To the Editor:
Elephantiasis nostras verrucosa (ENV) is a skin disorder caused by marked underlying lymphedema that leads to hyperkeratosis, papillomatosis, and verrucous growths on the epidermis. The pathophysiology of ENV relates to noninfectious lymphatic obstruction and lymphatic fibrosis secondary to venous stasis, malignancy, radiation therapy, or trauma. We present an unusual case of lymphedema and subsequent ENV limited to the arms and hands in a patient with scleroderma, an autoimmune fibrosing disorder.

A 54-year-old woman with a 5-year history of scleroderma presented to our dermatology clinic for treatment of progressive skin changes including pruritus, tightness, finger ulcerations, and pus exuding from papules on the dorsal arms and hands. She had been experiencing several systemic symptoms including dysphagia and lung involvement, necessitating oxygen therapy and a continuous positive airway pressure device for pulmonary arterial hypertension. A computed tomography scan of the lungs demonstrated an increase in ground-glass infiltrates in the right lower lobe and an air-fluid level in the esophagus. At the time of presentation, she was being treated with bosentan and sildenafil for pulmonary arterial hypertension, in addition to prednisone, venlafaxine, lansoprazole, metoclopramide, levothyroxine, temazepam, aspirin, and oxycodone. In the 2 years prior to presentation, she had been treated with intravenous cyclophosphamide once monthly for 6 months, adalimumab for 1 year, and 1 session of photodynamic therapy to the arms, all without benefit.
Physical examination showed cutaneous signs of scleroderma including marked sclerosis of the skin on the face, hands, V of the neck, proximal arms, and mid and proximal thighs. Excoriated papules with overlying crusting and pustulation were superimposed on the sclerotic skin of the arms (Figure 1).

A superinfection was diagnosed and treated with cephalaxin 500 mg 4 times daily for 2 weeks; thereafter, mupirocin cream twice daily was used as needed. She was prescribed fexofenadine 180 mg twice daily and doxepin 20 mg at bedtime for pruritus.

At 3-week follow-up, a trial of narrowband UVB therapy was recommended for control of pruritus. Two weeks later, a modified wet-wrap regimen using cloetasol ointment 0.5% twice daily covered with wet gauze followed by a self-adherent dressing was initiated only on the right arm for comparison purposes. This treatment was not successful. A biopsy taken from the left arm showed lymphedema with perivascular fibroplasia and epidermal hyperplasia consistent with ENV (Figure 2).

Two months after her initial presentation, we instituted treatment with tazarotene gel 0.1% twice daily to the arms as well as a water-based topical emulsion to the finger ulcerations and a healing ointment to the hands. A month later, the patient reported no benefit with tazarotene. She desired more flexibility in her arms and hands; therefore, after a discussion with her rheumatologist, biweekly psoralen plus UVA (PUVA) therapy was initiated. Five months after presentation, methotrexate (MTX) 15 mg once weekly with folic acid 1 mg once daily was added. The PUVA therapy and MTX were stopped 3 months later due to lack of treatment benefit.

The patient was referred to vascular medicine for possible compression therapy. It was determined that her vasculature was intact, but compression therapy was contraindicated due to underlying systemic sclerosis. She was subsequently prescribed mycophenolate mofetil 1000 mg twice daily by her rheumatologist. The options of serial excisions or laser resurfacing were presented, but she declined.

Elephantiasis nostras verrucosa is differentiated from elephantiasis tropica, which is caused by a filarial infection and is usually associated with infection.

Because options for medical treatment of severe ENV are limited, surgical debridement of the affected limb often remains the only viable option in advanced cases. A PubMed search of articles indexed for MEDLINE using the terms elephantiasis (MeSH terms) or elephantiasis (all fields) and scleroderma, systemic (MeSH terms) or scleroderma (all fields) and systemic (all fields) or systemic scleroderma (all fields) or scleroderma (all fields) or scleroderma, localized (MeSH terms) or scleroderma (all fields) and localized (all fields) or localized scleroderma (all fields) yielded only 1 other case report of lower extremity ENV in a patient with systemic sclerosis who ultimately required bilateral leg amputation. When possible, avoiding lymphostasis through compression and well documented. Characteristic features of systemic sclerosis include extensive fibrosis, fibroproliferative vasculopathy, and inflammation, which are all possible mechanisms for the internal lymphatic obstruction resulting in the skin changes observed in ENV. It seemed unusual that the fibrotic changes of lymphatic vessels in our patient were extensive enough to cause ENV of the upper extremities; lower extremity involvement is the more common presentation because of the greater likelihood of lymphedema manifesting in the legs and feet. Lower extremity ENV has been reported in association with scleroderma.

Regarding therapeutic options, Boyd et al reported a good response in a patient with ENV of the abdomen who was treated with topical tazarotene. Additionally, PUVA and MTX have been reported to be beneficial for the progressive skin changes of systemic sclerosis. Mycophenolate mofetil has been used in patients who fail MTX therapy because of its antifibrotic properties without the side-effect profiles of other immunosuppressives, such as imatinib. In our patient, skin lesions persisted following these varied approaches, and compression therapy was not advised due to the underlying sclerosis.

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control of any underlying infections is important in the treatment and prevention of ENV.³

REFERENCES