Scleral Fixation of a Hydrophobic Acrylic Intraocular Lens with Eyelets Using 8-0 Polypropylene Suture

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Abstract

**Purpose:** To report clinical outcomes of a scleral fixation technique of a hydrophobic acrylic intraocular lens (IOL) with eyelets using 8-0 polypropylene suture.

**Methods:** Nine eyes of nine patients who underwent combined pars plana vitrectomy and sclera fixation of an IOL using this technique were analyzed.

**Results:** The mean follow up period was 7.11 months (range 6-12 months), and there was a significant visual improvement at 6 months after surgery. The mean logarithm of the minimum angle of the resolution changed from 0.54 at baseline to 0.29 at postoperative 6 months (p=0.016). The mean postoperative spherical equivalent at 6 months was -0.90±0.90 D, and the mean predictive error was -0.49±0.62 D.

**Conclusion:** Postoperative visual and refractive outcomes were favorable, and the position of IOLs were well centered in all cases. This technique could be a useful alternative for surgeons without easy access to Gore-Tex suture.

**Keywords:** scleral fixation, hydrophobic acrylic intraocular lens, 8-0 polypropylene
Introduction

Secondary intraocular lens (IOL) implantation is often necessary in the absence of zonular support after successful cataract extraction. Various surgical techniques have been introduced including anterior chamber IOL, iris-sutured IOL, scleral-sutured IOL, sutureless intrascleral fixation, and iris-claw IOL. [1-8] The selection of surgical technique usually depends on each patient's ocular condition and surgeon preference. The advantage of scleral-sutured fixation over other techniques is placing the IOL in the anatomic position away from anterior chamber structures.

Recently, favorable outcomes have been demonstrated using Gore-Tex suture (W.L. Gore & Associates, Newark, DE) for IOL fixation to the sclera. [9-11] Although there are multiple advantages in using Gore-Tex suture including its long-term durability and positive outcomes, it is difficult to get the material in some countries because of importing issues. Therefore, we report the result of this scleral fixation technique using 8-0 polypropylene suture instead of Gore-Tex suture.

Methods

This scleral fixation technique was performed in nine eyes of nine patients between April 1, 2019 and October 31, 2019. All patients underwent combined PPV and scleral fixation of an IOL, and all surgeries were performed by a single surgeon (JHL). This retrospective, interventional case series followed the tenets of the Declaration of Helsinki and was approved by the institutional review board of Severance Hospital, Yonsei University College of Medicine.

 Conjunctival peritomy with a relaxing incision was performed at the temporal and nasal side. The horizontal meridians were marked using ASICO axis marker (ASICO, Westmont, IL) at 3- and 9-o’ clock positions of the limbus. An infusion cannula was inserted at the inferotemporal side, 3 mm away from the limbus. Four sclerotomy sites were marked 3 mm away from the limbus and 4 mm apart from each other. Two valved trocars were placed at the two superior marks. A standard pars plana vitrectomy (PPV) with scleral depression and shaving of vitreous base was performed. A clear corneal incision was made at the superior limbus using a 2.75 mm keratome. Two strands of 8-0
polypropylene sutures (Ethicon, Somerville, NJ) with the needle removed were used to form two loops on each side. The suture was passed through each eyelet of a MX60 IOL (Bausch and Lomb, Bridgewater, NJ) with both ends of the suture going over the haptic. Each end of the suture was placed into the anterior chamber and positioned at the pupillary area behind the iris plane. The suture was retrieved through the corresponding sclerotomy site using intraocular forceps. A similar pattern of suture passage was repeated with the other suture through the opposite sclerotomy sites. The IOL optic was folded in half using a Kelman-Mcpherson forcep, and the IOL was introduced in the anterior chamber. After confirming appropriate suture tension and centration of the IOL, the sutures were tied using a 3-1-1 technique. The knots were trimmed and buried into the sclerotomy site. The corneal incision and conjunctival peritomies were sutured (figure 1).

Preoperative lens calculations were performed using optical biometry (IOLMaster, Carl Zeiss Meditec, Dublin, CA). A routine in-the-bag calculation was used because sclerotomies were made 3mm posterior to the limbus. The formula used in each IOL calculation was based on axial length. For axial length less than 22.5 mm, the Hoffer Q formula was used. The Sanders-Retzlaff-Kraff theoretical (SRK/T) formula was used for axial lengths over 22.5 mm. Postoperative refractive error was measured at 6 months after surgery. Postoperative lens astigmatism was calculated using the method previously described by Munoz-Escriva and Furlan, in the standard notation spherical (S) / cylinder (C) x axis (α). [12,13]

Results

Baseline characteristics and clinical outcomes are summarized in table 1. The mean age was 57.56 years and the mean follow up period was 7.11 months (range 6-12 months). Indications for surgery included dislocated IOL (five eyes), postoperative aphakia (three eyes), and subluxated crystalline lens (one eye). There was a significant visual improvement at 6 months after surgery. The mean logarithm of the minimum angle of the resolution change from 0.54 at baseline to 0.29 at postoperative 6 months (p=0.016). The mean postoperative spherical equivalent (SE) at 6 months
was -0.86±0.82 D, and the mean predictive error (postoperative SE minus target refraction) was -0.49±0.62 D. The SRK/T formula was used in eight patients and the Hoffer Q formula was used in one patient with axial length 22.36 mm. Prediction error in this patient was -0.05 D, which was smaller than the mean prediction error of patients with SRK/T formula (-0.58±0.64 D). The IOLs were well centered in all cases without any tilt, subluxation, or dislocation. The mean calculated lens astigmatism at postoperative 6 months was 1.31±0.49 D, which was similar to that of conventional two-point fixation technique by the same surgeon (1.04±0.89D, p=0.394; n=7). No intraoperative complication was encountered. Postoperative complications included transient corneal edema in two eyes (22.2%), cystoid macular edema in two eyes (22.2%), and increased intraocular pressure controlled with medical therapy in one eye (11.1%). There was no case of postoperative hypotony, endophthalmitis, suture breakage, IOL dislocation, or retinal detachment during the follow up period.

Discussion

The selection of suture material is crucial to a long-term postoperative prognosis in cases of IOL scleral fixation. The use of 10-0 polypropylene suture is becoming less popular because of the suture breakage resulting in IOL subluxation or dislocation, which have been reported to be as high as 24% after 10 years. [14] In 2014, Kahn et al described a technique for scleral fixation of IOL using Gore-Tex suture. [9] Subsequent short-term and 1-year follow up outcomes demonstrated the effectiveness and safety of a trans-scleral IOL fixation using Gore-Tex suture. [10,11] Recently, a study comparing this technique with combined PPV with anterior chamber IOL revealed similar visual outcomes with reduced postoperative complications including early corneal edema. [15] Despite its high tensile strength and resistance to degradation, it is difficult to use Gore-Tex suture in some countries including South Korea because it is not commercially imported locally. In addition, Gore-Tex suture is not yet indicated for ophthalmologic use in the manufacturer’s label, and further studies regarding its long-term safety results are necessary. John et al recently described a technique of scleral IOL fixation using 8-0 polypropylene and Akreos AO60 IOL (Bausch and Lomb, Bridgewater, NJ) instead of 10-0 polypropylene or Gore-Tex suture, which showed good clinical outcomes with safety...
results. [16]

Our technique demonstrated favorable visual and refractive outcomes with acceptable short-term complications. Postoperative prediction error (-0.49±0.62 D) showed slightly myopic outcomes than targeted. Although transscleral fixation is known to have a higher refractive prediction error than other techniques of secondary IOL implantation, [17] the refractive outcomes in scleral fixation tend to vary from myopic [18,19] to hyperopic differences. [20] Recently, sutureless scleral fixation of IOL has been described, of which refractive outcomes were also varied. Whereas Todorich et al reported the mean final refraction as -0.565 D, [21] Abbey et al revealed that the prediction error was +0.41 D. [22] Combined PPV might be a possible factor accounting for the myopic shift, [23] but the exact mechanism is still unknown.

Our technique has the following advantages: first, a hydrophobic acrylic IOL was used to avoid the risk of lens opacification associated with gas or oil tamponade as reported previously; [24] second, 8-0 polypropylene suture has enough stiffness that the suture can stay stable in the pupillary area, allowing this technique to be also performed without needing to use the handshake maneuver; and third, we demonstrated acceptable refractive outcomes when suturing 3 mm behind the limbus (postoperative SE -0.86 at 6 months, prediction error -0.49 D). According to Su et al, the mean postoperative SE and the mean prediction error for fixation 3 mm posterior to the limbus were -0.82 D and -0.43 D, respectively. [25] Compared with sclerotomies placed 2 mm posterior to the limbus, creating sclerotomies 3 mm behind the limbus can result in less myopic shift and minimize iris chafing or vitreous hemorrhage.

There are several important points to be highlighted in this technique. First, consistency in sclerotomy site placement is crucial to prevent unpredictable refractive results. Every sclerotomy site should be measured and marked carefully to confirm the exact distance from the limbus and from each other. A slight anterior placement in sclerotomy may result in a myopic shift and vice versa. Second, suture tension is another important factor. If the suture is too tight, the lens may tilt and be decentered. If the suture is too loose, the lens may tremble along with eyeball movement, leading to unpredictable refractive outcomes. Third, careful preoperative assessment of the conjunctiva is crucial for
preventing postoperative suture exposure. The position of sclerotomy site should be made with consideration to previous surgical interventions on the conjunctiva such as glaucoma surgery. Lastly, smaller gauge instruments would be advantageous for preventing postoperative leakage from the sclerotomy site.

This report has several limitations, including its retrospective nature and small number of cases. Visual and refractive outcomes may be influenced by preoperative condition of the eye because various surgical indications and complicated preoperative histories were present. In addition, 23- or 25-gauge vitrectomy systems were used in this case series, which could have affected the refractive outcomes. However, we believe that these results revealed an evidence-based outcome in performing combined vitrectomy and scleral fixation of a hydrophobic acrylic IOL using 8-0 polypropylene suture. Further long-term outcome results with a larger number of cases would be necessary to confirm the surgical outcomes of this technique.

In summary, we report clinical outcomes of a scleral fixation technique of a hydrophobic acrylic IOL with eyelets using 8-0 polypropylene suture. Postoperative visual and refractive outcomes were favorable, and the position of IOLs were well centered in all cases. This technique could be a useful alternative for surgeons without easy access to Gore-Tex suture.

Conflicts of interest: No authors have any conflict of interest.

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Figure legends

**Figure 1.** Intraoperative findings of scleral fixation of a hydrophobic acrylic intraocular lens (IOL) with eyelets using 8-0 polypropylene suture. A. The horizontal meridians were marked at 3- and 9-o'clock positions at the limbus. B. Four sclerotomy sites were marked 3mm from the limbus and 4mm apart. Pars plana vitrectomy was performed through a sclerotomy site. C. An 8-0 polypropylene suture was passed through each eyelet of the IOL with both ends of the suture going over the haptic. D. Each end of the suture was placed into the anterior chamber and retrieved through the corresponding sclerotomy using intraocular forceps. E. A similar pattern of suture passage was repeated with the other suture through the opposite sclerotomy. F. The IOL optic was folded in half and inserted into the anterior chamber. G. The sutures were tied using a 3-1-1 technique. H. The knots were trimmed and buried into the sclerotomy site. I. The corneal incision and conjunctival peritomies were sutured.

**Figure 2.** Schematic drawing of scleral fixation of a hydrophobic acrylic intraocular lens (IOL) with eyelets using 8-0 polypropylene suture (surgeon’s view, superior position). A. After conjunctival peritomy was performed at the temporal and nasal side, four sclerotomy sites were marked 3mm from the limbus and 4mm apart. B. An 8-0 polypropylene suture was passed through each eyelet of the IOL with both ends of the suture going over the haptic. C. Each end of the suture was placed into the anterior chamber and retrieved through the corresponding sclerotomy using intraocular forceps. D. After inserting the IOL, the sutures were tied and buried into the sclerotomy site.
Table 1. Baseline characteristics and clinical outcomes of patients

<table>
<thead>
<tr>
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<th>n=9</th>
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<tbody>
<tr>
<td>Age, years</td>
<td>57.56±15.70</td>
</tr>
<tr>
<td>Sex (male/female)</td>
<td>7/2</td>
</tr>
<tr>
<td>Duration of follow up, months (range)</td>
<td>7.11±1.96</td>
</tr>
<tr>
<td>Indications for surgery, n (%)</td>
<td></td>
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<tr>
<td>Dislocated IOL</td>
<td>5 (55.6)</td>
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<tr>
<td>Aphakia (postoperative)</td>
<td>3 (33.3)</td>
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<tr>
<td>Crystalline lens subluxation</td>
<td>1 (11.1)</td>
</tr>
<tr>
<td>BCVA, logMAR</td>
<td></td>
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<tr>
<td>Preoperative</td>
<td>0.54±0.63</td>
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<tr>
<td>Postoperative 6 months</td>
<td>0.29±0.33</td>
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<tr>
<td>p-value</td>
<td>0.016</td>
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<tr>
<td>SE at postoperative 6 months, D</td>
<td>-0.86±0.82</td>
</tr>
<tr>
<td>Prediction error, D</td>
<td>-0.49±0.62</td>
</tr>
<tr>
<td>Calculated lens astigmatism at postoperative 6 months, D</td>
<td>1.31±0.49</td>
</tr>
<tr>
<td>Postoperative complications</td>
<td></td>
</tr>
<tr>
<td>Corneal edema</td>
<td>2 (22.2)</td>
</tr>
<tr>
<td>Cystoid macular edema</td>
<td>2 (22.2)</td>
</tr>
<tr>
<td>Increased intraocular pressure</td>
<td>1 (11.1)</td>
</tr>
</tbody>
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BCVA, best-corrected visual acuity; logMAR, logarithm of the minimum angle of resolution; SE, spherical equivalent.