Neodymium:yttrium-aluminum garnet capsulotomy rates after combined cataract surgery with implantation of a 4% water content hydrophobic acrylic intraocular lens and vitrectomy

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Introduction
X-70 (Eternity®, Santen Co, Ltd, Osaka, Japan) is a new three-piece, 4% water content, hydrophobic acrylic intraocular lens composed of hydroxylethyl methacrylate, polyethylene glycol phenyl ether acrylate, and styrene cross-linked with ethylene glycol dimethacrylate.1 X-70 has a square-edge design with an optic diameter of 7.0 mm and an overall diameter of 13.2 mm (Figure 1). The large optic diameter provides stability and visibility during vitreoretinal surgery. The X-70 is often used for combined cataract surgery and vitrectomy in Japan.2 In patients with diabetes mellitus, visualization and evaluation of the postoperative fundus is important in addition to visibility during surgery because of possible photocoagulation treatment requirements and reoperations.3
Posterior capsule opacification (PCO) is a common complication after cataract extraction with intraocular lens implantation, which can result in decreased fundus visualization postoperatively. Previous reports have shown that PCO rates can greatly vary depending on the intraocular lens material, and PCO rates after combined cataract surgery and vitrectomy are considered higher than those after cataract surgery alone.

Neodymium:yttrium-aluminum garnet (Nd:YAG) capsulotomy is an effective treatment for PCO. However, this therapy has rare but significant complications, so informed consent related to PCO is required prior to surgery.

We assumed that an investigation into Nd:YAG capsulotomy rates following implantation of the new acrylic intraocular lens, particularly in patients with diabetic retinopathy, would provide useful and important information. Thus, we conducted the current study to compare Nd:YAG capsulotomy rates in patients with diabetic and nondiabetic retinopathy after combined cataract surgery with implantation of the X-70 and vitrectomy.

Patients and methods

After obtaining institutional board approval, 153 eyes of 153 patients who underwent combined cataract surgery and vitrectomy at the Jikei University of Medicine Katsushika Medical Center between January 2009 and March 2011 were retrospectively reviewed. All patients were followed up for at least 2 years after surgery. Patients were divided into two groups, ie, a diabetic retinopathy group including 72 eyes of 72 patients and a nondiabetic retinopathy group including 81 eyes of 81 patients. Indications for surgery in the diabetic retinopathy group were proliferative diabetic retinopathy and those in the nondiabetic retinopathy group were retinal detachment (n=50), macular hole (n=13), epiretinal membrane (n=9), branch retinal vein occlusion (n=7), and uveitis (n=2). Exclusion criteria were previous intraocular surgery, trauma, silicon oil tamponade, scleral buckle, incomplete continuous curvilinear capsulorhexis, rupture of the posterior capsule, and further surgery.

All surgeries were performed by a single surgeon (YK). After a 5.5–6 mm continuous curvilinear capsulorhexis, a 3.2 mm wide and 1.5 mm long corneoscleral incision was created at the upper limbus, followed by phacoemulsification, cortex removal, and polishing of the posterior capsule. Vitrectomy was performed using a standard three-port pars plana vitrectomy technique with a 23-gauge system. The X-70 was implanted in a capsular bag after vitrectomy was completed.

PCO was identified using slit-lamp examination, grading density of the opacification from 0 to 4 (0, none; 1, minimal; 2, mild; 3, moderate; 4, severe) by the same surgeon (YK). An Nd:YAG capsulotomy was performed based on the presence of grade 2 opacity and subjective patient complaints of decreased visual acuity compared with best postoperative visual acuity.

Kaplan–Meier survival distribution plots were generated to display the time of Nd:YAG capsulotomy events, and the Mantel–Cox log-rank test was used. Comparisons were considered to be statistically significant if the P-value was less than 0.05. All analyses were performed with GraphPad Prism version 6 software (GraphPad Software, Inc., San Diego, CA, USA).

Results

Two years after the combination surgery, four of 72 eyes (5.6%) in the diabetic retinopathy group and four of 81 eyes (5.0%) in the nondiabetic retinopathy group (three eyes with retinal detachment and one eye with branch retinal vein occlusion) required Nd:YAG capsulotomies. In total, eight of 153 (5.2%) eyes required Nd:YAG capsulotomies after combination surgery and implantation of X-70. The average patient age was 61±11 years in the diabetic retinopathy group and 62±10 years in the nondiabetic retinopathy group. There were 53 males and 19 females in the diabetic retinopathy group, and 45 males and 36 females in the nondiabetic retinopathy group. No statistically significant difference was
observed between the two groups based on Kaplan–Meier survival estimates \( (P=0.30, \text{Figure 2}) \).

**Discussion**

Toda et al reported that the PCO rate did not significantly differ between diabetic and nondiabetic retinopathy groups after combined cataract surgery and vitrectomy. Similarly, our results did not suggest diabetes mellitus as a significant risk factor associated with a higher Nd:YAG capsulotomy rate. Previous reports have indicated that patients with diabetes mellitus have significantly higher Nd:YAG capsulotomy rates than those without diabetes mellitus after single cataract surgery. Blood-aqueous barrier breakdown is reportedly more severe in the eyes of patients with diabetes mellitus, and some chemical mediators stimulate the proliferation of lens epithelial cells that may lead to extensive PCO. A high rate of Nd:YAG capsulotomy after combined cataract surgery and vitrectomy has also been previously reported. Although we could not directly compare these reports because of the different types of intraocular lens, these accounts attributed the high rate of Nd:YAG capsulotomy to the use of silicone oil and further surgery, the absence of which may account for the fact that we found no statistically significant difference between diabetic and nondiabetic retinopathy groups.

Nishi et al reported that capsular bend formation by the intraocular lens with a square-edge design inhibits migration of lens epithelial cells, thereby decreasing the Nd:YAG capsulotomy rate. However, Toda et al reported that there is a slight posterior pressure in eyes with no vitreous after vitrectomy; thus, it is possible that the intraocular lens edge effect in reducing PCO is minimal. Previous reports have shown that the intraocular lens material plays a prominent role in preventing PCO, and the acrylic intraocular lens is associated with a significantly reduced degree of PCO. Our results suggest that the X-70 intraocular lens may have the potential for enhanced PCO prevention.

A limitation of our study was the use of Nd:YAG capsulotomy as a measurement of PCO. Criteria for Nd:YAG capsulotomy is subjective and largely based on patient complaints. If patients do not complain of low visual acuity (despite existence of severe PCO), Nd:YAG capsulotomy is not performed. Future studies with objective PCO measurement are definitely essential.

In conclusion, the rate of Nd:YAG capsulotomy did not significantly differ between the diabetic retinopathy group and the nondiabetic retinopathy group after combined cataract surgery with implantation of the X-70 and vitrectomy.

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**Figure 2** Kaplan–Meier survival distribution plots are generated to display the time of Nd:YAG capsulotomy events.

**Notes:** Two years after the combination surgery, four of 72 eyes (5.6%) of patients in the diabetic retinopathy group and four of 81 eyes (5.0%) of patients in the nondiabetic retinopathy group required Nd:YAG capsulotomies. The absence of statistically significant differences between the two groups \( (P=0.30, \text{Mantel–Cox log-rank test}) \).

**Abbreviation:** Nd:YAG, neodymium:yttrium-aluminum garnet.
The choice of an intraocular lens based on prevention of PCO is important. Our study suggests that the X-70 may have the potential for enhanced PCO prevention and be an acceptable intraocular lens for patients with different causes undergoing combined cataract surgery and vitrectomy.

Disclosure
No author has a financial and proprietary interest in this study.

References