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A case of WAGR syndrome in association with developmental glaucoma requiring bilateral Baerveldt glaucoma implants and subsequent tube repositioning

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Abstract: Glaucoma drainage device implantation is efficacious for the treatment of pediatric glaucoma patients when multiple angle surgeries fail. However, tube touching of the corneal endothelium is one of the major postoperative complications to deal with. A 15-month-old male patient with Wilms’ tumor, aniridia, genitourinary anomalies, and mental retardation (WAGR) syndrome was diagnosed with bilateral developmental glaucoma. He underwent Baerveldt glaucoma implant (BGI) surgeries in both eyes after multiple failed trabeculotomies. The tube in his right eye was touching the cornea 15 months after BGI surgery. To avoid corneal endothelium damage, BGI tube repositioning with scleral fixation was performed without serious complications. The bilateral BGI surgeries achieved successful intraocular pressure reduction for over 2 years and tube repositioning with scleral fixation of BGI tube was successful for BGI tube malposition. Although careful attention to intraocular pressure and tube malposition is essential after glaucoma drainage device implantation, especially in pediatric cases, BGI surgery is effective in the management of developmental glaucoma following unsuccessful angle surgeries.

Keywords: Baerveldt glaucoma implant, developmental glaucoma, WAGR syndrome, tube repositioning, glaucoma drainage device

Introduction

In eyes with developmental glaucoma, surgical therapies such as trabeculotomy or goniotomy are usually essential to control intraocular pressure (IOP). Although these angle surgeries have high success rates in primary developmental glaucoma, secondary developmental glaucoma, with associated ocular/systemic anomalies, including aniridia, has a poor prognosis, probably because of maldevelopment of the angle structures and Schlemm’s canal. Wilms’ tumor, aniridia, genitourinary anomalies, and mental retardation (WAGR) syndrome is caused by heterozygous contiguous gene deletions that involve at least Pax6 and Wt1, which has a risk of a developmental glaucoma.3

Even when the first trabeculotomy is unsuccessful, a subsequent trabeculotomy may be effective in lowering IOP.1 However, when multiple angle surgeries fail, further surgical procedures such as trabeculectomy with mitomycin-C, cyclodestructive surgery, and glaucoma drainage device (GDD) implantation are needed.2,4 Trabeculectomy in younger children has many problems: a rapid healing response, postoperative bleb management, and a high rate of bleb-associated endophthalmitis.2 Cyclodestructive
procedures require multiple retreatments and have a possible risk for a serious complication, phthisis bulbi. It has been reported that the success rate is significantly higher after Ahmed glaucoma valve or Baerveldt glaucoma implant (BGI) (87%) versus trabeculectomy with mitomycin-C (36%) in children.\(^4\) GDD implantation, such as BGI, is suggested as the most predictable and possibly the safest procedure after failed angle surgery.

Here, we report a case of WAGR syndrome in association with developmental glaucoma requiring bilateral BGIs and subsequent tube repositioning. This is the first report of tube repositioning with scleral fixation of BGI tube in a pediatric case.

**Case report**

A 15-month-old male patient was referred to Kyoto University Hospital. He was diagnosed with bilateral (OU) aniridia and developmental glaucoma in his right eye (OD) at 1 month of age and subsequently with WAGR syndrome. He had undergone three OD trabeculotomies before 1 year of age. At the initial visit to our hospital, he was using three types of glaucoma eyedrops; his IOP was 11 mmHg OD and 24 mmHg in the left eye (OS). Fundus examination showed macular hypoplasia OU; a tilted disc with 0.9 cup to disc ratio OD; and an oval disc with 0.6 cup to disc ratio OS. Figure 1 and 2 show pre- and postoperative photographs and IOP throughout the follow-up period, respectively. Briefly, after three failed OD and two failed OS trabeculotomies, BGI surgeries were performed in both eyes.

A BGI103-250 implant was placed in the superior temporal quadrant with a fornix based conjunctival flap (Figure 1A–D). The plate was secured with 9-0 nylon sutures 9 mm posterior to the limbus. The tube was ligated with 7-0 Vicryl and was slit to manage the transient hypertensive
Baerveldt glaucoma implants and tube repositioning for developmental glaucoma

Figure 2 The intraocular pressure (IOP) time course.

Notes: All IOP measurements were performed under sedation using a Perkins applanation tonometer (Clement-Clarke, Haag-Streit UK, Essex, UK). The patient underwent the first inferotemporal trabeculotomy OS (left eye) at 15 months (A). His IOP OS decreased to around the mid-teens once, but increased to over 30 mmHg and his IOP OD (right eye) gradually increased to over 40 mmHg. Therefore, a second inferonasal trabeculotomy OS and BGI OD were performed at 21 months (B). After 1 month, BGI OS was performed (C), because the second trabeculotomy OS was not sufficiently effective. The BGI tube repositioning with scleral fixation OS was performed (D).

Abbreviations: BGI, Baerveldt glaucoma implant; y, year(s); m, month(s).

Discussion

Tube positioning in the anterior chamber is especially important in pediatric cases due to the growth of the eyeball. Because tube retraction is a possible postoperative complication in pediatric GDD surgeries, we left a longer tube in the anterior chamber than in adults (Figure 1C and D). Tube touching of the corneal endothelium is another postoperative complication. Beck et al reported that tube repositioning was needed in 34.8% of 46 GDD-implanted pediatric eyes. Cutting the tube end or repositioning through a new scleral pathway is a possible choice. However, cutting the tube would be palliative, because it would not change the tube insertion angle. Reopening the conjunctiva and sclera for removal and tube reinsertion through a new pathway might cause insufficient wound closure and disturb the function of the BGI. Our choice was a novel technique of repositioning the tube with scleral fixation, which Ma et al reported in adult patients.

The bilateral BGI surgeries achieved successful IOP reduction for over 2 years. This is the first report of tube repositioning with scleral fixation of BGI tube in a pediatric case. This procedure was good for BGI tube malposition even in a pediatric case, at least for a short period. Although careful attention to IOP and tube malposition is essential after GDD implantation, especially in pediatric cases, BGI surgery is effective in the management of developmental glaucoma following unsuccessful angle surgeries.

Disclosure

The authors report no conflicts of interest in this work.
References


