Ectopic cilia associated with an orbital dermoid cyst and sinus tract: case report

David Krahalík, MD, PhD, Marta Karhanová, MD, PhD, Miroslav Vaverka, MD, PhD, Světlana Brychtová, MD, PhD, and Dagmar Pospišilová, MD, PhD

Departments of 1Neurosurgery, 2Ophthalmology, 3Clinical and Molecular Pathology, and 4Pediatrics, University Hospital Olomouc, Czech Republic

Ectopic cilia are extremely rare congenital anomalies in which eyelash follicles appear in an abnormal place on the eyelid, most typically on the lateral quadrant of the anterior surface of the upper eyelid. In the majority of cases, simple surgical excision of ectopic cilia is indicated because of its cosmetic aspect. There is usually no associated medical comorbidity with this anomaly. The authors report an unusual case of ectopic cilia associated with an orbital dermoid cyst and sinus tract. A 3-year-old boy was initially diagnosed with ectopic cilia on the left upper eyelid. There was no history of inflammation or swelling of the eyelid. An ophthalmological examination revealed only 1 mm of ptosis; no proptosis, inferior displacement, or palpable orbital mass was present. During surgical excision of the ectopic cilia, a thin sinus tract was identified, leading posteriorly to the orbit. Magnetic resonance imaging performed after the excision showed a supraorbital extraconal mass just below the roof of the left orbit. A supraorbital 2-piece craniotomy was performed with total extirpation of the dermoid cyst. The cyst was removed en bloc without damage to the extraocular muscles, but the sinus tract could no longer be identified. Follow-up MRI was performed 6 months after surgery and showed no evidence of recurrence. A follow-up ophthalmological examination showed no signs of inferior displacement or proptosis. To the best of the authors' knowledge, this case is the first reported instance of ectopic cilia associated with a dermoid cyst and sinus tract in which no typical clinical signs and symptoms of possible orbital pathology were present. This case highlights the value of radiological examination in all cases of ectopic cilia prior to surgical excision.


KEY WORDS ectopic cilia; dermoid cyst; orbit; sinus tract; congenital
normal, and no other associated abnormality was detected. The diagnosis of ectopic cilia was made and surgical excision was indicated specifically for cosmetic reasons. During the surgical excision of the ectopic cilia under general anesthesia, a thin fibrous band leading posteriorly to the orbit was detected. The band was excised after reaching the end of the band posteriorly. A small amount of oily liquid was released and a tiny sinus tract was clearly identified, but additional posterior dissection was not possible from the anterior approach. Histopathological examination of the excised material was completely consistent with the diagnosis of ectopic cilia when an abnormally arranged cluster of 12 hair follicles embedded in a desmoplastic stroma was found, without signs of inflammation. Four follicles had the appearance of an anagen phase of the hair growth cycle, and the remaining ones were formed by hair bulbs, with some of them miniaturized. There were also 2 small sebaceous glands in the lesion (Fig. 2).

Operation and Postoperative Course

Three months after excision of the ectopic cilia, a stiff edema of the upper eyelid, 3 mm of ptosis, and mild inferior displacement of the globe were present (Fig. 3). Magnetic resonance imaging showed a supraorbital extraconal mass just below the roof of the left orbit. The size of the mass was $30 \times 25 \times 15$ mm and the signal of the presented mass was heterogeneous on T1- and T2-weighted MR images, with slight enhancement of the margins after contrast agent administration (Fig. 4). On preoperative laboratory examination, a prolonged activated partial thromboplastin time was found and, in a detailed investigation, deficiency of von Willebrand factor was diagnosed. The patient was prepared for operation with an infusion of a high-purity factor VIII concentrate (antihemophilic factor/von Willebrand factor complex [Humate-P]; CSL Behring). A supraorbital 2-piece craniotomy was performed with total extirpation of the dermoid cyst. The cyst was removed en bloc without damage to the extraocular muscles, but the tiny sinus tract could no longer be identified. The specimen was 35 mm in length, and inside, beneath the membrane, yellow content was visible. Microscopically, a typical dermoid cyst was observed containing sebaceous glands, hairs, and hair follicles (Fig. 5).

Follow-up MRI performed 6 months after surgery showed no evidence of recurrence. A follow-up ophthalmological examination showed 1 mm of ptosis, but no signs of inferior displacement or proptosis.

Discussion

The first report of ectopic cilia was published by Wiegmann in 1936, in a 5-year-old girl. The presentations published in the literature fall into 2 distinct categories: congenital ectopic cilia situated externally protruding from the anterior surface of the tarsal plate away from the lid margin and posterior cilia acquired later in life emerging from the tarsal conjunctiva. Almost 20 cases of cutaneous ectopic cilia of the upper eyelid have been reported in the literature. Their origin is not well understood. In the past, they were believed to originate from meibomian glands, which had been completely or partially replaced. Current evidence has suggested their embryological origin, so they should be considered a congenital anomaly whose growth is influenced by various molecules released from the local microenvironment. This hypothesis is strongly supported by their anatomical distribution, because all described cases were situated...
Ectopic cilia associated with orbital dermoid cyst

J Neurosurg Pediatr Volume 16 • August 2015

205

consistently on the lateral part of the upper eyelid. This localization corresponds embryologically to the watershed area between 2 vascular supplies, the peripheries of the facial and superficial temporal arteries, where a Tessier Type 9 facial cleft is also known to occur.17 Morphologically, ectopic cilia are formed by hair with hair bulbs and should be accompanied by sebaceous and sweat glands of the large apocrine type.8

In humans, the association of ectopic cilia with only a small number of abnormalities and complications has been documented. Distichiasis and ectopic cilia were reported by Bader.2 Gordon et al.10 described a complex choristoma of the right eyelid containing ectopic cilia and a functioning aberrant lacrimal gland tissue intermittently producing tears. Sebum that had accumulated at the base of the cilia in 1 patient was found by Chappell et al.5 A case of ectopic cilia and atopic eczema in the periorbital region18 and a hypochromic nevus7 were also documented. Edmunds et al.9 presented the case of a 2-year-old girl with nail-patella syndrome and ectopic cilia.

Dermoid cysts are the most common orbital cystic tumor presenting in childhood and arise from ectodermal rests pinched off at suture lines during embryogenetic migration.1 A dermoid cyst should be differentiated from an atypical epidermoid cyst.22 Most orbital dermoid cysts present in childhood as a slow-growing subcutaneous mass near the superotemporal orbit and frontozygomatic suture. When the growth is outward into the eyelid, cysts present in early childhood, and when they grow inward into the orbit, they present later in life.21 A displacement of the globe could be found in some cases. Deep orbital dermoid cysts may present with proptosis and ocular motility disturbances or orbital nerve compression. Orbital dermoids only rarely form sinus tracts; they can be more commonly found in association with nasal dermoids.15,20 Bonavolontà and colleagues4 identified only 2 cases with sinus tracts from 145 cases of orbital dermoid cysts examined during a 16-year period. The tracts are not usually to the eyelids, but when they do occur, the sinus tract opening is typically situated in the frontotemporal or orbitotemporal region. Lacey et al.36 reported 3 cases of orbitotemporal dermoids with a sinus tract and bone invasion. When a sinus tract is present, recurrent infections or intermittent discharge of sebaceous material is common. Orbital cellulitis may also be the presenting sign of an orbital dermoid associated with a sinus tract. However, it is rare for eyelashes or hair to grow from the sinus tract opening. Wells and Harris24 reported an orbital dermoid cyst and sinus tract extending from a hypoplastic sphenoid wing, through the lateral orbit, to the skin surface, presenting with acute orbital cellulitis and discharge from the hair-bearing pit above the eyebrow in a 9-month-old infant. Wang et al.23 described an orbital dermoid cyst with a sinus tract and hairs mimicking ectopic cilia presenting with episodes of inflammation or infection of the pit, characterized by swelling, redness, a yellowish discharge with a putrid odor, and a slowly growing mass in the upper eyelid in a 2-year-old boy. The whole complex was removed via an infrabrow incision.

In our case, there were no clinical signs of a possible presence of an orbital dermoid or sinus tract before excision of the ectopic cilia, which, as we suppose, was due

FIG. 4. Sagittal (left) and coronal (right) contrast-enhanced T1-weighted MR images showing a supraorbital extracanal mass just below the roof of the left orbit (arrow).

FIG. 5. Gross specimen showing a typical dermoid cyst containing sebaceous glands, hairs, and hair follicles (upper) and a microphotograph of the dermoid cyst (lower) lined by keratinizing squamous epithelium, with multiple adnexal structures in the wall. H & E, original magnification ×40. Figure is available in color online only.
to the deep location of the dermoid cyst. The opening of the sinus tract during the first operation caused discharge of sebaceous material into the soft tissues of the eyelid. If the presence of the orbital dermoid was known, the whole complex could have been removed in a single step in cooperation with a neurosurgeon and an ophthalmologist. Based on our experience, we would like to emphasize that ectopic cilia should be viewed as a developmental disorder that might be associated with other abnormalities. We highly recommend careful clinical and radiological examination in cases of ectopic cilia prior to surgical excision to prevent unnecessary mutilation of the patient.

References


Author Contributions

Conception and design: Krahulík, Karhanová. Acquisition of data: Krahulík, Karhanová. Analysis and interpretation of data: Krahulík, Karhanová, Brychtová. Drafting the article: Krahulík. Critically revising the article: all authors. Approved the final version of the manuscript on behalf of all authors: Krahulík.

Correspondence

David Krahulík, Department of Neurosurgery, University Hospital Olomouc, I. P. Pavlova 6, 77900 Olomouc, Czech Republic. email: david.krahulik@fnol.cz.