



Factors affecting prediction error after cataract surgery with implantation of various multifocal IOLs in patients with previous refractive laser surgery

Young-Sik Yoo^{1,2#}, Min Chae Kang^{1#}, Jongyeop Park³, Hyung-Goo Kwon⁴, Eui-Sang Chung⁵, Dong Hui Lim^{1,6}, Tae-Young Chung¹

¹Department of Ophthalmology, Samsung Medical Center, Sungkyunkwan University School of Medicine, Seoul, South Korea; ²Department of Ophthalmology, College of Medicine, Uijeongbu St. Mary's Hospital, The Catholic University of Korea, Gyeonggi-do, South Korea; ³Department of Ophthalmology, Dongguk University Gyeongju Hospital, Dongguk University School of Medicine, Gyeongsangbuk-do, South Korea; ⁴Department of Ophthalmology, Keye Eye Center, Seoul, South Korea; ⁵Department of Ophthalmology, SNU Eye Center, Seoul, South Korea; ⁶Department of Health Sciences and Technology, Samsung Advanced Institute for Health Sciences and Technology, Sungkyunkwan University, Seoul, Republic of Korea

Contributions: (I) Conception and design: DH Lim, TY Chung; (II) Administrative support: DH Lim, TY Chung; (III) Provision of study materials or patients: TY Chung; (IV) Collection and assembly of data: J Park; (V) Data analysis and interpretation: HG Kwon, ES Chung; (VI) Manuscript writing: All authors; (VII) Final approval of manuscript: All authors.

[#]These authors contributed equally to this work.

Correspondence to: Tae-Young Chung, MD, PhD; Dong Hui Lim, MD, PhD. Department of Ophthalmology, Samsung Medical Center, Sungkyunkwan University School of Medicine, 81 Irwon-Ro, Gangnam-gu, Seoul 06351, Republic of Korea. Email: tychung@skku.edu; donghui.lim@samsung.com.

Background: This study aimed to compare the clinical outcomes of implantation of various multifocal intraocular lenses (mIOLs) and the prediction accuracy of two intraocular lens (IOL) power calculation formulas for eyes that underwent previous corneal refractive surgery.

Methods: Four types of mIOLs [TECNIS Symphony (Group I), AcrySof IQ PanOptix (Group II), LENTIS Mplus (Group III), and TECNIS ZLB00 (Group IV)] were used and the IOL power was calculated with the two no-history methods, Shammas-PL and Barrett True-K. Visual acuity and refractive outcomes including manifest refraction, prediction error (PE), absolute error (AE), and median absolute error (MedAE) were evaluated at three months after the cataract surgery.

Results: For all groups the Barrett True-K formula produced a narrower range of PEs and lower MedAE than Shammas-PL. Eyes of lower predictive accuracy (group B, AE >0.5D) showed weak uncorrected distance visual acuity resulting from myopic refractive error and target refraction when compared to that of higher predictive accuracy (group A, AE ≤0.5 D).

Conclusions: Targeting emmetropia using the Barrett True-K, which considers both anterior and posterior corneal curvature is recommended in patients undergoing mIOL implantation with prior corneal refractive surgery. Additionally, history of prior large amount of laser ablation seems to be an important factor related to low predictive accuracy.

Keywords: Multifocal intraocular lens; laser corneal surgery; intraocular lens implantation; cataract

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Introduction

With the popularization of refractive surgery, many patients have undergone refractive surgeries, and as the age of patients has increased, the number of cataract surgeries after refractive surgery has increased. There are many reports that intraocular lens (IOL) power calculation for eyes that have previously undergone refractive surgery is less accurate than that for virgin eyes (1-3). IOL power calculations for patients who have had corneal refractive surgery are clinically challenging because it is difficult to accurately predict corneal power and effective lens position (1,2). Many formulas are used to accurately calculate IOL power for eyes that have previously undergone corneal refractive surgery. When the refractive surgery parameters (pre-refractive surgery K-readings, amount of myopia corrected, or both) are known, many methods are available to correct the K-values and more accurately estimate the lens position (4,5). However, when the refractive surgery data is not available, the no-history method is a viable alternative for IOL power calculation after myopic refractive surgery (4).

In the absence of refractive surgery data, several formulas have been proposed to accurately calculate IOL power, including the Shammass post-LASIK (Shammass-PL) and Barrett True-K formulas (1,6). The Shammass-PL formula calculates IOL power based upon estimated postoperative anterior chamber depth (pACD), axial length (AL) and post-refractive surgery keratometry (1). The Barrett True-K formula (version 2.0) is based on the Barrett Universal II formula and is accessed online (7,8). Many studies have been published on monofocal IOL power calculation after refractive surgery, but to the best of our knowledge there have been few reports on multifocal IOL (mIOL) power calculation after refractive surgery. Although several studies have shown good refractive outcomes and visual acuity results, the use of mIOLs for eyes that have undergone corneal refractive laser surgery is controversial because of the assumption that the laser alteration of the cornea could cause vision to deteriorate after mIOL implantation (9,10).

To our knowledge, there are few studies that have evaluated refractive and visual outcomes and analyzed factors affecting prediction error (PE) for various mIOLs for patients who have had myopic refractive laser surgery. This retrospective study aimed to compare the clinical outcomes of four mIOLs and to evaluate predictive accuracy of two IOL power calculation formulas for patients that have previously undergone corneal refractive surgery. Additionally, preoperative characteristics that could lead

to low predictive accuracy of IOL power calculation were evaluated.

We present the following article in accordance with the STROBE reporting checklist (available at <https://dx.doi.org/10.21037/atm-21-3057>).

Methods

Subjects

This retrospective study included 131 eyes of 131 patients who underwent cataract surgery with mIOLs from January 2018 to July 2019 and had previously undergone myopic refractive surgery. Exclusion criteria were previous ocular surgeries except corneal refractive procedures such as LASIK or LASEK, corneal diseases, pseudoexfoliation, zonular weakness, corneal astigmatism greater than 1.00 diopter (D), glaucoma, macular disease, and amblyopia. Eyes with best-corrected distant vision less than 20/40 in the postoperative state were also excluded. The study was conducted in accordance with the Declaration of Helsinki (as revised in 2013). The Institutional Review Board (IRB #2020-09-079) for Human Studies at Samsung Medical Center approved this study and informed consent was exempted by IRB of Samsung Medical Center.

Surgical technique

All procedures were performed by one experienced surgeon (TYC) under topical anesthesia. Phacoemulsification was performed through a 2.75 mm temporal clear corneal incision. After performing continuous curvilinear capsulorhexis with intended diameter 5.0 mm and hydrodissection, phacoemulsification of the nucleus and bimanual aspiration of the residual cortex were performed using a cataract surgery phacoemulsification device (Centurion Vision System, Alcon, Fort Worth, TX, USA). The mIOL was implanted into the capsular bag using an injector and disposable cartridge system before removing the ophthalmic viscosurgical device (OVD). Four types of mIOLs were used: extended range of vision (Group I; TECNIS Symphony, ZXR00, Johnson & Johnson Vision Care, Inc. Santa Ana, CA, USA), quadrifocal (Group II; AcrySof IQ PanOptix, TNFT00, Alcon, Fort Worth, TX, USA), asymmetric zonal refractive bifocal (Group III; LENTIS Mplus, LS-313 MF30, LS-313 MF30, Oculentis GmbH, Berlin, Germany), and traditional diffractive bifocal (Group IV; TECNIS multifocal IOL, ZLB00, Johnson &

Johnson Vision Care, Inc., Santa Ana, CA, USA). Finally, a balanced salt solution was injected into the incision site to close the corneal incision, causing edema. After the surgery, postoperative antibiotic and corticosteroid eye drops were used four times daily and tapered over a month.

Patient examinations

Before cataract surgery, visual acuity was measured, including uncorrected distance visual acuity (UDVA) and uncorrected near visual acuity (UNVA). Manifest refraction (MR) testing was also conducted. To calculate mIOL power during cataract surgery, preoperative biometry of the eye was measured, namely the keratometry of the anterior surface, central corneal thickness, anterior chamber depth, and axial length, using a swept-source optical coherence tomography (SS-OCT) biometer (ARGOS, Suntec, Inc., Aichi, Japan). Keratometry of the posterior surface was measured with a Scheimpflug camera (Pentacam, Oculus, Wetzlar, Germany). Three months after the cataract surgery, visual acuity (UDVA and UNVA) and ocular refraction [MR and autorefractometry (AR) with autorefractor] were measured. UDVA and UCVA were measured at 4 m and 40 cm respectively.

Main measurement outcomes

Differences were compared between the predicted refractive value for each mIOL calculated by using different formulas and the refractive value measured three months after cataract surgery. The IOL power was calculated using two methods, the Shammas-PL formula, which is built into the biometer used, and the Barrett True-K formula (version 2.0, http://calc.apacrs.org/Barrett_True_K_Universal_2105/), which can be computed using a website. To calculate IOL power, the Shammas-PL uses only the anterior corneal curvature measured by the biometer, whereas the Barrett true-K formula uses both this curvature and the posterior corneal curvature. To calculate IOL power with both formula, preoperative biometry was used measured values from SS-OCT biometer except keratometry of the posterior surface. Posterior corneal curvature from Scheimpflug camera was used in the Barrett true-K formula. The IOL constants of each formula were as follows; In Shammas-PL [TECNIS Symphony (Group I); 118.8, PanOptix (Group II); 119.1, LENTIS Mplus (Group III); 118.5, TECNIS ZLB00 (Group IV); 118.8], in Barrett True-K formula [TECNIS Symphony (Group I); 119.39, PanOptix (Group

II); 119.1, LENTIS Mplus (Group III); 118.5, TECNIS ZLB00 (Group IV); 119.39].

Prediction error (PE) in refraction was defined as the predicted spherical equivalent (SE) from an IOL formula minus the actual postoperative SE, whereas the mean error (ME) was the mean of the PEs. Positive and negative signs of PE respectively represent myopic and hyperopic errors of postoperative refraction. The mean absolute error (MAE) and median absolute error (MedAE) were the mean and median of the absolute values of the MEs. We also calculated the percentage of eyes having MEs of ± 0.25 , ± 0.50 , and ± 1.00 D or less. Additionally, PE for each mIOLs were analyzed based on postoperative MR. After these refractive outcomes were calculated, the predictive accuracies of the two formulas were evaluated.

Statistical analysis

Visual acuity and refractive outcomes including MR, AR, and PE were compared between four mIOLs using the Kruskal-Wallis H test. The refractive outcome measures MAE and MedAE of the Shammas-PL and Barrett True-K formulas were compared for each of the four mIOL groups. Box plots were used to compare the PEs of the Shammas-PL and Barrett True-K formulas. Statistical analyses were performed using SPSS statistical software (version 23.0, SPSS, Inc., Chicago, IL, USA) and statistical significance was defined as $P < 0.05$.

Results

There was no difference in patient sex between the four groups (*Table 1*). Although the optical power of the implanted mIOLs was not different between the groups, group I showed younger age, flatter corneal curvature, and longer axial length than the other three groups (all $P < 0.05$). From the preoperative biometry of group I, it can be inferred that the extended depth of focus (EDOF) IOL was used in cataract patients with prior higher degrees of myopic laser correction.

Both postoperative UDVA and UNVA were improved in all groups compared with those before cataract surgery (all $P < 0.001$). Both postoperative mean UDVA and UNVA were logMAR 0.1 or less in groups II, III, and IV (*Table 2*). UDVA was better for groups II, III, and IV than for group I, and UNVA was better for groups II and III than for groups I and IV. Postoperative mean MR for four groups were -0.37 ± 0.64 , -0.14 ± 0.24 , -0.16 ± 0.26 , and -0.050 ± 0.44

Table 1 Subject demographics

Characteristics	Group I	Group II	Group III	Group IV	P value
Number	59	25	26	21	
Age	48.7±6.2 [§]	50.4±4.8	52.5±4.6	50.5±6.1	0.031 [†]
Female (%)	40 (67.8)	15 (60.0)	18 (69.2)	17 (81.0)	0.50 [‡]
Visual acuity (LogMAR)					
UDVA	0.56±0.37	0.29±0.20	0.25±0.33	0.37±0.30	0.23 [‡]
UNVA	0.40±0.26	0.34±0.28	0.41±0.21	0.32±0.24	0.45 [‡]
Manifest refraction (D)					
Spherical equivalent	-0.73±1.97	-0.49±0.92	-0.84±1.17	-0.75±1.36	0.56 [‡]
Refractive sphere	-0.43±2.02	-0.22±0.91	-0.35±0.49	-0.44±1.37	0.22 [‡]
Refractive astigmatism	-0.60±0.61	-0.53±0.71	-0.36±0.35	-0.61±0.59	0.48 [‡]
Keratometry (D)	39.92±2.47 [§]	41.30±1.65	41.66±2.16	40.61±2.52	0.010 [‡]
Central corneal thickness (μm)	483.0±41.4 [§]	514.5±33.2	482.4±42.4	479.8±34.0	0.007 [‡]
Anterior chamber depth (mm)	3.42±0.31	3.52±0.25	3.32±0.32	3.38±0.32	0.051 [‡]
Axial length (mm)	26.23±1.99 [§]	25.21±1.07	24.92±1.33	25.48±1.79	0.003 [‡]
IOL power (D)	20.11±4.64	20.28±3.52	19.98±2.07	21.12±2.78	0.46 [‡]

[†], P value by Kruskal-Wallis H test; [‡], P value by chi-squared test; [§], Group I showed statistically different values compared to each of groups II, III, and IV. UDVA, uncorrected distance visual acuity; UNVA, uncorrected near visual acuity; IOL, intraocular lens.

Table 2 Postoperative visual and refractive outcomes after cataract surgery in patients with prior refractive laser surgery

Outcome measures	Group I	Group II	Group III	Group IV	P value [†]
Number	59	25	26	21	
Visual acuity (LogMAR)					
UDVA	0.11±0.16 [‡]	0.076±0.14	0.043±0.10	0.072±0.10	0.047
UNVA	0.13±0.12 [§]	0.051±0.077	0.030±0.061	0.10±0.11 [§]	<0.001
Spherical equivalent (D)					
Manifest refraction	-0.37±0.64	-0.14±0.24	-0.16±0.26	-0.050±0.44	0.11
Auto refraction	-1.41±0.63 [¶]	-0.27±0.48	-1.50±0.45 [¶]	-0.51±0.49	<0.001

[†], P value by Kruskal-Wallis H test; [‡], Group I showed statistically different values compared to each of groups II, III, and IV; [§], Groups I and IV showed statistically different values compared to each of groups II and III; [¶], Groups I and III showed statistically different values compared to each of groups II and IV. UDVA, uncorrected distance visual acuity; UNVA, uncorrected near visual acuity.

D in groups I, II, III, and IV, respectively. There was no difference in SE based on MR between the four groups (P=0.106). Groups I and III showed myopic refractive error in mean AR compared to their mean MR. More than 60% of eyes in each group showed MR between -0.25 and +0.25 D (67.8%, 84.0%, 73.1%, and 66.7% in group I, II, III and

IV, respectively; *Figure 1*). More than 80% of eyes showed MR within 1 D in all groups (84.7%, 100.0%, 92.3%, and 95.2% in group I, II, III and IV, respectively).

Target refractions calculated from Barrett True-K showed no difference between all groups (P=0.355, *Table 3*). PEs based on Barrett True-K were closer to zero than those

based on Shammas-PL in groups I and II ($P=0.017$ and 0.007 , respectively; *Figure 2*). Although groups III and IV showed no difference in PE between Barrett True-K and Shammas-PL, mean PE values of Shammas-PL (0.050 ± 0.40 and -0.29 ± 0.65 D in groups III and IV, respectively) and Barrett True-K formulas (0.19 ± 0.22 and -0.012 ± 0.60 D in groups III and IV, respectively) were within 0.50 D of each other. For all groups, the Barrett True-K formula yielded lower MAE and MedAE than Shammas-PL. More eyes had MAE within 0.25 and 0.50 D when using the Barrett True-K formula (45.8%, 68.0%, 65.4%, 42.9% and 76.3%, 84.0%, 88.5%, 66.7% in group I, II, III, and IV, respectively) compared to Shammas-PL (23.8%, 44.0%, 50.0%, 38.1% and 72.9%, 84.0%, 76.9%, 66.7% in group I, II, III, and IV, respectively; *Figure 3*).

Postoperative mean UDVA in eyes with postoperative MR between 0.25 and -0.25 D were 0.053 ± 0.082 ,

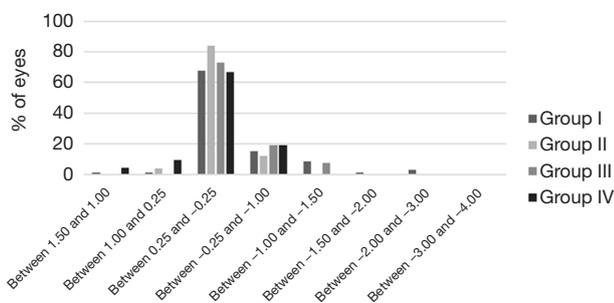


Figure 1 Postoperative manifest refractions of subjects implanted with various multifocal intraocular lenses.

0.044 ± 0.11 , 0.0072 ± 0.017 , and 0.058 ± 0.094 in group I, II, III, and IV, respectively (*Figure 4*). Although there was no difference in UDVA between four groups in case with postoperative MR between $+0.25$ and -0.25 D ($P=0.056$), UNVA at 40 cm was better in groups II and III than groups I and IV (all $P<0.001$). Group I achieved UNVA better than 0.1 logMAR in wider range of postoperative MRs than other groups, showing the EDOF properties. In subgroup analysis based on postoperative MR, Group II, III, and IV showed the best UDVA and UNVA in postoperative MR between $+0.25$ and -0.25 D.

Twenty-three of the 131 subjects (17.56%) showed an absolute value of PE (AE) of 0.5 D or more based on the calculation with Barrett True-K formula (*Table 4*); these subjects were grouped as Group B. Their UDVA was lower ($P<0.001$) and their postoperative refraction error was more myopic ($P=0.001$) than subjects having AE less than 0.5 D (Group A). Both SE and refractive sphere based on MR were closer to emmetropia for group A (-0.16 ± 0.36 and -0.0049 ± 0.38 D, respectively) than for group B [-0.58 ± 0.85 ($P=0.016$) and -0.40 ± 0.87 D ($P=0.028$), respectively]. Lower K-values ($P=0.002$) and longer axial length ($P=0.010$) were found in group B compared to group A. It is supposed that subjects who experienced more ablation during laser refractive surgery before cataract surgery fell into the higher-PE group B. Target refraction preoperatively calculated with Barrett-True-K was more myopic for group B than group A ($P<0.001$). However, target refraction calculated with Shammas-PL did not show any difference between

Table 3 Prediction accuracy of IOL power calculated using Shammas-PL and Barrett True-K formulas for four multifocal intraocular lenses

Prediction accuracy measures	Group I	Group II	Group III	Group IV	P value [†]
Number	59	25	26	21	
Target Refraction (D)					
Shammas-PL	-0.39 ± 0.34	0.067 ± 0.29	-0.13 ± 0.31	-0.34 ± 0.35	<0.001
Barrett True-K	-0.20 ± 0.65	-0.16 ± 0.37	0.027 ± 0.22	-0.062 ± 0.33	0.36
Absolute Error (D)					
Shammas-PL	0.44 ± 0.30	0.47 ± 0.33	0.32 ± 0.24	0.49 ± 0.50	0.28
Barrett True-K	0.41 ± 0.50	0.29 ± 0.19	0.24 ± 0.17	0.45 ± 0.39	0.16
MedAE (D)					
Shammas-PL	0.37	0.35	0.27	0.34	
Barrett True-K	0.28	0.28	0.22	0.29	

[†], P value by Kruskal-Wallis H test. MedAE, median absolute error.

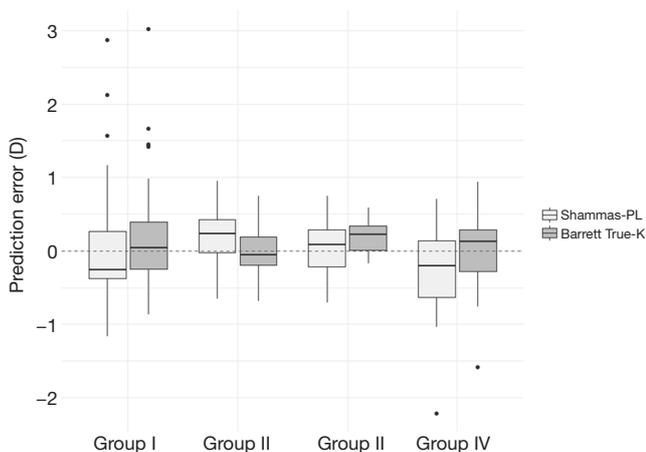


Figure 2 Prediction errors of Shammas-PL and Barrett True-K formulas [Prediction error = predicted spherical equivalent (SE) – postoperative SE].

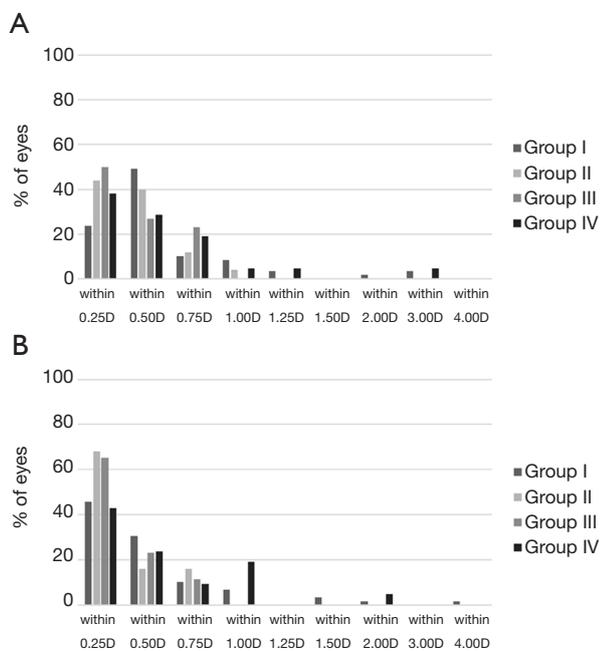


Figure 3 Mean absolute errors of (A) Shammas-PL and (B) Barrett True-K formulas.

group A (-0.22 ± 0.36 D) and group B (-0.36 ± 0.41 D, $P=0.090$). Regardless of which IOL formula was used, group B showed myopic refractive error compared to group A. For the high-AE group B, IOL power selection targeting myopic seems to have resulted in postoperative myopic PE after cataract surgery.

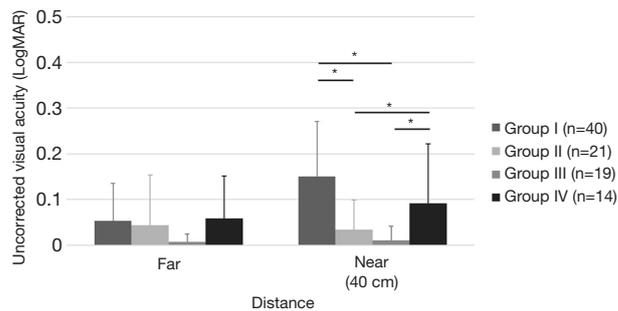


Figure 4 Postoperative LogMAR visual acuity of eyes having between 0.25 and -0.25 D of postoperative manifest refraction. Asterisks (*) indicate statistically significant differences ($P < 0.05$, Kruskal-Wallis H test).

Discussion

MIOL implantation yielded good distant and near visual acuity in post-refractive surgery eyes, and the emmetropia target is also recommended for good near and distance visual acuity. In calculating mIOL power after refractive surgery, the Barrett True-K formula, which considers posterior corneal curvature as well as anterior corneal curvature, was more accurate than the Shammas-PL formula. However, eyes with history of larger amount of laser ablation in corneal refractive surgery seems to show weaker predictive accuracy of IOL power calculation when using the Barrett True-K formula.

The predictive accuracy of the Barrett True-K formula was higher than that of the Shammas-PL formula, which is thought to be due to the fact that the Barrett True-K formula showed more hyperopic PE than Shammas-PL formula (6). Although conventional formulas such as Shammas-PL and Haigis-L calculate IOL power based upon preoperative keratometry measuring only the anterior curvature, supplemental factors have been added to these formulas to reduce PE for eyes with prior corneal refractive surgery (4,11). Specific methods for corneal power correction have been used with both Shammas-PL and Haigis-L to compensate for overestimated keratometry in eyes with prior corneal refractive surgery. However, the keratometry used in both formulas were not measured values, but expected values developed by using regression analysis (a no-history method). As a result, IOL power calculations with Shammas-PL and Haigis-L have shown more myopic PE in eyes with prior corneal refractive surgery than those preoperatively expected (7). Recently published meta-analysis concluded that the ASCRS average

Table 4 Factors related to absolute value of prediction error after cataract surgery in patients with prior refractive laser surgery. All subjects were divided into two groups: those having prediction error of 0.5 D or less (Group A) or prediction error more than 0.5 D (group B). Prediction error was calculated by using the Barrett True-K formula

Variables	Group A (AE ≤0.5 D)	Group B (AE >0.5 D)	P value
Number	108	23	
Postoperative values			
Visual acuity (LogMAR)			
UDVA	0.061±0.10	0.19±0.22	0.001 [†]
UNVA	0.090±0.11	0.10±0.12	0.65 [‡]
Manifest refraction (D)			
Spherical equivalent	-0.16±0.36	-0.58±0.85	0.016 [†]
Refractive sphere	-0.005±0.38	-0.40±0.87	0.028 [†]
Refractive astigmatism	-0.31±0.38	-0.35±0.32	0.37 [†]
Target refraction (D)			
Barrett true-K	-0.069±0.25	-0.36±1.05	<0.001 [†]
Preoperative values			
Age	50.1±5.8	50.0±5.8	0.70 [†]
Female (%)	40 (67.8)	15 (60.0)	0.92 [‡]
Keratometry (D)	40.96±2.27	39.13±2.30	0.002 [†]
Central corneal thickness (µm)	489.6±37.1	481.4±54.8	0.84 [†]
Anterior chamber depth (mm)	3.43±0.32	3.34±0.23	0.36 [†]
Axial length (mm)	25.46±1.66	26.59±1.97	0.010 [†]
IOL power (D)	20.3±3.4	20.3±5.1	0.23 [‡]

[†], P value by Man-Whitney test; [‡], P value by chi-squared test. AE, absolute value of prediction error; UDVA, uncorrected distance visual acuity; UNVA, uncorrected near visual acuity; IOL, intraocular lens.

based on ASCRS calculator (available at: <http://www.ascrs.org>) (6,12), Barrett True-K (7), or OCT formula (13) was recommended to calculate IOL power in eyes with prior laser refractive surgery for correcting myopia (14). Furthermore, the Barrett True-K (7), OCT (13), and optiwave refractive analysis formula (15) showed more accurate in prediction error than other formula including Haigis-L and Shammas formulas in eyes with previous myopic laser refractive surgery from Bayesian network meta-analysis (16).

The Barrett True-K formula more accurately calculates IOL power, representing the curvature of the entire cornea by including the posterior corneal curvature measurement (6,11,17,18). In previous studies reporting the results of cataract surgery after corneal refractive laser surgery, IOL power calculation using total keratometry which

considers both anterior and posterior corneal curvature showed better results than conventional calculation using only the anterior corneal curvature (19-24). Additionally, Shammas-PL based on total keratometry measured with an IOLMaster 700 biometer showed improved accuracy compared to that based on anterior keratometry in clinical studies by Lawless *et al.* and Yeo *et al.* (25,26). Recently, formulas using a ray-tracing method developed directly by the research institute were presented for more accurate IOL power calculation considering the anterior and posterior corneal curvature as well as the corneal thickness (20,27-29). Keratometry calculated from directly measured corneal curvatures of both the anterior and posterior surfaces would improve refractive outcomes due to the correction of myopic refractive error along with conventional IOL power calculations in eyes with prior

corneal refractive surgery.

In the present study, eyes with extreme biometry (mean K value of 39.13 D and mean AL of 26.59 mm) showed greater PE as calculated using Barrett True-K. IOL power calculation formulas for eyes with prior corneal refractive surgery are divided into history methods, which consider pre-refractive surgery data, and no-history methods, which are used when data from the refractive surgery are not available. Although there was no significant difference between the two methods (4,11,30), there has been no single formula showing high accuracy in various biometry conditions including eyes after corneal refractive surgery (31,32). Recently, Whang *et al.* reported in a retrospective study that the predictive accuracy of no-history IOL formulas depends on AL (33). Barrett True-K was most accurate for AL less than 28 mm, Triple-S was most accurate for AL between 28 and 30 mm, and Shammas-PL was most accurate for AL 30 mm or more. In the present study, mean preoperative AL was 25.84 ± 1.81 mm and only two eyes had a preoperative AL of more than 30 mm; thus, further evaluation of prediction accuracy in the case of extremely long eyes requires further study.

Eyes with high PE as calculated by Barrett True-K (group B in *Table 4*) showed preoperative characteristics of lower K-values and longer axial length. These biometry results correspond to eyes that experienced larger amount of ablation during refractive laser surgery. Additionally, both IOL power selection targeting myopic refraction and postoperative myopic refractive outcome resulted in PE being myopic. Previous studies with earlier formulas recommended targeting myopic due to hyperopic refractive surprise when calculating IOL power in eyes with previous corneal refractive surgery (32-36). Recent studies have reported that it is good to make the postoperative MR close to emmetropia in eyes having general corneal curvature, and the same results were reported for eyes which have undergone corneal refractive surgery (37,38). However, targeting slightly myopia may improve near vision without compromising distant vision, especially in high myopic patients, which is the case in the real world practice. Based on our result, we recommend aiming emmetropia when using Barrett True-K formula for implanting mIOL in eyes that have undergone corneal refractive surgery. Moreover, this study shows that predictive accuracy is more important than mIOL type for postoperative visual outcomes of patients with prior corneal refractive surgery.

To measure postoperative refractive power of the

eye, MR is recommended after mIOL implantation (39). Pseudophakia with a diffractive multifocal IOL (Tecnis ZM900 or ReSTOR) showed similar refraction between autorefractometry and MR measurements, including both spherical and astigmatic values (39,40). However, for eyes with a refractive multifocal IOL (ReZoom and Lentis LS-312 MF30), spherical values were found to be underestimated by autorefractometry compared to MR (39,41,42). Interestingly, in the present study both EDOF IOL (Tecnis Symphony) and rotationally asymmetric multifocal IOL (Lentis LS-312 MF30) also led to underestimation of spherical power by means of autorefractometry compared to MR. Postoperative refraction in patients with refractive or EDOF mIOL should be measured by MR and AR cannot be a replacement for MR.

However, there were some limitations in this study. IOL power calculation formulas with no-history methods were not compared to history methods in this retrospective study. Type of corneal refractive surgery was not identified due to a lack of information in some medical records. SS-OCT biometer used in the present study (ARGOS) could not be the same one with other biometer which other clinician uses (IOL Master 700; Carl Zeiss Meditec AG, Jena, Germany). As there is a difference in axial length measurement between two biometer, the results from this study cannot be applied to the cases with IOL Master 700. The adequate values for keratometry and axial length to estimate the lower predictive accuracy for postoperative refraction error could not be concluded as only two eyes had an axial length of >30 were included into the present study. Therefore, further prospective study based on a larger number with same type of mIOL is needed in order to improve the predictive accuracy of each IOL power calculation, especially in eyes that have undergone a large amount of corneal ablation.

In conclusion, MIOL implantation after corneal refractive laser surgery showed good refractive and visual outcomes. The Barrett True-K formula, which considers both anterior and posterior corneal curvature, was more accurate than the Shammas-PL formula, which considers only anterior corneal curvature. Targeting emmetropia is recommended when implanting mIOL in eyes that have undergone corneal refractive surgery. Especially in eyes that have undergone extensive corneal ablation, the previously large amount of laser ablation for cornea seems to be an important factor causing low predictive accuracy of IOL power calculation in cataract surgery.

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Footnote

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Ethical Statement: The authors are accountable for all aspects of the work in ensuring that questions related to the accuracy of integrity of any part of the work are appropriately investigated and resolved. The study was conducted in accordance with the Declaration of Helsinki (as revised in 2013). The Institutional Review Board (IRB #2020-09-079) for Human Studies at Samsung Medical Center approved this study and informed consent was exempted by IRB of Samsung Medical Center.

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