

Chromoblastomycosis Masquerading as Tuberculoid Leprosy

TO THE EDITOR:

It is not uncommon for the skin lesions of leprosy to be mistaken for other skin diseases but only rarely does the reverse occur. There are occasional reports of diseases such as histiocytic lymphoma, pachydermoperiostosis, Melkersson-Rosenthal syndrome, amniotic band syndrome, and lymphocytic infiltration of the skin being misdiagnosed as leprosy (1-5).

A 21-year-old male developed an asymptomatic, well-circumscribed, slightly erythematous, dry, flat, hypoesthetic, hypoallergic, oval plaque of 3 × 4 cm on the lumbar region (Fig. 1). There was no thickening of the cutaneous or peripheral nerves. The routine laboratory tests on blood and urine were normal. Blood VDRL and Mantoux tests were negative. Slit-skin smears from the earlobes and from the plaque did not show any acid-fast bacilli. Histopathological study of the biopsy specimen taken from the edge of the plaque revealed mild atrophy of the epidermis and a granulomatous infiltration in the dermis consisting of many "tubercles," some eroding the basal-cell layer of the epidermis. There were also a few Langhans'-type giant cells (Fig. 2). A clinico-histopathological diagnosis of tuberculoid leprosy was made, and he was treated with dapsone 100 mg daily and rifampin 600 mg once a month for 6 months. Since there was no therapeutic response even at the end of 6 months and there was periph-

eral extension of the lesion, a skin biopsy was repeated. Histopathology this time revealed tuberculoid granuloma in the dermis and numerous dark brown, thick-walled, rounded spores (sclerotic bodies) inside the Langhans' giant cells (Fig. 3), suggesting a diagnosis of chromoblastomycosis. There was no associated epidermal hyperplasia or neutrophilic abscesses in the epidermis or dermis. Culture of the biopsy specimen in Sabouraud's agar yielded growth which was identified to be *Cladosporium carrioni*. He was later treated by surgical excision of the lesion followed by skin grafting.

Cutaneous chromoblastomycosis, being a verrucous lesion, usually does not come under the differential diagnoses of leprosy. The well-defined, dry, hypoesthetic plaque in our patient simulated a tuberculoid leprosy lesion. However, the lack of therapeutic response to antileprosy drugs prompted us to repeat the biopsy and histopathological study which revealed "sclerotic cells," sug-

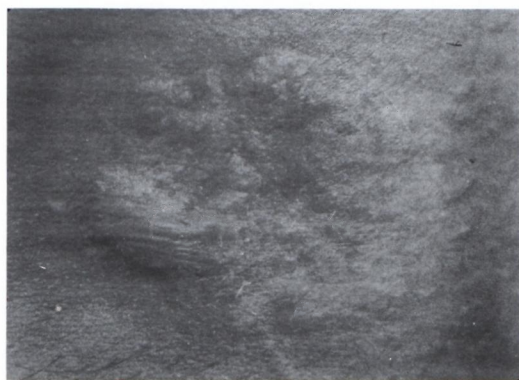


FIG. 1. Well-defined, dry, flat plaque of 4 × 3 cm on the lumbar area.

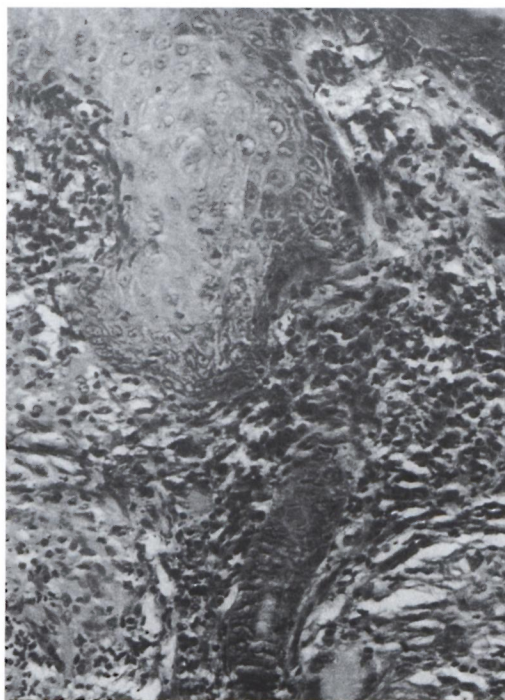


FIG. 2. Histology (H&E ×100) revealed tuberculoid granuloma in the dermis. The granuloma eroded the basal-cell layer of the epidermis.

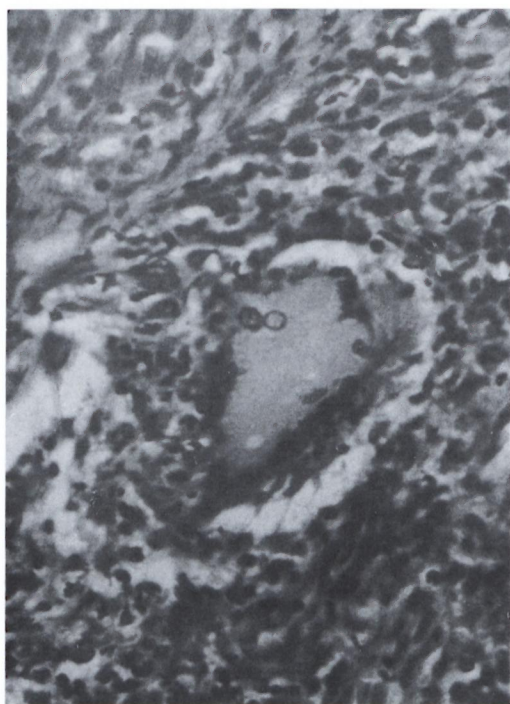


FIG. 3. Note two "sclerotic cells" within a Langhans' giant cell (H&E $\times 400$).

gesting that the previous leprosy diagnosis was wrong.

This report emphasizes the need for considering chromoblastomycosis also in the

differential diagnoses of tuberculoid leprosy and for a thorough search for "sclerotic cells" in all sections of biopsy specimens from skin lesions suspected to be tuberculoid leprosy, especially in areas where chromoblastomycosis is endemic. A tuberculoid granuloma in the dermis is only a tissue reaction to different etiologic agents that include bacteria, viruses, fungi, parasites, and foreign bodies.

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Suitability of a Skin-smear Examination Needle for Leprosy Screening by PCR

TO THE EDITOR:

Although leprosy has already been eradicated from some parts of the world, the disease still remains a major health problem, especially in developing countries. Recently, some investigators have reported the detection of *Mycobacterium leprae*, the causative agent of leprosy, by the polymerase chain reaction (PCR) (¹⁻³). We synthesized the PCR primers used in these reported studies, and compared their sensitivity in an attempt to use PCR as a practical screening test for leprosy in regions where the disease is endemic, and where

such trials are most needed. The subjects were patients in a leprosy sanatorium in Japan. The needle used for conventional skin-smear examination was used to obtain material from which DNA was extracted. In our experiments, although most patients were negative for acid-fast bacilli in the skin-smear test because all of them had been treated, DNA amplification was observed (The Table). This result demonstrated the sensitivity of PCR, since it was capable of detecting partially digested DNA in dead bacteria from the treated patients.

The primers reported by Woods, *et al.* (³) showed the highest sensitivity, but accuracy