Background: To compare refractive outcomes after phacoemulsification and toric IOL implantation using two different toric calculators for initial astigmatism assessment in a real-world setting.

Methods: This was a retrospective, comparative, interventional case series. Patients over 30-year-old who underwent phacoemulsification and toric IOL implantation (AcrySof® Toric IOL) by the same surgeon between 2017 and 2018 were included. Eyes with irregular astigmatism, previous corneal refractive surgery, intraocular surgery, corneal pathology, macular pathology and pupil abnormalities were excluded. IOL toricity was determined by using a calculator provided by the AcrySof Toric calculator before 2018 and Barrett Toric Calculator after 2018. Patient demographics, corneal topography, vector and preoperative and postoperative refraction were collected and analyzed at three months postoperative.

Results: Thirty-two eyes of 32 patients were included in the final analysis. 0.1D for surgically induced astigmatism was used. Group 1 included 14 eyes assessed with the original (AcrySof) toric IOL calculator, and group 2 included 18 eyes assessed with the Barrett toric IOL calculator. In group 1, postoperative astigmatism less than −1.00D, −0.75 D, and −0.5D was achieved in 88.2%, 76.1% and 53.7% of eyes, respectively, while, in group 2, 89% eyes achieved postoperative residual astigmatism less than 0.5D and all eyes achieved postoperative residual astigmatism less than 0.75D. The proportion of patients with lower postoperative astigmatism was significantly higher in Group 2 (p< 0.05 by chi-square test), a pattern that still held when we divided patients into multiple groups. Vector analysis with the Alpins methods also supported better outcomes in the Barrett group (0.71 D vs 0.35 D).

Conclusion: The Barrett Toric calculator resulted in better results in the prediction of residual astigmatism than original (AcrySof) toric calculators.

Keywords: astigmatism, cataract surgery, toric intraocular lens, posterior corneal astigmatism

Background

In cataract patients with astigmatism, using toric IOLs in cataract surgery treats cataracts and astigmatism in one procedure and results in better postoperative visual quality. Nevertheless, they require additional effort, including preoperative measurement, planning, and modifications in intraoperative techniques. Previous studies have reported good visual and refractive outcomes, which are an important part of the standard of care for astigmatism correction with cataract surgery, especially in higher degrees of astigmatism. This procedure is also ideal for
patients with regular astigmatism who hope to reduce their dependence on glasses for distance vision.

Unfortunately, in real-world practice, this is not always the case, with “refractive surprise” still present from time to time. Variations in refractive outcomes of toric IOLs stem from several factors, including imperfections in anterior corneal surface measurement and alignment of toric IOL, corneal surgical-induced astigmatism (SIA), and disparities in the formulas used to calculate IOL power. Early toric calculators (eg, AcrySof Toric calculator) led to variable results, with only 50–75% of eyes exhibiting postoperative astigmatism of −0.50D or less. The Barrett Toric IOL calculator theoretically accounts for posterior corneal astigmatism (PCA) and effective lens position (ELP), which may lead to more accurate preoperative prediction of residual astigmatism. It is the first formula to provide non-inferior and reliable predictions based on measurements of posterior corneal curvature.

Therefore, we conducted a retrospective, comparative, interventional case series to compare refractive outcomes after standard 2.2 mm clear corneal phacoemulsification and toric IOL implantation using either the original (AcrySof) Toric calculator or Barrett Toric IOL calculator.

Methods

A retrospective, comparative, interventional case series was conducted. The study was approved by an institutional review board Institutional review board approval was obtained from the China Medical University Hospital. This study was conducted in compliance with Good Clinical Practices (GCPs) and was consistent with the 1996 version of the Declaration of Helsinki. Inclusion criteria were patients over 30-years-old with a follow-up period of more than three months. The subjects were diagnosed with cataracts but otherwise had good ocular health. Detailed preoperative evaluation including dilated indirect fundoscopy and OCT were performed in all patients to screen out those with retinal pathology. Patients also had regular corneal astigmatism between 1.00 D and 5.00D. Eyes with irregular astigmatism identified by corneal topography were excluded. To limit confounding variables, all eyes with previous corneal refractive surgery, intraocular surgery, corneal pathology, maculopathy and pupil abnormalities were excluded. Any subjects who experienced complications during surgery were also excluded. All eyes underwent standard 2.2 mm clear corneal phacoemulsification and toric IOL implantation (AcrySof® Toric IOL, Alcon Laboratories, Inc., Fort Worth, TX) between July, 2017 and June, 2018 by the same surgeon (Lin CJ). Automated keratometry, IOL Master 700 biometry (Carl Zeiss Meditec, Jena, Germany), and Pentacam (OCULUS, Wetzlar, Germany) were used as measurement methods in undilated eyes. Repeated measurements were performed and the value with repeatability was chosen. Also, patients with dry eye were pre-treated before biometry measuring.

The A standardized corneal refractive index 1.3375 was used. SIA magnitude value was set at 0.10 diopter (Table 1). The IOL toricity was determined with the AcrySof Toric calculator for patients treated before 2018 and Barrett Toric Calculator for patients treated after 2018. The online AcrySof–Barrett Toric Calculator was used to plan the axis of placement and cylinder power for the toric IOL with a printout of the toric calculator results made available at operation room.

The eye was marked with the patient in the upright position; this was performed with the patient fixating with the contralateral eye at a distant target at head height to avoid cyclotorsion from near-fixation under topical anesthesia, and the limbus was then marked at the desired angle of alignment using a needle and a surgical marking pen.

All operations were performed by the same surgeon using a two-plane, 2.2-mm incision placed just anterior to the limbus in the axis of the clear cornea. Following a divide-and-conquer nucleofractis and irrigation / aspiration for cortex removal, a toric IOL was implanted. The surgical wound was closed with stromal hydration. Levofloxacin 5mg/mL (Cravit, Santen Pharmaceutical Co., Osaka, Japan) and prednisolone acetate 1.0% eye drops were given 4 times daily for 2 weeks.

Patient demographics, corneal topography, and preoperative and postoperative refraction were collected before.

### Table 1 Centroid Surgically Induced Astigmatism Used in Barrett Toric Calculator

| Centroid | OS | OD |
|----------|----|----|
| **Incision Location** | **SIA (D)** | **Number** | **SIA (D)** | **Number** |
| Nasal | 0.17 | 3 | 0 | 0 |
| Superior | 0 | 0 | 0 | 0 |
| Superior temporal | 0 | 0 | 0.28 | 1 |
| Temporal | 0 | 0 | 0.17 | 6 |
| Inferior temporal | 0 | 0 | 0 | 0 |
| Inferior | 0 | 0 | 0 | 0 |
| Overall SIA | 0.17 | 3 | 0.14 | 7 |
and 3 months after surgery. The results from using the original (AcrySof) and Barrett Toric IOL calculator were compared.

**Vector Analysis**

The overall accuracy of astigmatism correction was calculated with the Alpins vector analysis method at 3 months after operation. The software included three astigmatism parameters: preoperative, target (set as 0), and achieved astigmatism. Four standard graphs were produced: 1) Target-Induced Astigmatism Vector (TIA), 2) Surgically Induced Astigmatism Vector (SIA), 3) Difference Vector (DV), and 4) Correction Index (CI). 17

**Statistical Analysis**

For statistical evaluation, SPSS for Windows (Version 13.0; SPSS Inc., Chicago, Illinois, USA) was used. Descriptive statistics on univariate data were performed in histogram. Chi-square test was used for residual astigmatism comparison between the two different toric calculators.

**Results**

In total, 32 eyes of 32 patients were collected. All operations were performed by the same surgeon. Group 1 included 14 eyes from 14 patients (4 males and 10 females) assessed with the original (AcrySof) Toric IOL calculator. The mean age was 65.2 ± 12.2 years (range, 37–80). Figure 1 showed the cumulative result of the two group. 14.3% of eyes achieved postoperative residual astigmatism less than −0.25D. 57.2% of eyes achieved less than −0.5D and 64.3% of eyes achieved less than −0.75D; only 78.6% of eyes had residual astigmatism less than −1.00D.

Group 2 included 18 eyes from 18 patients (7 males and 11 females) assessed with the Barrett Toric IOL calculator. The mean age was 67.6 ± 7.7 years (range, 53–84). 50% of eyes achieved postoperative residual astigmatism less than −0.25D. 88.9% of eyes achieved postoperative residual astigmatism less than −0.5D, and 100% of eyes achieved postoperative residual astigmatism less than −0.75D and −1.00D. (Figure 1).

The proportion of patients with lower postoperative astigmatism was significantly higher in Group 2 (p< 0.05 by chi-square test), a pattern than held when we divided patients into multiple groups (postoperative astigmatism cylinder ≥ −0.25D, −0.25D > cylinder ≥ −0.5D, −0.5D > cylinder ≥−0.75D, −0.75D > cylinder ≥ −1D) and into binary groups (postoperative astigmatism cylinder ≥ −0.5 D and < −0.5D; cylinder ≥ −0.25D and < −0.25D) (Figure 2).

Figures 3 and 4 show the results of the vectorial astigmatism analysis at 3 months of follow-up. Figure 3 displays the vector analysis in Group 1 (AcrySof Toric IOL calculator) while Figure 4 demonstrates the analysis in Group 2 (Barrett toric IOL calculator.) Mean TIA were similar in both groups (2.39 D vs 2.43 D). Mean DV, which represents the remaining astigmatism and provides

![Figure 1](cumulative.png) Cumulative percentage of postoperative residual astigmatism analysis using original and Barrett toric IOL calculator.
The proportion of patients with lower postoperative astigmatism is significantly higher in Group 2 (p< 0.05 by chi square test), no matter we divided patients in multiple groups (postoperative astigmatism cylinder ≥ −0.25D, −0.25D > cylinder ≥ −0.5D, −0.5D > cylinder ≥ −0.75D, −0.75D > cylinder ≥ −1D) or in binary groups (postoperative astigmatism cylinder≥ −0.5D and < −0.5D; cylinder ≥ −0.25D and < −0.25D).

Figure 2 The proportion of patients with lower postoperative astigmatism is significantly higher in Group 2 (p< 0.05 by chi square test), no matter we divided patients in multiple groups (postoperative astigmatism cylinder ≥ −0.25D, −0.25D > cylinder ≥ −0.5D, −0.5D > cylinder ≥ −0.75D, −0.75D > cylinder ≥ −1D) or in binary groups (postoperative astigmatism cylinder≥ −0.5D and < −0.5D; cylinder ≥ −0.25D and < −0.25D).
a summary of the astigmatic error, showed better results in the Barrett group (0.71 D vs 0.35 D). In CI, which demonstrates the under- (<1) or overcorrection (>1) of astigmatism treatment, revealed both these two calculators were slightly under-corrected (0.99 vs 0.93).

Discussion
With the recognition of the importance of posterior corneal astigmatism, new toric calculators were developed and have greatly helped with lowering post-operative astigmatism. In our study, the differences we found between the older calculator and the Barrett Toric calculator were larger than expected, with statistical significance reached in all parameters (p < 0.05 by chi-square test) (Figure 2). The old AcrySof calculator used simulated K values generated by a fixed cornea model of 500 µm thickness and anterior/posterior radius of 0.82,18 which was shown in our study to be less precise in reflecting the overall corneal astigmatism.

The posterior surface of most corneas is steep vertically, approximately 0.26–0.78D in previous studies, causing against-the-rule (ATR) refractive astigmatism.13,19 Previous generations of toric calculator (eg, AcrySof Toric calculator), which ignored the posterior corneal surface, would over-correct eyes with with-the-rule (WTR) astigmatism and under-correct eyes with ATR astigmatism.

Due to person-to-person variability, it is important to measure PCA individually so for a more precise astigmatism correction. This could be done by the incorporation of measured PCA from swept-source OCT such as with IOL Master 700 (Carl Zeiss Meditec, Jena, Germany), or topographic Scheimpflug analysis, such as with Pentacam (OCULUS, Wetzlar, Germany). However, whether measured PCA is significantly better than predicted PCA is still worth further investigation. Skrzypecki et al did not find any statistically significant difference in residual astigmatism when using the Barrett Toric Calculator incorporating either the measured PCA (with IOL master 700) or predicted PCA,20 but Kingrey et al found that 14% of cases ended up with a different toric IOL selection with the Barrett toric calculator.21

Our study showed comparable results with previous studies. Generally, prospective studies reveal relatively better
outcomes then retrospective studies due to its nature. But retrospective designs were much closer to daily clinical practice from some aspects. A recent large dataset came from a retrospective 823 cases series by Kane and Connell, revealing 88.2%, 76.1% and 53.7% of eyes had a post absolute prediction error less than −1.00 D, −0.75 D, and −0.50 D, respectively, using Barrett Toric calculator. In other prospective studies the results were slightly better, with 96.7 to 100%, 83.9% to 100% and 75 to 96% of eyes with a post absolute prediction error less than −1.00 D, −0.75 D, and −0.50 D, respectively. In our study, 100%, 100% and 88.9% of eyes had postoperative residual astigmatism less than −1.00 D, −0.75 D, and −0.50 D, respectively.

With the Alpins vector analysis method, our studies showed slightly larger values of under-correction in the Barrett calculator group (Correction index 0.99 vs 0.93). In further analyses, we found that there was one outlier value in this group causing the deviation. It was −0.75 D in before operation and −0.5 D after operation. Although the definite value was small, it accounted for a large proportion of the CI value.

The regression algorithms based on population mean values in the Barrett Toric calculator estimates posterior corneal astigmatism from anterior corneal astigmatism. As the result, the best possible anterior surface astigmatism measurements should be emphasized. We suggest the following clinical practice: first, perform at least two measurements with different devices (in our practice: IOL Master 700 and Pentacam). Then, repeat topography and ensure that all measurements closely match. Finally, re-evaluate the corneal surface if there are ambiguous results. We demonstrate that good preoperative measurements, combined with an advanced toric calculator such as the Barrett Toric calculator, are capable of achieving −0.50 D or less of residual refractive astigmatism 90% of the time in real-world clinical practice.

The limitations of this study include its retrospective nature and the small sample size. In Taiwan, patients pay extra fees of astigmatism-correcting intraocular lenses out of pocket. Nevertheless, overlooking the contribution of the posterior cornea might not make a huge difference in the outcome for the patients. However, selection of an IOL calculator that considers posterior corneal astigmatism and SIA may improve refractive outcomes.

Figure 4 Vector analysis by the Alpins method of astigmatism changes in Group 2 (Barrett toric IOL calculator). Abbreviations: Arith mean, arithmetic mean; Ax, Axis.
Conclusions
In conclusion, the Barrett Toric calculator, which includes consideration of PCA and SIA, was found to have lower median absolute errors in the prediction of residual astigmatism than the original toric calculator in our present study. The Barrett Toric calculator helps minimize postoperative variation after toric IOL implantation and a high percentage of eyes had a residual refractive astigmatism of −0.5D or less. Prospective randomized studies are necessary to further validate our findings.

Abbreviations
ATR, Against-the-rule; BCVA, Best-corrected visual acuity; ELP, Effective lens position; IOL, Intraocular lens; OCT, Optical coherence tomography; PCA, Posterior corneal astigmatism; SIA, Surgically induced astigmatism; WTR, With-the-rule.

Data Sharing Statement
The datasets used and/or analysed during the current study are available from the corresponding author on reasonable request.

Ethics Approval and Consent to Participate
The study protocol described here was approved by the Institutional Review Board of China Medical University Hospital and conformed to the tenets of the Declaration of Helsinki.

Consent for Publication
Owing to the retrospective design of the study and the use of deidentified patient information, the review board waived the need for written informed consent.

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