Alternative topical therapy for post-Scabietic Mastocytoid nodules using pimecrolimus cream 1%

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INTRODUCTION

Post scabietic nodules are one of the commonest manifestations that are seen after the cure and recovery of scabies. A recent review by Sharquie et al showed that post scabietic nodules are non-excoriated, pigmented dusky red in color and had a tendency to occur in specific predilection areas like the scrotum and penile shaft. Darier sign similar to that observed in mastocytosis was reported in 8.82% and hence the term “post scabietic mastocytoid nodules” was recently applied [1].

The histopathology of post scabietic nodules is a dynamic reaction as the previous studies showed that the main cells in the inflammatory infiltrate are eosinophils and lymphocytes and hence the terms pseudolymphoma and eosinophiloma were used respectively in the past, however, no stain was carried out for the detection of the mast cells [2,3]. While a recent study that used H&E stain and Giemsa stains revealed that there were three patterns of the inflammatory infiltrate in the post scabietic nodules: eosinophils predominant variant in 36.36%, mast cells predominant variant in 27.27% and a mixed cellular variant of eosinophils and mast cells in 36.36% [1].

The already known modalities for the treatment of the post-scabietic nodules including the use of topical and oral corticosteroid, but the drawbacks of these treatments are the plenty side effects [4]. The current study applied pimecrolimus cream 1% as an alternative modality for the treatment of the post-scabietic nodules relying on its pharmacological actions through stabilizing the cell membranes, inhibiting the release of mediators from the mast cells and basophils and reducing the activation of eosinophils [5].

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MATERIALS AND METHODS

This observational therapeutic study was done in the center of Dermatology and Venereology -Baghdad Teaching Hospital, Medical City; Baghdad, Iraq during the period from April 2017- November 2019.

Thirty-four patients with post-scabietic nodules were included in this study, based on the medical history of scabies and clinical examination. Formal consent was taken from all enrolled patients after the nature of the current study was fully explained and ethical approval was performed by the Scientific Council of Dermatology and Venereology-Arab Board for Medical Specialization. The patients were treated with pimecrolimus cream 1% applied to the nodules twice daily for one month.

Assessment of the patient response to treatment was based on the clinical judgment (change in the itching subjectively by the patients, size and the number of the nodules objectively and by photograph assessment). The response was classified into A- Complete response (complete disappearance of the nodules and itching). B- No response (no change in the size, number of the nodules and the severity of itching).

C- Partial response (any response that is less than the complete response).

Ethics Statement

Formal consent was taken from all enrolled patients after the nature of the current study was fully explained and ethical approval was performed by the Scientific Council of Dermatology and Venereology-Arab Board for Medical Specialization.

RESULTS

Following the four-week daily application of pimecrolimus cream 1% to the post scabietic nodules, the response of the patients was as the following:

a- Seventeen patients (50%) showed a complete response as there was complete disappearance of itching and the nodules at the end of the treatment period (Fig. 1).

b- Nine patients (26.47%) showed a partial response as there was partial reduction in the size, number of the nodules and itching (Fig. 2).

c- Eight patients (23.52%) showed no response as no change in the size and number of the nodules and in the severity of itching.

Side effects were not detected in any of the treated patients. Regarding the relapse rate after cessation of topical pimecrolimus, it could not be determined as the patients did not attend the follow-up visit after they got relief from their disease.

DISCUSSION

Treatment of post scabietic nodules is often problematic as they respond variably to topical, oral steroid +/- antihistamines that are usually followed by a high relapse rate after stopping the therapy [4].

Most recent histopathological evaluation of the post scabietic nodules have shown that the mast cells are...
the predominant cells either alone or admixed with eosinophils in the inflammatory infiltrate of post-scabietic nodules. These findings have led us to the use of pimecrolimus cream 1%, a calcineurin inhibitor that has specific effects on the mast cells and eosinophils mainly by stabilizing the cell membrane and preventing their rupture and release of granules [5]. Two previous case reports have evaluated the use of tacrolimus ointment 0.1%, another calcineurin inhibitor, for the treatment of post-scabietic nodules, while only a single case report of successful use of pimecrolimus cream in a single case of steroid-resistant nodular scabies have been reported [6-8].

Hence we planned to use topical pimecrolimus 1% twice a day for one month as a therapy for these post-scabietic nodules and the results were encouraging as 50% of the patients had complete response with complete disappearance of itching and nodules and 26.47% had partial response with reduction in itching and nodular number and size while 23.52% had no response after one month of treatment. Notably, no side effects were reported during the treatment period.

Unfortunately, the limitation of the current study was the difficulty in assessing the relapse rate following the treatment cessation, as most of the treated patients were reluctant to attend the follow-up sessions.

**CONCLUSION**

The present work showed that pimecrolimus cream 1% is an effective alternative treatment for the post-scabietic mastocytoid nodules acting through its pharmacological action in the suppression of the inflammatory process in these nodules.

**Statement of Human and Animal Rights**

All procedures followed were in accordance with the ethical standards of the responsible committee on human experimentation (institutional and national) and with the Helsinki Declaration of 1975, as revised in 2008.

**Statement of Informed Consent**

Informed consent was obtained from all patients for being included in the study.

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