ORIGINAL RESEARCH

Refractive Outcomes in Fuchs' Endothelial Corneal Dystrophy: Conventional and Femtosecond Laser-Assisted Cataract Surgery

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Correspondence: Ellen H Koo Bascom Palmer Eye Institute, Department of Ophthalmology, Miller School of Medicine at the University of Miami, Miami, FL, USA Tel + 1 561-515-1544 Email exk126@med.miami.edu **Purpose:** To investigate the refractive outcomes of eyes with Fuchs' endothelial corneal dystrophy (FECD) following phacoemulsification.

Methods: This is a retrospective chart review of patients with FECD who underwent phacoemulsification. Manifest refraction at the early postoperative period (1–6 weeks) and late postoperative period (3–12 months) was collected. The spherical equivalent (SE) and variance of SE from target in diopters (D) were analyzed.

Results: A total of 219 eyes from 175 FECD patients (73 FLACS, 146 conventional phacoemulsification) were included. In the early postoperative period, when comparing variance from intended target, 62% (n=126) had a SE variance of ≤ 0.5 D, 22% (n=44) >0.5 D and ≤ 1 D, and 17% (n=34) >1 D. In the late postoperative period, 62% (n=85) had a SE variance of ≤ 0.5 D, 19% (n=26) >0.5 D and ≤ 1 D, and 20% (n=27) >1 D. There was no difference in the variance of SE comparing FLACS versus conventional phacoemulsification in either the early postoperative period (p=0.78) or the late postoperative period (p=0.29).

Conclusion: Patients with mild-to-moderate FECD had favorable refractive outcomes with phacoemulsification. There was no difference in refractive outcomes in eyes with FECD between the group that underwent FLACS versus the group that underwent conventional phacoemulsification.

Keywords: femtosecond laser assisted cataract surgery, Fuchs' dystrophy, Fuchs' endothelial corneal dystrophy, refractive error, refractive outcome

Introduction

One of the many challenges associated with cataract surgery in Fuchs' endothelial corneal dystrophy is predicting the refractive outcome. Factors, such as fluctuations in pre-operative data, central corneal thickness, manifest refraction, topography and tomography, as well as biometric data, serve to influence this difficulty with predictability.^{1–4}

Femtosecond-laser assisted cataract surgery (FLACS) has been proposed to confer several advantages compared to standard phacoemulsification including shorter phacoemulsification time and less ultrasound energy usage, both of which have been shown to reduce endothelial cell loss and postoperative corneal edema in the general cataract population.^{5–11} In addition, it has been shown FLACS may result in a more predictable Effective Lens Position (ELP), and thereby aid in targeting the refractive outcome.^{12,13}

As modern phacoemulsification has become more advanced, we have seen an expansion in the recommendation criteria for cataract surgery in FECD.^{14–16}

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Existing literature evaluating the refractive outcomes of patients with FECD after phacoemulsification—whether conventional or FLACS—is extremely sparse.

Methods

This retrospective cohort study was approved by the Institutional Review Board (IRB)/Ethics Committee at the University of Miami. The patient informed consent waiver was obtained for this retrospective study. The study was carried out with patient data confidentiality and compliance in accordance with the Declaration of Helsinki. A chart review of all patients with a prior diagnosis of FECD who underwent either conventional phacoemulsification cataract surgery or FLACS between January 1, 2014 and January 1, 2017 at Bascom Palmer Eye Institute was performed. The cases that underwent FLACS all underwent femtosecond laser capsulorrhexis formation, nucleus fragmentation, as well as corneal wound creation. All procedures were performed at the Bascom Palmer Eye Institute by the surgeons with the Comprehensive and Cornea services. Surgery was performed using either the Centurion Vision (Alcon, Irvine, CA) or Whitestar Signature (Johnson & Johnson, New Brunswick, NJ) phacoemulsification system and any of the following femtosecond laser platforms: LenSx (Alcon, Irvine, CA), Catalys (Johnson & Johnson, New Brunswick, NJ), or Victus (Bausch & Lomb Surgical, Rancho Cucamonga, CA).

Inclusion and Exclusion Criteria

A search for charts within the electronic medical record system with concurrent CPT procedural codes for cataract surgery (66,982 and 66,984) and ICD-10 (H18.5) or ICD-9 (371.57) diagnostic codes for FECD yielded a total of 753 cases. Cases with prior keratoplasty or glaucoma surgery, concurrent cataract extraction with keratoplasty, complications related to cataract surgery or laser application, incomplete records, and follow-up time less than 3 months were excluded. Eyes that required keratoplasty in the 12-month period following cataract surgery as a result of corneal decompensation were also excluded. A total of 219 eyes (73 FLACS and 146 conventional phacoemulsification) from 175 patients were evaluated in this study.

Chart Review and Data Collection

The charts were evaluated for demographic and clinical variables including age, gender, date, type of cataract surgery (conventional vs FLACS), preoperative bestcorrected visual acuity (BCVA), and central corneal thickness (CCT) by ultrasound pachymetry. Nuclear sclerosis was assessed by slit-lamp examination and graded based on the Wisconsin Cataract Grading System.¹⁷ The clinical grade for FECD was assigned according to the degree of corneal guttae and edema documented upon slit-lamp exam as previously described.¹⁸ All cases of FECD were categorized as mild (grades 1–2) or moderate (grades 2.5– 4). Advanced cases of FECD (grades 5–6) were excluded as they underwent concurrent endothelial keratoplasty. The BCVA, CCT (when available), and corneal edema findings documented at each postoperative visit were analyzed. The spherical equivalence (SE) and variance of SE from target refraction in diopters (D) were analyzed.

Follow-Up Periods

Due to the retrospective nature of the study, postoperative visits were not available to all patients at consistent follow-up times. Therefore, we divided the follow-up into broad windows: the early postoperative period (including visits from 1 to 6 weeks following surgery) and the late postoperative period (including visits from 3 to 12 months following surgery). If a patient had more than one visit per period, the visit closest to the center of the window was selected for analysis.

Statistical Analyses

Categorical and ordinal variables were summarized with counts and percentages, while continuous variables were summarized with means and standard deviations (SD). Due to the inclusion of both eyes of some patients, statistical analyses comparing the femtosecond group to the conventional treatment group, as well as assessing the influence of risk factors, were conducted using Generalized estimating equation (GEE) models (IBM SPSS Statistics version 22, Armonk NY) employing exchangeable correlation structure and interval, binomial, or multinomial links as appropriate.

Results

Baseline Patient Characteristics

A total of 175 patients were identified. Out of the 175 patients, 52 patients had undergone conventional phacoemulsification, and 123 had undergone FLACS. Two eyes were included from a single patient as long as the study eye met the Inclusion and Exclusion criteria. A total of 219 eyes were included in the study, with 73 eyes in the

Laser Treatment	Femtosecond Laser	Conventional	Generalized Estima			
Table I Baseline Characteristics in Risk Factor Analysis° for Fuchs Endothelial Corneal Dystrophy Patients						

Laser Treatment	Femtosecond Laser	Conventional	Generalized Estimating Equation (GEE) Model P- value
Number of Patients	52	123	
Number of Eyes for Analyses	73	146	
Age (y), mean age ± SD	74.1 ± 7.5	75.2 ± 8.5	0.58
Best-corrected visual acuity, Mean logMAR (BCVA) ± SD [Range]	0.231 ± 0.163	0.331 ± 0.264	0.001
Visual Acuity Group, N (%) ≥ 20/30 20/30 – 60 20/70 – 5/200	40 (56) 30 (42) 2 (3)	60 (41) 62 (43) 24 (16)	0.024
Central corneal thickness (μm) ^{Φ,} mean ± SD [Range] Not collected N, (%) Collected N, (%)	576 ± 39 [496, 623] 55 (75) 18 (25)	574 ± 39 [496, 648] 83 (57) 63 (43)	0.81 0.038
Gender N, (%) Male Female	21 (29) 52 (71)	48 (33) 98 (67)	0.83
Corneal guttata grade N, (%) 0.5–1.0 1.5 –2.0 2.5–3.0 3.5–4.0	16 (24) 24 (36) 22 (33) 5 (8)	42 (33) 38 (30) 35 (28) 12 (9)	0.67
Cataract grade N, (%) 0.5–1.0 1.5–2.0 2.5–3.0 3.5–4.0	7 (10) 34 (49) 29 (41) 0	5 (4) 84 (58) 52 (36) 3 (2)	0.16
Target refractions Mean (SD) [Range]	-0.3 (0.3) [-2, 0]	-0.1 (0.1) [-3, 0]	0.005

Notes: ^{Φ}Preoperative central corneal thickness was not available for every patient; values were calculated from n = 18 (femtosecond laser) and n = 63 (conventional); ^{ε}Tabled samples sizes are based on all eyes with early follow-up data at 1–6 weeks and/or late follow-up data at 3–12 months.

conventional phacoemulsification group, and 146 eyes in the FLACS group. Demographics were similar between treatment groups (Table 1). Eyes with pre-operative CCT measurement were more common in the conventional group (43%, n=63) than in the FLACS group (25%, n=18, p=0.038). However, there was no significant difference between the mean thickness values (p=0.81). Conventionally treated eyes also had visual acuity of 0.1 logMAR (approximately one Snellen line on average) worse than the FLACS eyes (p=0.001). Notably, there was no statistically significant difference in cataract grade (p=0.16) and corneal guttata grade (p=0.67).

Spherical Equivalent Outcomes

Early and late follow-up period Spherical Equivalent (SE) outcomes for both the FLACS and conventional groups were investigated and compared (Tables 2 and 3, Figure 1). The mean target SE correction for eyes in the FLACS group was -0.3 diopter (D) (range of -2 D to plano). The mean target SE correction for eyes in the conventional group was -0.1 diopter (D) (range of -3D to plano). The mean deviation from the target SE correction was -0.10 D \pm 0.83 in the early postoperative period for all eyes. The mean deviation from target SE correction was -0.14 D \pm 0.85 in the late follow-up group for all

Table 2 Eyes with Early	(I-6 Weeks)	Postoperative Follow-Up
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Eyes with Early (I–6 Week) Postoperative Follow-Up						
	Femtosecond	Conventional	Total	GEE p-value		
N Eyes	70	134	204			
Preoperative Spherical Equivalent, Mean ± SD [range]	N/A	N/A	N/A	N/A		
Preoperative logMAR BCVA Mean ± SD [range]	0.23 ± 0.17 [0, 1.0]	0.32 ± 0.26 [-0.1, 1.6]	0.29 ± 0.24 [-0.1, 1.6]	0.003		
Preoperative Acuity Group, N (%)				0.042		
≥ 20/30	38 (55%)	57 (43%)	95 (47%)			
60	29 (42%)	56 (42%)	85 (42%)			
20/70 – 5/200	2 (3%)	21 (16%)	23 (11%)			
	Early Postoperative Spherical	Equivalent, target corrected (SE	тс)			
SE _{TC} , Mean ± SD [range]	-0.07± 0.89 [-2.00, 2.75]	-0.11 ± 0.80 [-2.63, 3.25]	-0.10 ± 0.83 [-2.63, 3.25]	0.78		
Absolute value SE _{TC} , Mean ± SD [range]	0.66 ± 0.60 [0, 2.75]	0.56 ± 0.58 [0, 3.25]	0.60 ± 0.59 [0, 3.25]	0.25		
Outcome classification by deviation from target refraction, N (%)				0.24		
Good (≤0.5D)	39 (56%)	87 (65%)	126 (62%)			
Fair (>0.5 to ≤1D)	19 (27%)	25 (19%)	44 (22%)			
Poor (>ID)	12 (17%)	22 (16%)	34 (17%)			
	Early Postop	erative Visual acuity		_		
logMAR BCVA Mean ± SD [range]	0.14 ± 0.14 [0, 0.7]	0.17 ± 0.25 [-0.1, 2.0]	0.16 ± 0.22 [-0.1, 2.0]	0.18		
Visual Acuity Group				0.29		
≥ 20/30	57 (81%)	97 (73%)	154 (76%)			
20/30 - 60	12 (17%)	28 (21%)	40 (20%)			
20/70 – 5/200	I (1%)	8 (6%)	9 (4%)			

eyes. In the group that underwent FLACS, the mean deviation from target SE correction was -0.07 D \pm 0.89 in the early postoperative period and $-0.10 \text{ D} \pm 0.68$ in the late postoperative period. In the conventional group, the mean deviation from target SE correction was $-0.11 \text{ D} \pm$ 0.80 in the early postoperative period and $-0.15D \pm 0.94$ in the late postoperative period. In both the early postoperative period and the late postoperative period, there was no statistical difference (p > 0.005) when comparing the mean deviation from target SE between FLACS and conventional groups (p=0.78 and 0.47, respectively for the two follow-up periods). For the early period, the corresponding 95% CI for the regression coefficient 0.039 (representing the difference between the mean deviation from target SE in the FLACS group and the mean deviation from target SE in the conventional group) was -0.23, 0.31. For the later period, the corresponding 95% CI for the regression coefficient 0.12 (representing the difference between the mean deviation from target SE in the FLACS group and the mean deviation from target SE in the conventional group) was -0.21, 0.45. There was no difference between the groups with respect to the absolute value of the target corrected SE (Tables 2 and 3).

The refractive outcomes of groups in both the early and late postoperative periods were further divided into Good (defined as having a mean deviation of ≤ 0.5 D from target SE), Fair (defined as having a mean deviation of >0.5 D and ≤ 1.0 D from target SE) and Poor (defined as having a mean deviation of >1.0 D from target SE). In the FLACS group, the percentages to achieve a SE that was within ± 0.5 D from the intended target (Good refractive outcome) for the early and late postoperative periods were 56%

	Femtosecond	Conventional	Total	GEE p-value
N Eyes	50	88	138	
Preoperative Spherical Equivalent, Mean ± SD [range]	N/A	N/A	N/A	N/A
Preoperative logMAR BCVA Mean ± SD [range]	0.21 ± 0.16 [0, 1.0]	0.40 ± 0.29 [0, 1.6]	0.33 ± 0.27 [0, 1.6]	P<0.001
Preoperative Acuity Group, N (%) ≥ 20/30 20/30 – 60 20/70 – 5/200	31 (63%) 17 (35%) 1 (2%)	24 (27%) 44 (50%) 20 (23%)	55 (40%) 61 (45%) 21 (15%)	0.001
	Postoperative Spherical Eq	uivalent, target corrected (SE _{TC})		
SE _{TC} , Mean ± SD [range]	-0.10 ± 0.68 [-2.00, 2.25]	-0.15 ± 0.94 [-2.13, 2.75]	-0.14 ± 0.85 [-2.13, 2.75]	0.47
Absolute value SE _{TC} , Mean ± SD [range]	0.51 ± 0.46 [0. 2.25]	0.70 ± 0.64 [0, 2.75]	0.63 ± 0.59 [0, 2.75]	0.29
Outcome classification by deviation from target refraction, N (%)				0.11
Good (≤0.5D)	34 (68%)	51 (58%)	85 (62%)	
Fair (>0.5 to ≤1D)	11 (22%)	15 (17%)	26 (19%)	
Poor (>ID)	5 (10%)	22 (25%)	27 (20%)	

Table 3 Eyes with Late Postoperative Follow-Up (3-12 Months)

(n=39) and 68% (n=34), respectively (Tables 2 and 3). In the conventional group, the percentages to achieve a SE that was within ± 0.5 D from the intended target (Good refractive outcome) for the early and late postoperative periods were 65% (n=87) and 58% (n=51), respectively (Tables 2 and 3). There was no statistical significance in the mean deviation from target SE between the two groups in either the early postoperative period (p=0.78) or the late postoperative period (p=0.29). For all FECD eyes, in the early postoperative period, 62% (n=126) of eyes were found to be in the Good category, 22% (n=44) in the Fair category, and 17% (n=34) in Poor category (Table 2). In the late postoperative period, 62% (n=85) of eyes were found to be in the Good category, 19% (n=26) in the Fair category, and 20% (n=27) in the Poor category (Table 3).

There were no statistically significant differences in outcome classifications (Tables 2 and 3) between the FLACS and conventional groups at either early follow-up (p=0.24) or late follow-up (p=0.11). Of note, at late follow-up 15% more (total n=22) conventionally treated eyes were found to have poor outcomes than FLACS treated eyes.

Drift in Spherical Equivalent Refraction from Early-to-Late Postoperative Period

Comparison of outcome classifications within the two treatment groups reveals some shift in outcomes between early and late postoperative periods (Tables 2 and 3). However, these rates cannot be directly compared because the two follow-up periods included different subsets of eyes. Therefore, we examined the drift in SE from early-to-late period in the 47 FLACS and 76 conventionally treated eyes who contributed data for both follow-up periods. Figure 2 presents SE drift from early-to-late follow-up by treatment group. There was no statistically significant difference between the groups in drift categories (p=0.45, multinomial GEE); however, of note, 10 (13%) of eyes receiving conventional treatment experienced a change in SE of 2 or more diopters from early to late follow-up compared to none of the FLACS treated eyes. These large drifts did not always result in a worse outcome. Of these 10 eyes, 5 of 7 with a Poor early outcome achieved a Fair or Good outcome at late follow-up; nevertheless, 3 eyes with Good early follow-up outcome had Poor late follow-up outcomes.

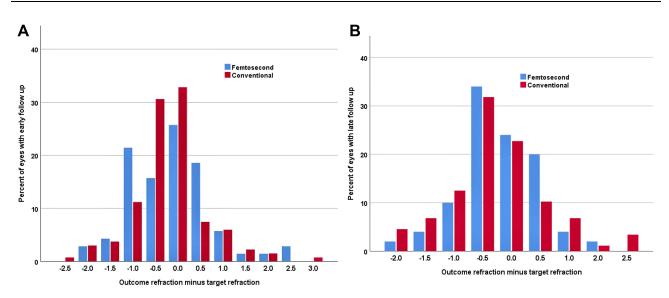


Figure I Spherical equivalent outcome by type of cataract surgery for early follow period (A) and late follow-up period 9 (B).

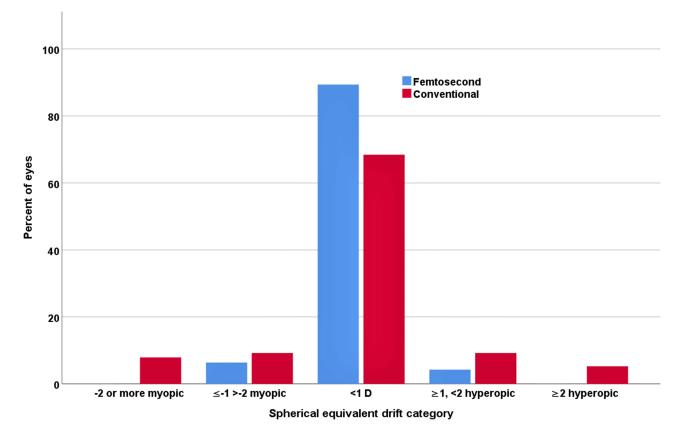


Figure 2 Spherical Equivalent drift from early-to-late follow-up period by cataract surgery type.

Risk Factors for Poor Refractive Outcome

The study population was quite heterogeneous with respect to preoperative ocular characteristics, some of which differed between the FLACS and conventional groups (Table 1). Therefore, we assessed the influence of these characteristics on Poor early and late spherical equivalent outcomes (Table 4). With the exception of preoperative cataract status, none of the putative risk factors was statistically significantly related to Poor outcome at either early or late follow up. The influence of cataract was

Table 4 Preoperative Risk Factors for Poor Outcome (>ID from Target Refraction)

	Follow Up Period	
	Early	Late
Number of Eyes	204	138
N (%) of eyes with poor outcome	34 (17%)	27 (20%)
Risk factor	Odds ratio	(95% CI) * [†]
Age (10 years older age)	0.97 (0.62, 1.51) P=0.88	0.75 (0.46, 1.22) P=0.24
Best-corrected visual acuity (0.3 worse logMAR acuity	1.17 (0.74, 1.85) P=0.50	1.46 (0.92, 2.31) P=0.11
CCT (if measured, 50um thicker)	1.09 (0.46, 2.59) P=0.84	0.84 (0.32, 2.18) P=0.84
Gender	P=0.52	P=0.89
Male	0.76 (0.33, 1.76)	0.94 (0.38, 2.34)
Female	Reference	Reference
Visual Acuity Group, N (%)	P=0.98	P=0.13
≥ 20/30	Reference	Reference
20/30 – 60	0.94 (0.44, 2.01)	1.23 (0.43, 3.52)
20/70 – 5/200	1.02 (0.31, 3.33)	3.32 (0.87, 11.3)
Central corneal measured	P=0.66	P=0.86
Yes	1.19 (0.55, 2.57)	1.08 (0.46, 2.56)
No	Reference	Reference
Corneal guttata grade N, (%)	P=0.97	P=0.84
0.5–1.0	Reference	Reference
1.5–2.0	1.16 (0.43, 3.15)	1.96 (0.44, 8.69)
2.5–3.0	0.95 (0.34, 2.61)	1.25 (0.32, 4.88)
3.5-4.0	0.92 (0.18, 4.78)	1.40 (0.44, 4.46)
Cataract grade N, (%)	P<0.001	0.90
0.5–1.0	Reference	Reference
1.5–2.0	1.32 (0.29, 6.05)	1.21 (0.25, 5.95)
2.5–3.0	0.89 (0.17, 4.69)	1.44 (0.26, 8.17)
3.5-4.0	5.73 (1.32, 24.9)	None present
Target refraction (D)	P=0.75	P=0.47
>-0.5	Reference	Reference
-0.75, -0.5	0.94 (0.34, 2.60)	0.45 (0.13, 1.60)
≤ −1.0	1.58 (0.46, 5.38)	(0.87 (0.17, 4.40)

Notes: *Univariate GEE, not adjusted for type of laser treatment; [†]Reference = Odds ratio reference category.

limited to grades >3 (3 eyes) which had odds of Poor refractive outcome 5.7 times greater than those of eyes with less severe cataract. None of these patients returned for examination in the later follow-up period.

The presence of postoperative corneal edema was examined in relation to the refractive outcome categorization. There was no statistical significance in the refractive outcome distribution when comparing absence versus presence of corneal edema in both the early (Table 5, p=0.15) and late postoperative periods (Table 6, p=0.27).

The fluctuation in SE from early to late period was found to be 0.02 ± 0.59 in the FLACS group versus -0.12 ± 1.20 in the conventional group. There was no statistical difference between the two groups (P=0.3). However, a wider range was observed in the conventional group [-3.50D to +3.25D] compared to the

Refractive Outcome	Absence of Postoperative Edema	Presence of Postoperative Edema
Good	61 (66%)	23 (51%)
Fair	14 (15%)	10 (22%)
Poor	18 (19%)	12 (27%)

Table 5 Refractive Outcomes and Postoperative Edema, EarlyPostoperative Period

Table 6 Refractive Outcomes and Postoperative Edema, LatePostoperative Period

Refractive Outcome	Absence of Postoperative Edema	Presence of Postoperative Edema		
Good	72 (63%)	12 (52%)		
Fair	20 (17%)	4 (17%)		
Poor	23 (20%)	7 (30%)		

FLACS group [-1.25D to +1.75D] (Table 7). The regression coefficient was found to be 0.18 with a hyperopic increase in FLACS at the late time-point.

Discussion

Fuchs' Endothelial Corneal Dystrophy (FECD) is a disease known to adversely affect the outcome of phacoemulsification surgery due to its direct risks of subsequent corneal decompensation, and authors have previously described strategic approaches to cataract surgery in this patient population.^{14–16,19} As modern phacoemulsification has become more advanced, studies have demonstrated that phacoemulsification can be safely done in FECD.^{14–16}

Refractive outcomes of FECD patients undergoing phacoemulsification remain sparsely documented, however. Fritz et al have shown via Scheimpflug image studies that the curvature of the cornea affects the refractive outcome in their study, due to the oblate posterior profile of FECD corneas.³ Others have proposed that in FECD corneas, the presence of stromal edema affects the refractive state. $^{\rm 20,21}$

In the past few decades, there have been advancements in phacoemulsification equipment and in surgical technique, and improved accuracy of biometric measurements and of intraocular lens (IOL) power formulas.^{22,23} Patient expectations have also risen with cataract surgery, and thus the demand for a desirable refractive outcome makes up a large part of patient expectations.^{23–25}

Femtosecond Laser Cataract Surgery

FLACS is one of the newest technological advancements in modern-day phacoemulsification surgery. The clinical advantages of FLACS over conventional phacoemulsification are continually being investigated and debated, although for many parameters, the technique seems to yield similar refractive and safety outcomes relative to conventional phacoemulsification.^{6,26–29} In studies comparing refractive outcomes, FLACS was shown to offer a more favorable outcome; this has been attributed to intraocular lens stability, and more predictable effective lens position (ELP) and capsulorrhexis size.^{30,31}

The study of FLACS outcomes in FECD patients has garnered increasing attention in the last few years, as this group of patients are particularly vulnerable to corneal decompensation with cataract surgery.¹⁴ Previous studies show mixed results in terms of clinically significant advantage in FECD when comparing rates of corneal edema and rates of corneal decompensation.^{32–35}

FECD Cornea and Refractive Challenges

In FECD patients undergoing cataract surgery, there is the additional challenge of predicting the refractive outcome in these eyes. In particular, a hyperopic trend has been described, and this is thought to be due to the posterior corneal curvature changes from edema,^{3,20} as well as the large change in refractive index at the anterior corneal surface.³⁶

As cataract surgery has evolved, so have patient expectations. Most patients desire a reasonably low residual

Table 7 Spherical Equivalence Change from Early to Late Period

	Femtosecond	Conventional	Total	GEE P-value
N SE change from early to late Mean ± SD [range]	47 0.02 ± 0.59 [-1.25, 1.75]	76 -0.12 ± 1.20 [-3.50, 3.25]	123 -0.06 ± 1.01 [-3.50, 3.25]	0.30

refractive error after cataract surgery. Based on the severity of the Fuchs' Dystrophy, an appropriate refractive target is determined by the surgeon; in particular, with more severe disease, mild residual myopia is targeted with the expectation that the patient may eventually undergo endothelial keratoplasty with an anticipated hyperopic shift.^{34,37,38}

When comparing the fluctuation of SE from the early to late period, there was no statistical difference between the two groups (P=0.3). Interestingly, a wider range was observed in the conventional group [-3.50 D to +3.25 D]compared to the FLACS group [-1.25 D to +1.75 D]. The wide fluctuation range from the early to late period in both groups is noteworthy—this may be clinically relevant to the surgeon during the early postoperative period when addressing a patient who is anxious or unhappy about the refractive results of cataract surgery. Our study suggests that it may be prudent to wait longer in order to assess the true refractive results in the FECD patient.

Limitations

The retrospective nature of this investigation means there is inherent bias. In addition, the procedures were performed by different surgeons with varying phacoemulsification techniques. There was also variability amongst femtosecond laser platforms and phacoemulsification systems. In addition, the CCT was not consistently recorded for every eye. While the Spherical Equivalent was our main refractive outcome measure, we did not include astigmatism or astigmatism-correcting treatments because the disease progression of FECD is known to create posterior corneal changes and induced astigmatism.³⁹ In addition, the retrospective nature of the study would mean that there would be variances in the surgeons' individual clinical practice preferences for astigmatism correction. Furthermore, in-depth tomographic study of posterior corneal changes as well as astigmatism in these eyes was not possible as tomographic imaging was not consistently performed. The limitation of examining the presence or absence of corneal edema was that we relied on the clinician's subjective documentation in the charts. Another limitation was the lack of IOL formula documentation by the surgeons. However, these are factors common to clinical retrospective studies like ours. A controlled prospective study would be warranted to further study these areas.

Summary

In this study, we sought to investigate the overall refractive outcomes in this population. Despite the afore-mentioned challenges presented by the FECD corneas, patients with mild-to-moderate FECD could still expect favorable refractive outcomes (the patients with severe classification were omitted per our exclusion criteria as these eyes either underwent concurrent keratoplasty or experienced corneal decompensation). We did not find statistically significant differences in outcome classifications (Tables 2 and 3) between the FLACS and conventional groups at either early follow-up (p=0.24) or late follow-up (p=0.11). We did not find a statistical significance when looking at the absence or presence of postoperative corneal edema and the effects on refractive outcomes.

Disclosure

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