Image-guided system versus manual marking for toric intraocular lens alignment in cataract surgery

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Image-guided system versus manual marking for toric intraocular lens alignment in cataract surgery

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Purpose: To compare the accuracy of toric intraocular lens (IOL) alignment using the Verion Image-Guided System versus a conventional manual ink-marking procedure.

Setting: University Eye Clinic Maastricht, Maastricht, the Netherlands.

Design: Prospective randomized clinical trial.

Methods: Eyes with regular corneal astigmatism of at least 1.25 diopters (D) that required cataract surgery and toric IOL implantation (Acrysof SN6AT3-T9) were randomly assigned to the image-guided group or the manual-marking group. The primary outcome was the alignment of the toric IOL based on preoperative images and images taken immediately after surgery. Secondary outcome measures were residual astigmatism, uncorrected distance visual acuity (UDVA), and complications.

Results: The study enrolled 36 eyes (24 patients). The mean toric IOL misalignment was significantly less in the image-guided group than in the manual group 1 hour (1.3 degrees ± 1.6 [SD] versus 2.8 ± 1.8 degrees; \( P = .02 \)) and 3 months (1.7 ± 1.5 degrees versus 3.1 ± 2.1 degrees; \( P < .05 \)) postoperatively. The mean residual refractive cylinder was −0.36 ± 0.32 D and −0.47 ± 0.28 D in the image-guided group and manual group, respectively (\( P > .05 \)). The mean UDVA was 0.03 ± 0.10 logarithm of minimum angle of resolution (logMAR) and 0.04 ± 0.09 logMAR, respectively (both \( P > .05 \)). No intraoperative complications occurred during any surgery.

Conclusion: The IOL misalignment was significantly less with digital marking than with manual marking; this did not result in a better UDVA or lower residual refractive astigmatism.


In most patients who have cataract surgery, emmetropia can be achieved by correcting for refractive errors by implantation of a monofocal intraocular lens (IOL). However, approximately 20% to 30% of patients having cataract surgery have preexisting corneal astigmatism of 1.25 diopters (D) or more, which, when uncorrected during surgery, will result in spectacle dependency postoperatively. In patients without comorbidities and a desire to achieve postoperative freedom from spectacles for distance vision, the correction of astigmatism could be addressed at the time of surgery by using toric IOLs.

Numerous studies have shown that the implantation of a toric IOL is safe and effective. After toric IOL implantation, 70% to 85% of all patients are spectacle-independent for distance visual acuity. One factor that determines the effectiveness of the astigmatism correction is the accuracy of toric IOL alignment; every 5 degrees of misalignment will decrease the anticipated effect by 17%. Therefore, accurate alignment of the toric IOL is essential to achieve excellent postoperative outcomes and reduce residual refractive astigmatism.

Since the introduction of the toric IOL, many manual marking techniques have been used for the alignment of toric IOLS. However, with a recently introduced digital marking system (Verion Digital Marker M, Alcon Laboratories, Inc.) toric IOLs can be aligned without preoperative manual marking.

The purpose of this study was to compare the accuracy of a new digital marking technique with a conventional manual marking technique for toric IOL alignment.

PATIENTS AND METHODS

In this randomized controlled trial, patients with cataract and concomitant corneal astigmatism were recruited from the
Maastricht University Eye Clinic. All patients provided informed consent. The study was performed in compliance with the tenets of the Declaration of Helsinki and good clinical practice guidelines and was registered in a clinical trial register. Approval of an investigational review board was obtained before the start of this study.

The main inclusion criteria were unilateral or bilateral preexisting corneal astigmatism of 1.25 D or more and age of 18 years or older. Exclusion criteria were irregular corneal astigmatism, previous intraocular or corneal surgery, Fuchs endothelial dystrophy (stage 2 or higher), extensive age-related macular degeneration, glaucoma or diabetic macular disease, or other contraindications to cataract surgery and toric IOL implantation.

**Toric Intraocular Lens Selection and Calculation**

In all cases, an Acrysof aspheric toric IOL (model SN6AT3-9, Alcon Laboratories, Inc.) with cylindrical powers in 0.75 D steps ranging from +1.50 D (T3) to +6.00 D (T9) and spherical powers ranging from +6.00 to +30.00 D was implanted. To calculate the spherical power of the toric IOL, optical biometry (Carl Zeiss Meditec AG) and the Sanders-Retzlaff-Kraff formula (SRK/T, A-constant 118.9) were used. The cylindrical power of the toric IOL was determined by transferring into an online toric calculator the keratometry reading obtained by an optical biometer based on partial coherence interferometry (PCI) (IOLMaster, Carl Zeiss Meditec AG). The web-based toric calculator takes into account mandatory data input, such as the expected surgically induced astigmatism (SIA) and the position of the main corneal incision. The expected SIA was 0.30 D for a 2.2 mm superior incision.

**Surgical Technique**

All surgeries in this study were performed by 1 of 2 experienced surgeons (R.N., N.B.). The surgical technique, except the method of marking for toric IOL alignment, was standardized in each patient. In all cases, a standard divide-and-conquer phacoemulsification technique was performed through a superior 2.2 mm clear corneal incision.

The manual marking technique consisted of a 3-step procedure. After the eye was marked at 0 degree, 180 degrees, and 270 degrees using the Nuijt-Lane preoperative toric reference corneal marker (AE-27935, Asico LLC), with the patient seated, the desired implantation axis was marked intraoperatively using a Mendez ring and a Nuijt toric axis marker (AE-2740, Asico LLC). Implantation and alignment of the toric IOL were performed until the IOL marks were in line with the manually placed ink alignment marks.

The Verion Image-Guided System, which consists of a measurement module and digital marker, was used to perform digital marking for toric IOL alignment. Preoperatively, a high-resolution color reference image of the patient’s eye was obtained using the measurement module. These images were transferred to the digital marker. Using multiple reference points on the conjunctiva and limbus, a digital overlay of the imported preoperative image and live-surgery image was created. Because of the eye-tracking navigation of the system, cyclotorsion and eye movements are eliminated, allowing the desired implantation axis of the toric IOL to be accurately projected in the right ocular of the surgeon’s microscope.

**Randomization**

Consecutive patients were randomly assigned to have digital marking or manual marking. Blocked randomization was performed to reduce bias and achieve balance in the allocation of participants to both treatment arms. The assigned randomization was e-mailed to the surgeon 1 day before surgery. The patient and investigator performing the postoperative examinations were masked to the treatment allocation. At the 3-month follow-up, the randomization was revealed.

**Preoperative Assessment**

Preoperatively, each patient had a full ophthalmologic evaluation using slitlamp biomicroscopy, including funduscopy and intraocular pressure measurement. Automated keratometry (PCI-based and the image-guided system’s reference unit) and corneal topography (Pentacam, Oculus Optikgeräte GmbH) were performed before surgery.

**Determination of Misalignment and Rotation Stability**

At all postoperative visits (1 hour, 1 week, 1 month, and 3 months), slitlamp photography in full mydriasis (phenylephrine hydrochloride 100 mg/mL, tropicamide 5 mg/mL) was performed. All slitlamp photographs were examined by the same researcher (V.W.). Misalignment was defined as the difference between the desired toric IOL axis and the achieved axis 1 hour postoperatively. To eliminate cyclotorsion or an in-habitual head position, the postoperatively obtained slitlamp photographs were compared with preoperative images obtained with the image-guided system’s measurement unit; these images were obtained while the patient’s eyes were in a horizontal position. The preoperative image-guided system measurements were performed according to the manufacturer’s directions. These include having patients place their chin in the center of the chinrest and affirming constant contact of the forehead with the horizontal supporting band. In addition, head tilt was prevented during the measurements by ensuring that an imaginary line connecting the outer canthi was running parallel with the supporting band.

Reference spots on the conjunctiva and limbus were marked in the preoperative images and postoperative images using Photoshop CS6 (version 13.01.3, Adobe Systems, Inc.). After an overlay of both images was created by matching all reference spots, the achieved toric IOL axis was determined using its marks, thereby excluding changes in eye or head position.

A comparable technique was used to determine rotation stability postoperatively. Rotation was defined as the difference between the implanted toric IOL axis 1 hour postoperatively and the toric IOL axis at consecutive follow-up visits. The rotation of the toric IOLs was determined using Adobe Photoshop CS6 and the horizontal reference markings obtained from the preoperative image (as described above).

**Visual and Refractive Outcomes**

At each follow-up visit, monocular uncorrected distance visual acuity (UDVA) and corrected distance visual acuity (CDVA) were measured using Early Treatment Diabetic Retinopathy Study charts at 4 m. Patients were asked to start reading the charts at the smallest row on which all letters were easily distinguishable. Patients were asked to continue to read rows with smaller letters until no letters on a row were read correctly. The correct number of letters was noted and transferred into logarithm of minimum angle of resolution (logMAR) values for use in the analyses.

**Sample Size**

A sample-size calculation was performed using an α of 0.05 and a power of 80%. A previous study examining manual eye marking found a mean toric IOL misalignment of 5.0 degrees ± 2.1 (SD) in pseudophakic eyes. In a pilot study, the mean misalignment of the image-guided system was 2.5 degrees. Based on these assumptions and assuming a dropout of 15% on the primary outcome measure, the study would have to include a minimum of 36 eyes.

**Statistical Analysis**

All data were collected in an electronic data-capture system for medical research (Castor EDC, CIWIT B.V., Amsterdam) and transferred to SPSS for Windows software (version 23.0, 2010, International Business Machines Corp.) for data analysis. An intention-to-treat analysis was performed. Differences in
alignment between groups were analyzed using the independent Student t test. If the distribution of variables was not normal, the nonparametric Wilcoxon rank-sum test was used. Paired t tests were used to analyze the changes in visual acuities and residual refractive cylinder between preoperative visits and postoperative visits. Vector analysis according to the Alpin method was used to translate corneal and refractive astigmatism into Cartesian coordinates (x and y). In all tests, a threshold of statistical significance was assumed equal to a P value of 0.05.

RESULTS
Thirty-six eyes were randomly assigned to the manual group or the image-guided group (Figure 1). The image-guided group comprised 18 eyes of 17 patients (9 women [50%]) with a mean age of 68 years (range 23 to 89 years). The manual group comprised 18 eyes of 17 patients (6 women [33%]) with a mean age of 70 years (range 23 to 89 years). Preoperatively, there were no statistically significant differences in age, sex distribution, refractive, and anterior or posterior corneal astigmatism between the 2 groups (Table 1). A statistically significant difference was noted in preoperative CDVA favoring the image-guided group (Table 1). One patient in the image-guided group was lost to follow-up as a result of a systemic health disorder. Intraoperative digital marking failed in 1 patient, resulting in conversion to an intraoperative manual marking technique.

Misalignment
Table 2 shows the mean misalignment 1 hour postoperatively in both groups. Of the 36 anterior segment slitlamp photographs, 35 could be used for analysis of misalignment 1 hour postoperatively. Misalignment was significantly lower in the image-guided group than in the manual group (P = .02). The misalignment ranged from 0.0 to 6.1 degrees in the image-guided group and from 0.3 to 6.5 degrees in the manual group. Postoperative toric IOL misalignment of 5 degrees or less occurred in 17 patients (94%) in the image-guided group and 14 patients (81%) in the manual group. Counterclockwise rotation compared with the desired implantation axis was seen in 10 patients (57%) in the image-guided group and in 11 patients (64%) in the manual group.

Rotational Stability
Table 2 shows the mean rotation of the toric IOLs measured at consecutive follow-up visits. In 27 of 36 cases, anterior slitlamp photographs of good quality were obtained at all 4 follow-up visits. The mean rotation during the follow-up period (1 hour to 3 months) was comparable between the 2 groups (P = .25). In terms of rotation over time (1 hour to 3 months), a statistically significant decrease in rotation was measured (P = .04, image-guided group; P = .03, manual group); however, no statistically significant differences were seen between the 2 groups at any of follow-up timepoints (P = .13, P = .71, and P = .25 for 1 week, 1 month, and 3 months, respectively). The percentage of clockwise rotation during follow-up was 78% (14 patients) in the image-guided group and 82% (14 patients) in the manual group. Three months postoperatively, the mean error in alignment (misalignment and rotation combined) was significantly less in the image-guided group than in the manual group (P = .05).

Visual Acuity and Refraction
Three months postoperatively, the mean UDVA was 0.03 ± 0.10 logMAR in the image-guided group and 0.04 ± 0.09 logMAR in the manual group (P = .74). The mean CDVA was −0.05 ± 0.08 logMAR and −0.04 ± 0.07 logMAR, respectively (P = .82).

A statistically significant reduction between preoperative corneal astigmatism and postoperative refractive astigmatism occurred in both groups (both P < .0001) (Figure 2). The mean residual refractive astigmatism 3 months postoperatively was −0.36 ± 0.32 D in the

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**Figure 1.** Patient screening and follow-up.

- **93 Eyes (67 patients) assessed for eligibility**
- **57 eyes (43 patients) excluded**
  - 53 eyes (41 patients) did not meet inclusion criteria
  - 4 eyes (2 patients) refused to participate

- **36 Eyes (24 patients) randomized**
  - **18 Eyes randomized to manual marking**
  - **18 Received assigned intervention**
  - **0 Lost to follow-up**
  - **1 Unusable for primary outcome determination**

- **18 Eyes randomized to digital marking**
  - **17 Received assigned intervention**
  - **1 Lost to follow-up after primary outcome was assessed**

- **17 Eyes included for primary outcome analysis**
- **18 Eyes included for primary outcome analysis**
image-guided group, representing a reduction of 84% of the preexisting astigmatism magnitude, and $-0.47 \pm 0.28$ D in the manual group, representing a reduction of 81% of the preexisting astigmatism magnitude. Three months postoperatively, the mean SIA was $-0.27 \pm 0.44$ @ 113 in the image-guided group and $-0.21 \pm 0.51$ @ 121 in the manual group. The mean magnitude of SIA was $0.64 \pm 0.38$ D and $0.58 \pm 0.33$ D, respectively. Twenty-one patients (60%) had a magnitude of SIA greater than $0.50$ D.

Figure 3 shows the distribution of UDVA and residual refractive astigmatism in both groups 3 months postoperatively. There were no statistically significant differences in terms of UDVA, CDVA, or mean residual refractive astigmatism between the 2 groups at any time during the follow-up ($P > 0.05$).

### Vector Analysis

Table 3 shows the astigmatism parameters 3 months postoperatively obtained through vector analyses. No statistically significant differences were seen in the correction index, index of success, magnitude of error, or angle of error between the 2 groups ($P > 0.05$). The mean correction index in the image-guided group was slightly higher than in the manual group ($P = 0.18$) (Figure 4), representing a general overcorrection in both groups. In the image-guided group, 9 patients (53%) had an overcorrection of $0.25$ D or more and 5 patients (29%) had $0.50$ D or more. In the manual group, the overcorrection was $0.25$ D or more in 7 patients (39%) and $0.50$ D or more in 4 patients (22%). The success of astigmatism correction was 73% in the image-guided group and 78% in the manual group ($P > 0.05$).
Complications and Adverse Events

No complications occurred during any surgery. Postoperatively, a macular pucker was seen in 1 patient in the manual group. Neodymium:YAG laser capsulotomy was performed in 1 eye in both groups to treat posterior capsule opacification. No toric IOL had to be repositioned.

**DISCUSSION**

The purpose of this study was to compare the accuracy of a new digital marking technique with a commonly used manual marking procedure in the alignment of toric IOLs during cataract surgery. Using slitlamp photography, the misalignment 1 hour after surgery was determined. In our study, there was a significant difference in misalignment between the groups favoring the Verion image-guided system over the manual marking procedure. However, the difference in misalignment did not result in significant differences in UDVA or residual refractive astigmatism between the 2 groups 3 months postoperatively.

Sources of residual refractive astigmatism after toric IOL implantation include variations in SIA, the effects of posterior astigmatism, and misalignment of the toric IOL. A reduction of 3.3% in astigmatism correction for every degree of misalignment of the toric IOL axis to its desired axis reflects the importance of perfect intraoperative alignment and excellent postoperative rotation stability. Developments in toric IOL design and material and marking techniques for toric IOL alignment have resulted in improved outcomes after toric IOL implantation for the correction of preexisting corneal astigmatism.29–34

Various methods to mark the eye before toric IOL implantation have been described; of them, manual marking is considered the gold standard at present. In a study by Visser et al.,22 the use of a 3-step ink-marker procedure led to a mean error in alignment of 4.9 ± 2.1 degrees. Popp et al.24 compared several manual methods for marking the eye (slitlamp, pendulum, bubble-marker, and tonometer). The mean errors in toric IOL alignment ranged from 1.8 ± 2.2 degrees (pendulum) to 4.7 ± 2.9 degrees (tonometer), indicating that the pendulum method achieved the highest accuracy.

Several studies evaluated other approaches, such as eye mapping, using a corneal analyzer system and using a 3-dimension (3-D) imaging system. Cha et al.23 compared 3 methods; that is, marking the eye with a toric reference marker, using the slit beam, and using a mapping method.

**Table 3. Alpins vector analysis** of the effectiveness of astigmatism correction using toric IOLs.

<table>
<thead>
<tr>
<th>Parameter</th>
<th>Image-Guided Group</th>
<th>Manual Group</th>
</tr>
</thead>
<tbody>
<tr>
<td>Target induced astigmatism (D)</td>
<td>2.10 ± 0.80</td>
<td>2.24 ± 0.83</td>
</tr>
<tr>
<td>Total surgically induced astigmatism (D)</td>
<td>2.39 ± 0.90</td>
<td>2.42 ± 0.99</td>
</tr>
<tr>
<td>Difference vector (D)</td>
<td>0.51 ± 0.29</td>
<td>0.48 ± 0.27</td>
</tr>
<tr>
<td>Magnitude of error (D)</td>
<td>0.29 ± 0.40</td>
<td>0.18 ± 0.31</td>
</tr>
<tr>
<td>Angle of error (°)</td>
<td>−2 ± 16</td>
<td>−6 ± 14</td>
</tr>
<tr>
<td>Absolute angle of error (°)</td>
<td>11 ± 12</td>
<td>9 ± 11</td>
</tr>
<tr>
<td>Correction index*</td>
<td>1.16</td>
<td>1.07</td>
</tr>
<tr>
<td>Index of success*</td>
<td>0.27</td>
<td>0.22</td>
</tr>
</tbody>
</table>

*Mean only
The mapping method consisted of preoperatively obtained slitlamp photography to identify several reference vessel points. During surgery, these reference points were used to mark the desired toric IOL axis using calipers. To evaluate the accuracy of the 3 axis-marking methods, anterior segment photographs were taken. This new mapping method led to statistically significant smaller error in marking the eye than when conventional marking methods were used (2.3 ± 1.1 degrees versus 3.7 ± 1.5 degrees versus 3.1 ± 1.6 degrees) (all P < .05). Carey et al.21 analyzed postoperative misalignment after manual intraoperative marking using a 4-point procedure. After the eye was marked at 12 o’clock and 6 o’clock using a Codman surgical pen, the slit beam was rotated 90 degrees to mark the eye at 9 o’clock and 3 o’clock. The 3-month postoperative misalignment was measured using 2 techniques, a slitlamp approach and a corneal analyzer. Both methods showed a high correlation and comparable means of misalignment (2.7 ± 2.0 degrees versus 2.6 ± 2.8 degrees for corneal analyzer and slitlamp approach, respectively) (r = 0.99, P < .001). Montes de Oca et al.25 analyzed the accuracy of toric IOL axis alignment using a 3-D computer-guided visualization system compared with the accuracy of a manual marking procedure to mark the eye at 0 degree and 180 degrees. The digital method used a preoperative high-resolution photograph and intraoperative registration of the patient’s eye based on scleral and limbal vessels to allow digital surgical guidance and alignment of the toric IOL. The mean error of 3.0 ± 2.5 degrees and 2.9 ± 2.2 degrees in the digital group and manual group, respectively, was not significantly different between the 2 methods.

Table 4 shows an overview of the findings in previous studies, reflecting a wide range in misalignment between different marking methods. In the current study, significantly less misalignment immediately after surgery was seen with the image-guided system than with the manual marking procedure. The findings in the manual group are comparable to those seen in previous studies. However, the mean misalignment in the image-guided group was less than that reported in previous studies of manual marking, indicating the former is more accurate than other manual marking methods. However, there are limitations to comparing previously published studies21–25 and the current study. In many of these studies, postoperative misalignment was not measured immediately after surgery but rather several weeks later. It is well known that toric IOLs can rotate in the early postoperative period. Therefore, measuring misalignment after a few weeks can distort the true misalignment because of the possible rotation of the IOL. Furthermore, most studies used a slitlamp orientation to determine misalignment, while we created an overlay between preoperative images and postoperative images to eliminate cyclotorsion and differences in the patient’s head position.

At 3 months, 82% in the image-guided group and 72% in the manual group achieved a postoperative residual refractive cylinder of 0.50 D or less. This appears to be a higher proportion than in most recent studies, in which residual astigmatism of 0.50 D or less occurred in approximately 50% of cases.11,33 In our study, there were no statistically significant differences in terms of the mean UDVA and mean residual refractive astigmatism between the 2 groups. However, 3 months postoperatively the mean error in alignment (misalignment and rotation combined) was significantly less in the image-guided group than in the manual marking group (1.7 ± 1.5 degrees and 3.1 ± 2.1 degrees, respectively) (P = .05). These findings could be explained by different causes. First, although the difference in misalignment immediately after surgery between the 2 groups was significant, it was rather small (ie, 1.5 degrees). This difference in alignment remained stable during the 3 months of follow-up. At 3 months, the mean absolute difference between the intended axis and the achieved axis was 1.7 ± 1.5 degrees in the image-guided group and 3.1 ± 2.1 degrees in the manual group (P < .05). In the current study the mean preoperative corneal astigmatism was relatively low (2.42 ± 0.95 D and 2.29 ± 0.90 D, respectively). Therefore, the impact of a 1.5-degree difference in alignment error might not be clinically relevant in this population. However, this difference might become clinically relevant in patients with a higher level of preexisting corneal astigmatism. Second, in the current study, we used a first-generation toric calculator.9 This calculator does not take posterior corneal astigmatism and effective lens position (ELP) into account.

It is well known that posterior astigmatism plays an important role in the calculation of cylindrical power of a toric IOL.41–44 Because the posterior side of the cornea acts as a minus lens, neglecting this part of the cornea will result in overcorrection of the preexisting corneal astigmatism in a patient with with-the-rule astigmatism, whereas an undercorrection will occur in patients with against-the-rule astigmatism. In the current study, a mean

<table>
<thead>
<tr>
<th>Study*</th>
<th>Mean Misalignment (°)</th>
</tr>
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<tbody>
<tr>
<td>Visser22</td>
<td>4.9 ± 2.1</td>
</tr>
<tr>
<td>Popp24</td>
<td>2.3 ± 1.8</td>
</tr>
<tr>
<td>Sliitlamp Pendulum Bubble marker Tonometer</td>
<td>1.8 ± 2.2 2.9 ± 1.9 4.7 ± 2.9</td>
</tr>
<tr>
<td>Cha23</td>
<td>3.7 ± 1.5</td>
</tr>
<tr>
<td>Reference marker Sliitlamp Mapping method</td>
<td>3.1 ± 1.6 2.3 ± 1.1</td>
</tr>
<tr>
<td>Carey23</td>
<td>2.6 ± 2.8</td>
</tr>
<tr>
<td>Sliitlamp Corneal analyzer</td>
<td>2.7 ± 2.0</td>
</tr>
<tr>
<td>Montes de Oca25</td>
<td>2.9 ± 2.2</td>
</tr>
<tr>
<td>Reference marker 3-D imaging</td>
<td>3.0 ± 2.5</td>
</tr>
<tr>
<td>Current</td>
<td>2.8 ± 1.8</td>
</tr>
<tr>
<td>Reference marker Image-guided system</td>
<td>1.3 ± 1.6</td>
</tr>
</tbody>
</table>

*First author
overcorrection occurred in both groups (1.16 D and 1.07 D in image-guided group and manual group, respectively) \((P = .18)\). The higher amount of overcorrection in the image-guided group might be explained by the lower misalignment 3 months postoperatively. The cylindrical power of the toric IOL calculated by a toric calculator that uses input from anterior keratometry and neglects posterior astigmatism was on average 0.40 D and 0.41 D in the image-guided group and manual group, respectively, and will usually result in overcorrection. Because of the higher misalignment in the manual group, a reduction of the anticipated correction of the preexisting corneal astigmatism resulted in less overcorrection than in the image-guided group. Therefore, it is essential in the near future to implement second-generation toric calculators that take posterior astigmatism and ELP into account to prevent residual refractive astigmatism in cases in which the toric IOL is or nearly is perfectly aligned.38,42

In conclusion, both the digital and manual marking methods showed high accuracy in aligning toric IOLs intraoperatively. Although this study did not show significant advantages in terms of UDVA and residual refractive astigmatism using the digital marking system, IOL misalignment was significantly less in this group than in the manual marking procedure. Further studies will be required to determine whether the current difference might be more clinically relevant in a patient population with higher levels of preexisting corneal astigmatism. Future studies should also assess the effect on residual refractive astigmatism of updated toric calculators that account for posterior refractive astigmatism in combination with digital marking technology.

WHAT WAS KNOWN
• Adequate alignment of toric IOLs is essential to achieve excellent postoperative results.

WHAT THIS PAPER ADDS
• The new digital approach that uses eye-tracking navigation to eliminate cyclotorsion and eye movements reduced misalignment immediately after surgery by 50% compared with a commonly used 3-step manual marking procedure.

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C. Webers VSC, Bauer NJC, Visser N, Berendschot TTJM, van den Biggelaar FJHM, Nuijts RMMA (unpublished data)

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