Extratarsal chalazia

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SUMMARY Three cases of atypical, extratarsal chalazia are presented. All three patients initially presented with a history of an inflammatory mass close to the lid margin. Inferior migration subsequently occurred, with loss of connection to the tarsus, causing confusion in the diagnosis. The basis for this migration is postulated to be the anatomical relationships of the tarsus, postorbicular fascia, and lower eyelid retractors. A good response was obtained with standard surgical therapy.

Although it is one of the commonest benign tumours of the eyelid,12 the lowly chalazion does not stir much interest or excitement in most ophthalmologists. Because the presentation is usually common and characteristic, of more importance are other lesions, such as sebaceous gland carcinoma, masquerading as chalazia.34 The following three cases represent the opposite circumstance, where an atypical presentation of a chalazion with an extratarsal location caused concern.

Case reports

CASE 1
A 4-year-old girl developed a small mass slightly inferior to the margin of her left lower lid. She was seen by her ophthalmologist and the diagnosis of a chalazion was made. Treatment consisted of application of warm soaks. Over the ensuing three months the lesion enlarged and appeared to migrate inferiorly. It became tender, and stabilised in size.

When the child was referred for examination, she presented with a $5 \times 10$ mm violaceous mass located at the level of the inferior orbital rim, at the junction of the medial and middle thirds of the left lower lid (Fig. 1). The lesion was covered by thin, atrophic skin, with scale formation and dilatation of surrounding capillaries. There was no evidence of fixation to deeper tissues. The rest of the ocular examination was normal.

At surgery a large collection of necrotic-appearing tissue was located within the substance of the orbicularis muscle anterior to the orbital septum. Frozen-section examination revealed florid granulation tissue. No connection to the tarsus was seen. The abnormal tissue was removed and the wound closed with advancement flaps.

Examination of fixed tissue showed a mixture of diffuse acute and chronic inflammatory cells. Neutrophils, lymphocytes, plasma cells, fibroblasts, and histiocytes were seen (Fig. 2). A granulomatous reaction corresponding to probable lipid globules was also present.

The patient’s postoperative course was uneventful. When seen eight months after surgery the lid was healed with no evidence of recurrence.

CASE 2
A 39-year-old woman developed a lesion she described as a ‘sty’ three months prior to examina-
Fig. 2 Case 1. Diffuse acute and chronic inflammation with granulomatous reaction. Haematoxylin and eosin.

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ation. It was treated with a combination of oral penicillin and topical sulphacetamide eye drops, with the application of moist heat. Over the next three months the lesion remained relatively stable in size, but proceeded to migrate inferiorly from its initial location near the lid margin. The lesion would periodically drain purulent material.

When examined the patient had a 15x7 mm mass located in the mid portion of the right lower lid, at the level of the inferior orbital rim. The overlying skin was thin and of a violaceous hue. A central ulcer or fistula was present, with a dried purulent plug. With the exception of mild follicular hypertrophy of the inferior tarsal conjunctiva in both lower lids nothing else abnormal was noted.

At surgery a friable necrotic-appearing mass was found within the substance of the preseptal orbicularis muscle. The mass extended to the level of the orbital septum but did not penetrate it. No connection with the tarsus was found. Frozen sections at the time of surgery showed an inflammatory mass. It was excised and the resulting defect closed with a small rotational flap. Betamethasone sodium phosphate/betamethasone acetate suspension was injected into the area.

Examination of the permanent sections showed a chronic granulomatous response, with giant cell formation. The postoperative course was uneventful, and when the patient was seen seven months after surgery no recurrence was noted.

Case 3

A 47-year-old man gave a history of surgical incision of multiple lid lesions consistent with chalazia in the past. Four months prior to examination he developed an apparently similar lesion on his left lower lid. It arose approximately 3 mm inferior to the lower lid margin and was treated initially with warm compresses. When it did not disappear, he saw his ophthalmologist, who added topical antibiotics to the treatment regimen. There was still no improvement, and oral antibiotics were prescribed. Finally the lesion was incised and drained. The patient was referred for evaluation when there was still no improvement.

On examination a 10x24 mm mass was seen to occupy the medial two-thirds of the left lower lid (Fig. 3). The central part of the mass was violaceous, contained a fistula, and was located at the level of the inferior orbital rim. Medially the lesion extended up to the level of the inferior border of the tarsus.

Fig. 3 Case 3. 10x24 mm mass occupying the medial two-thirds of the left lower lid in a 47-year-old man.
However, no abnormality of the pretarsal part of the lid was present. The lesion was limited to the superficial part of the lid, with no deep fixation. Inspissated material was seen in the orifices of the meibomian glands on both lower lids. The rest of the ocular examination showed nothing of note.

At surgery a quantity of friable material was found to occupy the plane of the preseptal orbicularis muscle, extending down to but not through the orbital septum. The septum itself was moderately thickened. The lesion was followed up to the inferior border of the tarsus, but no direct connection with the tarsus could be seen. Frozen-section examination was consistent with chronic inflammatory tissue. The necrotic tissue was excised and the wound closed with small advancement flaps after the injection of betamethasone sodium phosphate/betamethasone acetate suspension.

Permanent sections showed chronic inflammatory cells with focal granulomatous reaction and aggregated histiocytes. Giant cells were seen, along with intra- and extracellular empty spaces consistent with lipid material (Fig. 4). When the patient was seen two months following surgery, no recurrence was noted.

**Discussion**

A chalazion is a chronic lipogranulomatous inflammation of the eyelid associated with retention of meibomian gland secretions. Histologically it is characterised by granulation tissue composed of lymphoid, epithelioid, and plasma cells, with the addition of giant cells with intra- and extracellular lipid.

The exact pathogenesis of this lesion is still uncertain. A chalazion normally presents as a firm nodular mass extending from the tarsus either anteriorly towards the skin or posteriorly towards the conjunctiva. Groenvall, in a study of 1693 chalazia, found a predilection for the lesion to occur more often in adults than in children and most often to involve the upper lid. Unfortunately in terms of the present report his study did not consider distance from the lid margin.

A variety of therapies have proved to be effective against this lesion. Conservative therapy, consisting of the application of moist heat either alone or in conjunction with lid scrubs or topical antibiotics, has been associated with cure rates of 25–80%. Persistent lesions respond to either surgical removal or intralesional steroid injection.

The lesions presented in this paper are somewhat unusual. Although when referred none of the patients showed any abnormality of the tarsal portion of the eyelid, in all the cases a history of the antecedent presence of a more typical chalazion was given. All the patients described an inferior migration of the lesion until the mass assumed its final location at the level of the inferior orbital rim. At surgery no connection with the tarsus was seen, though the lesions extended up to the level of the inferior border of the tarsus in the preseptal plane. Pathological examination showed typical mixed inflammatory cells with granuloma formation.

The mechanism for the inferior migration and final location is most probably explained by anatomical considerations of the lower eyelid. Hawes and Dortzbach have described the anatomy of the lower eyelid retractors. In the lower eyelid the orbital septum fuses to the capsulopalpebral fascia some
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distance from the inferior tarsal border. The main
terception of this complex is the inferior border of the
tarsus, though there is some extension on to the
inferior one-third of the posterior aspect of the tarsus
and a rudimentary extension for a very short distance
on to its anterior inferior surface.

The apparent sequence in these cases would seem
to be the following: (1) The process began in the
usual fashion in the tarsus. (2) The inflammatory
mass extended anteriorly and inferiorly, beneath the
junction of the pretarsal and preseptal parts of the
orbicularis. Although the pretarsal segment of the
orbicularis is tightly adherent to the tarsus, less of an
adherence is present between the preseptal portion of
the muscle and the underlying septum.17 A
potential space thus exists between the preseptal
orbicularis and the orbital septum, normally
occupied by a loose layer of suborbicular fascia.18
(3) With little fibrous barrier anterior to the tarsus
because of the limited anterior insertion of the orbital
septum/capsulopalpebral fascia complex, the inflam-
matory process easily entered the potential space
between the preseptal orbicularis and the orbital
septum. There the inflammation expanded with
subsequent loss of the original connection to the
tarsus. Although the resulting atypical appearance
delayed the diagnosis, the lesions responded to
standard therapy with good results.

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