CASE REPORT

Membranous lipodystrophy: case report and review of the literature

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Abstract: Membranous lipodystrophy is a distinct type of membranocystic fat necrosis. It is associated with many local and systemic diseases, including vascular disorders. The histopathological changes which characterize this phenomenon are variably sized cysts in the fat lobules of the subcutaneous tissue, which are surrounded by eosinophilic membranes projecting into the cystic space. We report a case of secondary membranous lipodystrophy associated with both hypertension and venous insufficiency.

Keywords: Cysts; Fat necrosis; Ischemia; Lipodystrophy; Venous insufficiency

INTRODUCTION

Membranous lipodystrophy is membranocystic fat necrosis. Initially, it was described as polycystic lipomembranous osteodysplasia. Various clinical conditions, such as lupus erythematosus, diabetes mellitus, erythema nodosum, morphea, as well as trauma and vascular disorders have been found to exhibit similar membranous lipodystrophies and lipodystrophy-like changes. We report a case of secondary membranous lipodystrophy associated with hypertension and venous insufficiency.

CASE REPORT

A 60-year-old man presented with dyspnea and painful plaques on the lower extremities, which progressed slowly within two months. He was using antihypertensive drugs and had never consulted a physician for his varicose leg veins. The patient was a smoker and a habitual drinker. He had no history of connective tissue disease, Behçet’s disease, infection or trauma.

On physical examination, expirium was prolonged. Blood pressure was 180/100 mmHg. There were multiple, 2x4 cm erythematous indurated plaques on his distal lower legs. Multiple varices located bilaterally on the posterior aspect of his knee, around his ankle and on the medial side of his feet were identified (Figure 1). Laboratory findings were in normal ranges. The patient was diagnosed with chronic obstructive pulmonary disease by the specialist in chest diseases. Superficial venous insufficiency at the saphenofemoral junction was observed by colored Doppler ultrasonography.

Histopathology of the punch biopsy revealed slight infiltration of mononuclear cells and fibrosis in the dermis, septa and fat lobules. The walls of the vessels were thickened and infiltrated by lymphohistocytes. There were multiple cysts of irregular shape and variable size in the adipose tissue: the cysts were lined by an amorphous eosinophilic material (Figure 2A). Eosinophilic membranes were observed projecting...
into the cystic spaces (Figures 2B and 2C). The patient was diagnosed with membranous lipodystrophy due to vascular ischemia.

**DISCUSSION**

Membranous lipodystrophy was described by Nasu et al in 1973 as a genetic entity with membranocystic lesions of fat occurring in the long bones and associated with sudanophilic leukoencephalopathy. Later, membranocystic changes were noted in morphea profunda, lupus panniculitis and factitial ulcer, and the term “secondary membranous lipodystrophy” was proposed. Similar changes were also detected in diabetes mellitus, multiple myeloma, atypical mycobacterial infection, erythema nodosum, and vascular disorders, including arterial ischemia of lower limbs, varicose veins, thrombophlebitis and deep venous thrombosis.

The pathogenesis of membranous lipodystrophy is unknown. Idiopathic processes, enormous proliferation of the fat cell membranes, physiochemical interaction between the ground substance in connective tissue and fat droplets, free fatty acids released from degenerated fat cells processed by macrophages, metabolic disorders of lipids in mesenchymal cells, loops and fold in the basal laminae of fat cells in varying stages of lipid depletion at the time of necrosis; and ischemic injury of adipose tissue resulting from venous insufficiency are various mechanisms that have been proposed. Our case had hypertension and superficial venous insufficiency, which were thought to be the underlying causes. Several clinical presentations comprising tender or nontender nodules, ulcerated or atrophic plaques, depressed lesions, diffuse swelling, and echymotic lesions may occur. Our patient had tender, indurated plaques suggesting panniculitis, probably erythema nodosum or thrombophlebitis. The condition can only be diagnosed histopathologically by the presence of multiple, variably sized cysts in the panniculus, lined by an amorphous, homogeneous, eosinophilic membrane. This membrane may be flat and, in some cysts, be replaced by granulomatous inflammation. It often projects into cysts, creating a pseudopapilla or arabesque appearance. Our histopathological examination revealed similar findings, as well as dermal vascular hyperplasia.
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