An Unusual Presentation of Squamous Cell Carcinoma as a Subcutaneous Nodule Without Overlying Skin Changes: A Case Report

Steven L Chang, Michael L Chang, Alyssa L Chang, Brian L Chang and Lawrence D Chang

Introduction

Cutaneous squamous cell carcinoma (SCC) arises from the squamous epithelia of the epidermis and is the second-most common skin cancer with a lifetime risk of about 10% [1-4]. Cutaneous SCC can arise de novo or evolve from a precursor lesion, such as actinic keratosis, Bowenoid papulosis, or epidermotropism verruciformis [2,5-8]. While the appearance the appearance of cutaneous SCCs varies greatly, they generally present as firm, hyperkeratotic, ulcerated, skin-colored papules or plaques found on sun-exposed skin and may be itchy or painful [2,3].

Nodular types of SCC have also been described and are frequently confused with other types of nodules, including nodular basal cell carcinoma, fibroxanthoma, Merkel cell carcinoma, nodular melanoma, seborrheic keratosis, lipomas, etc. [8]. The nodular type of SCC is commonly characterized by a keratin crust or scale, a central keratin mass, prominent vascularity or haemorrhage, and multiple colors. As with most other types of SCC, nodular SCC arises in the epidermis and then invades the dermis [9-10].

While nodular SCCs typically have some form of epithelial change, there have been a few reports of SCCs that present as subcutaneous nodules without significant epithelial manifestations [11-16]. Currently, it is believed that these SCCs are not primary skin cancers but rather metastatic: either from a visceral malignancy or as an in-transit metastasis from a prior SCC, similar to melanoma [11-16]. Regardless of the origin, the subcutaneous SCCs have been reported to be more aggressive and recur more frequently. It is therefore important to be aware of this type of SCC and treat it appropriately.

Here, we present a case of subcutaneous SCC and review the existing literature on disease course and management. The patient initially presented with a subcutaneous nodule without overlying skin changes and was subsequently diagnosed with SCC limited to the dermis without involvement of the epidermis. We hope that this case report will help raise awareness of this atypical presentation of SCC so that it can be properly managed.

Case Report

KF is a 72-year-old Caucasian male with a history of BCC, SCC, type II diabetes mellitus, and Parkinson’s disease who presented with a tender lump in his left outer cheek that had been slowly growing over the past two months. Exam revealed a 2 x 1 cm subcutaneous mass with tenderness, erythema, and warmth but no drainage or overlying skin changes (Figure 1). KF underwent routine skin surveillance examination 9 months prior to presentation; at that time, there was no sign of a lump or other skin changes (Figure 2).

At this time, it was thought that KF likely had an infected sebaceous cyst, so he was empirically treated with oral amoxicillin. However, 2 weeks later, the subcutaneous growth remained unchanged, though the symptoms of tenderness and erythema resolved. He was then referred to plastic surgery.

Figure 1: Image of KF’s left cheek at initial presentation that demonstrated a 2x1 cm subcutaneous mass that was tender, erythematous, and warm to the touch though lacked significant epithelial changes.

Keywords: Cutaneous squamous cell carcinoma; Nodular squamous cell carcinoma; Dermal squamous cell carcinoma; In-transit squamous cell carcinoma metastasis

Abbreviations: Squamous Cell Carcinoma (SCC), Basal Cell Carcinoma (BCC)
Upon initial presentation to plastic surgery, we were unsure what KF’s lesion was but could not rule out malignancy, so we performed incisional biopsy to get a tissue diagnosis. Pathology revealed deeply-invasive, moderately-differentiated SCC within the dermis but without a direct origin from the epidermis (Figure 3). KF underwent formal surgical excision with 4 mm margins and resection down to the subcutaneous layer (Figure 4). Final pathology revealed invasive SCC with clear peripheral and deep margins. 3-month surveillance follow-up revealed no recurrent SCC, so we thought KF was similar to all our other SCC patients and should be simply monitored with routine skin checks (Figure 5).

Unfortunately, 7 months after surgical resection, KF represented to plastic surgery with a rapidly growing subcutaneous mass without overlying skin changes of his left outer cheek in the same area as his prior SCC (Figure 6). Suspecting that the lesion was a recurrent SCC, we biopsied the lesion, which revealed well-differentiated SCC. Because KF’s SCC recurred so quickly and aggressively, we recommended consultation with radiation oncology for post-resection radiation therapy in an effort to prevent recurrence and optimize KF’s prognosis.

KF underwent surgical excision with 1 cm margins including superficial fascia, followed by skin graft reconstruction. Final pathology revealed clear peripheral and deep margins. 1 month after surgical resection to allow for stable soft tissue healing, KF underwent a 6-week course of 6000 cGy radiation to the skin cancer bed. 6 months later, KF is cancer-free and will undergo surveillance every 3 months for the next 2 years (Figure 7).

Discussion

This case report presents a 72-year-old male with a history of skin cancer who presented with a subcutaneous nodule without overlying skin changes and was subsequently diagnosed with invasive SCC that recurred despite complete surgical excision. This was the first time in our experience in which we managed a patient with SCC confined to the dermis without involvement of the epidermis that presented as a subcutaneous...
nodule. We hoped to present this case and review the existing literature to increase awareness of this more aggressive type of SCC.

Subcutaneous SCC has not been extensively studied and has only been presented in small case series. Three case series by Howe and Lang, Copcu et al., and Carucci et al. present a combined total of 28 patients with subcutaneous SCC confined to the dermis without an apparent epidermal origin [11-13]. The patients reported were very similar to KF: prior history of SCC and/or BCC, dermatologic findings of a subcutaneous nodule without significant superficial changes, pathology findings of invasive SCC in the dermis without clear origin in the epidermis, and aggressive or recurrent disease.

The authors individually reviewed the literature on subcutaneous SCC and proposed what they thought was the cause of these lesions. Prior studies have reported SCC presenting as subcutaneous nodules to occur as cutaneous metastases of primary visceral malignancies.14-16 However, despite extensive workup for visceral malignancies, the case series could not consistently identify a potential primary cancer in the patients [11-13]. Therefore, the authors hypothesized that the subcutaneous SCCs may actually represent in-transit cutaneous metastasis from prior SCC, similar to melanoma [11-13]. In one of the series, re-review of the pathology slides revealed dystrophic dermal-epidermal bridges, which the authors concluded supported their belief that the subcutaneous SCCs were in-transit metastases [11].

While there is no basic science to truly support the theory of in-transit metastasis of SCC, we believe that, if anything, it is a reasonable theory. These subcutaneous SCCs have no overlying epidermal origin, which casts doubt on them being primary cutaneous malignancies. In addition, many patients with this type of SCC do not have an identifiable visceral malignancy, and there must be some origin of these cancers.

In reviewing KF's case, as well as those presented by the few existing case series on subcutaneous SCCs, it is important to remember that SCCs have incredibly diverse presentations, and providers should operate with a low threshold when examining a new skin lesion in the elderly, sun-exposed patient. Because SCCs presenting as subcutaneous nodules can represent either cutaneous or extra-cutaneous metastases, it is important to consider the possibility that the patient has other malignancies. Because this was our first experience with a subcutaneous SCC, we were unaware of the existing thought that these could arise from primary visceral malignancies, so we did not pursue any additional diagnostic workup. In hindsight, KF never complained of any other symptoms that would suggest a visceral malignancy, which raises the question as to which patients, especially otherwise asymptomatic ones, should be worked up aggressively for visceral malignancy. Finally, the subcutaneous SCCs are often more aggressive with high rates of recurrence, necessitating adjuvant radiation therapy, as in KF's case and the patients of the three-case series.

Conclusion

This case report presents a 72-year-old male with a history of skin cancer who presented with a subcutaneous nodule, later found to be invasive SCC of the dermis that recurred despite complete surgical excision. SCCs are a very common, diverse, and potentially dangerous skin cancer, so it is important to have a low index of suspicion when evaluating new skin lesions in elderly patients. As subcutaneous SCCs may be a sign of cutaneous or extra-cutaneous metastasis and are particularly aggressive and recurrent, it is important to treat such cancers aggressively to optimize patient outcomes.

Conflict of Interest

The authors declare that they have no conflict of interest.

References