Letter to the Editor

Solitary Reticulohistiocytoma (SRH) of the skin [1,2] is a benign mesenchymal tumor with histiocytic proliferation of unknown etiology, belonging to reticulohistiocytosis (RH) [3,4]. In contrast to multicentric reticulohistiocytosis (MRH) [1,3,5], the other clinical setting of RH, SRH has no associations with arthritis or systemic lesions. It usually presents an asymptomatic papule or nodule, often sited on the trunk, although any site may be affected. Different from the necessity of systemic therapy for the challenging clinical process of MRH [6,7], the only curative therapy for SRH is surgical removal, or it may even regress over the course on observation. On the other hand, however, it often needs to be distinguished from so many skin tumors including some kind of malignancies [2,3,8]. Among such differential diagnosis, the most important problem is its discrimination from melanoma in view of the unfavorable prognosis for the latter. Here we report a case of SRH mimicking clinical appearance of amelanotic melanoma. A 29-year-old male visited our outpatient office presenting a surface-eroded nodule 7 mm in diameter on his rt. lateral abdomen (Figure 1). He was otherwise healthy. He had had that lesion for 2 month before his visit. Relatively rapid growth of the tumor and its peculiar appearance at the gross level led us to raise amelanotic malignant melanoma as the diagnostic possibility. The pathological findings of hematoxylin and eosin stained sections of the excisional biopsy revealed large hematoma just beneath the epidermis, and circumscribed dermal nodule (Figure 2). The nodular lesion was composed of multinucleate epitheloid giant cells with an eosinophilic, glassy appearance-cytoplasm, histiocytes and inflammatory cells mainly by neutrophilic infiltrate (Figure 3). Additional findings include scattered xanthomatized cells. These were histologically pathognomonic characteristics of reticulohistiocytoma. Immunohistochemistry exhibited that such massive distribution of epitheloid histiocytes were immunoreactive for anti-CD68 antibody, in contrast to the negative stains for S-100, HMB-45 and melan-A, allowing distinction from malignant melanoma (Figure 4). Our patient has been disease-free for a year. An asymptomatic skin nodule of subtle appearance should prompt consideration of reticulohistiocytoma in the differential diagnosis.

Figure 1. Clinical appearance of a solitary reticulohistiocytoma in the present patient.

Solitary reticulohistiocytoma of the skin mimicking amelanotic malignant melanoma.

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**Figure 2.** Histological findings from the surgical specimen. (25X).

**Figure 3.** Characteristic intralesional multinucleate giant cells, histiocytes and inflammatory cells. (200X).

**Figure 4.** Immunoreactivity of multinucleate giant cells for CD68. (200X).
Conflict of Interest

The author has nothing to disclose.

References


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