Case Report

Microcystic adnexal carcinoma of the centrofacial region: a case report

Carcinoma annessiale microcistico centrofacciale: presentazione di un caso

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SUMMARY

Microcystic adnexal carcinoma is a rare, locally aggressive neoplasm with both eccrine and follicular differentiation and a high probability of perineural invasion of the centrofacial region. Given the histopathological features of this tumour, early diagnosis is essential for adequate management. This report refers to a case of microcystic adnexal carcinoma of the nasogenial region, with infiltration of the deep planes extending to the anterior wall of the maxillary sinus. Surgical treatment involved wide demolition of the centrofacial region followed by reconstruction using four locoregional flaps: an Indian flap and a Mustardé flap were used for cutaneous reconstruction; a septal flap to support the maxillogenous region; a mucosal flap to separate the nasal cavities.

KEY WORDS: Microcystic adnexal carcinoma • Mustardé flap • Forehead flap

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Microcystic adnexal carcinoma (MAC) is a rare eccrine gland tumour with low-grade malignancy and a low probability of metastasis. It generally affects the centrofacial region in older individuals. A characteristic histological feature is epithelial proliferation that progressively involves microcysts, strands, and cords as the tumour invades the deeper tissues from the skin surface. It has a severe tendency to neurotropic spread and presents with widely infiltrated margins. These characteristics make demolition surgery, especially in the centro-facial region, particularly challenging.

In this report, a case is presented of MAC of the centrofacial region, with involvement of the deep planes extending to the anterior wall of the maxillary sinus. The tumour was excised with wide demolition which was reconstructed using 4 local flaps: 2 chondro-mucosal flaps and 2 fascio-cutaneous flaps for skeletal support and restoration of the facial integument with good matching of skin colour.

Case report

In January 2007, an 81-year-old female was seen for an extensive, hard-rubbery neoplasm (~3 cm in diameter) fixed to the deep planes and involving the ala nasi and right nasogenial sulcus (Fig. 1). The case history revealed that a basal cell of the right genial region, arising in a setting of actinic keratosis, had been removed several years previously.

Contrast computed tomography (CT) and magnetic resonance imaging (MRI) revealed a neoplasm involving the nasogenial region, ala nasi, and right lip, with infiltration into the deep planes towards the anterior wall of the max-
illary sinus, without actually invading the sinus (Fig. 2). A biopsy specimen demonstrated the epithelial origin of the lesion and its proliferative nature.

Surgery included demolition of the right half of the nasal pyramid to the cartilage septum and part of the right nasal bone, with excision of the cheek skin and anterior wall of the maxillary sinus to the orbital rim, molar pillar, alveolar crest, and lip skin (Fig. 3).

Intra-operative frozen section analysis of the resection margins and the infra-orbital nerve was performed.

Reconstruction was carried out using a fasciocutaneous cheek advancement (Mustardé) flap to restore the aesthetic subunit of the cheek. The nasal pyramid was reconstructed using a left para-median forehead flap rotated 180° to restore the half dorsum, tip of the nose, and ala nasi. Cutaneous lining was achieved with a flip-flap harvested from the septum (Fig. 4). A superiorly pedicled flap, composed of septal cartilage and mucosa of the septum contralateral to the lesion, was rotated and sutured to the orbital rim to support the skin flaps; an inferiorly pedicled flap composed of the septal mucosa of the affected side was fashioned to create a septum to separate the nasal cavities (Figs. 5, 6).
Macroscopic examination showed contraction of the upper lip (1.4 cm in diameter); on incision, a hard-rubbery neoplasm with indistinct margins was observed in the dermohypodermal layer, with infiltration of the soft tissues extending to the maxillary bone. Histopathological examination of the uppermost part of the reticular derma showed the proliferation of micronests and solid cords consisting of mildly atypical monomorphic cuboid cells embedded in an abundant desmoplastic stroma alternating with keratinic cysts (Fig. 7). In the deeper plane, there were ductal structures with two layers of cuboid epithelial cells gradually replaced by microcysts and prevalently neurotropic monofilament cords (Fig. 8). Based on the histologic features, a
diagnosis of MAC with eccrine differentiation was established.
The differential diagnosis included several other neoplasms: benign adnexal tumours (syringoma and desmoplastic trichopithelioma), locoregional malignant tumours (malignant mixed tumour of skin appendages, sclerodermiform basal cell carcinoma of the skin, and neoplasms of the minor salivary glands), and metastasis of carcinomas occurring elsewhere, such as breast cancer 4.

Discussion
The classification of cutaneous adnexal tumours, especially eccrine tumours, continues to give rise to difficulties on account of the number of adnexal tumours variously identified and the variety of names used to describe them 5. The basic organising principle is to classify adnexal tumours according to histologically distinguishable differentiation (follicular, sebaceous, apocrine, and eccrine) and their degree of maturity (hamartoma, adenoma, benign, primitive, and malignant) 6. Once thought to be derived from mature adnexal cells, rather than via the transformation of genetically altered skin cells, the current most-accredited hypothesis is that adnexal tumours arise from adnexal differentiation of pluripotent cutaneous stem cells. This explains why some adnexal tumours can present concomitantly with three different modes of differentiation: pilosebaceous, eccrine, and apocrine 7,8.
Micro-cystic adnexal carcinoma is classified as a malignant neoplasm with prevalently eccrine differentiation, which in some cases manifests with pilosebaceous or apocrine differentiation 9. Other names used in the histopathological diagnosis of this tumour are sclerosing sweat duct carcinoma, eccrine epithelioma, and syringomatous carcinoma.
Excision of the neoplasm requires wide tissue demolition, because of the wide resection margins necessary. Repairing the resulting defect poses a considerable reconstructive challenge, particularly in the centrofacial region. While free flaps may meet tissue requirements, this cannot be said for their quality. In addition, as skin is lost due to the excision, reconstruction of the facial region with free flaps represents a limitation due to dyschromia after healing. Therefore, we chose local cutaneous, mucosal, and cartilage flaps to support the classic Mustardé and Indian flaps and to achieve an aesthetically acceptable outcome with more uniform facial skin chromatics 10-12.

References
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