Verrucous sarcoidosis associated with human papillomavirus infection: A case report

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Verrucous sarcoidosis (VS) is a rare variant of cutaneous sarcoidosis that most often appears on the lower extremities. It could represent a localized hypertrophic response over an area with underlying noncaseating sarcoi dal granulomas or a response secondary to a viral wart overlying a sarcoi dal plaque. A case of annular VS on the face is reported in the setting of widespread background cutaneous papular and plaque sarcoidosis.

CASE REPORT

A 51-year old African-American woman with a history of pulmonary sarcoidosis for more than a decade presented for evaluation of a facial rash. The rash had been present for several years and failed to respond to hydrocortisone 2.5% cream twice daily during this period. A laceration on the nasal bridge induced by blunt trauma from the nose pad of her eyeglasses occurred immediately before the appearance of expanding verrucous papillomatous changes in this area. She had no personal or family history of warts.

An annular plaque with verrucous surface features and central clearing was present on the bridge of the nose (Fig 1, A and B). Numerous skin-colored to violaceous papules and plaques, all demonstrating a smooth surface, were distributed on the forehead, temples, chin, and cheeks (Fig 1, A and B).

A 4-mm punch biopsy specimen from a dermal plaque on the right cheek showed sarcoi dal granulomas. Periodic acid-Schiff and Acid-Fast Bacilli stains showed no offending organisms, and no polarizable foreign material was present. A 4-mm punch biopsy from the verrucous lesion on the bridge of the nose demonstrated hyperkeratosis, acanthosis, hypergranulosis, and papillomatosis with parakeratosis overlying digitate projections of epithelium. Mild vacuolization of the granular layer was noted between the projections. Sarcoi dal granulomas, composed of histiocytes including scattered multinucleated giant cells, were present in the underlying dermis surrounded by a few scattered lymphocytes (Fig 2, A and B).

Sections from the punch biopsy of the verrucous lesion were sent for DNA quality assessment by β-globin reference gene polymerase chain reaction (PCR). DNA quality assessment was positive, and human papillomavirus (HPV) typing by PCR using the nested primer system was performed.1 HPV-PCR product was detected by the nested

Abbreviations used:
HPV: human papillomavirus
PCR: polymerase chain reaction
VS: verrucous sarcoidosis

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forslund antonsson primer (FAP) system in DNA extracted from the tissue sample, and the HPV-PCR product obtained from the sample was cloned and sequenced. The National Center for Biotechnology Information Basic Local Alignment Search Tool (NCBIBLAST) analysis of the sequence information obtained from the clones found the presence of HPV type candidate FA14 (Fig 3, A-C).

Imiquimod 5% cream 3 times a week was initiated but then discontinued because of severe irritation after 2 weeks. The cutaneous sarcoidosis was treated with hydroxychloroquine, 200 mg orally twice a day. Three months later, the patient returned after being lost to follow-up. The original annular verrucous lesion expanded, and 2 new annular verrucous lesions appeared overlying areas in which previous sarcoideal papules and plaques had been present (Fig 4, A-C). Despite the new and expanding verrucous lesions, the background smooth-surfaced sarcoideal papules and plaques had improved 80% to 90% since her previous visit. Treatment with imiquimod 5% cream was restarted once weekly to verrucous areas, and hydroxychloroquine was continued.

DISCUSSION

Sarcoidosis is known as one of the great imitators, with skin findings occurring in 20% to 35% of cases. In some cases, skin lesions are the only manifestation of sarcoidosis.2,3 These cutaneous findings are classified as specific when they are associated with cutaneous granulomas (Table I). The maculopapular form is the most common specific dermatologic manifestation of sarcoidosis.3 Erythema nodosum is the most common nonspecific dermatologic manifestation of sarcoidosis.3

VS is a rare, specific form of cutaneous sarcoidosis usually occurring in association with severe pulmonary disease.1,5 It most commonly affects the lower extremities.4,6 The differential diagnoses for verrucous sarcoidosis include verruca vulgaris, hypertrophic lichen planus, prurigo nodularis, and

![Fig 1](image1.png) Fig 1. A. An annular verrucous plaque with central clearing slowly expanded over several years after blunt trauma to the bridge of the nose. B. Scattered 1- to 8-mm hyperpigmented dermal papules and plaques without scaling on the cheeks.

![Fig 2](image2.png) Fig 2. A. Epidermis shows papillomatous architecture with hyperkeratosis and acanthosis. Parakeratosis overlies digitate projections (left pointing arrow) of epithelium. Hypergranulosis with vacuolization presents between the projections. A sarcoideal granuloma (downward arrow) is noted in the upper dermis. B. Sarcoideal granulomas with multinucleated giant cells are surrounded by a sprinkling of lymphocytes. (A and B, Hematoxylin-eosin stain; original magnifications: A, ×40; B, ×100.)
infectious processes such as blastomycosis and mycobacterial diseases.2,4

The pathophysiologic basis of VS is not clear. The verrucous epidermal response seen on histopathology could represent an alteration in cutaneous response to persistent mechanical irritation and, therefore, characterizes a type of hypertrophic localized neurodermatitis.7 A similar mechanism has been proposed for the epidermal change in psoriasiform sarcoidosis.8 The patient in this case did not complain of pruritus and denied rubbing the lesions. Furthermore, HPV was identified in this case supporting viral etiology of the verrucous hyperplasia.

Second, the HPV could serve as a sarcoideal priming antigen initiating the underlying cutaneous sarcoidosis.9-11 Of course, in this case, most facial cutaneous sarcoideal papules and plaques were not associated with overlying verrucous change. It is unlikely that 2 priming antigens were present.

A third explanation appears to be the most likely. The verrucous change may have been initiated by local inoculation of HPV after eyeglass trauma. The HPV then preferentially spread and rapidly grew in previously formed sarcoideal plaques. These plaques may represent immunocompromised districts.12

First, the granulomatous inflammatory process in the dermis may suppress immune mechanisms in the epidermis potentiating the growth of the virus. Second, the immunosuppressive effects of topical steroid treatment may have played a role in viral propagation and spread. This explanation is
supported by the occurrence of new warts only in areas overlying previous cutaneous sarcoidal granulomas without interval trauma at these sites.

A limitation of the positive HPV tissue sample is the finding of HPV FA14 in healthy skin and in the skin of immunocompromised renal transplant patients. On the other hand, the clinical appearance, expansive growth, and Hematoxylin-eosin stain findings support the viral etiology of these verrucous lesions. To the best of our knowledge, this is the first time HPV has been confirmed in verrucous sarcoidosis. Studies should be performed to determine if this is an unusual finding or if HPV is present in many cases of verrucous sarcoidosis. In addition to better understanding the pathophysiologic basis of this condition, the presence of HPV-associated verrucous sarcoidosis in some patients would lead to more focused prevention and treatment options.

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