The mechanism of intraocular pressure rise after keratoplasty

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ABSTRACT

Purpose: To determine the pathomechanism of increased intraocular pressure after penetrating and lamellar keratoplasty based on clinical observation and literature data.

Materials and methods: Morphometric analysis of the anterior segment of the eye using spectroscopic optical coherence tomography, gonioscopy and biomicroscopy.

Results: Pre- and postoperative spectroscopic optical coherence tomography scans in patients after keratoplasty reveal deformation of the peripheral cornea, narrowing of the iridocorneal angle, presence of anterior synechiae and various types of pupillary block.

Conclusions: The mechanism of elevated intraocular pressure after corneal grafting is complex and depends on pre-, intra- and postoperative factors, such as anatomy changes of the iridocorneal angle, presence of anterior synechiae as well as anterior and posterior pupillary block. Spectroscopic optical coherence tomography plays an important role in diagnosis and further treatment management.

Key words: keratoplasty, keratoplasty postoperative complications, secondary glaucoma
INTRODUCTION

Nowadays, keratoplasty is a well known ophthalmic surgery and the most common procedure in transplantology. It dates back to 1905, when dr Eduard Zirm performed the first successful penetrating keratoplasty (PKP) in patient with corneal lime burns [1]. In Poland number of performed keratoplasties is comparable with total number of all other transplantations [2]. Popularity of keratoplasty is based on its high efficacy consisting of immune privilege of cornea and anterior chamber as well as improvement in surgical techniques and tissue storage procedures. Nevertheless, keratoplasty bears a risk of postoperative complications, such as increased intraocular pressure (IOP) and glaucoma. Uncontrolled elevated IOP not only decreases eye function and graft survival but can also lead to blindness. The risk of transplant rejection is almost 3 times higher in patients with preoperative glaucoma than in patients without glaucoma [3].

Increased IOP after keratoplasty was described in 1969 for the first time [4]. Nowadays, it is observed after penetrating [5–10] as well as lamellar keratoplasty [11–13] and its frequency oscillates around 27–36%. Elevated IOP depends on pre-, intra- and postoperative factors. Among preoperative factors glaucoma plays a main role and can increase the risk of elevated IOP after Descemet’s stripping endothelial keratoplasty (DSEK) by 10% [11]. Intraoperative factors include not properly aspirated viscoelastic agent, using too small corneal grafts (under 7 mm) or suturing corneal grafts too „tight” [14]. Long-lasting steroid therapy as well as presence of blood or fibrin in iridocorneal angle can lead to elevated IOP postoperatively.

Increased IOP depends on type of keratoplasty and its indication. It occurs most often (60%) after PKP and DSEK [15], whereas considerably rarely after Descemet’s membrane endothelial keratoplasty (DMEK) (6.5%) [16]. Penetrating keratoplasties „à chaud” in keratitis or corneal perforation also carry a higher risk of elevated IOP [17].

Glaucoma neuropathy after keratoplasty is diagnosed mostly in the first postoperative months [17]. The lack of early diagnosis of glaucoma is due to inaccurate tonometry as a result of changed pachymetry measures and corneal biomechanics. It is suggested that precise pachymetry relied on spectroscopic optical coherence tomography (OCT) seems to be a reliable method of iridocorneal angle and corneal thickness assessment in patients after keratoplasty.

The paper presents mechanisms of increased IOP during early period after keratoplasty, based on clinical evaluation of the eye morphology as well as literature review. There are three types of morphometrical changes observed within the anterior segment of the eye that seem to be a reason of increased IOP after keratoplasty, e.g. narrowing of the iridocorneal angle, presence of peripheral anterior synechiae and presence of pupillary block.

NARROWING OF THE IRIDOCORNEAL ANGLE AS A CAUSE OF INCREASED INTRAOCULAR PRESSURE AFTER KERATOPLASTY

There are some factors that influence postoperative width of the iridocorneal angle, e.g. diameter of the recipient’s cornea, size of the corneal graft and graft suturing method [13]. Based on the AS-OCT imaging the following parameters can be measured to assess important morphometrical changes after keratoplasty: AOD500 (angle opening distance) – which is a distance between cornea and iris perpendicular to the cornea 500 μm from the scleral spur, TISA500 (trabecular iris space area) – trapezoidal area between cornea and iris 500 μm from the scleral spur into the anterior chamber and SSA (scleral spur angle) – representing iridocorneal angle width starting in the scleral spur [21].

Case reports

Case 1.

Penetrating keratoplasty combined with cataract surgery and IOL implantation in patient with bullous keratopathy, pachymetry > 1000 μm and mature cataract (fig. 1A). Before surgery iridocorneal angle was 41° wide in the OCT scans (fig. 1B). In the early postoperative period, correctly sutured corneal graft and narrowed iridocorneal angle (38°) was found (fig. 1C, 1D). There was no other potential cause of elevated IOP.
Case 2.
A patient with ocular pain and reduced visual acuity due to Fuchs’ corneal dystrophy (fig. 2A). Two days after endothelial keratoplasty (fig. 2B) elevated IOP ≥ 30 mmHg was observed. In AS-OCT narrowing of the iridocorneal angle from 54° preoperatively (fig. 2C) to 42° postoperatively was found. Additionally, bulging of the central and peripheral iris as well as air bubble in the anterior chamber were still present (fig. 2D).

Case 3.
A patient suffering from central corneal scarring and significantly reduced visual acuity, who underwent PKP with 8 mm corneal graft. During first postoperative weeks an increased IOP > 30 mmHg and narrow iridocorneal angle were observed (fig. 3A). Applied laser iridotomy widened iridocorneal angle sufficiently (fig. 3B), however the IOP stayed unchanged. Finally, a non-penetrating deep sclerectomy was performed (fig. 3C), which resulted in lowering the IOP to 19 mmHg without additional drug treatment.

Anterior synechiae as a cause of increased intraocular pressure after keratoplasty
Among numerous morphological changes of the iridocorneal angle, anterior synechiae seem to play an important role in pathomechanism of increased IOP after keratoplasty. They can be present in the anterior chamber before the surgery or appear as a consequence of postoperative complication [22]. It is suggested that anterior synechiae observed preoperatively should be treated as a routine surgical step. It has been proved that synechiolysis itself, the use of corneal graft 1 mm bigger than trephinated recipient's cornea, filling the anterior chamber with viscoelastic substance during surgery or performing oculopression before surgery can effectively reduce synechiae. Moreover, the use of steroid drops after keratoplasty reduces inflammatory response and efficiently prevents formation of anterior synechiae in early postoperative period [23].

Case reports

Case 1.
A patient 2 years after previously performed penetrating keratoplasty due to acute corneal hydrops complications. In the follow-up constantly elevated IOP ≥ 25 mmHg as well as intumescent cataract were observed (fig. 4A). OCT scans revealed presence of peripheral anterior synechiae in the huge area (fig. 4B). The patient was qualified to cataract surgery and IOL implantation combined with intraoperative mechanical synechiolysis (fig. 4C). The procedure resulted in significant widening of the iridocorneal angle, change of iris shape (fig. 4D) and gradual drop in IOP. Eventually the level of IOP of 18 mmHg was achieved that enabled withdrawing anti-glaucoma drugs.
Case 2.
A patient with diagnosed rheumatoid arthritis and glaucoma, who underwent PKP due to corneal perforation. One-year follow-up observation showed thinning of the corneal graft, descemetocele, shallowing of the anterior chamber, anterior synechiae formation and rise in IOP ≥ 25 mmHg (fig. 5A, B). Decision of repeated PKP combined with synechiolysis was made, which resulted in significant widening of the iridocorneal angle and consequently gradual decrease in the IOP to 15 mmHg (fig. 5C, D).
Pharmacological synechiolysis including mydriatics or miotics as well as oculopression is an advisable method before keratoplasty in some cases. When the surgery is finally planned it should be completed with intraoperative synechiolysis.

**PUPILLARY BLOCK AS A CAUSE OF INCREASED INTRAOCULAR PRESSURE AFTER KERATOPLASTY**

Pupillary block is commonly observed as a consequence of graft air tamponade after DSEK/DMEK [24]. Depending on location and volume, air bubble can induce: anterior pupillary block (when located in the anterior chamber and pushing iris backwards) or posterior pupillary block (when the air is dislocated behind the iris, pushing the iris anteriorly making peripheral iridocorneal adhesion and closing the iridocorneal angle).
Case reports

Case 1.
A patient after DSEK combined with cataract surgery and IOL implantation due to bullous keratopathy associated with Fuchs’ corneal dystrophy. Postoperatively a significant increase in IOP > 40 mmHg was observed and did not normalize after anti-glaucoma drops and intravenous osmotic agents. Slit lamp and OCT examination revealed air bubble located behind the iris, leading to iris bombe and posterior pupillary block (fig. 6A, B). Taking all these information into account, the patient was qualified to YAG-iridotomy, which resulted in iris flattening and fast lowering of the IOP < 20 mmHg. Interestingly, the procedure did not relocate the air bubble to the anterior chamber.

Case 2.
A patient, who underwent endothelial keratoplasty with basal iridectomy due to corneal edema (fig. 7A, B). Directly postoperatively elevated IOP > 35 mmHg was observed. Biomicroscopy and OCT scans presented a “mixed” type of pupillary block, in which one air bubble deepened the upper half of the anterior chamber leading to iridolenticular adhesion and anterior pupillary block (fig. 7C). Whereas the second air bubble dislocated behind the iris moved lower part of the iris anteriorly inducing posterior pupillary block (fig. 7D). Because YAG-iridotomy was not effective, the treatment included anti-glaucoma eye drops, mydriatics and intravenous osmotic agents for the next 2 days. Finally, spontaneous decrease in air bubble volume together with its relocation into the anterior chamber led to breaking the posterior pupillary block improving the iridocorneal angle anatomy and eventually lowering IOP < 20 mmHg.
It is worth mentioning, that DSEK/DMEK procedure requires iridotomy that should be performed preoperatively using NdYAG/argon laser on intraoperatively with scissors excision. It is well known that postoperative iridotomy is difficult to perform, as it has been shown in our cases. Other possible option is anterior chamber puncture with aspiration of some air volume, which can be a risky maneuver for corneal graft adhesion or anterior chamber stability.

OTHER CAUSES OF INCREASED INTRAOCULAR PRESSURE AFTER KERATOPLASTY

While the iridocorneal angle seems to be normal, the mechanism of elevated IOP after keratoplasty may be not so well defined. A meticulous examination of patient’s medical history is necessary to find any sign of glaucomatous neuropathy, method of previous treatment (conservative or surgical) or long lasting steroid therapy.

Case report

A patient after endothelial (DSEK) surgery due to corneal edema associated with Fuchs’ corneal dystrophy (fig. 8A). Postoperatively a significant increase in IOP > 30 mmHg was observed. Iridocorneal angle in gonioscopy and on OCT scans was wide and open (fig. 8B, C). However, patient’s history revealed long-term drug therapy of glaucoma with two medications. Reintroducing of previously used drugs did not normalize IOP and glaucoma progression was observed. A non-penetrating deep sclerectomy was performed (fig. 8D, E), which finally resulted in decrease in IOP < 20 mmHg preventing further glaucoma progression.
CONCLUSIONS
Keratoplasty is one of the most effective types of transplantation procedure with relatively good postoperative prognosis. Immunological graft rejection and increased IOP or glaucoma neuropathy are the most significant reasons of decreased visual functions and graft lost. Pathomechanism of elevated IOP is complex and includes pre-, intra- and postoperative factors. Moreover, the assessment of increased IOP in eyes after keratoplasty is burdened with error, mainly due to changed corneal biomechanics leading to inaccurate IOP measurements. To find out potential cause of increased IOP, gonioscopy and OCT with iridocorneal angle assessment are crucial, revealing deformation of peripheral cornea with narrowing of the iridocorneal angle, presence of anterior synechiae and pupillary block. It is worth mentioning that an increase in IOP is possible independently of above mentioned factors, therefore long-term observation after keratoplasty is recommended.

Figures: from the author’s own materials.

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References

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Piotr Jurowski: 25% – substantive consultation; Zofia Pniakowska: 25% – acquire of the photograph.

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