Case Report

Bilaterally symmetrical lupus profundus with livedo reticularis as a sole cutaneous manifestation in a case of systemic lupus erythematosus

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INTRODUCTION

Lupus erythematosus profundus (LEP) is a rare variant of lupus erythematosus. It may occur as a separate disease or coexist with systemic lupus erythematosus (SLE) or discoid lupus erythematosus (DLE). Here we report a case of SLE with lupus nephritis with lupus profundus and livedo reticularis.

CASE REPORT

A 31-year-old female presented with the hardness of skin over the inner aspect of both thighs for four months associated with mild pain and pruritus. There was no history of preceding trauma. She had joint pains. She was diagnosed as SLE, without cutaneous lesions, with lupus nephritis and anti-ds DNA positivity two years back. She had been treated with steroids and other immuno suppressants. She underwent bilateral core decompression of femoral head six months back for avascular necrosis, probably induced by steroids. She had been under treatment with mycophenolate mofetil (MMF) for 6 months.

Examination revealed hard, indurated, slightly pigmented, non-tender plaques with ill-defined margins, shiny, irregular surface over the medial side of both lower thighs, measuring 10x10 cm over right and 20x10 cm over the left side (Figure 1).

No associated specific cutaneous lesions of SLE/DLE were found. Livedo reticularis over the left leg was present (Figure 2). There was no lymphadenopathy.

Routine laboratory tests like complete blood picture, erythrocyte sedimentation rate, blood sugar, liver function tests were within normal limits. X-ray chest was normal. Connective tissue profile showed positive antinuclear antibodies with 1:320 titer by indirect immune-fluorescent assay method. Anti-double stranded DNA was positive. The rheumatoid factor was negative. Incisional biopsy from the lesion over the right thigh showed the histopathology of fat necrosis, calcification,
perivascular infiltration of plasma cells, lymphocytes, eosinophils, and histiocytes (Figure 3). Direct immuno fluorescence was focally positive for IgA, IgG& IgM at DEJ, and around blood vessels in the subcutaneous tissue. Clinical, histopathological, and immunological features are suggestive of Lupus profundus.

Treatment was planned to increase the MMF dose from 500 mg 12th hourly to 500 mg 8th hourly and hydroxychloroquine sulphate from 200 mg 24 hourly to 200mg 12th hourly.

DISCUSSION

Kaposi first described this entity in 1883 as an association of subcutaneous nodules with lupus erythematosus.1 Subsequently, this finding has been recognized as lupus erythematosus (LE) profundus.2 The inflammatory reaction in LE prof takes place primarily in the deep corium and the subcutaneous tissue leading to deep indurated nodules or sharply defined plaques. The overlying skin usually appears normal, but there may be erythema, atrophy, ulceration, or poikilo-dermatous or hyperkeratotic changes.2 The lesions are most frequent on cheeks, but other predilection sites are upper arms, face, hands, chest, buttocks, and thighs. The histology is that of lobular panniculitis, and the direct immunofluorescence test shows a linear pattern at the dermo-epidermal junction. The overlying epidermis may or may not show evidence of changes in typical LE.2 MMF had been continued as therapy with increased dosage as it is accepted as a treatment modality for cutaneous lupus.3,4,5 Lupus profundus used to be a specific cutaneous criterion as per American College of Rheumatology criteria 1999 and Systemic Lupus Erythematosus International Consensus Conference 2012 but in the latest 2019 EULAR/ACR criteria for SLE, this was not mentioned.6

Livedo reticularis (LR) is a cutaneous physical sign characterized by transient (or) persistent, blotchy, reddish-blue to the purple net like discoloration in a cyanotic pattern. It is a manifestation of cutaneous blood flow disturbance that may occur in various physiological and pathological states. LR may be benign as in physiological cutis marmorata of infancy or severe as in the vasculitis of LE.7 The appearance of livedo reticularis in patients with SLE may herald central nervous system involvement. In this case, the patient does not have any symptoms or signs of CNS involvement. But the patient and her family members were educated about the CNS symptoms and informed them to meet the physician as soon as they first noticed such symptoms.

Bilaterally symmetrical location of lupus profundus over both medial thighs and association of lupus profundus with livedo reticularis had not been mentioned in the literature till now as per our knowledge.8

CONCLUSION

Lupus profundus can occur as sole specific cutaneous manifestation even two years after the diagnosis of SLE, as has happened in this case. The bilaterally symmetrical presentation of lupus profundus over both medial thighs, and association of lupus profundus with livedo reticularis,
as per our knowledge, had not been mentioned till now. As livedo reticularis can precede cerebral vasculitis, we should look for neurological symptoms and signs so that initiation of early treatment in case of positive symptoms can prevent mortality.

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