New Topical Treatment for Atopic Dermatitis

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ABSTRACT

The primary approach for managing atopic dermatitis (AD) involves the use of topical corticosteroids as the first-line treatment. While high-potency topical corticosteroids have shown to be effective, they come with an increased risk of local and, rarely, systemic adverse effects. Additionally, patients often experience a relapsing and remitting course. A revolutionary topical treatment for psoriasis and AD has recently received patent approval from the Spanish Ministry of Industry, Trade, and Tourism. This innovative treatment, presented in the form of a lotion, includes a combination of clobetasol, papaverine hydrochloride, spironolactone, a milk-peptide complex, and propylene glycol. An 18-year-old female presented with AD on the back of her neck and scalp. The patient had no significant past medical history and primarily complained of intense pruritus in the AD lesions. The patient received guidance to apply our recently patented lotion, Psorisbye, once a day for 5 days. In total, 50 ml of Psorisbye was utilized over 4 days. On the fifth day, the patient underwent an examination at the outpatient clinic. The patient reported a significant improvement in pruritus sensations and observed a reduction in scaled lesions. Upon evaluating our patient, a comparison of the lesions before and after applying the topical treatment for 4 days revealed a notable improvement in the SCORAD index, decreasing from 49.95 to 0. While the results of Psorisbye in this case show promise, it is crucial to conduct further studies with larger sample sizes and extended follow-up periods to validate the findings presented in our case report.

Keywords: Atopic dermatitis, SCORAD, SCORing Atopic Dermatitis, Topical treatment.

1. Introduction

Atopic dermatitis (AD) stands as the prevalent chronic inflammatory skin ailment, impacting approximately 230 million individuals globally. In developed nations, its prevalence varies, with rates ranging from 10% to 25% in children and 7%–10% in adults [1].

Clinically, AD is identified by severe itching, dry skin, and eczematous lesions, primarily affecting flexures, of the head, neck, and hands [1], [2]. Furthermore, disruptions in attention and low sleep quality, likely associated with itching, can manifest, impacting both academic and professional performance [3].

The primary approach for managing AD involves the use of topical corticosteroids as the first-line treatment [3]. As per a recent comprehensive review and meta-analysis, the treatments found to be most effective for AD comprise pimecrolimus, tacrolimus, and topical corticosteroids with moderate potency [4]. While high-potency topical corticosteroids are more effective, they come with an increased risk of local and, rarely, systemic adverse effects. Additionally, patients often experience a relapsing and remitting course [5].

The Spanish Ministry of Industry, Trade, and Tourism has recently granted patent approval (Invention patent reference number 202030824) for a ground-breaking topical treatment designed for psoriasis and AD. This innovative treatment, presented in the form of a lotion, includes a combination of clobetasol, papaverine hydrochloride, spironolactone, a milk-peptide complex, and propylene glycol.

We showcase a case of AD that was treated using our newly patented lotion, Psorisbye.

2. Case Report

An 18-year-old female presented with AD on the back of her neck. The patient had no significant past medical
history and primarily complained of intense pruritus in the AD lesions.

During the examination, a red rash on swollen skin, accompanied by crusting and scaling, was observed on the back of the neck and scalp. Additionally, a few abrasions from scratching were identified, primarily located on the back of the neck.

She received the diagnosis during infancy and has undergone treatment involving a combination of corticosteroid creams and moisturizers to address dry skin.

The patient received guidance to apply our recently patented lotion, Psorisbye, once a day for 5 days. In total, 50 ml of Psorisbye was utilized over 4 days. Throughout this duration, the patient did not undergo antihistamine treatment in conjunction with the application of the lotion.

On the fifth day, the patient underwent an examination at the outpatient clinic. The patient reported a significant improvement in pruritus sensations and observed a remission of the scaled lesions.

The gold standard for assessing AD is the SCORing Atopic Dermatitis Index (SCORAD). SCORAD incorporates both objective symptoms (extent and intensity) and subjective criteria (daytime pruritus and sleep loss). Clinical features considered for evaluating disease severity encompass erythema, swelling, oozing/crusting, scratch marks, skin thickening, and dryness. Each characteristic is assigned a severity rating of 0, 1, 2, or 3. The extent of the disease is estimated by the percentage of affected skin with eczema. SCORAD assesses subjective symptoms, such as itch intensity and sleep disturbance, on a scale from 0 (no symptoms) to 10 (maximum), reflecting the preceding three days. Numerical data for all three aspects are tabulated and summed, with a maximum score of 103 [6].

Upon evaluating our patient, a comparison of the lesions before and after applying the topical treatment for 4 days revealed a notable improvement in the SCORAD index, decreasing from 49.95 to 0 (Fig. 1).
more efficient absorption of moisture. This concerted action not only fosters heightened hydration levels but also serves as a proactive agent in the cultivation and maintenance of skin suppleness, resulting in an overall improvement in the skin’s tactile and visual qualities.

- Excipients encompass a hydroalcoholic solution carefully formulated in precisely measured proportions, a strategic composition designed to achieve the utmost solubility and stability of the formula.

In our case report, we did not observe any adverse reactions or interactions among the different molecules in the new topical treatment.

Our previously published case reports on Psorishbye have recently demonstrated the efficacy of the new foaming lotion in addressing moderate psoriasis in the short term [7], [8].

4. CONCLUSIONS

The effectiveness of the new topical treatment was evident in the significant improvement noted, demonstrated by the disappearance of the pruritus sensations and a notable difference in the pre- and post-SCORAD index assessment.

While the preliminary results exhibited by Psorishbye in the context of this specific case are indeed promising, the imperative to establish its efficacy and reliability necessitates the initiation of further investigations characterized by more expansive sample sizes and prolonged follow-up periods. Conducting such comprehensive studies will not only serve to validate the preliminary findings articulated in our current case report but also contribute to a more robust understanding of the potential long-term effectiveness and safety profile of the aforementioned topical treatment.

AUTHOR CONTRIBUTIONS

The study design was conducted by José Miguel Ingelmo Calvo, José Ruiz Cobo, and Mohamed Farouk Allam. Analysis and interpretation of data were performed by José Miguel Ingelmo Calvo and José Ruiz Cobo. Manuscript writing was carried out by José Ruiz Cobo and Mohamed Farouk Allam. The collection of data was managed by José Miguel Ingelmo Calvo and José Ruiz Cobo. Critical revision was conducted by José Ruiz Cobo and Mohamed Farouk Allam.

ETHICAL STATEMENT

Written informed consent was obtained for the patient, treatment, and publication of this case report and the accompanying image.

CONFLICT OF INTEREST

Authors declare that they do not have any conflict of interest.