A Rare Case of Acute Bilateral Endothelial Decompensation after Prophylactic Nd:YAG Laser Iridotomy Requiring Endothelial Keratoplasty

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ABSTRACT

Aim: To describe a case of acute bilateral endothelial decompensation following prophylactic Nd:YAG laser iridotomy (LI) for occludable angles. Despite regarded safe, LI is occasionally a source of various ocular complications, including corneal endothelial damage. In the herein case, we describe the first case of acute bilateral endothelial decompensation after Nd:YAG LI.

Case description: A 63-year-old man was referred for consultation due to visual acuity deterioration in both eyes 2 weeks after undergoing an uneventful prophylactic LI for occludable angles. On examination, bilateral corneal edema with Descemet’s membrane folds was observed.ROB

Conclusion: Subacute endothelial dysfunction should be considered as a possible adverse event following Nd:YAG LI and patients should be advised accordingly.

Clinical relevance: Surgeons should be aware of the potentially devastating complication of bilateral corneal decompensation following routine Nd:YAG LI, even in patients without preexisting corneal injury. Patients should be advised accordingly.

Keywords: Angle-closure glaucoma, Corneal edema, Corneal transplant, Laser iridotomy, Nd:YAG laser.

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BACKGROUND

Laser iridotomy (LI) is a well-established procedure for the treatment and prevention of angle-closure glaucoma (ACG). Although both argon and neodymium-doped yttrium aluminum garnet (Nd:YAG) LI are regarded as being safe, they carry the risks of causing various ocular complications, including elevated intraocular pressure (IOP), iritis, hemorrhage, lens opacity, and corneal burns.1 Decreased endothelial density after LI has been reported in a few long-term studies.2 Late endothelial decompensation was documented in patients after argon LI, eventually necessitating corneal transplantation years after the procedure.2 To the best of our knowledge, acute endothelial decompensation after Nd:YAG LI has not been reported before.

CASE DESCRIPTION

A 63-year-old man was referred to our corneal unit 2 weeks after undergoing an uneventful prophylactic bilateral Nd:YAG LI for occludable angles. He complained of bilateral visual deterioration of decreased vision before the Nd:YAG LI procedure, and there was no personal or familial history of fluctuating vision, episodes of ocular pain, corneal dystrophy, or corneal transplantation.

He underwent an ophthalmic examination in our clinic, and it revealed a corrected VA of RE 20/60, LE 20/40 (glasses +7.50 D-0.75 D × 21 and +8.50 D–1.75 D × 92 right eye and left eye, respectively). The IOP was 12 mm Hg in both eyes. Slit-lamp examination revealed bilateral diffuse corneal edema with Descemet’s membrane folds and a few epithelial bullae. The corneal edema was worse in the right eye, and inferior peripheral iridotomies, round and reactive pupils, and mild nuclear sclerosis cataract. The fundus examination ruled out any definitive changes in the retina.

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Non-touch specular microscopy (EM-3000, Tomey, Japan) failed to measure endothelial density in both eyes. Anterior segment spectral-domain optical coherence tomography (OCT) (Heidelberg-Spectralis, version 6.9.4.0, Heidelberg Engineering, Germany) demonstrated an attached Descemet’s membrane with prominent folds in each eye (Fig. 2), and ultrasound biomicroscopy confirmed the presence of narrow angles. The axial length was 20.45 and 20.17 mm, the anterior chamber depth was 2.10 and 2.25 mm, and the central corneal thickness was 736 and 689 μm in the right and left eyes, respectively (Swept-Source OCT Biometer, IOLMaster 700, Carl Zeiss, Germany).

The patient was diagnosed as having acute corneal decompensation following Nd:YAG LI. He was treated with topical 0.1% dexamethasone and NaCl 5% ointment for 2 months without improvement and was subsequently referred for bilateral endothelial keratoplasty.

**DISCUSSION**

Peripheral LI is a well-established procedure for ACG and one that is widely performed worldwide. Potential cornea-related complications following LI are numerous, and they may include endothelial damage, which ultimately can lead to endothelial decompensation necessitating corneal transplantation. Several mechanisms of LI-related endothelial damage have been proposed, including direct corneal burn, thermal and shockwave injury, shear stress, iris pigment dispersion, and blood–aqueous barrier disruption. Although some risk factors predicting significant endothelial cell loss after LI have been hypothesized, no direct association has been found between any single factor and the development of corneal decompensation.

In the few studies that described endothelial decompensation after LI, the symptoms had appeared years after the procedure, and argon laser rather than Nd:YAG laser had been applied. Argon LI is achieved by means of a photoagulation mechanism, while Nd:YAG laser pierces the iris through photodisruption. Endothelial cell density was shown to decrease after both procedures, although to a less extent after NG:YAG laser. Part of the reason for that difference may be that penetration of the iris with the Nd:YAG laser generally requires fewer laser spots than an argon laser due to its high-power density.

In the current case report, our patient developed bilateral endothelial decompensation 2 weeks after a documented uneventful prophylactic Nd:YAG LI for occludable angles. This sequela is extremely unusual in light of the rapid progression of corneal decompensation, the application of an Nd:YAG laser rather than an argon laser, and the fact that the patient had no history of corneal pathology. The underlying etiology for the presented rare occurrence was not determined. Acute inflammation, a high IOP, or pigment dispersion were not likely to contribute to the corneal damage since the IOP was normal at presentation as well as after the procedure, the anterior chamber was clear, and no pigment deposition was observed on the endothelium. Direct corneal damage was also ruled out since none was documented following the LI, no signs of damage were observed on presentation, and because the most injured area did not correspond to the LI location. Lastly, the absence of corneal dystrophy findings, as well as the lack of prior visual symptoms, indicates that it was highly unlikely for preexisting corneal dystrophy to contribute to the acute corneal decompensation. Although we cannot rule out non-guttate Fuchs endothelial corneal dystrophy (FED) in our patient, the absence of any clinical signs or symptoms before and following the LI and the...
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lack of predominant central corneal edema makes FED an unlikely diagnosis.

Recent studies have questioned the suitability of LI as a first-line treatment for primary ACG (PACG) in a patient with normal IOP and suggested that cataract extraction with IOL implantation may serve as a superior alternative.8,9 While LI solely resolves the relative pupillary block component in ACG, phacoemulsification with IOL implantation deepens the anterior chamber by a posterior repositioning of the ciliary processes due to a debulking effect, resulting in widening of the anterior chamber angle.10 It has also been suggested that cataract extraction may have lower failure rates in lowering IOP in the long-term.2 The recent randomized controlled EAGLE study concluded that clear lens extraction had better efficacy and was more cost-effective than LI for PACG.8,9 Another possible reason for favoring phacoemulsification in PACG patients is the effect of the procedures on the corneal endothelium, with several studies have demonstrated a lower rate of endothelial cell loss after phacoemulsification.2 Although larger trials are still needed to better define the comparative effect of these two interventions on corneal endothelium health, performing phacoemulsification rather than LI in our patient may have prevented the devastating corneal injury.

CONCLUSION

Laser iridotomy can impair VA due to rapid corneal endothelial decompensation, in our case without a detectable etiology. To the best of our knowledge, this is the first to report acute bilateral corneal decompensation following Nd:YAG LI necessitating keratoplasty. Although more cases are needed to reinforce this observation, physicians should be aware of this potential sight-threatening complication, and patients should be informed accordingly.

CLINICAL SIGNIFICANCE

Although considered a simple and straightforward procedure, Nd:YAG LI can be a cause of various ocular complications. In this context, surgeons should be aware of the rare and potentially devastating complication of bilateral corneal decompensation following routine Nd:YAG LI, necessitating keratoplasty, even in patients without preexisting corneal injury, and advise their patients accordingly.

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