Similar findings have been reported in the context of keratoconus post-crosslinking treatment.⁵¹

Study C	Complications T	otal		Proportion	95%-CI	Weight (common)	•
Salgado 2011 (33)	0	20 -		0.00	[0.00; 0.17]	1.6%	1.6%
Li G 2012 (34)	0	11		- 0.00	[0.00; 0.28]	0.5%	0.5%
Richoz 2014 (35)	0	26 🛉		0.00	[0.00; 0.13]	2.7%	2.7%
Yildirim 2014 (36)	0	20 🛉		0.00	[0.00; 0.17]	1.6%	1.6%
Tong 2017 (37)	0	14 🛉		0.00	[0.00; 0.23]	0.8%	0.8%
Chanbour 2021 (39)	0	54		0.00	[0.00; 0.07]	11.1%	11.1%
Tian 2021 (38)	0	25 🛉		0.00	[0.00; 0.14]	2.5%	2.5%
Margines 2023 (41)	0	82 ⊦		0.00	[0.00; 0.04]	25.1%	25.1%
Nasef 2022 (40)	0	21		0.00	[0.00; 0.16]	1.8%	1.8%
Hersh 2017 (30)	1	91		0.01	[0.00; 0.06]	15.2%	15.2%
Khairy 2019 (31)	0	90 F		0.00	[0.00; 0.04]	30.2%	30.2%
Marino 2015 (42)	0	40		0.00	[0.00; 0.09]	6.2%	6.2%
Vinciguerra 2010 (43)	0	13 -		0.00	[0.00; 0.25]	0.7%	0.7%
Common effect model		507	>	0.00	[0.00; 0.01]	100.0%	
Random effects model Heterogeneity: $I^2 = 0\%$, $\tau^2 =$	0 = 1.00	Ċ	>	0.00	[0.00; 0.01]		100.0%
Helefogeneity: $I = 0\%, \tau =$	ο, <i>ρ</i> – 1.00	0	0.05 0.1 0.15 0.2 0.25				

Figure 15 Serious Complications rates.

Abbreviations: I², heterogeneity; p, p-value; Pre-CXL, before crosslinking; Post-CXL, after crosslinking.

SE analyses are essential for evaluating refractive outcomes following corneal cross-linking in post-refractive surgery corneal ectasia. The results show a statistically significant decrease in SE after CXL, indicating a beneficial impact on refractive error, similar to other systematic reviews of CXL.⁵² This reduction in SE suggests that CXL contributes to the correction of defocus and possibly higher-order aberrations due to improvements in the corneal surface, which are critical for achieving better visual acuity. This can be explained by corneal flattening resulting from CXL. This flattening is associated with potential myopia correction. However, its effect on hypermetropia remains a subject of ongoing debate.⁵³ Furthermore, the heterogeneity was low, unlike in other studies that compared SE with CXL treatment.²⁵

Analysis of the topographic results further supports the positive impact of CXL on corneal anatomy. Significant reductions in parameters, such as Kmax, indicate flattening of the corneal curvature and might improve refractive and visual outcomes.⁵⁴ One possible explanation for these differences in the outcomes is the specific focus and sensitivity of each parameter. Parameters such as Kmax are directly related to corneal curvature and astigmatism; thus, changes in these parameters are likely to be clinically significant and readily detectable.⁵⁵ In contrast, other indices such as ISV and IVA may be more subtle and less responsive to the effects of CXL, particularly in cases where extreme changes in surface variance or vertical asymmetry do not primarily characterize ectasia. Additionally, individual patient variability and heterogeneity of corneal ectasia patterns could contribute to the lack of significant changes in specific parameters. Corneas with different ectatic patterns may respond differently to CXL, and this diversity could attenuate the observed effects on specific indices.⁵⁶ Furthermore, because simulated keratometry does not consider the total corneal surface, we considered the Klyce^{57–60} and Ambrósio⁶¹ indices of keratoconus. The CKI decreased with decreasing severity of central keratoconus. This augmentation may be consistent with the fact that ectatic corneas after excimer laser refractive surgery may have the most pronounced curvature changes near the center and that CXL may result in improvements in this central corneal area, where the deepest ablation and tissue removal are found. Therefore, there seemed to be a stabilization of corneal curvature and a trend towards central flattening, leading to an enhanced refractive outcome.

In contrast to the previous meta-analysis, this analysis revealed a statistically significant decrease in CCT after CXL. This occurs because of the regularization of epithelium and compaction of the collagen lamellae in the corneal stroma. Similar results have been reported for keratoconus after postoperative crosslinking.^{62,63} The lack of statistical heterogeneity in this outcome enhanced the reliability of this finding, indicating consistency across the included studies. Conversely, the evaluation of the EEC did not show a statistically significant change after CXL. The stability in EEC implies that the CXL procedure does not harm endothelial cells, which are essential for maintaining corneal deturgescence and transparency, as well as in previous studies.^{35,64} The moderate heterogeneity observed in this result suggests that variations in study methodologies and patient populations may play a role in changes in EEC.

The observed results in parameters like Kmax value (showing tissue applanation) are likely due to the improved biomechanical stability of the cornea after CXL, a trend consistent with findings in primary keratoconus patients undergoing similar treatment.^{23,34} Another study noted corneal changes after CXL in a comparable manner, though less pronounced, in eyes with keratoconus.^{43,46} However, contrasting results emerged from a different study, where the progression rate post-CXL was notably higher in patients with ectasia compared to those with keratoconus over three years.³⁹ Additionally, in another investigation, CDVA and Kmax were inferior in patients with ectasia compared to keratoconus patients after CXL.⁶⁵ This discrepancy may be attributed to heightened variability in the timeline of changes in ectasia patients compared to those with keratoconus. The underlying cause of these potential differences between keratoconus and post-RSL ectatic corneas remains unclear. Various factors, such as biomechanical disparities introduced by the LASIK flap, possible differences in riboflavin diffusion rates in post-LASIK corneas (especially at the flap interface), and intrinsic pathophysiologic distinctions between keratoconus and post-RSL ectasia may contribute to the varied responses to CXL observed in these groups.⁶⁵ To gain a more comprehensive understanding of these differences and determine which is more stable between eyes with keratoconus and eyes with corneal ectasia after RLS, further research involving a more significant number of eyes is essential.

Some investigations of ectasia risk factors have shown it is necessary. LASIK surgery may compromise the cornea's structural integrity by diminishing the total load-bearing tissue available and reallocating the load-bearing duty to the structurally less robust posterior stromal layer of the cornea.⁶⁶ The eye asymmetry scoring⁶⁷ and the presence of abnormal topography are the other risk factors for ectasia.^{68–70} The Belin-Ambrosio Enhanced Ectasia Display (BAD) uses nine specific parameters, including elevation and pachymetry data, to generate a "D value". This value identifies patients at risk for keratoconus or conditions like keratoconus who might not be suitable for refractive surgery or may have a poor prognosis.⁷⁰ The BAD provides an exhaustive perspective on the corneal tomography architecture, thereby enabling the identification of such patients via a holistic examination. Thus, factors including the topographic abnormality, percentage tissue altered, residual stromal thickness, inferior-superior index of the sagittal map, Bellin Ambrosio display data, and information from the thickness curve map are acknowledged risk factors for ectasia development.^{68,69,71} However, our investigation into this topic was constrained by inconsistent data availability across the analyzed studies. The lack of detailed information about this data in many of the included studies limited our ability to delve deeper into understanding these risk factors' role in CXL treatment. Due to this, it is necessary future research that consistently incorporates these metrics, allowing for more robust analyses and more definitive conclusions about the causes of ectasia after RLS and different prognoses after CXL.

CXL for corneal ectasia after RLS proved to be a safe procedure, with an extremely low complication rate. The absence of statistically significant variance among the studies suggests that complications after CXL are relatively rare and are not significantly affected by variations in the study methodologies. Most CXL complications are caused by corneal epithelial removal. Innovative protocol modifications are used to improve safety and efficacy without the need for epithelial removal and to expand indications in thinner corneas.^{64,72–77} We look forward to reviewing the results of these new techniques and technologies shortly.

Our meta-analysis had limitations and biases due to the quality of the selected studies. While the results of this metaanalysis provide valuable insights into the safety and efficacy of corneal collagen cross-linking for treating post-refractive surgery corneal ectasia, it is essential to consider the limitations of this study. As corneal ectasia after refractive surgery has a low occurrence, our analysis included a relatively small number of studies and patients. Although the inclusion of 15 studies is a strength, a larger dataset would provide more robust and generalizable conclusions. The presence of heterogeneity in some outcomes, such as CKI, IHD, and EEC, suggests variability in treatment responses across studies. This variability could be influenced by factors, such as variations in CXL protocols, patient populations, and study methodologies. Many studies included in this analysis reported follow-up periods ≥ 12 months. However, long-term outcomes beyond the follow-up period were not assessed. It is essential to evaluate the stability of CXL effects over an extended period. The possibility of publication bias cannot be ruled out, as studies with positive or significant results may be more likely to be published, whereas studies with negative or non-significant findings may not. This bias may have affected the overall conclusions of the meta-analysis. Different studies may have employed variations in CXL protocols, such as the type of riboflavin used, ultraviolet light intensity, and duration of treatment. These variations can influence treatment outcomes and introduce heterogeneity. The meta-analysis relied on aggregated data from published studies. Individual patient data were not available for analysis, which could have provided a more detailed understanding of the treatment effects and potential predictors of response. The included studies were conducted in various countries, potentially introducing variability in the patient demographics, surgical techniques, and follow-up protocols. This may limit the generalizability of the findings to a broader population. This meta-analysis primarily focused on assessing the outcomes of CXL treatment compared with the preoperative baseline values. It did not include a direct comparison with alternative treatments for post-refractive surgery corneal ectasia, limiting its ability to determine the relative efficacy of CXL.

Conclusions

In conclusion, this systematic review and meta-analysis suggest that CXL for managing cases of corneal ectasia after surgery is a safe and effective technique, successfully arresting or stopping the process of corneal ectasia, improving topographic indices, and providing good refractive and visual outcomes with a very low risk of severe complications. There is a need to standardize protocols for riboflavin concentration, duration of UV exposure, and classification of corneal ectasia severity before treatment for better understanding and indication of this treatment and patient counseling. Moreover, conducting a new meta-analysis to validate these findings is essential, especially as more randomized, more significant, and longer clinical trials become available for publication.

Data Sharing Statement

All relevant data are within the paper.

Author Contributions

All authors made substantial contributions to the conception and design, acquisition of data, or analysis and interpretation of data; took part in drafting the article or revising it critically for important intellectual content; gave final approval of the version to be published; have agreed on the journal to which the article has been submitted; and agree to be accountable for all aspects of the work.

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