







Visual Outcomes After Sequential Posterior Chamber Phakic IOL with Corneal Refractive Surgery (Bioptics) for the Treatment of Myopic Astigmatism

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Purpose: To report the outcomes of sequential posterior chamber phakic intraocular lens (PC-pIOL) with corneal refractive surgery in conventional (PC-pIOL prior to refractive surgery) and reverse (refractive surgery prior to PC-pIOL) biptics for treating high myopic astigmatism.

Setting: Tertiary refractive center, Draper, Utah, USA.

Design: Retrospective case series.

Methods: Medical records of patients who underwent planned biptics were reviewed. Surgery involved PC-pIOL placement using an implantable collamer lens (ICL) with preceding or subsequent LASIK or PRK. Pre- and postoperative manifest spherical equivalent (SEQ), visual acuity, and PC-pIOL vault were analyzed.

Results: Of the 51 eyes present at 12 months postoperatively, 49 eyes (96%) achieved target SEQ within ± 1.00 D and an identical amount achieved refractive astigmatism ≤ 1.00 D. Post-biptics eyes achieved a postoperative UDVA equal to or better than preoperative CDVA in 45 eyes (88%). Efficacy and safety indices were 1.08 ± 0.20 (41 eyes) and 1.13 ± 0.22 (44 eyes) for conventional biptics and 0.99 ± 0.42 (7 eyes) and 1.15 ± 0.38 (7 eyes) for reverse biptics eyes at 12 months. The maximum PC-pIOL vault of conventional biptics eyes (27 eyes) within 6 months before and after LASIK/PRK was 385 ± 159 μ m and 377 ± 135 μ m, respectively ($P = 0.71$).

Conclusion: Biptics for high myopic astigmatism was safe and effective. Reverse biptics, although not as traditional, could provide similar results. Additionally, the PC-pIOL vault does not appear to be affected by LASIK/PRK.

Keywords: biptics, LASIK, PRK, posterior chamber phakic IOL, high myopic astigmatism, visual outcomes

Introduction

Biptics, originally described by Zaldivar et al,¹ utilizes the combination of a posterior chamber phakic intraocular lens (PC-pIOL) with excimer laser corneal ablation to split the optical correction between two planes, the corneal plane and the ciliary sulcus plane. Consequently, biptics can treat complex and high refractive errors while maintaining a large optical zone with minimal induced spherical aberrations. As of today, the term has expanded to encompass phakic, pseudo-phakic, and clear lens extraction cases combined with several forms of corneal refractive procedures (LASIK, PRK, intracorneal rings, and RK).²⁻⁵ The

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combination of intraocular implant and corneal curvature modification expands the scope of treatment for patients with significant refractive errors that fall outside the recommended treatment range of excimer laser refractive surgery as well as patients with unfavorable corneal thicknesses that prevent full treatment of their refractive error.^{6–9} Only bioptics procedures involving PC-pIOL utilizing the Visian Implantable Collamer Lens V4b (ICL; STAAR Surgical, Monrovia, CA), and to a lesser extent, the Visian Toric ICL (TICL; STAAR Surgical, Monrovia, CA), will be discussed here.

Previously, PC-pIOL implantation could only address the spherical component of refractive errors, relying on a separate corneal procedure to correct the astigmatic element. With the advent of the TICL, coexisting astigmatism can now be treated using a single procedure, and several studies have already demonstrated the safety and efficacy of TICL implantation.^{10–14} However, TICLs are currently limited to four diopters of astigmatic correction. As a result, bioptics remains a useful treatment option in patients with large amounts of astigmatism.

The majority of reported bioptics cases involve using a PC-pIOL to correct most of the patient's refractive error followed by a corneal refractive procedure several months later to reduce any residual refractive error. The literature on planned bioptics procedures performed in the reverse order, with corneal surgery prior to intraocular lens implantation, for the primary treatment of high refractive errors is not as extensive. This reverse order of procedures, termed by Leccisotti⁷ as “reverse bioptics,” could offer similar results to conventional bioptics.

This retrospective study aims to assess the safety, efficacy, and predictability of PC-pIOL in combination with LASIK and PRK bioptics procedures in both conventional and reverse orders as performed at one site.

Methods

This study followed the tenets of the Helsinki Declaration and was approved by the Hoopes Vision Ethics Board. All patients were adequately informed prior to obtaining their signed consent. Only conventional bioptics or reverse bioptics cases with planned, consecutive procedures were included in this study. Patients undergoing unplanned LASIK or PRK enhancements after or before PC-pIOL implantation and patients receiving corneal refractive surgery in addition to cataract surgery were excluded. The preoperative evaluation included uncorrected distance visual acuity (UDVA), corrected distance visual acuity

(CDVA), manifest refraction, tonometry, keratometry, pachymetry, and anterior chamber depth (ACD) measurements. Adverse events and complications were recorded at all visits.

Surgeries were performed by two surgeons (MM and PH). At least 1 week prior to PC-pIOL implantation, YAG peripheral iridotomies were performed at the 11 and 1 o'clock positions. The selection of proper Visian ICL size was based upon the Parkhurst nomogram utilizing measurements of sulcus-to-sulcus dimension, lenticular rise, and aqueous depth obtained by VUmax (Sonomed Escalon, New Hyde Park, NY) ultrasound biomicroscopy. The PC-pIOL was inserted through a temporal 3.0 mm clear corneal incision and placed into the sulcus. Postoperatively, patients received ofloxacin 0.3% or moxifloxacin 0.5% four times a day for 1 week. Prednisolone acetate 1% was also applied topically four times daily for the first week then tapered weekly.

For LASIK procedures, the AMO iFS (Abbott Medical Optics, Inc., Santa Ana, CA) femtosecond laser system was used for flap creation and the WaveLight EX500 excimer laser system (Alcon Laboratories, Inc., Fort Worth, TX) was used for stromal ablation with a 6.5 mm central optical zone and blend zone to 9.0 mm. Flap diameter was between 8.5 and 9.0 mm, and flap thickness was between 100 and 115 μ m with the creation of a superior hinge. The postoperative treatment protocol included ofloxacin 0.3% or moxifloxacin 0.5% four times a day for 1 week. Patients were instructed to apply prednisolone acetate 1% every hour while awake for the first 24 hours. On postoperative day 1, the prednisolone was decreased to four times daily for 1 week and subsequently stopped.

With PRK procedures, alcohol debridement of the corneal epithelium was performed, then excimer laser was used for stromal ablation as previously discussed. Postoperative treatment included ofloxacin 0.3% or moxifloxacin 0.5% four times a day for 1 week. Ketorolac 0.5% was used twice daily for 3 days. Prednisolone acetate 1% was also applied topically four times daily for the first month, then stopped. Fluorometholone 0.1% was then used three times daily for 3 weeks, then twice daily for 3 weeks, then once daily for 3 weeks. For treatments involving an ablation depth of greater than 64 μ m, a topical application of 0.02% mitomycin-C was used to decrease the risk of corneal haze. Lastly, a bandage contact lens was inserted postoperatively and removed at the 1 week post-op visit.

Table 1 Preoperative Baseline Characteristics for Eyes That Underwent Conventional Bioptics and Reverse Bioptics

	Conventional Bioptics	Reverse Bioptics	P value
Number of eyes (patients)	87 (49)	13 (7)	–
M/F (eyes)	41:46	4:8	–
PRK/LASIK (eyes)	61:26	2:11	–
Mean age (range) (y)	33.1 (21 to 56)	33.7 (21 to 45)	0.82
Time between procedures (month)*	3.8 ± 2.1 (1 to 10)	2.67 ± 2.1 (1 to 7)	0.07
UDVA*	1.56 ± 0.32 (1.00 to 3.00)	1.56 ± 0.12 (1.48 to 1.90)	0.97
CDVA*	0.03 ± 0.09 (0.00 to +0.40)	0.11 ± 0.10 (0.00 to +0.30)	<0.01*
SEQ (D)*	−11.12 ± 3.39 (−19.88 to −3.38)	−12.42 ± 1.81 (−15.38 to −9.13)	< 0.01*
Refractive cylinder (D)*	−2.49 ± 1.16 (−6.00 to −0.50)	−3.88 ± 1.43 (−6.75 to −2.00)	< 0.01*

Notes: Visual acuity reported as LogMAR. *Values reported as mean ± SD (range). Monovision eyes were excluded from UDVA analysis.

Abbreviations: UDVA, uncorrected distance visual acuity; CDVA, corrected distance visual acuity; SEQ, spherical equivalent refraction.

Follow-up after the initial procedures for both groups was performed at 1 day, 1 week, 1 month, 3 months, 6 months, and 12 months unless the second procedure occurred before the 6-month or 12-month appointment. The perioperative evaluation included UDVA, CDVA, manifest refraction, tonometry, keratometry, pachymetry, and anterior segment ocular coherence tomography (OCT). Postoperative follow-up after the final procedures occurred at 1 day, 1 month, 3 months, 6 months, and 12 months and included the above-mentioned parameters.

Statistical analysis was performed using IBM SPSS Statistics 2020 v27.0 (IBM corp., Armonk, NY). Wilcoxon rank-sum test was applied to compare outcomes in conventional and reverse bioptics. A paired *t*-test was used to compare changes in PC-pIOL vault prior to and after LASIK/PRK in conventional bioptics. $P < 0.05$ was considered statistically significant.

Results

In this retrospective study using de-identified data, the medical records of 100 eyes of 56 patients (46 male and 54 female) undergoing sequential, planned PC-pIOL implantation and corneal refractive surgery at one site between January 2012 and December 2019 were evaluated (Table 1). The mean age of patients was 33.5 ± 7.6 years (SD) (range 21 to 57).

In 87 eyes, PC-pIOL implantation was followed by either PRK for 61 eyes (70%) or LASIK for 26 eyes (30%). Three eyes of two patients, whose procedures occurred after FDA approval of TICL in

September 2018, received TICL due to high astigmatic errors (≥ 4.00 D). The TICL provided its maximum of four diopters of astigmatic correction and PRK corrected the remaining astigmatic (0.00 to 1.25 D) and spherical refractive error. Thirteen eyes of eight patients received PRK for two eyes (15%) or LASIK for eleven eyes (85%) prior to PC-pIOL implantation. Time between procedures was 3.6 ± 2.1 months (range 1 to 10).

Follow up was 83 eyes at 1 month, 77 eyes at 3 months, 61 eyes at 6 months, and 51 eyes at 12 months (Table 2). Although not all patients completed a 1-month postoperative visit, all patients were evaluated during at least one of the future time points analyzed. Four eyes that were purposely corrected to be myopic to achieve monovision were excluded from the postoperative UDVA and manifest spherical equivalent (SEQ) results. Refractive surgery results were reported via the standard graphs¹⁵ using data collected from the 3- and 12-month follow-ups.

Preoperative UDVA was worse than 20/200 in all 100 eyes. Visual outcomes at 3 and 12 months postoperatively are presented in Figure 1. Only one eye (1.3%) experienced a transient loss of 2 lines of CDVA at 3 months. However, this loss in CDVA was likely due to the initial post-PRK recovery phase and the steroid taper, as the patient's BCVA and UCVA recovered to 20/20 at the 6-month visit. The mean preoperative SEQ was -11.30 ± 3.25 D (range -19.88 to -3.38) improving to 0.01 ± 0.35 D (-0.50 to 1.13) for the 77 eyes at 3 months. At the 12-month follow-up visit, the mean SEQ was -0.08 ± 0.31 D (-0.88 to 0.75) for 51 eyes.

Table 2 Pre- and Postoperative Visual Outcomes at Postoperative Time Points

	Preop	1mo	3mo	6mo	12mo
Number of eyes (patients)	100 (56)	83 (47)	77 (44)	61 (36)	51 (30)
UDVA	1.56 ± 0.30 (1.00 to 3.00)	0.09 ± 0.13 (−0.12 to +0.48)	0.03 ± 0.11 (−0.12 to +0.40)	0.04 ± 0.13 (−0.12 to +0.40)	0.02 ± 0.12 (−0.12 to +0.30)
CDVA	0.04 ± 0.09 (0.00 to +0.40)	0.03 ± 0.10 (−0.12 to +0.40)	−0.01 ± 0.10 (−0.12 to +0.30)	−0.01 ± 0.10 (−0.12 to +0.30)	−0.01 ± 0.10 (−0.12 to +0.30)
SEQ (D)	−11.30 ± 3.25 (−19.88 to −3.38)	−0.15 ± 0.46 (−1.50 to +1.13)	0.01 ± 0.35 (−0.50 to +1.13)	−0.03 ± 0.34 (−0.63 to +1.13)	−0.08 ± 0.31 (−0.88 to +0.75)
Refractive cylinder (D)	−2.67 ± 1.29 (−6.75 to −0.50)	−0.49 ± 0.36 (−1.50 to 0.00)	−0.41 ± 0.32 (−1.50 to 0.00)	−0.41 ± 0.31 (−1.25 to 0.00)	−0.42 ± 0.34 (−1.25 to 0.00)

Notes: Visual acuity is reported as LogMAR. Values reported as mean ± SD (range). Monovision eyes were excluded from UDVA analysis.

Abbreviations: UDVA, uncorrected distance visual acuity; CDVA, corrected distance visual acuity; SEQ, spherical equivalent refraction.

Pre- and postoperative outcomes for refractive astigmatism are displayed in [Figure 2](#). The mean preoperative refractive astigmatism was -2.67 ± 1.29 D (range -6.75 to -0.50) improving at 3 months postoperatively to -0.41 ± 0.32 D (-1.50 to 0.00). At 12 months postoperatively, the mean refractive astigmatism was -0.42 ± 0.34 D (-1.25 to 0.00).

At 3 months, the surgically induced astigmatism vector (SIA) was within ± 1.00 D of target induced astigmatism vector (TIA) for 76 eyes (99%) (one eye overcorrected) and within ± 0.50 D for 70 eyes (91%) (three over- and four under-corrected). At 12 months, SIA was within ± 1.00 D of TIA for 49 eyes (96%) (one over- and one under-corrected) and within ± 0.50 D of TIA for 43 eyes (84%) (one over- and seven under-corrected). Linear regression curves for SIA as a function of TIA demonstrated equal distribution of over- and under-corrections and mild amount of scatter (R^2 of 0.92) for both 3 and 12 months postoperatively.

The mean SEQ was utilized to evaluate stability in all patients which is displayed in [Figure 3](#). The preoperative mean SEQ of the 100 eyes was -11.30 ± 3.25 D (range -19.88 to -3.38). Mean SEQ of 83 eyes at 1 month was -0.15 ± 0.46 D (-1.50 to 1.13); 77 eyes at 3 months was 0.01 ± 0.35 D (-0.50 to 1.13); 61 eyes at 6 months was -0.03 ± 0.34 D (-0.63 to 1.13); and 51 eyes at 12 months, was -0.08 ± 0.31 D (-0.88 to 0.75).

[Table 3](#) compares the change in efficacy index (mean postoperative UDVA/mean preoperative CDVA) and safety index (mean postoperative CDVA/mean preoperative CDVA) for conventional and reverse bioptics. At 1

month, the differences between the efficacy and safety indices of each group were statistically significant ($P < 0.01$); however, this significance was not present at any of the follow-up visits. At 12 months, 38 eyes (93%) after conventional bioptics and 4 eyes (57%) after reverse bioptics achieved a postoperative UDVA equal to or better than preoperative CDVA.

[Table 4](#) compares the change in mean preoperative SEQ and refractive cylinder in both groups. Monovision eyes were excluded from the SEQ results. The difference in preoperative values and refractive cylinders at 3 months was statistically significant ($P < 0.01$). Differences between the two groups at all other time points were no longer significant. At 12 months postoperatively, all eyes after conventional bioptics were within ± 1.00 D of target SEQ and 36 eyes (88%) were within ± 0.50 D. Seven eyes (100%) after reverse bioptics achieved ± 0.25 D. The astigmatic error was ≤ 1.00 D for all but one eye (2.3%) after conventional bioptics, and ≤ 0.50 D for 33 eyes (75%) and 5 eyes (71%) after conventional and reverse bioptics, respectively.

PC-pIOL vault depth was recorded using the RTVue-XR Avanti OCT system (Optovue, Inc., Fremont, CA) prior to and following LASIK/PRK for 27 eyes (31%) in the conventional bioptics group to evaluate the effect LASIK/PRK had on the PC-pIOL vault ([Figure 4](#)). The vault prior to LASIK/PRK was 385 ± 159 μ m (44 to 672) shifting to 377 ± 135 (130 to 610) after the ablative procedures. The difference was not significant ($P = 0.71$). Of the 27 available, 15 eyes (58%) had an average decrease in vault of 67 ± 88 μ m (2 to 350), while the

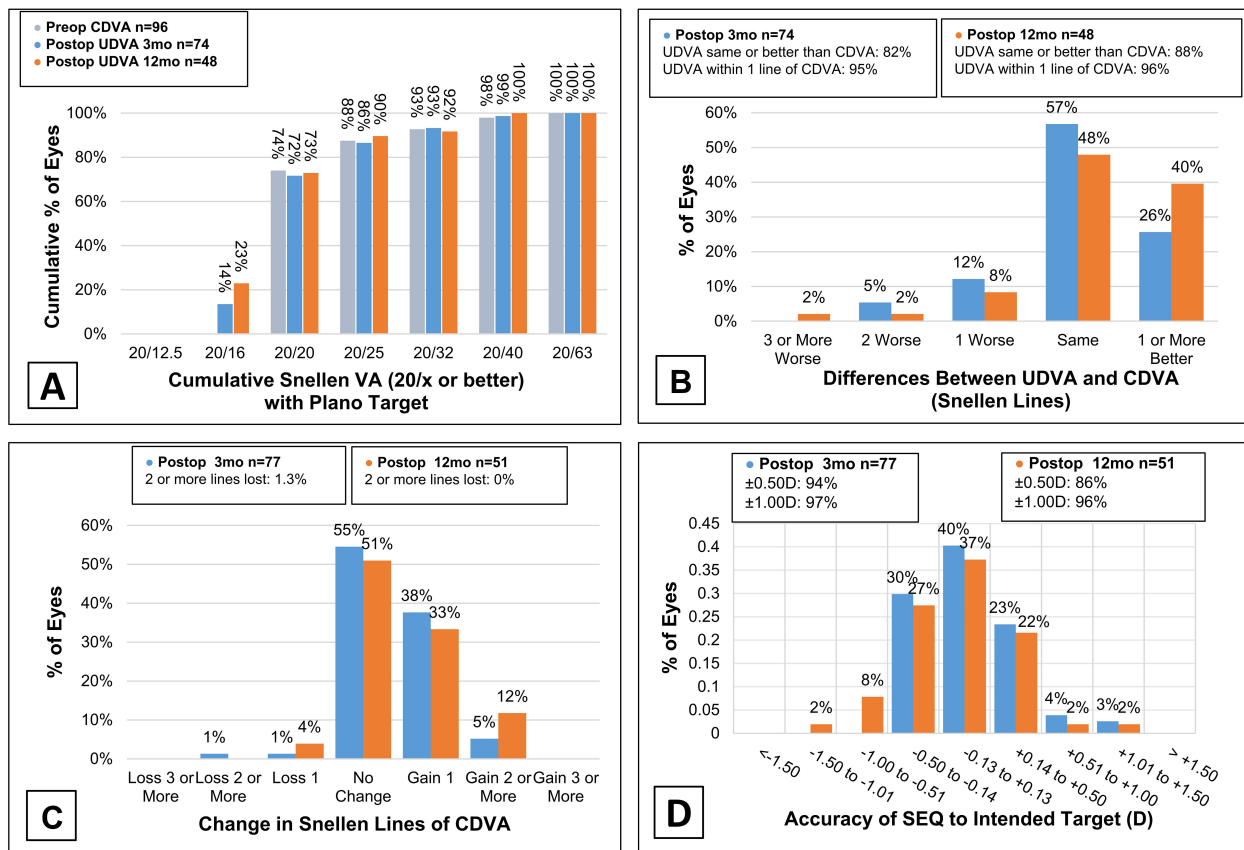


Figure 1 Visual and refractive outcomes for eyes at 3 and 12 months postoperatively following all bioptics procedures. UDVA values of monovision eyes were not included. **(A)** Cumulative postoperative UDVA and preoperative CDVA. **(B)** Difference in postoperative UDVA and CDVA. **(C)** Change in CDVA. **(D)** Attempted vs achieved SEQ. **Abbreviations:** UDVA, uncorrected distance visual acuity; CDVA, corrected distance visual acuity; D, diopters; SEQ, manifest spherical equivalent.

remaining 12 eyes (46%) had an average increase of $67 \pm 46 \mu\text{m}$ (10 to 166).

The most common adverse event was dry eyes requiring punctal plugs, which occurred in three eyes (3%) of two patients after PC-pIOL implantation and six eyes (6%) of three patients after laser ablation surgery. Two patients had bilateral flap dislocation detected on postoperative day one requiring flap irrigation and refloat; however, neither patient experienced a loss in UDVA nor in CDVA at the final follow-up visits.

Discussion

This study examines the visual outcomes after planned bioptics procedures with a sub-analysis of eyes that underwent reverse bioptics. As reported in previous studies of planned and unplanned bioptics,^{1-9,16-19} our results further support bioptics as a safe, effective, and stable option for correcting myopic astigmatism.

Data on the outcomes of planned reverse bioptics in the literature is limited in comparison to that of conventional bioptics. Reverse bioptics is typically unplanned, occurring in patients who have previously undergone a corneal refractive procedure and request an enhancement but are subsequently poor corneal surgery candidates.^{7,20} In our study, patients receiving planned reverse bioptics showed similar visual improvements when compared to patients after conventional bioptics as well as in comparison to studies of unplanned reverse bioptics.²⁰⁻²³ However, due to the limited sample size ($n = 13$), additional studies with longer follow-up and larger sample sizes are needed to make a definitive conclusion about the outcomes of planned reverse bioptics for high myopic astigmatism. The small sample size of reverse bioptics was in part due to patient preference being the main determinant of whether or not reverse or conventional bioptics was performed. Further studies could reveal unique potential benefits of planned reverse bioptics as well.

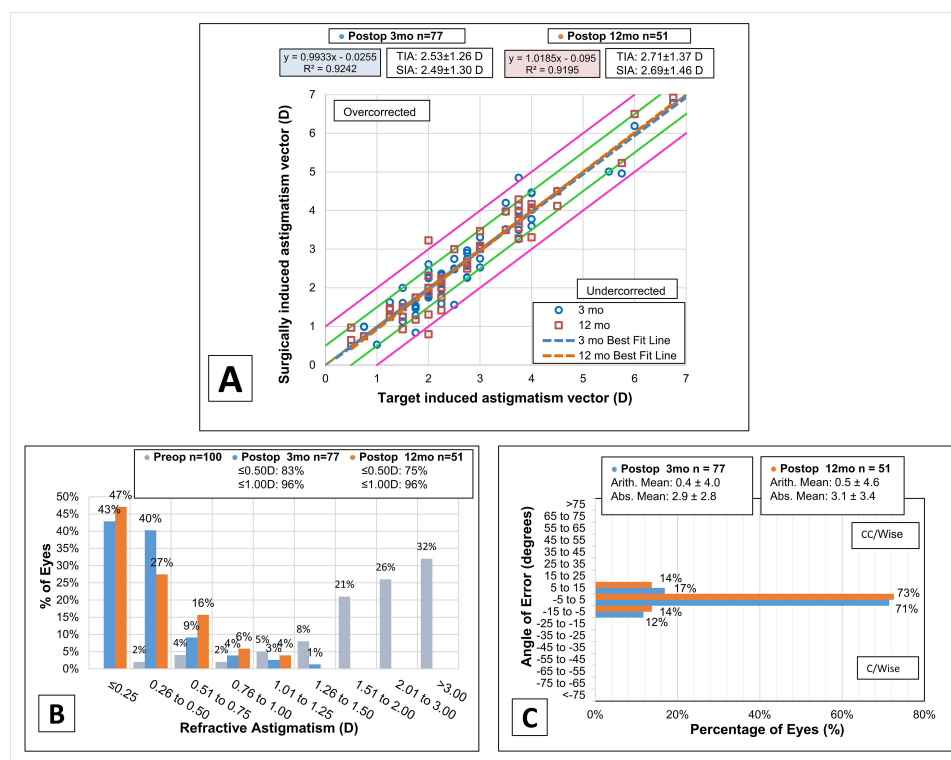


Figure 2 Outcomes of refractive astigmatism for myopic eyes at 3 and 12 months postoperatively following all bioptics procedures. **(A)** Magnitude of TIA versus the magnitude of SIA. **(B)** Change in refractive astigmatic magnitude. **(C)** Postoperative angle of error.

Abbreviations: TIA, target induced astigmatism; SIA, surgically induced astigmatism; D, diopters.

Unlike the IOL power calculations for cataract surgery, PC-pIOL calculations are less dependent on corneal refractive power and therefore less affected by preceding corneal refractive surgery in reverse bioptics patients.^{23,24} Additionally, because PC-pIOL calculations are partly based on subjective refraction, reducing myopia provides a more accurate calculated PC-pIOL power.¹ This in turn could lead to a larger optical zone along with diminished glare and halos in reverse bioptics patients. Reducing the power of the PC-pIOL also lessens the thickness of the lens. A thinner PC-pIOL implant would be less likely to contact and abrade the anterior crystalline lens, theoretically leading to decreased risk of anterior subcapsular cataract formation.

Due to the time required for PC-pIOL incisions to heal, LASIK is typically delayed at least 3 months to avoid potential complications from the suction ring applied in both microkeratome- and femtosecond-assisted LASIK. In reverse bioptics, prolonging the delay before the last procedure is not necessary. In our study, the average time between procedures for conventional (14 patients) and reverse bioptics (6 patients) that received LASIK was 15 and 8 weeks, respectively, demonstrating the potential decrease in inter-procedural delay.

The effects of LASIK/PRK on the PC-pIOL vault has rarely been examined in the literature. Vault narrowing has been directly linked to adverse events such as pigment dispersion syndrome and anterior subcapsular cataract formation.^{24,25} The multiple forces applied to the cornea during LASIK/PRK could theoretically lead to vault narrowing. However, our study found that LASIK/PRK did not cause acute vault narrowing. Future research should confirm these initial findings.

While bioptics was previously the principle method for correcting high astigmatic refractive errors, several studies have demonstrated the safety and efficacy of TICL implantation.^{10–14} When Choi et al²⁷ compared 29 eyes which received TICL with 26 eyes that underwent bioptics procedures to treat moderate to high myopia with astigmatism, the two groups had similar visual outcomes at 12 months postoperatively, with quicker visual improvement and fewer postoperative higher-order aberrations in the TICL group. In a later study by Alfonso et al,²⁸ patients who underwent TICL versus bioptics had comparable outcomes; however, the bioptics group achieved a slightly higher efficacy index (1.09 ± 0.19 compared to 0.98 ± 0.20) and refractive

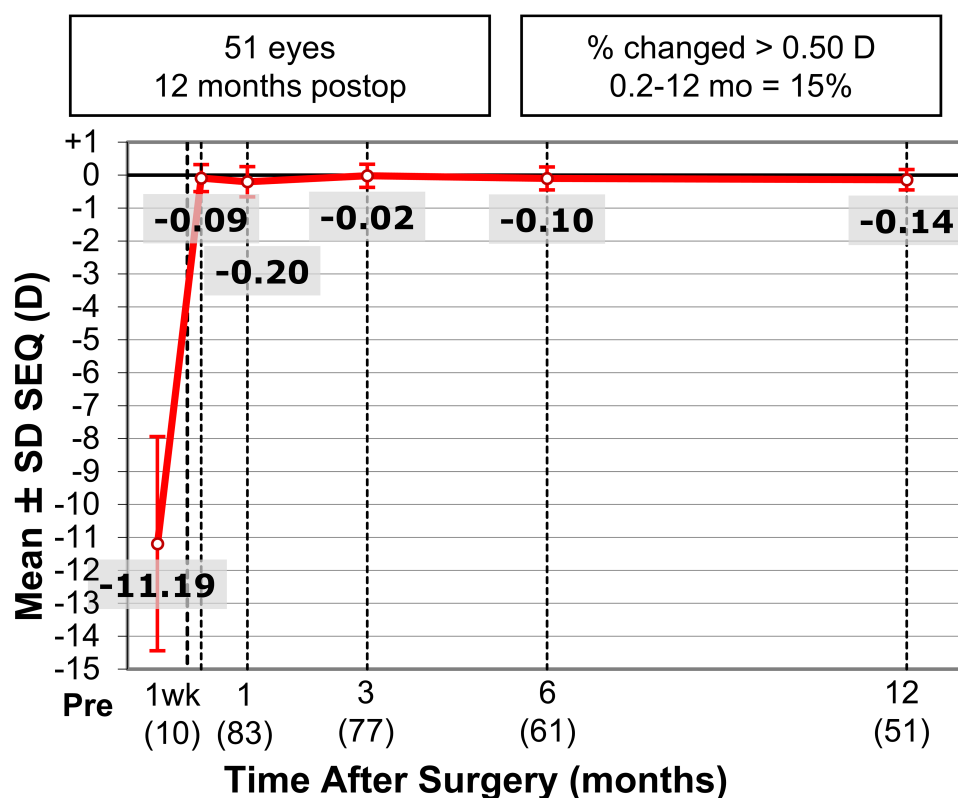


Figure 3 Stability of bioptics procedures reported as mean SEQ at 1 week and 1, 3, 6, and 12 months postoperatively.
Abbreviations: SEQ, manifest spherical equivalent; D, diopters.

predictability (77 of eyes [94%] within ± 0.50 D of emmetropia compared to 66 eyes [81%] in the TICL group). Regardless, TICL may overall be the preferred treatment. With a lower risk of surgical complications, decreased cost to the patient, quicker results, and the convenience of a single surgery; TICL is superior to bioptics in several regards.²⁷

However, bioptics may yet still play a useful role, as TICL alone may be unable to address cases of high myopia (> -19.00 D) and significant astigmatism (> 4.00 D). In our study, three eyes of two patients with astigmatic errors ranging from -4.00 to -5.25 D received TICL implants that provided a maximum astigmatic correction of 4 diopters followed by PRK 4.5 ± 2.6 months (3 to 9) later to

Table 3 Efficacy and Safety Index for Eyes That Underwent Conventional Bioptics and Reverse Bioptics

Postop	Efficacy Index			Safety Index		
	C Bioptics [Eyes]	R Bioptics [Eyes]	P value	C Bioptics [Eyes]	R Bioptics [Eyes]	P value
1 mo	0.87 ± 0.23 (0.40 to 1.50) [69]	1.24 ± 0.32 (1.00 to 2.00) [11]	$<0.01^*$	1.00 ± 0.20 (0.50 to 1.67) [72]	1.39 ± 0.24 (1.25 to 2.00) [11]	$<0.01^*$
3 mo	1.01 ± 0.20 (0.67 to 1.67) [67]	1.28 ± 0.24 (0.67 to 1.33) [7]	0.06	1.11 ± 0.19 (0.67 to 1.67) [70]	1.25 ± 0.12 (1 to 1.33) [7]	0.09
6 mo	1.02 ± 0.22 (0.50 to 1.60) [49]	1.10 ± 0.30 (0.63 to 1.67) [9]	0.95	1.12 ± 0.18 (0.80 to 1.67) [52]	1.30 ± 0.20 (1.00 to 1.67) [9]	0.16
12 mo	1.08 ± 0.20 (0.67 to 1.67) [41]	0.99 ± 0.42 (0.50 to 1.60) [7]	0.56	1.13 ± 0.22 (0.80 to 1.67) [44]	1.15 ± 0.38 (1.00 to 2.00) [7]	0.49

Notes: Values reported as mean \pm SD (range) [number of eyes]. Monovision eyes were excluded from efficacy index analysis. *Statistically significant.

Abbreviations: C Bioptics, conventional bioptics; R Bioptics, reverse bioptics.

Table 4 Postoperative Spherical Equivalent Refraction and Refractive Cylinder in Eyes That Underwent Conventional Bioptics and Reverse Bioptics

Spherical Equivalent Refraction (D)				Refractive Cylinder (D)		
	C Bioptics	R Bioptics	P value	C Bioptics	R Bioptics	P value
Preop	-11.12 ± 3.39 (-19.88 to -3.38) [83]	-12.42 ± 1.81 (-15.38 to -9.13) [13]	<0.01*	-2.49 ± 1.16 (-6.00 to -0.50) [87]	-3.88 ± 1.43 (-6.75 to -2.00) [13]	<0.01*
Postop 1 mo	-0.16 ± 0.49 (-1.50 to +1.13) [69]	-0.15 ± 0.22 (-0.38 to +0.38) [11]	0.96	-0.50 ± 0.37 (-1.50 to 0.00) [72]	-0.43 ± 0.32 (-0.75 to 0.00) [11]	0.70
3 mo	0.01 ± 0.37 (-0.50 to +1.13) [67]	0.00 ± 0.19 (-0.25 to +0.38) [7]	0.57	-0.38 ± 0.31 (-1.50 to 0.00) [70]	-0.71 ± 0.25 (-1.25 to -0.50) [7]	<0.01*
6 mo	-0.05 ± 0.33 (-0.63 to +1.13) [49]	0.06 ± 0.41 (-0.63 to +0.88) [9]	0.47	-0.38 ± 0.32 (-1.25 to 0.00) [52]	-0.56 ± 0.23 (-1.00 to -0.25) [9]	0.23
12 mo	-0.08 ± 0.33 (-0.88 to +0.75) [41]	-0.11 ± 0.18 (-0.25 to +0.25) [7]	0.69	-0.40 ± 0.33 (-1.25 to 0.00) [44]	-0.50 ± 0.38 (-1.00 to 0.00) [7]	0.53

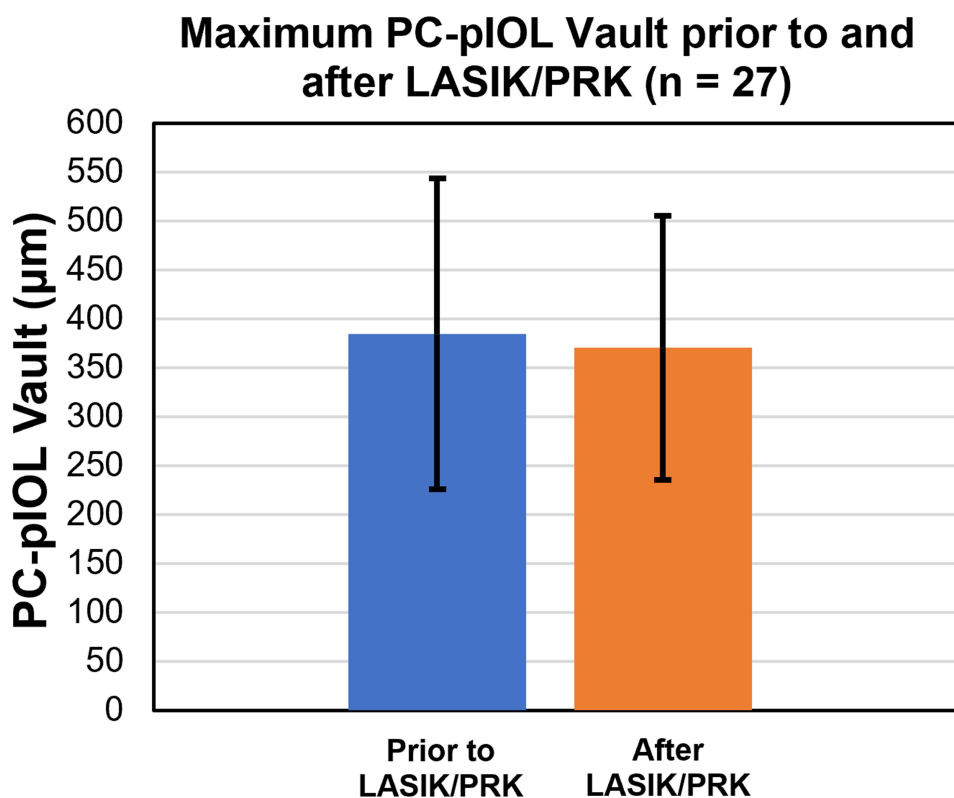
Notes: Values reported as mean ± SD (range) [number of eyes]. Monovision eyes were excluded from spherical equivalent refraction analysis. *Statistically significant.

Abbreviations: C Bioptics, conventional bioptics; R Bioptics, reverse bioptics.

reduce the remaining astigmatic and spherical error. All three eyes achieved an SEQ ± 0.50 D of emmetropia and ≤1.00 D of astigmatism with no loss in CDVA at their final follow-up visits. Our results indicate that TICL bioptics

could be a viable treatment option for high astigmatic patients.

In addition to the limitations inherent in retrospective studies, this study was limited, from a statistical

**Figure 4** PC-plIOL vaults recorded within 6 months prior to and after LASIK/PRK in conventional bioptics. (P = 0.71).

Abbreviation: PC-plIOL, posterior chamber phakic intraocular lens.

standpoint, by the relatively small sample size of the reverse bioptics group ($n = 13$). Additionally, in most of the patients, eyes were treated bilaterally rather than randomly assigning treatment to a single eye. Deciding which patients received bioptics versus reverse bioptics as well as LASIK versus PRK was left to the discretion of the surgeon and patient preference and was not randomized. Lack of stratification of the changes in PC-pIOL vault size between LASIK and PRK due to the limited sample size was an additional limitation. Specular microscopy data were not collected; however, this was not the objective of this work. The study was also limited due to a lack of follow-up beyond 12 months. Due to the higher risk of cataract formation and endothelial cell density loss after PC-pIOL implantation, the importance of long-term follow-up should be emphasized when counselling patients. Despite the limitations, this study provides useful information currently missing in the scientific literature as data regarding planned reverse bioptics is notably sparse.

In summary, bioptics procedures are safe and effective. While TICL is superior to bioptics in several regards, bioptics may still serve a purpose for treating cases of high myopia or myopia with significant astigmatism that fall outside the effective treatment ranges of PC-pIOLs and TICLs alone. The present findings suggest that PRK/LASIK does not appear to cause substantial PC-pIOL vault narrowing, though further research is needed to confirm this conclusion. Lastly, in addition to treating patients who are poor corneal enhancement surgery candidates, planned reverse bioptics could be a useful, primary treatment option for astigmatic and myopic refractive errors.

Ethics and Consent Statement

This retrospective study using de-identified data has been approved by the Hoopes Vision Ethics Board and conforms with the Helsinki Declaration of 1964, as revised in 2013, concerning human and animal rights. The patients signed informed consent.

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Author Contributions

All authors take responsibility for the integrity of the work, and have given final approval to the version to be published. All authors made substantial contributions to 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; agreed to submit to the current journal; gave final approval of the version to be published; and agree to be accountable for all aspects of the work.

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Disclosure

All authors declare that they have no conflicts of interest.

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