Clinical Image

Volume 10 (2024) Issue 15

ISSN: 2379-1039

Trichoscopic features in scalp dermatomyositis

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Abstract

Scalp involvement in Dermatomyositis (DM) represents a diagnostic challenge. Its clinical presentation is often atypical, and can be observed in other collagenosis. In this report, we outline the trichoscopic signs of two cases of DM of the Scalp (SDM).

Trichoscopy revealed cicatricial alopecia (absence of follicular openings with white structureless areas), enlarged tortuous capillaries, linear branched vessels, curved branched vessels, lake-like vessels, perifollicular and interfollicular scales. We observed an uncommon sign that was the «large yellowish clods», corresponding to giant follicular plugs.

In the field of connective tissue diseases, dermoscopy gained popularity for the evaluation of inflammatory skin conditions. The clinico-dermoscopic correlation seems to be an accessible substitute for biopsy. Trichoscopy is a rapid, easy to apply, non-invasive technique that enables the diagnosis and should be performed routinely in all patients with dermatomyositis.

Keywords: Trichoscopie; Alopecia; Dermatomyositis.

Introduction

Scalp involvement in Dermatomyositis (DM) represents a diagnostic challenge. Its clinical presentation is often atypical, and can be observed in other collagenosis, which can delay the diagnosis and consequently the treatment of this affliction. Its dermoscopic characteristics have been poorly described. In this report, we outline the trichoscopic signs of two cases of DM of the Scalp (SDM).

Case Presentation

A 56-year-old female patient was diagnosed with amyopathic dermatomyositis since 1998. She had a characteristic rash of dermatomyositis consisting in heliotropic periorbital erythema, Gottron's papules, violaceous scaling patches in her upper extremities, and periungual erythema with telangiectasias. During the follow up, she never developed an inflammatory myopathy but had several cutaneous flare-ups. Repeated explorations did not show evidence of malignancy, infection, or association with another collagenosis. Topical corticosteroids, methotrexate, antimalarials lead to partial improvement. Recently, the patient developed persistent scalp itchiness. The scalp examination showed a large erythematous, squamous and telangiectasic alopecic plaque of the vertex (Figure 1). Trichoscopy revealed cicatricial alopecia (absence of follicular openings with white structureless areas), enlarged tortuous capillaries, linear branched vessels, curved branched vessels, lake-like vessels, perifollicular and interfollicular scales (Figure 2).

The second patient was a 65-year-old female patient who had been treated for DM since 2013. She presented with several erythematous and pruritic, alopecic plaques in the frontal, parietal and occipital areas. These lesions were associated with poikiloderma and had been evolving for 1 year. Trichoscopic examination revealed signs of scarring alopecia, blood vessels and perifollicular skin similar to the first patient, accompanied by a perifollicular hyperpigmentation.

In both our patients, we observed «large yellowish clods» (Figure 2), corresponding to giant follicular plugs. Mycological samples were negative, and scalp biopsies ruled out other diagnoses and confirmed the diagnosis of SDM.



Figure 1: Alopecic plaque on the vertex.



Figure 2: Linear branched vessels (green arrows), curved branched vessels (bleu arrows), large yellow clods (yellow arrows), white structureless area (white star), yellow structureless area (black star), lake-like vessels (red arrows).

Discussion/Conclusion

SDM mainly affects women. Clinically, it is characterized by alopecia, pruritus and/or a burning sensation [1]. These chronic scalp symptoms impair the patient's quality of life and require adequate treatment. Trichoscopy is a non-invasive and helpful tool for the diagnosis of SDM. Jasso-Olivares and al. reported the biggest study describing clinical and trichoscopic features of 31 cases of SDM [1]. The most common

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findings included enlarged tortuous capillaries and peripilar scales. Tufting with three or more hair shafts emerging together, interfollicular scales, bushy capillaries, interfollicular and perifollicular pigmentation and vascular lake-like structures were least common. Zychowska et al. described other trichoscopic features of SDM such as yellow dots, hair diameter diversity, and interfollicular honeycomb pigment pattern [2]. The main differential diagnosis is discoid lupus with some dermoscopic similarities such as white dots and patches, brown honeycomb pattern, interfollicular scales, dotted vessels, arborizing vessels. However, enlarged capillaries with a tortuous shape, linear and curved branched vessels, and lake-like vessels seen in our patients seems to be more specific of SDM [3-5]. For our patients, trichoscopy showed specific findings of SDM. We also observed "large yellow clods" which were not previously described in the literature and they correspond to giant follicular plugs.

In the field of connective tissue diseases, dermoscopy gained popularity for the evaluation of inflammatory skin conditions [5]. The clinico-dermoscopic correlation seems to be an accessible substitute for biopsy. Trichoscopy is a rapid, easy to apply, non-invasive technique that enables the diagnosis and should be performed routinely in all patients with dermatomyositis.

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Manuscript Information: Received: May 15, 2024; Accepted: June 07, 2024; Published: June 10, 2024

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Citation: Lahouel M, Mabrouk RSE, Mokni S, Fetoui NG, Thabouti M, Saad S, Rejeb MB. Trichoscopic features in scalp dermatomyositis. Open J Clin Med Case Rep. 2024; 2253.

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