Combined Descemet-stripping automated endothelial keratoplasty and phacoemulsification with toric intraocular lens implantation for treatment of failed penetrating keratoplasty with high regular astigmatism

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We present the case of a 57-year-old woman who had combined Descemet-stripping automated endothelial keratoplasty (DSAEK) and phacoemulsification with implantation of a toric intraocular lens (IOL). Surgery was intended to treat a cataract developing in a post-penetrating keratoplasty (PKP) eye with high astigmatism and endothelial decompensation. Six months after uneventful surgery, the cornea was clear and the corrected distance visual acuity was 20/20 with a refraction of +0.25 –1.00 × 10 (from –3.00 –8.50 × 12 preoperatively). The internal topography map (OPD-Scan) showed an IOL rotation of 4 degrees. The endothelial cell loss was 15% of the eye-bank value. Descemet-stripping automated endothelial keratoplasty combined with phacoemulsification and toric IOL implantation is a relatively simple and very effective procedure for eyes with endothelial failure and high post-PKP astigmatism. The speed of visual rehabilitation and final visual acuity achieved with this approach was superior to that obtained with other surgical procedures.

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Currently, Descemet-stripping automated endothelial keratoplasty (DSAEK) is the surgical procedure of choice for the treatment of endothelial failure after penetrating keratoplasty (PKP).1,5 However, if high astigmatism is present as a result of the PKP, it remains substantially unaffected by DSAEK and visual acuity is unlikely to improve postoperatively despite the recovery of corneal clarity.6,7 Recently, moderate or high astigmatism resulting from PKP has been successfully corrected by implantation of a toric intraocular lens (IOL) alone (phakic IOL) or in combination with phacoemulsification.8,9 We report the outcome of a combined procedure of DSAEK and phacoemulsification with implantation of a hydrophilic acrylic toric IOL to treat high regular corneal astigmatism in a patient with cataract in a failed PKP.

CASE REPORT

A 57-year-old woman presented to our institution complaining of a progressive decrease in visual acuity in her left eye. Penetrating keratoplasty for keratoconus had been performed in that eye in 1995. Since then, the acuity had been affected by anisometropia and high regular astigmatism, which could be only partly corrected by spectacles. The patient was intolerant to hard contact lenses.

On presentation, the uncorrected distance visual acuity (UDVA) and corrected distance visual acuity (CDVA) were 20/40 and 20/20, respectively, with a manifest refraction of –3.00 –8.50 × 12. Slitlamp examination showed a full-thickness graft (8.5 mm in diameter) with mild edema and a dense nuclear cataract (Figure 1, A). Endothelial cell...
density could not be evaluated as the noncontact specular microscope (Topcon SP200F, Topcon Corp.) could not obtain a useful picture of the endothelial surface. Corneal topography (OPD-Scan, Nidek Co. Ltd) showed high astigmatism with a regular asymmetrical bowtie pattern (Figure 1, B); keratometry values (K1/K2) were 48.91/40.55 (102).

Before surgery, the patient was asked to sit in the slit lamp. After the beam was set horizontally, a sterile ink pencil was used to mark the 0 and 180-degree positions, thus compensating for cyclorotation. Local anesthesia was administered with a peribulbar injection of a mixture of lidocaine hydrochloride 2.0% and hyaluronan hydrochloride 0.5%. Standard photomicrofiltration was performed through a 2.2 mm clear corneal tunnel centered on the steepest corneal axis previously marked using a Meizer ring.

The customized hydrophilic acrylic bitoric IOL (ATTOR-BI 70PMA-Ak, Contour 64E TLC, Carl Zeiss Meditec AG) was manufactured based on the biometric data obtained by partial coherence interferometry (PCI) (IOLMaster, Carl Zeiss Meditec AG). The IOL power was +12.5 +8.0, which included an intended overcorrection of 0.75 D to compensate for the mild hyperopic shift usually seen after DSAREK surgery. The IOL was implanted in the capsular bag and then rotated until proper alignment with the planned axis was achieved. The pupil was constructed with acetylcholine chloride (Möc-E), and the clear corneal tunnel was widened to 4.0 mm. The anterior chamber was filled with air, and the Descemet membrane and endothelium were peeled off en masse. At this point, a small peripheral iridectomy was performed at the 6 o'clock position with a vitrectome to prevent the possible onset of pupillary block.

The donor lenticule, prepared using the automated lamellar therapeutic keratoplasty system (Moria), was then placed on a Barron punch (Keravision Products, Inc.), endothelial side up, and trimmed to 9.0 mm in diameter. An anterior chamber maintainer was placed at the 12 o'clock position for continuous irrigation, and the donor tissue was inserted into the anterior chamber with the pull-through technique using the Busin glide (Moria). Both the clear corneal tunnel and the side entries were sutured tightly with interrupted 10-0 nylon sutures, and the anterior chamber was filled with air injected through the temporal side entry. Triamcinolone acetonide and gentamicin were injected subconjunctivally at the end of the procedure. The patient was pressure-patched overnight and instructed to lie supine for 6 to 8 hours. Postoperatively, dexamethasone (0.1%) and tobramycin antibiotic eye drops were administered initially every 2 hours and then tapered over 3 to 4 months. All sutures were removed 3 weeks after surgery. No intraoperative or postoperative complications occurred.

By 1 month postoperatively, the UDVA was 0.30 and the CDVA was 0.25 with a refraction of +0.75 -1.50 × 10. At the last follow-up, 6 months postoperatively, the corneal graft was transparent and the central endothelial cell density, measured with a noncontact specular microscope, was 2210 cells/mm², representing a 15% cell loss from the initial eye bank count (2600 cells/mm²). The CDVA had improved to 0.25 with a minimal change in refraction (+0.25 ± 1.00 × 10) compared with the 1-month measurement. The keratometry readings had not changed substantially from the preoperative values. The alignment of the toric IOL was evaluated by slitlamp examination and further confirmed by the wavefront analysis and internal refractive map obtained with the topography system that showed a postoperative IOL rotation of 4 degrees from the intended axis (Figure 2).
DISCUSSION

Several authors have recently reported that a triple procedure combining DSAEK, phacoemulsification, and IOL implantation allows rapid visual rehabilitation with minimal surgically induced astigmatism and/or ametropia. However, performing this triple procedure in an eye with a full-thickness edematous graft has the potential limitation of not correcting the high astigmatism that may be present in up to 20% of cases. In these cases, after the cornea has cleared, additional surgery is required to allow spectacle correction. Incisional surgery and/or excimer laser procedures (both photorefractive keratectomy and laser in situ keratomileusis) have been used in the past but have variable refractive results, i.e., undercorrection, overcorrection, and/or regression of effect over time, as well as relatively high rates of other complications, including perforation, wound gaping, infection, and loss of CDVA.

Another option would be to combine cataract surgery with a repeat PKP, but this approach would expose the patient to a renewed higher immunologic risk as well as other PKP-related possible complications. In addition, at least 1 year would be required before sutures could be removed and final refraction achieved.

Recently, phacoemulsification and implantation of a customized toric IOL have been suggested as a safe and effective procedure for the correction of high astigmatism resulting from PKP, a peripheral marginal degeneration, and keratoconus. Our case demonstrates that toric IOLs have an additional indication in eyes with full-thickness corneal grafts that are visually impaired due to the development of cataract as well as the presence of endothelial failure and high post-PKP astigmatism. In these eyes, a toric IOL can be implanted at the time of combined DSAEK and phacoemulsification, thus effectively treating all conditions negatively affecting vision with a single procedure. As with the procedures reported in the past, the toric IOL in our case exhibited minimal rotation from the intended position and good stability despite the additional maneuvers required during DSAEK, particularly the complete air fill of the anterior chamber and its consequent pronounced deepening.

A potential limitation of our approach concerns the calculation of the toric IOL. The accuracy of this calculation may have been negatively affected by 2 factors. First, the corneal edema may have prevented the PCI from measuring as precisely as in the presence of a clear cornea. Second, the 4.0 mm wide clear corneal tunnel that we routinely use for DSAEK may have caused an unpredictable change in corneal curvature in this post-PKP eye compared with the effect in an unoperated eye. Therefore, we decided not to consider the theoretical calculations of surgically induced astigmatism during the IOL calculation, as to date these have not been validated in the setting of PKP. However, in cases such as this, the aim of surgery is to obtain a debulking effect, that is to achieve a postoperative refractive error that can be corrected with spectacles, and the above-mentioned variables have had a negligible effect in this respect. Minor residual ametropia, including astigmatism, could then be addressed, if required, with more accurate procedures involving an excimer laser treatment.

In conclusion, this case describing implantation of a toric IOL at the time of combined cataract surgery and DSAEK in a post-PKP eye with high astigmatism shows that a 1-stage approach is at least as effective as the 2-stage or even 3-stage approaches. This procedure reduced postoperative recovery to a period of several weeks and allowed excellent final CDVA.

REFERENCES


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