Simultaneous Occurrence of Two Squamous Cell Carcinomas Developing in a Nevus Sebaceous

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Abstract
Nevus sebaceous (NS), also known as organoid nevus, is a congenital skin hamartoma involving pilosebaceous follicle, epidermis and adnexal structures, which usually occurs in the head and neck region. During the first stage, which is seen in infancy and childhood, the lesion remains static, but can subsequently grow during puberty to become more evident and verruvious or nodular. Depending on the location and extent, nevus sebaceous causes only a cosmetic problem during childhood; however, the development of benign and malignant neoplasms on the lesion is the most severe complication during adulthood. The association between nevus sebaceous and various benign and malignant cutaneous neoplasms is well documented. Basal cell carcinoma (BCC) is the most common malignant lesion, and syringocystadenoma papilliferum (SP) is the most common benign tumor associated with nevus sebaceous. Development of squamous cell carcinoma within NS is very rare. The literature contains only a few reports of simultaneous multiple malignant tumors developing on nevus sebaceous. We present a review of the literature and report the case of a 42-year-old female patient with two squamous-cell carcinomas (SCC) simultaneously within a single NS.

Keywords: Malignant cutaneous neoplasms, nevus sebaceous, organoid nevus, squamous cell carcinoma

Introduction
Nevus sebaceous (NS) affecting ectoderm- and mesoderm-derived structures occurs predominantly on the head, neck and hairy skin. Although it remains stable throughout childhood, it grows and obtains a verruvious appearance as a result of the development of pilosebaceous-apocrine units due to the effect of androgens during puberty.1

The association of NS with many kinds of benign and malignant skin tumors is documented in detail by many cases and series within the literature. The global incidence of benign or malignant lesions on nevus sebaceous is approximately 15%.2 The most commonly observed benign tumors are syringocystadenoma papilliferum, apocrine cystadenoma, apocrine carcinoma, adnexal carcinoma, tubuloglandular sweat gland carcinoma, leiomyoma, metastasizing adenocarcinoma, and sebaceous carcinoma, whereas the most commonly observed malignant tumors are basal cell carcinomas (BCCs), trichoblastomas, squamous cell carcinomas (SCCs), and basosquamous carcinomas.1–3 The incidence of BCC, the most common form of malignant tumor, is less than 2%.4 Other malignant tumors such as SCC and adnexal tumors are rarely observed.

This study presents the case of a 42-year-old female with SCC development in two areas on NS, and attention is drawn to malignant neoplasms developing on NS background by reviewing the existing literature.

Case Report
A 42-year-old female patient presented to our clinic with increasingly growing lesions with occasional bleeding, located on hairy skin, the left ear and neck. The patient indicated that verruvious areas, which did not include yellowish hair in the background, had been present since childhood and had grown after puberty. She also stated that over the previous six months, indolent and wet nodular lesions had developed on this lesion, and that these bled spontaneously or following minor trauma (Figure 1). The patient did not respond to various topical treatments and systemic antibiotic treatment. Similar lesions were not observed in close relatives of the patient who was otherwise healthy. The patient’s medical history did not include previous skin cancer, immunosuppression, trauma or radiation exposure.

Dermatological examination revealed a yellow plaque lesion with occasional verruvious appearance, located on the hairy occipital area of scalp, neck and left ear region. In addition, two ulcerated nodules (22 × 34 mm and 25 × 43 mm) were detected on the left ear tragus helix and the left occipital region, respectively (Figure 1). No pathological symptoms were detected on regional lymph node examination. Magnetic resonance imaging detected no regional metastasis in lymph nodes or surrounding tissue. No pathological symptoms were detected in laboratory analysis of the patient’s samples (hemogram, biochemical, immunoglobulin levels). No metastatic symptoms were detected on abdominal ultrasonography or chest radiography.

The patient was referred to Plastic and Reconstructive Surgery; her lesions were completely excised with wide surgical margin and the defects were closed with a split-thickness skin graft. On histopathological examination of the excised material, hyperkeratosis, papillomatosis and irregular acanthosis were observed in non-ulcerated areas of the epidermis. On the dermis, a large
number of irregularly distributed mature sebaceous glands, apocrine glands and occasional immature hair follicles were observed (Figure 2). On sections of ulcerated areas of the epidermis, differentiated squamous cell proliferation was observed where dyskeratotic and atypical keratinocytes were found. Moderately differentiated cells were also observed to invade the papillary and reticular dermis (Figure 3). The clinical and histopathological findings are consistent with NS settled in a large area and presence of SCC in two distinct areas on this nevus. On a control examination after one week, sutures were removed and the patient was transferred to periodic follow-up examination. No infection or surgical complication was observed. No recurrence was observed during an 18-month follow-up period.

Discussion

NS is a complex hamartomatous skin malformation affecting 0.3% of newborns. This tumor belongs to the congenital lesions group that originates from primary epithelial embryonic germ cells—also called organoid nevus. Clinically, it is a well-demarcated and verrucous papillomatous lesion of yellowish color. Histopathologically, it consists of epithelial hyperplasia, papillomatosis, ectopic apocrine glands, mature hypoplastic and superficially localized sebaceous glands. It is often observed on hairy skin and less commonly on the neck and body. It was first defined by Jadassohn in 1895, and a three-stage process of nevus was defined by Mehregan and Pinkus. Papillomatous epithelial hyperplasia is the early infantile stage, in which hair development in the lesion area is low. During the pubertal period, overdevelopment of sebaceous glands occurs and epidermal hyperplasia and apocrine glands undergo maturation. In the third stage, during the 4th to 8th decades, neoplasms of epidermal, adnexal or mesenchymal origins develop on the nevus. On the other hand, there are several case reports in literature which indicate BCC development on childhood sebaceous nevus.
While complaints are generally only of an aesthetic nature during childhood, benign and malignant neoplasms develop in 15% of cases during adulthood.2 Benign or malignant neoplasm development is the most commonly observed and potentially serious complication of NS, and was reported to have a global incidence of 14.4% in a large case series (13.6% benign; 0.8% malignant, all of which were basal cell carcinomas).2

The most commonly observed biding neoplasm on NS is syringocystadenoma papilliferum.14 Apart from this, trichoblastoma, trichilemmoma, sebaceous, keratoacanthoma, seborrheic keratosis, syringoma, follicular poroma, nodular hyadentoma and apocrine cystadenoma may be observed.2,8,10,15 While BCC may, trichilemmoma, sebaceoma, keratoacanthoma, seborrheic keratosis, syringoma, follicular poroma, nodular hyadentoma and apocrine cystadenoma may be observed.2,8,10,15 While BCC is the most commonly developing malignant tumor on nevus sebaceous, widely differing rates of development are reported. In recent papers, the incidence of BCCs on nevus sebaceous is lower compared to those reported in earlier papers. The rate of BCCs reported until 1985 is 6%–50%.16 However, an 18-year study by Rosen et al. showed that many cases were actually trichoblastoma misdiagnosed as BCC. In addition, it was reported that the incidence of BCCs on nevus sebaceous was less than 2%.7

Cases of malignant tumors other than BCCs developing on NS are very rare. Those rare malignant neoplasms comprise SCC, adnexal carcinoma, apocrine carcinoma and eccrine poroma. A study of 997 cases by Domingo et al. reported nine aggressive carcinomas detected on nevus sebaceous, two of which were BCC, whereas seven were adnexal carcinoma.1 In two different nevus sebaceous series consisting of 180 and 140 cases, there was one case in each series.8,18 In addition, verrucous hyperplasia was seen in the epidermis of one quarter of the NS cases, and it was reported that, unless it is well-documented, the condition could be mistaken for well-differentiated SCC. Wilson et al. concluded that some reports within the literature, of SCC developing on NS, had histopathology representing pseudoepitheliomatous hyperplasia, and might have been misdiagnosed.8

In the literature review, we identified 22 cases in which malignant neoplasms other than BCC developed on the nevus sebaceous. Of those, 12 were BCC and 10 were adnexal carcinoma. Three of 11 case reports were adenocarcinoma and eight were SCC. In four cases where patients were defined with clinic and pathological details, there was more than one malignant neoplasm on nevus sebaceous.1,6,19,12 A case report documented the development of many benign neoplasms (trichoedanoma, trichoblastoma, syringocystadenoma papilliferum) in addition to two malignant neoplasms (sebaceous carcinoma, BCC).20 There is only one case reported in literature in which there were two different SCC lesions on nevus sebaceous.7

In our case, there were two different SCC developments on the same nevus sebaceous lesion; there were SCC developments on NS of hairy skin, neck and tragus, as well as on normal skin of hairy skin and tragus. Cribier et al. stated that in 596 cases, 79% of NS that showed tumor development were located on hairy skin.21 In our case, there was SCC development on NS located on both hairy and normal skin. Our case, which was clinically diagnosed as nevus sebaceous and SCC development, was referred for plastic and reconstructive surgery; lesions were excised with wide surgical boundary and the defects were closed with a partial-thickness skin graft. The diagnosis of NS and SCC was verified via histopathological examination of excised material.

There are differing opinions about the recurrence of malignant lesions on NS. A previous study reported no recurrence in a case of invasive SCC and BCC on NS during four years following total excision2 whereas there was one fatal case of recurrence 11 months after diagnosis.3 There was no recurrence in our case during 18 months of follow-up.

Longstanding opinion recommended early excision of NS seen in childhood, due to the perceived risk of malignant tumor development.15,21 In recent years, some authors have objected to this “prophylactic excision” because malignant tumors are rarely seen in children.22 According to these authors, malignant tumors are rare even among adults, and most BCCs are non-fatal when treated.

Nevertheless, the majority of authors defend the view that aggressive neoplasms developing on NS could cause important morbidity and, in rare cases, death, and therefore early excision is the safest course.23 In the literature, since there are many studies that report malignant neoplasms developing on NS, early excision has become a popular approach—not in all, but in cases with atypical clinical properties.23,24

Since literature contains limited cases of malignant neoplasms developing on NS, it is difficult to determine predicting factors or progression. Prophylactic excision of pre-puberty nevus sebaceous is a widely accepted approach. On the other hand, some studies contend that the rate of neoplasms developing on NS is low, and therefore patients could instead be followed and offered early excision in cases displaying atypical clinical properties.25 Although early treatment is still debated, once surgical removal of a tumor is decided, this can be achieved via simple excision or Mohs micrographic surgery method.26 In wider lesions, the excision area can be closed with a partial-thickness graft.26

Declaration of Interest

The authors report no conflict of interest.

References


