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This department is for the publication of informal communications that are of interest because they are informative and stimulating, and for the discussion of controversial matters. The mandate of this Journal is to disseminate information relating to leprosy in particular and also other mycobacterial diseases. Dissident comment or interpretation on published research is of course valid, but personality attacks on individuals would seem unnecessary. Political comments, valid or not, also are unwelcome. They might result in interference with the distribution of the Journal and thus interfere with its prime purpose.

Erythema Nodosum Leprosum in a Case of Histoid Leprosy

TO THE EDITOR:

The occurrence of erythema nodosum leprosum (ENL) is an uncommon phenomenon in patients with histoid leprosy. While some authors postulate that patients with histoid leprosy do not develop the ENL type of reaction (2), others have reported the occurrence of ENL in histoid leprosy (1,3-6). We recently saw a patient with histoid leprosy who developed ENL lesions simultaneously.

A 45-year-old male patient presented with diminished sensation over his hands and feet for 15 months and fever, joint pain, testicular pain, and the development of recurrent crops of erythematous tender nodules, subsiding within 3-5 days, over the forearms and shins. Examination revealed a generalized infiltration of the skin with infiltration of the face and earlobes and moderately thickened, extremely tender, bilateral symmetrical ulnar, median and the terminal branches of radial and lateral popliteal nerves. In addition to the nodules arising from infiltrated skin, there were multiple skin-colored, dome-shaped, shiny, firm nodules of 2-5 mm arising from normal-looking skin over the lower back on both flanks. He also had bilateral epididymo-orchitis and bilateral palpable, tender, inguinal lymph nodes.

Investigations showed a slit-skin smear result of 5+ 0% from the infiltrated skin and 5+ 1% from the histoid nodules. Acid-fast bacilli from the histoid nodules were larger and did not form any globi. A skin biopsy of a histoid nodule showed localized collections of histiocytes, foam cells and lymphomononuclear cells aggregated in the mid- and lower dermis with unremarkable epidermis. Early and late reactions of lepromin were negative. His total leukocyte count was 24,800/cmm; differential count

THE FIGURE. Photomicrograph showing focal leukocytoclastic vasculitis consistent with ENL (H&E x140).
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was: polymorphs 71%, lymphocytes 26%,
eosinophils 0%, monocytes 3%, T and B
lymphocytes were 77% and 23%, CD4+ and
CD8+ were 40% and 37%, respectively. The
ENL histology was consistent with ENL (The
Figure).

The patient was started on the World
Health Organization multibacillary regi-
men with dapsone, rifampin and clofazim-
ine along with prednisolone 1 mg/kg/day.
Within 48 hr there was symptomatic im-
provement and the ENL lesions disap-
peared.

Why patients with histoid leprosy are rel-
atively immune to the ENL reaction is not
known exactly, but it is believed to be be-
cause histoid leprosy is an immunologically
relatively stable form in the multibacillary
spectrum of leprosy (6).

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Necrotic Erythema Nodosum Leprosum;
A Presenting Manifestation of Lepromatous Leprosy

TO THE EDITOR:

Usually lepromatous leprosy presents with
multiple hypopigmented macules, papules,
nodules or plaques, symmetrically distrib-
uted over the face, trunk and extremities (3).
Sometimes the presentation is unusual, in
the form of a single nodule, spontaneous
ulceration, histoid nodule or Lucio leprosy
(1, 2, 4, 5). But, to the best of our knowledge,
lepromatous leprosy presenting as necrotic
erythema nodosum leprosum (NENL) le-
sions has not been reported in the literature
so far.

We report here a patient with leproma-
tous leprosy who presented with necrotic
and pustular lesions. Demonstration of acid-
fast bacilli (AFB) from skin and pus smears
by Ziehl-Neelsen staining and histopathol-
ogy confirmed the diagnosis of lepromatous
leprosy with ENL.

A 48-year-old male presented with painful,
tender nodules and pustules with ne-
ecrotic ulcers on the extremities of 2 weeks’
duration. Some of the initial lesions were
nodules which had turned into pustules in
2–3 days’ time; other lesions were first no-
ticed as pustules. The pustules had been
covered with crusts after ulceration. There
were neither any constitutional symptoms,
such as fever and joint pain, nor any evi-
dence of iridocyclitis, conjunctivitis, neur-
ritis or orchitis. He had two similar epi-
isodes; one in January 1989 and one in
August 1990, each lasting for about 2 months
and subsiding with oral antibiotics and cor-
ticosteroids. In each episode multiple nodu-
lo-pustular lesions appeared in crops. These
episodes were associated with fever without
any history suggestive of systemic involve-
ment. There was neither a history nor clinical
evidence of hypopigmented, hypoesthetic
skin patch(es), paresthesia, glove-and-
stocking anesthesia or motor weakness. A
cutaneous examination revealed multiple,