

Giant Plaques on the Back with Gyrus-like Alteration: A Quiz

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A 21-year-old man presented with skin lesions. Six years ago, scattered skin-coloured papules were observed on the upper back with no obvious cause or significant subjective symptoms. The number and size of the papules gradually increased and became fused. Physical examination revealed 2 large skin-coloured plaques on the back and waist, measuring approximately $55 \times 50 \times 0.5$ cm and $16 \times 8 \times 0.2$ cm, respectively. The surface exhibited gyrus-like alteration with a soft texture and clear boundaries. Scattered skin-coloured papules with a tendency to fuse were observed on the left shoulder, sacrum, and buttocks (Fig. 1). Histopathological

examination revealed a slightly pigmented basal epidermal layer with a small amount of mature adipose tissue in the superficial dermis (Fig. 2).

What is your diagnosis?

1. Giant skin tag
2. Lipofibroma
3. Nevus lipomatosus cutaneus superficialis
4. Focal dermal hypoplasia

See next page for answer.



Fig. 1. Skin-coloured plaques on the back and waist, whose surface exhibited gyrus-like alterations with clear boundaries. Scattered skin-coloured papules with a tendency to fuse were observed on the left shoulder, sacrum, and buttocks.

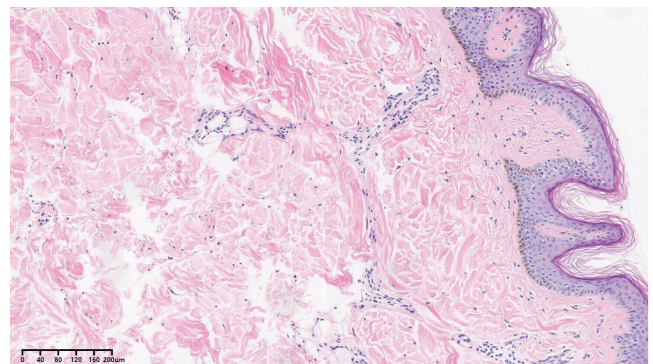


Fig. 2. Histopathological examination revealed a slightly pigmented basal epidermal layer with a small amount of mature adipose tissue in the superficial dermis (HE $\times 100$).

Giant Plaques on the Back with Gyrus-like Alteration: A Commentary

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Diagnosis: Nevus lipomatosus cutaneus superficialis

Nevus lipomatosus cutaneus superficialis (NLCS) is a rare cutaneous hamartoma characterized by the presence of mature ectopic adipose tissue within the dermis. The pathogenesis of this disease remains unclear, with the main theories including: (i) lipomatous metaplasia caused by degenerative changes in dermal connective tissue; (ii) a true nevus formed by developmental displacement of adipose tissue; (iii) differentiation of perivascular adipoblasts into adipocytes (1).

In 1921, Hoffmann and Schuhelle were the first to report this disease and classified it into 2 clinical types: classical and solitary. The classical type mostly manifests at birth or before the age of 30, preferentially involving the lower back and buttocks. The skin lesions present as multiple soft papules or nodules with skin-coloured or light-yellow hues; the overlying skin is smooth or wrinkled, occasionally exhibiting a cerebriform appearance. Meanwhile, the solitary type is characterized by a single skin-coloured papule or nodule, with no obvious age predilection or anatomical site preference. Most cases are asymptomatic, although a few might show abnormal growth or morphological changes, including giant NLCS, comedo-like alterations, foul-smelling secretions, and ulcerative lesions (2).

The characteristic histopathological features of NLCS include the proliferation of mature ectopic adipose tissue within the dermis, accounting for 10–50% of the lesion. Adipocytes are often distributed in clusters around blood vessels or eccrine glands; they may also appear as single cells between collagen bundles. Increased density of collagen fibres and fibroblasts, along with perivascular infiltration of mononuclear cells and spindle cells, is occasionally observed. Additionally, the epidermis shows acanthosis, basket-weave hyperkeratosis, and increased basal pigmentation, as well as focal elongation and fusion of epidermal ridges (3).

This disease should be differentiated from disorders such as giant skin tag, lipofibroma, and focal dermal hypoplasia. While giant skin tag is clinically similar to the solitary type of NLCS, histopathological examination

can confirm the absence of adipocytes in the dermis of giant skin tag cases. Meanwhile, lipofibroma typically presents as a pedunculated mass, characterized by a mixture of adipose tissue and dense collagen fibres around skin appendages. Focal dermal hypoplasia (Goltz syndrome) shares similar histopathological features with NLCS; however, in addition to ectopic adipose tissue proliferation in the dermis, it exhibits widespread collagen fibre rarefaction and absence of skin appendages. In cases where adipocyte components in NLCS lesions contain numerous spindle cells (representing immature adipocytes), they should be differentiated from other dermal spindle cell lesions such as neurofibroma, leiomyoma, and dermatofibroma (4).

Treatment is typically not required for NLCS except for cosmetic purposes. Surgical excision is curative for this condition, and no recurrence has been reported to date. Other treatment modalities include electrocautery, cryotherapy, phosphatidylcholine injection, and CO₂ laser therapy (5). Notably, the patient in this case had giant skin lesions, and reconstructing tissue defects after lesion excision was challenging. Thus, early surgical excision is necessary for such cases to achieve a favourable prognosis. Furthermore, potential abnormal morphological changes should be vigilantly monitored.

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