Sarcoatosis is a complex, multisystem disease with an unclear cause. Research suggests that the pathogenetic mechanism of sarcoidosis is dysregulation of the immune system in individuals with a genetic predisposition who are subsequently exposed to inciting environmental agents.

Cutaneous sarcoidosis is often one of the earliest clinical signs of the disease and can be divided into 2 subclasses: specific and nonspecific lesions. Specific lesions are characterized by granulomas identified histologically and include macules, papules, plaques, annular lesions, lupus pernio, infiltration of scars, and subcutaneous nodules. Alternatively, nonspecific sarcoid lesions, including erythema nodosum, prurigo, or calcifications, are reactive inflammatory processes.

**Report of a Case** | A woman in her 40s presented with erythematous papules with perinasal and periorcular distribution involving both the upper and lower eyelids and an atrophic plaque on the dorsal surface of her neck. The patient complained of vaginal changes including itching, burning, tearing, pain with intercourse, and a painful lesion in the perianal area. Examination revealed an erythematous scaly plaque on the mons pubis with atrophic, white, discolored patches in the vaginal area without evidence of tearing (Figure 1).

The patient presented with a 5-week history of productive cough with exertional dyspnea, and her chest radiograph revealed upper-lobe calcified granulomas but no bilateral hilar lymphadenopathy. The patient’s pulmonary function tests revealed no airflow limitations, normal lung volume, and a mild reduction in the diffusing capacity of the lungs for carbon monoxide. The results of blood tests, including complete blood cell count, comprehensive metabolic panel, angiotensin-converting enzyme level, and erythrocyte sedimentation rate, were within normal limits.

Biopsies of the vulva and mons pubis revealed pauci inflammatory nodular granulomas consistent with sarcoidosis. On examination of the epidermis, vaguely psoriasiform epidermal acanthosis with hyperkeratosis was found. Within the dermis, tissue biopsy specimens demonstrated numerous, well-circumscribed nodular granulomas with multinucleated giant cells and lacking significant numbers of neutrophils (Figure 2A). Most of the granulomas were pauci inflammatory, but some had a cuff of lymphoplasmacytic inflammation. Finally, in areas of the epidermis there was transepidermal elimination (TEE) of the granulomas, a unique finding in the vulvar area of a patient with sarcoidosis.
All specimens tested negative for microorganisms by Grocott-Gomori methenamine silver and acid-fast bacilli stains and were not consistent with lichen sclerosus or lichen simplex chronicus. Hydroxychloroquine therapy had previously failed for this patient, and she was therefore prescribed mycophenolate (500 mg, twice a day) by her pulmonologist. In addition, we recommended 3-times-daily application of topical clobetasol ointment and nightly hydrocortisone suppositories.

Discussion | Although it is rare, sarcoidosis may occur in other regions of the female genital tract, including the uterus, ovaries, fallopian tubes, cervix, placenta, and vaginal wall.4 The differential diagnosis of granulomatous disease of the vulva includes tuberculosis, Crohn disease, syphilis, foreign body reactions, and lymphogranuloma venereum.5

Our case demonstrates TEE, a histologic phenomenon that, to our knowledge, has never been reported in a case of vulvar sarcoidosis. During TEE, the epidermis or the follicular epithelium undergoes a process of pseudohyperplasia to encompass the targeted material, which is eventually eliminated by maturing keratinocytes.6 Importantly, the epithelium does not undergo major structural remodeling and returns to normal after the targeted material is successfully removed.6 In the dermis of our patient’s vulva, the granulomas were the targeted material to be removed.

In summary, our patient presented with sarcoidosis of the vulva with histologic evidence of TEE. This case highlights that sarcoidosis can have an uncommon presentation, warranting extensive physical examination of patients presenting with possible sarcoidosis.

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Published Online: March 5, 2014. doi:10.1001/jamadermatol.2013.7204.

Conflict of Interest Disclosures: None reported.


Hypertensive Emergency, Matlike Telangiectasias, and Calciphylaxis in POEMS Syndrome

POEMS syndrome (polyneuropathy, organomegaly, endocrinopathy, monoclonal gammopathy, and skin changes) is a paraneoplastic syndrome caused by plasma cell dyscrasias. We report a case of POEMS syndrome presenting with a hypertensive emergency and acral matlike telangiectasias as novel manifestations of POEMS syndrome (POEMS). Also, to our knowledge, this is the first reported case of calciphylaxis and POEMS occurring in the absence of sclerotic bone lesions or a peripheral monoclonal gammopathy, highlighting the diagnostic challenges posed by this disease.

Report of a Case | A woman in her 30s with a 2-year history of distal polyneuropathy and matlike telangiectasias presented with a hypertensive emergency and peak blood pressure of 247/154 mm Hg. Workup revealed a serious pericardial effu-