SOLITARY KERATOACANTHOMA INVOLVING UPPER LIP: A DIAGNOSTIC DILEMMA - CASE REPORT AND A BRIEF REVIEW

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Abstract
Keratoacanthoma (KA) is a benign epithelial proliferative lesion which frequently occurs on the sun exposed areas of the skin. KA originates within the pilosebaceous apparatus of the skin and may be solitary or multiple. Solitary KA can be difficult to differentiate from squamous cell carcinoma (SCC) both clinically and microscopically. However, the clinical course of the lesion and its ability to self involute makes it a distinct clinical entity. Solitary KA appears on the vermilion border of the lower lips with some frequency. A case of keratoacanthoma involving the upper lip is reported presenting as an exophytic growth that resolved after excisional biopsy. Photographic documentation of the case along with relevant management protocol is discussed. This article emphasizes the significance of recognizing such lesion and discriminating it from SCC thus carrying diagnostic and therapeutic implications. However, in case of dilemma it is prudent to assume that the lesion is SCC unless proved otherwise clinically or histologically.

Key Words
Keratoacanthoma, squamous cell carcinoma, upper lip, self involution, regression after biopsy.

Introduction
Keratoacanthoma is defined as a benign and usually rapidly growing epithelial proliferation originating in the supra sebo-glandular portion of hair follicles (1-3). Lesions typically present as a solitary, firm, skin colored reddish papules that rapidly progress to dome shaped sessile, nodule with a central crateriform ulceration or keratin plug. The lesion shows a male preponderance and occurs frequently on sun exposed areas of face, neck, forearms and in older age group (2,4). KA can appear frequently on the vermilion border of lips thus mandating the dental professionals to be aware of the lesion. Etiology of KA still remains obscure; however actinic rays, HPV, trauma, genetic factors and immunocompromised status have been implicated (2-6). Solitary KA is often difficult to differentiate from well-differentiated SCC both clinically and histologically. The difference in diagnosis between these two lesions is of paramount importance since it carries therapeutic implications, thus prompting us to report this case. The case is unique in that KA is reported in a young individual involving upper lip contrary to the previous reported cases occurring on the lower lip and in older age group usually above 40 years of age (1-3, 5-8).

Case Report
A 26 year old male was referred from Department of Dermatology to the outpatient Department of Oral Medicine and Radiology for evaluation of an exophytic growth on the upper lip with one and half years of duration. There was no associated history of previous local trauma, but the patient recalls occurrence of acute onset ulcers involving mouth and lips before one and half years which subsided following local therapy. The lip lesion did not resolve and persisted with intermittent variation in the size. Patient was a smoker since 6-7 years and occasional gutkha chewer. Facial examination revealed well-demarcated sessile exophytic nodular growth located at the junction of vermilion border and the skin of upper lip (Fig.1). The lesion was non-tender, firm with no discharge. Palpation of neck did not disclose any suspicious lymph nodes and medical history was negative. An initial differential hypothesis of actinic cheilo-
sis, keratoacanthoma, squamous cell carcinoma, bacterial or mycotic infection was made and the patient was subjected for complete hemogram along with x-ray chest and Mantoux test. The test results showed no admissible changes. The need for biopsy was discussed with the patient but could not meet his approval.

Patient reported back after 2 months from his initial visit with no change in the size of the lesion, however, the lesion exhibited a yellowish white central zone of plug which was likely to be keratinous mass with raised edges. Subsequently the lesion was completely excised with a wedge excision followed by primary closure and the tissue was sent for histopathologic examination. The histopathology report showed proliferating parakeratinized stratified squamous epithelium with thickened stratum granulosum and slightly proliferative spinous and basal cell layers. At numerous places, epithelium showing downward growth forming budding islands with central portion of the island showing keratin (Fig.2 and 3). These down growth islands were surrounded by intense chronic inflammatory infiltrate predominantly comprising of lymphocytes, histiocytes and macrophages. The pathological findings were considered compatible with keratoacanthoma. The patient reported seven days after the biopsy with favorable healing observed.

Association of this observation with clinical course of the lesion coupled with the histopathological presentation confirmed the hypothesis of keratoacanthoma. The patient was regularly followed up at every two weeks of interval. There was no recurrence of the lesion after 6 months of follow up.

Discussion
KA is a well known benign squamous epithelial neoplasm with a strong clinical and histological similarity to well-differentiated SCC (2,3,5,6). Few authors are of the opinion that solitary KA represents an extremely low grade SCC (5,6). If so it fails to explain the static or regressive behavior of KA. In an attempt to discriminate the two lesions a comparative chromosomal aberration analysis study by Clausen et al. (9) showed that chromosomal instability was significantly higher in cutaneous SCC than in KAs. This difference in genomic expressivity of the two lesions has been correlated to their different biological behavior and pathogenicity. Irritation in the form of actinic rays, trauma, tars, virus which are common implications for both KA and SCC, the difference in the type of tumor produced was explained by experimental work by Ghadially (10) showing that the timing when the irritant took effect whether
Cases of solitary KA have been reported to persist for 6-8 weeks after undergoing spontaneous involution with good prognosis. Solitary lesions approximating 1-1.5 cm have been traditionally excised with a wedge excision (3, 5, 6). Such an approach has the advantage of obtaining entire lesion for histological examination and also the resulting scar is usually less disfiguring than the scar remaining after self involution of the lesion. In our case the patient recalls the static or regressive behavior of the lesion more suggestive of KA than a true neoplasm. Further, complete healing was observed after excisional biopsy which is unusual to be seen in a true neoplasm. Either spontaneous regression or healing by surgical intervention is considered as definitive evidence of KA (7). Thus the present case documents complete resolution of the lesion after surgical intervention indicative of KA. It is thus rationale to include KA in the diagnostic hypothesis list of every solitary nodular or ulcerative condition of lip. Further studies may be required to better understand the mechanisms involved in tumor proliferation, growth controlling factors, and involution of KA.

References
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