Case Report

Atypical Presentation of a Rare Eyelid Tumor: A Case Report

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Introduction

Superficial Angiomyxomas (SA) are a group of uncommon soft tissue, myxoid mesenchymal tumours, found especially on the trunk. They mostly occur in people between 40 and 60 years of age and affect equally men and women. The eyelid is an uncommon site of SA. Only two more cases have been reported in the past. We present a case of Superficial Angiomyxoma of the eyelid in an adult male patient.

Case Report

A 39-year-old male presented with edema in the right upper eyelid (Figure 1), with no cutaneous lesions, which appeared the year prior to the hospital presentation and did not respond to cortisone therapy. He had no personal or family history of ophthalmic disorders, skin tumours or other types of cancer.

The MRI scan (Figure 2) revealed a heterogeneous tumoural mass located in the superior temporal quadrant of the orbit, that was not separated from the lacrimal gland. The preoperative diagnosis, based on the clinical findings and the MRI examination, was that of lacrimal gland tumour.

Incisional biopsy was performed (Figure 3), under local anaesthetic, in order to get a sample for the histological diagnosis. Intraoperative, when the orbital septum was incised, through the incision herniated a well circumscribed, nonencapsulated, red-pinkish mass, 1 × 0.5 cm in size, of mucinous consistency unseparated from the palpebral lobe of the lacrimal gland.

The first histological diagnosis was that of a Lymphangioma with the further recommendation of immunohistochemical tests to confirm it. The microscope slides (Figure 4) were reexamined and revealed a lesion consisting of spindle-shaped cells, distributed in a prominent myxoid matrix, with numerous thin-walled blood vessels. The immunohistochemical (Figure 5) analysis showed CD 34 intense positive cells and negative CD 31, EMA, D2-40, S100, MITF, SMA, GFAP, CD10, resulting in a final diagnosis of Superficial Angiomyxoma.

After the final diagnosis was made, the patient underwent a new surgical intervention in order to complete the excision of the tumour. The post-operative evolution was good and the patient remained tumour free for approximately one year after surgery when he developed right upper eyelid edema. The consult revealed the recurrence of the tumour and a new surgical intervention was scheduled. Intraoperative, the tumour invaded the orbital lobe of the lacrimal gland, but we managed to perform a complete excision (Figure 6).

Discussion and Conclusion

Superficial Angiomyxomas are rare, benign myxoid tumours, described for the first time by Allen et al. [1]. The word “superficial” was used in order to distinguish them from aggressive angiomyxomas which are highly locally aggressive tumours occurring preferentially in the female genital region [6,7].

Superficial Angiomyxomas are usually located on the trunk, lower limbs, head, neck or the genitals (penis and vagina) [2,3,6,8]. The present case, hence, is unusual, because the eyelid is a highly uncommon site for its occurrence [2,5,9].

The diagnosis is histological-characteristic features include extensive myxoid change with spindle or stellate shaped stromal cells that show no nuclear atypia or hyperchromasia. The presence of neutrophils is a diagnostic clue, as these are not present in other superficial myxoid lesions, such as digital myxoma or nerve sheath myxomas [3,8,10].

The treatment is surgical, total excision being curative [8,11,12]. This type of tumours do not metastasize, but local recurrence is possible (30%-40%), due to their infiltrative growth and the difficulties of a complete excision [5,8,10,12].
Figure 1: Patient presenting right upper eyelid edema.

Figure 2: MRI scan revealing a tumoural mass located in the superior temporal quadrant of the orbit.
   a) Coronal view
   b) Axial view

Figure 3 (a,b): Intraoperative aspect of the tumour.

Figure 4: Histological appearance of the tumour - a) H&E magnification 5x, b) H&E magnification 10x and c) H&E magnification 20x.

Figure 5: Immunohistochemical analysis – CD 34 intense positive cells.

Particularities of this Case

Atypical clinical presentation - no cutaneous lesions

Localization - only two more cases of superficial angiomyxoma in the eyelid are reported in the literature

First case of superficial angiomyxoma encountered by us in over 20 years of ophthalmic practice

First case of superficial angiomyxoma published by romanian ophthalmologists

Conflict of Interest

The authors declare no conflict of interest.

Patient consent

Obtained.

Ethics approval

All procedures performed in our case involving the human participant were in accordance with the ethical standards of the institutional and national research committee and with the 1964 Helsinki Declaration and its later amendments or comparable ethical standards. Informed consent was obtained from the participant included in the study, which is retrospective.
References


