Surgery of congenital and juvenile cataracts: a pars plicata approach with the vitrophage

GHOLAM A. PEYMAN, MOTILAL RAICHAND, AND MORTON F. GOLDBERG
From the Department of Ophthalmology, University of Illinois Eye and Ear Infirmary, Chicago, USA

SUMMARY Eighteen eyes with congenital cataracts were surgically managed by pars plicata lensectomy and vitrectomy using the vitrophage. No major intraoperative or postoperative complications were encountered. All eyes have maintained extremely clear media, attached retinas, and absence of pupillary block during a follow-up period ranging from 12 to 36 months.

In the last two decades numerous advances have been made in the management of congenital cataracts. The intraoperative and postoperative complications associated with discission, linear extraction, and intracapsular lens extraction have been reduced with the re-introduction of the aspiration technique (Scheie, 1960). Development of phacoemulsification instruments (Kelman, 1967; Shock, 1974; Girard, 1974) greatly simplified the aspiration procedure. However, such postoperative complications as secondary membrane formation continue to occur with these techniques and require additional surgery (Hiles and Wallar, 1974; Hiles and Hurite, 1973).

With the use of automated vitrectomy instruments a new dimension has been added to the treatment of congenital cataracts. Surgical approach through the limbus (Calhoun and Harley, 1975) and pars plicata (Peyman et al., 1977) has been described previously.

This report indicates our moderately long-term results in patients with congenital and juvenile cataracts who were managed with a pars plicata approach by combined lensectomy and vitrectomy using the vitrophage (Peyman and Sanders, 1975).

Material and methods

Seventeen patients (18 eyes) were referred for the management of congenital cataract to the Vitreous Service of the University of Illinois Eye and Ear Infirmary. In 14 patients the contralateral eye had been managed with other surgical techniques such as phacoemulsification and aspiration. In 1 patient both eyes were managed by pars plicata approach as described below. The other patients had no previous surgery, and only 1 eye of each was operated on. In 1 of these the cataract was unilateral, and the other patient refused operation on the fellow eye.

The patients’ ages ranged from 6 months to 37 years. Three patients were 25 years or older and the remainder were 15 or younger (Table 1). Patients 3 and 5 are twins. In 4 patients the cataracts were due to maternal infection with rubella during pregnancy; in 2 patients the family history suggested autosomal dominant mode of inheritance; and in the remainder the aetiology was unknown. In all the eyes combined lensectomy and vitrectomy was performed by means of the vitrophage (Peyman and Sanders, 1975).

In children surgery was performed with the patient under general anaesthesia. In adults local (retrobulbar) anaesthesia was used. Initially an examination of both eyes is done before the standard surgical preparation is made. Traction sutures are then passed through both eyelids, followed by traction sutures under the insertion of the 4 rectus muscles, 4-0 black silk being used. After adequate exposure is obtained, a limbal peritomy, usually in the inferotemporal quadrant, is made. In children a 4-mm-long sclerotomy is made 2-5 mm posterior and parallel to the limbus, a No. 15 Bard-Parker blade being used. In adults, however, the sclerotomy can be made 3-5 mm from the limbus.

The underlying ciliary body is cauterised or diathermised gently. A mattress suture of 5-0 polyglactin 910 or 4-0 Supramid is passed through the lips of the sclerotomy, and a double throw-knot is applied loosely. The suture material is then looped out of the sclerotomy. A 52-s Beaver blade is inserted through the sclerotomy into the lens and then removed with a slicing motion to widen the
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Table 1  Patients treated with pars plicata lensectomy and vitrectomy

<table>
<thead>
<tr>
<th>Case no.</th>
<th>Age</th>
<th>Aetiology and remarks</th>
<th>Complications</th>
<th>Approximate clarity of fundus view</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>14 yr</td>
<td>Unknown</td>
<td>None</td>
<td>20/20</td>
</tr>
<tr>
<td>2</td>
<td>5 yr</td>
<td>Unknown</td>
<td>None</td>
<td>20/20</td>
</tr>
<tr>
<td>3</td>
<td>4 yr</td>
<td>Rubella syndrome, esotropia</td>
<td>None</td>
<td>20/20</td>
</tr>
<tr>
<td>4</td>
<td>4½ yr</td>
<td>Rubella syndrome</td>
<td>None</td>
<td>20/20</td>
</tr>
<tr>
<td>5</td>
<td>4 yr</td>
<td>Rubella syndrome</td>
<td>Transient corneal oedema post-operatively</td>
<td>20/20</td>
</tr>
<tr>
<td>6</td>
<td>1½ yr</td>
<td>Unknown, esotropia</td>
<td>Self-limited bleeding from iris intra-operatively</td>
<td>20/20</td>
</tr>
<tr>
<td>7</td>
<td>6 mo</td>
<td>Autosomal dominant congenital cataract</td>
<td>Transient slight corneal haze postoperatively</td>
<td>20/20</td>
</tr>
<tr>
<td>8</td>
<td>6 mo</td>
<td>Autosomal dominant congenital cataract</td>
<td>None</td>
<td>20/20</td>
</tr>
<tr>
<td>9</td>
<td>4 yr</td>
<td>Rubella syndrome</td>
<td>None</td>
<td>20/20</td>
</tr>
<tr>
<td>10</td>
<td>10 mo</td>
<td>Unknown</td>
<td>None</td>
<td>20/20</td>
</tr>
<tr>
<td>11</td>
<td>2½ yr</td>
<td>Unknown</td>
<td>None</td>
<td>20/20</td>
</tr>
<tr>
<td>12</td>
<td>15 yr</td>
<td>Aniridia and nystagmus</td>
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<td>20/20</td>
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<td>13</td>
<td>25 yr</td>
<td>Chorioretinal scars</td>
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<td>34 yr</td>
<td>Unknown</td>
<td>None</td>
<td>20/20</td>
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<td>None</td>
<td>20/20</td>
</tr>
<tr>
<td>16</td>
<td>37 yr</td>
<td>Unknown, nystagmus</td>
<td>None</td>
<td>20/20</td>
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<tr>
<td>17</td>
<td>6 mo</td>
<td>Autosomal dominant congenital cataract</td>
<td>None</td>
<td>20/20</td>
</tr>
<tr>
<td>18</td>
<td>7 mo</td>
<td>Unknown</td>
<td>None</td>
<td>20/20</td>
</tr>
</tbody>
</table>

tract for insertion of the lens fragmentor. A lens fragmentor (Peyman and Sanders, 1975) is introduced into the lens through the sclerotomy, and the lens material is fragmented for about 60 seconds.

After this the lens fragmentor is removed and replaced with a wide-angle cutter vitrophage (Peyman, 1975). The mattress scleral suture is tightened round the vitrophage. The cortical and nuclear fragments are removed first, followed by the lens capsule. An anterior and a central vitrectomy is performed to the optic disc, after which the vitrophage is removed and the scleral mattress suture tied. The conjunctiva is sutured with 6-0 plain catgut. At the conclusion of the operation, antibiotic and cycloplegic eye drops are instilled.

In cases when the pupil does not dilate well sector iridectomy is performed with the vitrophage. The infusion fluid used for replacement of vitreous is 5% dextrose in 0.45% normal saline containing 2-2 g sodium bicarbonate and 0.2 g/litre calcium chloride. We routinely add 4 µg/ml of gentamicin of infusion fluid.

Results

No operative complications were encountered in 17 eyes. One eye (Patient 6, Table 1) had minimal bleeding from the iris, which stopped spontaneously.

During the early postoperative period transient corneal oedema, noticed in 2 patients’ eyes, cleared over a period of 2 to 3 weeks. There was no evidence of intraoperative or postoperative bleaching at the site of entry into the pars plicata or development of retinal dialysis. Surgery was tolerated well, and there was no obvious difference between the eyes with rubella- and non-rubella-induced cataracts during the postoperative period.

Postoperatively the media were extremely clear (approximately 20/20 in clarity) in all eyes. As a majority of these patients have associated disorders such as amblyopia, microphthalmia, retinal scarring, or mental retardation, the final visual acuity was often difficult to assess.

No increase in intraocular pressure was noticed, and no severe or persistent intraocular inflammation appeared. No retinal detachment has been evident during the follow-up, ranging from 12 to 36 months. No case of pupillary block with shallow anterior chamber occurred.

Discussion

Management of congenital cataracts, a matter of great controversy, can be separated into three main areas of debate: the indication and timing of surgical intervention; the technical procedures of choice; and the management of amblyopia and postoperative optical correction. Our discussion will be confined to the technical aspects of congenital cataract surgery, although it is recognised that restoration of good visual function often depends on postoperative management and correction of amblyopia.

Despite the availability of various surgical techniques no one procedure has gained wide acceptance because of inherent complications associated with each. The simple discission technique had an unacceptable rate of complications such as swelling or retained lens matter, glaucoma, and retinal detachment (Barkan, 1932; Chandler, 1951; Cordes, 1956). Although better results have been reported with linear extraction (Owens and Hughes, 1948; Bagley, 1949), this technique was also...
associated with complications such as delayed formation of anterior chamber, anterior synechiae and updrawn pupil, occlusion of pupil, membrane formation, vitreous loss, glaucoma, and retinal detachment (Cordes, 1961). Similarly, intracapsular extraction produced complications in a very high percentage of cases (Ryan et al., 1965) and was advocated by only a few surgeons (Escapini, 1968).

The aspiration techniques have greatly reduced the complications associated with older methods (Sheppard and Crawford, 1973). Although the incidence of postoperative glaucoma and retinal detachment was reduced significantly (Parks and Hiles, 1967; Ryan et al., 1965; Ryan and von Noorden, 1971), secondary membrane formation and vitreous loss have continued to be major problems. Parks and Hiles (1967) reported a 73% incidence of membrane formation and 6% vitreous loss in a review of 52 eyes that underwent discission and aspiration. Ryan and von Noorden (1971) reported 23% incidence of secondary membrane and 8% vitreous loss in 75 cases by using the discission and the aspiration technique. The incidence of postoperative secondary membrane formation with aspiration techniques ranged from 18% (Sheppard and Crawford, 1973) to 73% (Parks and Hiles, 1967).

Although phacoemulsification has greatly facilitated the surgical management of congenital cataract surgery, complications such as vitreous loss and secondary membrane formation are still major problems that often require additional surgery (Hiles and Hurite, 1973). Hiles and Wallar (1974) reviewed their experiences with phacoemulsification and aspiration in infantile cataract surgery in 390 eyes. They reported secondary or delayed posterior capsulotomies in 66% of their aspiration group and 39% in their phacoemulsification group. A total of 70% of the eyes in the aspiration series required a capsulotomy either at the end of the operation or later. Of those eyes undergoing phacoemulsification 58% required a capsulotomy. Hiles (1977) states that as many as 92% of these children will need capsulotomies if enough time has elapsed after the original cataract surgery.

Automated vitrectomy instruments have greatly aided our ability to deal with secondary membrane and vitreous loss. Although we have advocated the pars plana approach for adult patients, we suggest the pars plicata approach for children. Anatomically the pars plana region is smaller in children than in adults. By making the sclerotomy 2.5 mm from the limbus and entering the eye through the pars plicata a safe distance from the ora serrata is maintained. Transillumination through the pupil is sometimes helpful in identifying the ciliary body.

The pars plicata approach offers many advantages over the limbal approach. With the limbal approach Calhoun and Harley (1975) reported unplanned iridectomies in 14% of cases and a similar incidence of unexplained prolonged intraocular inflammation. The pars plicata route avoids direct manipulation of the corneal endothelium, and, similarly, contact with the iris is kept to a minimum. Any accidental bleeding from the iris into the deep vitreous can be treated immediately with the pars plicata approach, because a fundus contact lens can immediately be placed on the unmanipulated cornea. Moreover, lens particles that may fall into the vitreous can also be retrieved, so that the chance of postoperative inflammation is thus reduced. With the pars plicata approach the tip of the vitrophage can easily reach behind the iris and remove most of the lens material by applying only suction, a procedure which is rather difficult through the limbal approach, particularly in the axis of the shaft of the instrument.

In our series no patient has had severe intraocular inflammation postoperatively. We routinely remove the anterior vitreous with a central core of vitreous down to the optic disc. This technique has eliminated the development of pupillary block glaucoma.

Our patients have been followed up for periods ranging from 12 to 36 months. All eyes have maintained extremely clear media postoperatively. Using serial refraction measurements, we have been able to observe normal growth and development in these eyes when compared with contralateral eyes. With the evidence at hand we can postulate that pars plicata lensectomy and vitrectomy has no immediate untoward effect on the development of infant eyes. Although additional long-term results of this technique are clearly desirable, the present data are encouraging.

Combined pars plicata lensectomy and vitrectomy offers a 'one procedure' technique for the surgical management of congenital cataracts with minimal complications.

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References


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