Congenital intrascleral cyst

Müslime Akbaba1
Güllühan Hacıyakupoğlu2
Aysun Uğuz3
Şafak Karslioğlu1
Zeynel Karcioğlu4

1Istanbul Oculoplastic and Orbital Surgery and Ocular Oncology Center, Şişli/Istanbul, Turkey; 2Department of Ophthalmology of Faculty Medicine, Çukurova University, Balcali/Adana, Turkey; 3Department of Pathology of Faculty Medicine, Çukurova University, Balcali/Adana, Turkey; 4Department of Ophthalmology, University of Virginia Health Sciences Center, Charlotte, NC, USA

Abstract: Congenital intrascleral cysts are rare. They are mostly located at the limbus with corneal involvement. We report a case of a 30-month-old boy with a bulbar conjunctival cyst noticed at birth. The lesion enlarged over the following months but did not involve the cornea. During surgery the cyst proved to be intrascleral and a complete excision was carried out. The remaining defect was repaired with banked fascia lata. The histopathology revealed a scleral cyst wall lined by nonkeratinizing squamous epithelium with no goblet cells. We conclude that congenital intrascleral epithelial cysts are rare but should be considered in differential diagnosis of external eye cystic lesions. In our case, early excision and repair with fascia lata led to an uncomplicated postoperative course of 6 years.

Keywords: intrascleral cyst, epithelial cyst, banked fascia lata

Introduction

Congenital intrascleral cysts are unusual clinical conditions with uncertain etiology. The lesions are usually located at the limbus with corneal and scleral components. In this report, we present a case with progressively enlarging congenital intrascleral cyst without corneal involvement in a 30-month-old boy, and describe its clinical and histopathologic features and surgical management.

Case report

A 30-month-old boy presented to the clinic with a bulbar subconjunctival cystic mass of the right eye (Figure 1A). The lesion was present at birth and had been gradually enlarging for 1 year. Delivery was uneventful and there was no history of ocular trauma or amniosynthesis.

A thin-walled, multilocular scleral cyst, measuring 5 mm × 8 mm in its largest dimensions, was identified at the inferior nasal quadrant, extending from the medial canthus to the limbus of the right eye (Figure 1A, B, and C). There was no conjunctival congestion or vascularization, and the cornea was clear. Visual acuity was 1.0 with a “tumbling E” chart. The rest of the ocular examination in the right eye and the left eye was normal.

During surgery, saline hydrodissection was carried out with a 30-gauge needle to separate the underlying cyst wall from conjunctiva, confirming that the lesion was intrascleral. A conjunctival incision was made through the upper margin of the mass from the 3 o’clock position to the inner canthus to expose the anterior portion of the...
lesion by blunt dissection without perforating the cyst. The anterior portion of the cyst containing transparent fluid was exposed and removed carefully. The posterior aspect of the cyst was then gently cleaned with a Weck-cell sponge to be sure that no residual epithelium was left behind. After the total removal of the cyst, the sclera at the posterior aspect was noted to be very thin and the underlying choroid was clearly visible. This thin area was patched with preserved fascia lata using 8.0 vicryl sutures to prevent staphyloma formation (Figure 1B). Conjunctiva above the patch was also closed with 8.0 vicryl sutures. At the end of the surgery, 0.5 mL dexamethasone and 0.5 mL gentamicin were injected subconjunctivally. Topical tobramycin and prednisolone acetate were used postoperatively. Histopathologic examination revealed that the cyst wall was lined with nonkeratinized epithelial cells without goblet cells. The presence of goblet cells is said to favor congenital origin. Rao et al reported the presence of glandular structures simulating lacrimal gland tissue and stated that this indicates the developmental nature of the lesion.

These cysts may recur postoperatively because of residual epithelium. Careful removal of the cyst wall, denaturation of remnants with saline and distilled water, and chemical coterization with trichloroacetic acid and tetracyclin injection are advised to prevent recurrence. In limbal cysts, peripheral lamellar keratoplasty is known to be effective to prevent the development of staphyloma and recurrence. In order to prevent staphyloma formation in our case, we utilized banked fascia lata, which worked well without complications. Based on our experience in this case, we would suggest that banked fascia lata may be considered as an alternative grafting material in defect restoration when banked human sclera is not readily available.

In conclusion, we report our experience with a congenital intrascleral cyst that was surgically repaired with the use of banked fascia lata and had an uncomplicated postoperative course of 6 years.

Disclosure
The authors report no conflict of interest with this work.

References
