

VERRUCOUS SARCOIDOSIS OF THE SKIN SIMULATING SQUAMOUS CELL CARCINOMA

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ABSTRACT. Here, we report a case of a 38-year-old woman with a history of systemic sarcoidosis who developed cutaneous verrucous sarcoidosis simulating a squamous cell carcinoma. This modality of presentation is unusual in both caucasian patients and in woman and may represent a diagnostic challenge for dermatologists. (*Sarcoidosis Vasc Diffuse Lung Dis* 2013; 30: 70-72)

KEY WORDS: skin sarcoidosis, squamous cell carcinoma, granulomatous disease

INTRODUCTION

Sarcoidosis is a systemic disease of unknown etiology characterized by the formation of non-caseating granulomas in several organs. Lungs, mediastinal and peripheral lymph nodes, liver, spleen, skin, eyes, and parotid glands may be involved; central nervous system, heart, upper respiratory tract, bones, and joints are less frequently but usually more seriously affected (1, 2).

Cutaneous manifestations of sarcoidosis occur in approximately 25% of patients and may indicate onset of the disease (3, 4). Sarcoidosis of the skin usually peaks between the ages of 25 and 35 years and a second peak occurs in women from 45 to 65 years old (5). Various skin lesions associated with sarcoidosis have been described (6, 7). Most authors

divide these lesions into specific skin lesions, where histologic examination shows the typical sarcoid granulomas, and nonspecific skin lesions (6-8). Specific lesions are lupus pernio, infiltrated plaques, maculopapular eruptions, subcutaneous nodules and scars. The most important nonspecific skin lesion in sarcoidosis is erythema nodosum⁷. Many other clinical presentations of chronic cutaneous lesions have been reported, that's why cutaneous sarcoidosis is known as one of the "great imitators" in dermatology (9).

Here, we report a case of a 38-year-old woman with a history of sarcoidosis who had an unusual cutaneous presentation.

CASE REPORT

A 38-year-old woman presented with an asymptomatic nodular verrucous lesion on the forehead, surrounded by an erythematous halo, which had appeared during her second pregnancy (Fig. 1-2). The patient had been diagnosed systemic sarcoidosis two years before, when she had complained about respiratory and neurologic disorders and afterwards ematochemical examinations and instrumen-

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Fig. 1. Nodular verrucous lesion, surrounded by an erythematous halo, on the forehead of our patient

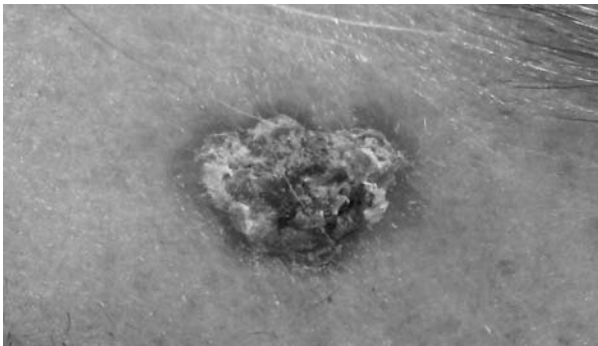


Fig. 2. Higher magnification of the nodular verrucous lesion

tal tests (chest TC, bronchoscopy, mediastinoscopy) had resulted positive. In the same year she had also been diagnosed cutaneous sarcoidosis after a biopsy from an erythematous patch appeared on her face during the first pregnancy. The biopsy obtained from the nodular lesion on the forehead was submitted for histological evaluation. This analysis revealed hyperkeratosis with parakeratosis of the corneum stratum and presence of dermal noncaseating granulomas consisting of epithelioid and giant cells surrounded by a lymphoplasmacytic infiltrate (Fig. 3). Stains and culture for acid-fast bacilli and fungi were negative. The clinical and histological characteristics were consistent with the diagnosis of verrucous sarcoidosis. The patient was treated with hydroxychloroquine and low doses of oral steroid with an adequate response.

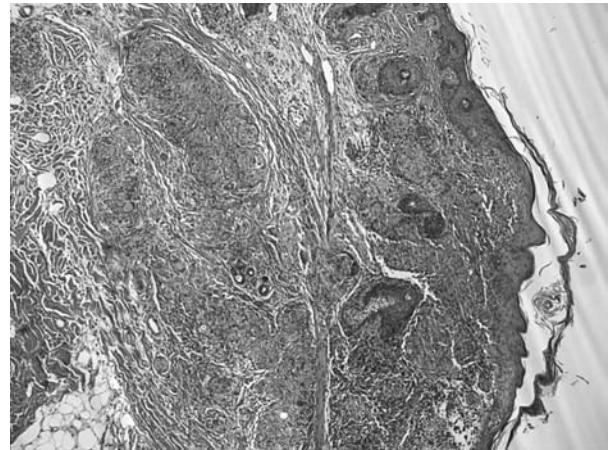


Fig. 3. Epidermal hyperkeratosis and parakeratosis and non-caseating granulomas in the dermis consisting of epithelioid and giant cells surrounded by a lymphoplasmacytic infiltrate

DISCUSSION

Cutaneous sarcoidosis is known as one of the “great imitators” in dermatology, because lesions may assume a vast array of morphologies (9). Some uncommon presentations include morpheaform plaques (10), lower extremity swelling (11), lichen nitidus-like papules (12), gyrate erythema (13), verrucous lesions (6), genital lesions (14, 15), palmar erythema (16), discoid lupus-like plaques (17), lesions mimicking polymorphous light eruption (6), plaques resembling lichen sclerosus (18) and pustular lesions. Therefore cutaneous sarcoidosis frequently represents a diagnostic challenge for physicians, which often results in a diagnosis of exclusion, established by the histological findings of non-caseating granulomas and after eliminating other pathological causes of skin lesions such as fungus, mycobacteria, and polarizable foreign bodies (19). Verrucous sarcoidosis is a rare specific cutaneous lesion. Our case is unusual because verrucous lesions in sarcoidosis are uncommon (20–26). Moreover, our case occurred in a Caucasian woman whereas all the cases described in literature were generally observed in male black patients and only one case was observed in a black female patient (20). Most patients had a systemic involvement principally of the respiratory system, as observed in our case.

In conclusion, we describe a case of cutaneous verrucous sarcoidosis simulating a squamous cell carcinoma. This modality of presentation is unusual in

both caucasian patients and in woman and may represent a diagnostic challenge for dermatologists.

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