Comparing the Therapeutic Effects of Aloe vera and Olive Oil Combination Cream versus Topical Betamethasone for Atopic Dermatitis: A Randomized Double-blind Clinical Trial

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Objectives: Atopic dermatitis (AD) is a prevalent and chronic, pruritic inflammatory skin condition that can influence all age groups. AD is associated with a poor health-related quality of life. This randomized clinical trial was performed to compare the effectiveness of Olivederma (combination of aloe vera and virgin olive oil) or betamethasone regarding disease severity, quality of life, serum IgE and eosinophil count.

Methods: Thirty-six AD patients were randomly allocated to topical Olivederma or betamethasone, and were followed for 6 weeks.

Results: Total SCORAD severity scores showed significant decrease in both groups, while it was more prominent in Olivederma group (64.5% improvement in Olivederma vs. 13.5% improvement in Betamethasone, p-value < 0.001). Quality of life (DLQI questionnaire) of AD patients was significantly improved after 6 weeks treatment with Betamethasone (22.3%, p < 0.001) and Olivederma (60.7%, p-value < 0.001). Olivederma group showed a significantly lower DLQI score in comparison with Betamethasone treated patients after 6 weeks of therapy (p < 0.001). Improvements in eosinophil count and serum IgE was observed.

Conclusion: In summary, this study shows that Olivederma is superior to topical Betamethasone after 6 weeks of therapy with regard to disease severity, quality of life and eosinophil count.

Keywords: atopic dermatitis, Aloe vera, olive oil, corticosteroid, topical administration

INTRODUCTION

Atopic dermatitis (AD) is a common chronic inflammatory genetically determined disease of the skin which is marked by increased ability to form regain (IgE), with increased susceptibility to allergic rhinitis and asthma, and hereditary disposition to a lowered threshold for pruritus. It affects up to 20% of children and 1% to 3% of adults [1]. AD is associated with a decrement in health-related quality of life [2].

Topical medications are the mainstay of AD treatment. Even in more severe cases needing systemic or phototherapy, they are often used in combination with these interventions [3]. Topical corticosteroids have been used to treat AD for more than 60 years [4]. The effectiveness of topical steroids is demonstrated [3, 4].

Topical corticosteroids exert various cutaneous side effects including purpura, telangiectasia, striae, focal hypertrichosis, and acneiform or rosacea-like eruptions [5]. Skin atrophy is our
greatest concern, which can be induced by any TCS, though high-potent steroids are more hazardous [4, 5]. Due to these adverse reactions and considering the fact that AD patients may have multiple flares during their life, it is quite necessary to develop no-steroidal anti-inflammatory and disease modifying agents.

Olivederma is a combination of virgin olive oil (VOO) and aloe vera gel. VOO has 82% unsaturated fatty acids [6], which helps in moisturizing dryness and removing Staphylococcus aureus from colonized AD skin [6]. Moreover, the anti-inflammatory activity of aloe vera has been demonstrated [7].

There are few randomized clinical trials for comparing the effects of topical corticosteroids and natural topical products like Olivederma on the severity of AD and the quality of life of patients. Therefore, we conducted this randomized clinical trial for this purpose.

### MATERIALS AND METHODS

#### 1. Trial design

This randomized clinical trial was performed during July 2009-July 2010 in Dermatology clinics of Tehran Baqiyatallah University. Patients satisfying inclusion criteria were randomly allocated in two groups receiving Olivederma or betamethasone, in a parallel group design (1:1 ratio) using block randomization (with a block size of 4). The specialists and staffs were blinded through trial.

#### 2. Participants and study setting

We enrolled 36 patients with atopic dermatitis for this randomized, double-blind placebo- clinical trial. Patients were diagnosed by the U.K. Working Party’s Diagnostic Criteria for atopic dermatitis [8].

Inclusion criteria were defined as: age OF 17-70 years, confirmed diagnosis of atopic dermatitis, not having allergy to herbal products, and providing the written informed consent. These patients were excluded: patients who do not show adherence to intervention (based on weekly follow-up calls; defined as refusing to consume utmost 10% of recommended treatments), and those with any adverse reactions to the medications.

Ethical Committee of Tehran Islamic Azad University approved the study.

#### 3. Interventions

Intervention group received Olivederma topical (Kimi Daru Pharmaceutical Co, Tehran, Iran) for 2 times a day for 6 weeks and the control group received topical betamethasone (Kimi Daru Pharmaceutical Co, Tehran, Iran) 2 times a day for 6 weeks.

#### 4. Randomization

We used permuted block randomization sequences (created using Microsoft Excel version 2010 software; the block sizes were considered fixed as 4). A blinded statistician generated the sequentially numbered containers for concealment. The blinded secretary of outpatient dermatology clinic enrolled participants using sealed sequential numbers with attached pockets containing medications (identical betamethasone and Olivederma topical medications). Blinded dermatologists of clinic explained the interventions to the participants and checked their adherence by weekly calls and monthly visits. All adverse events were recorded by them.

#### 5. Sample size

This primary aim of this study was to compare the effects of topical Olivederma and betamethasone on SCORAD severity index of atopic dermatitis. Moreover, the quality of life (DLQI), serum IgE and eosinophil count in the peripheral blood were also compared. The sample size was obtained regarding the primary outcome of SCORAD scores. The effect size was selected as 15 scores mean difference between intervention groups (SD = 15). Also 5% type I error and 80% power were considered.

Formula written here was used:

\[ n = \left( \frac{Z_{1-\alpha/2} + Z_{1-\beta}}{d} \right)^2 \]

where \( d = (D_1 - D_2) / (\sigma\sqrt{2}) \)

Considering 5% loss to follow-up, sample size per each group was determined as 17 atopic dermatitis patients.

#### 6. Outcomes and measurements

The primary outcome variables were SCORAD severity index of atopic dermatitis [9]. Moreover, the validated Persian version of Dermatology Life Quality Index (DLQI) questionnaire was measured for assessing the quality of life.

Blood samples of patients were obtained (10 cc) to measure...
the serum IgE and peripheral blood eosinophil count.

7. Statistical methods

Quantitative data are shown as mean ± SD. Within-group (before/after intervention) differences were tested by paired t-test. Between-groups differences were analyzed by independent t-test and the time×intervention interaction was analyzed by repeated measures ANOVA. Categorical data are reported as numbers (percentages) and were tested by \( \chi^2 \), Fisher’s exact, or Monte Carlo tests; where indicated. p-values less than 0.05 were considered statistically significant.

RESULTS

79 patients with AD were recruited for initial assessment of eligibility. Twenty-four patients were excluded due to not meeting the inclusion criteria and 19 ones refused to participate. The remaining 36 atopic dermatitis patients were randomly allocated to topical Olivederma or betamethasone, and were followed for 6 weeks.

The studied AD patients had a Mean (Range) age of 33.41 (13-57) years. They were 56% male. We did not observe significant differences for demographic characteristics and past medical history between intervention groups (Table 1). Overall, 36 T2DM patients (19 in the Olivederma group; 16 patients in the Betamethasone group) successfully finished the trial and were analyzed.

Regarding the severity of AD (Table 2), total SCORAD scores showed significant decrease in both groups, while it was more prominent in Olivederma group (64.5% improvement in Olivederma vs. 13.5% improvement in Betamethasone, p-value < 0.001).

More interestingly, we observed that quality of life (DLQI questionnaire, Table 3) of AD patients was significantly improved after 6 weeks treatment with Betamethasone (22.3%, p < 0.001) and Olivederma (60.7%, p-value < 0.001). Olivederma group showed a significantly lower DLQI score in comparison with Betamethasone treated patients after 6 weeks of therapy (p < 0.001).

We found that eosinophil count (Table 4) of Betamethasone group was significantly increased (p = 0.011), while it was significantly decreased after 6 weeks of therapy with Olivederma (36%, p = 0.003).

We observed that Olivederma did significantly decrease serum IgE level (Table 4), while it was remained constant in Betamethasone group. However, the between-group difference of Betamethasone and Olivederma was non-significant.

DISCUSSION

In this randomized, double blind clinical trial we observed that topical Olivederma is superior to topical Betamethasone after 6 weeks of therapy with regard to disease severity, quality of life and eosinophil count of AD patients.

Table 1. Baseline characteristics of patients with atopic dermatitis randomly allocated in topical Betamethasone and Olivederma interventions

| Patients’ characteristics | Group | Difference | p-value |
|--------------------------|-------|------------|---------|
| Age (years)              | Betamethasone | 30.44 ± 9.65 | 36.37 ± 12.45 | 0.123 |
| Sex                      | Olivederma   | 0.296       |         |
| Male                     | 6 (35.3%)    | 10 (52.6%)  |         |
| Female                   | 11 (64.7%)   | 9 (47.4%)   |         |
| Allergic rhinitis        | 4 (23.5%)    | 8 (42.1%)   | 0.238   |
| Asthma and allergic rhinitis | 13 (76.47%) | 9 (47.4%) | 0.074 |
| Positive family history of atopic dermatitis | 17 (100%) | 16 (84.2%) | 0.087 |

Table 2. Comparing topical Betamethasone and Olivederma on SCORAD severity index of patients with atopic dermatitis randomly allocated in interventions

| Indices | Phase | Betamethasone | Olivederma | p-value* |
|---------|-------|---------------|------------|----------|
| SCORAD-A | Baseline | 44.4 ± 8.7 | 75.1 ± 7.2 | < 0.001 |
|          | 6th week | 35.5 ± 9.9 | 23.3 ± 13.6 | 0.006 | < 0.001 |
| SCORAD-B | Baseline | 9.6 ± 1.5 | 9.4 ± 1.7 | < 0.001 |
|          | 6th week | 8.1 ± 2.3 | 3.2 ± 1.8 | 0.004 | < 0.001 |
| SCORAD-C | Baseline | 15.8 ± 2.7 | 19.5 ± 0.9 | < 0.001 |
|          | 6th week | 15.2 ± 3.4 | 8.2 ± 5.7 | 0.045 | < 0.001 |
| SCORAD total | Baseline | 58.4 ± 7.3 | 67.3 ± 6.3 | < 0.001 |
|          | 6th week | 50.5 ± 10.7 | 23.9 ± 13.3 | 0.003 | < 0.001 |

*Within-group (time effect) significance.
*Between-group (intervention * time effect) significance.
Traditionally, topical corticosteroids are pivotal therapeutic agents for patients with AD [3-5]. Considering the chronic nature of AD, its multiple recurrences throughout life, and the disadvantageous effects of topical corticosteroid; it is warranted to conduct studies aimed to develop non-steroidal agents and with minimal harms [6]. Natural products have always been a potential resource to discover and develop drugs. Combinations of such products have been introduced in literature [3] to conduct therapeutic effects. However, the evidence is insufficient to support the superiority of such natural products for treatment of chronic inflammatory dermatitis like AD. We conduct this study to compare the effectiveness of Olivederma (combination of aloe vera and virgin olive oil) and Betamethasone for improving the quality of life of AD patients and decreasing the severity of disease.

Several studies support the role of Aloe vera in wound healing [10-19]. It has been found to decrease the healing time of burn wound patients in comparison with petroleum gel [18]. Its effects have been postulated to be exerted through its anti-inflammatory effects by prostaglandins and bradykinin [7, 14, 16, 17]. However, aloe vera was not better that placebo in the treatment of slight to moderate psoriasis vulgaris [20]. To the best of our knowledge, there were no randomized controlled trials comparing the efficacy of aloe vera and betamethasone in treatment of AD patients. We conduct this study to compare the effectiveness of Olivereda (combination of aloe vera and virgin olive oil) and Betamethasone for improving the quality of life of AD patients and decreasing the severity of disease. Several studies support the role of Aloe vera in wound healing [10-19]. It has been found to decrease the healing time of burn wound patients in comparison with petroleum gel [18]. Its effects have been postulated to be exerted through its anti-inflammatory effects by prostaglandins and bradykinin [7, 14, 16, 17]. However, aloe vera was not better that placebo in the treatment of slight to moderate psoriasis vulgaris [20]. To the best of our knowledge, there were no randomized controlled trials comparing the efficacy of aloe vera and betamethasone in treatment of AD patients. We conduct this study to compare the effectiveness of Olivederma (combination of aloe vera and virgin olive oil) and Betamethasone for improving the quality of life of AD patients and decreasing the severity of disease.

Emollients are a standard of care for inhibition of dryness, steroid-sparing effect, and maintenance therapy in AD [4, 6]. Virgin olive oil coat the skin, occluding and protecting it by slowing down transepidermal water loss and increasing hydration within the stratum corneum [6, 21]. Also, it makes the skin look less rough and scaly [21]. The superiority of Olivederma in comparison with Betamethasone regarding the severity of dryness reported by our patients might be due to such moisturizing effects of olive oil.

AD patients have a decreased quality of life [2]. Evidence reveals that they have inferior mental health scores compared with diabetes or hypertension patients, and inferior social functioning scores compared with hypertension patients. In comparison with psoriasis patients, AD patients are reported to have inferior scores in the role-physical, vitality, social functioning, role-emotional, and mental health SF-36 domains [2]. In this study we observed that all of patients treated with topical Olivederma showed improvements in DLQI scores, while only 53% of AD patients treated with Betamethasone have mild improvements regarding the quality of life.

The association between the severity of AD symptoms and serum IgE level has been observed [22, 23]. Also, to our knowledge, the disease severity and personal or family history of respiratory atopy are associated with higher eosinophil count [24]. We found that both eosinophil count and serum IgE level are significantly decreased after 6 weeks of therapy with Olivederma, however eosinophil count of Betamethasone treated patients were found to be increased. Moreover, betamethasone did not show significant effects on serum IgE. Now, we do not know the exact cause of such advantageous effects of Olivederma in comparison with Betamethasone. Further studies are required to confirm this finding.

This study had some limitations, as we could not follow our patients for long-term and examine the effectiveness of Olivederma in severe flare-ups. However, we showed the superiority

| Indices | Phase | Betamethasone | Olivederma | p-value* |
|---------|-------|---------------|------------|----------|
| DLQI score | Baseline | 22 ± 3.2 | 18.3 ± 2.8 | < 0.001 |
| 6th week | 17.1 ± 5.4 | 7.2 ± 1.4 | < 0.001 | < 0.001 |
| p-value** | 0.011 | 0.003 |
| DLQI change | Excellent improve | 0 (0%) | 6 (31.6%) | < 0.001 |
| Mild improve | 7 (41.2%) | 13 (68.4%) | 0.23 | 0.007 |
| No changes | 9 (52.9%) | 0 (0%) | 0.23 | 0.007 |
| Exacerbated | 1 (5.9%) | 0 (0%) | 0.23 | 0.007 |

*Within-group (time effect) significance.
*Between-group (intervention * time effect) significance.
of Olivederma to Betamethasone by various subjective and objective markers in short-term. The considerable improvement of AD patients after treatment with a natural combination of Virgin olive oil and aloe vera is quite encouraging to conduct more trials investigating its superiority to other topical regimens especially corticosteroids.

In conclusion, this randomized double-blind clinical trial shows that Olivederma, a combination of aloe vera and Virgin olive oil, is superior to topical Betamethasone after 6 weeks of therapy with regard to disease severity, quality of life and eosinophil count. Further clinical trials are required.

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CONFLICT OF INTEREST

We declare no conflicting financial or non-financial interests.

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