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Effect of Pharmacist's Intervention in the Quality of Life of Patients with Atopic Dermatitis - A Pilot Study



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ABSTRACT

Background: Atopic dermatitis is a chronic skin condition that is variable in age and onset. Therefore, in order to improve the quality of life of patients, it is essential to provide counseling so as to improve compliance with the medications and hence disease control. Methods: This study was carried out in 28 patients who were diagnosed with atopic dermatitis from the Dermatology Department of a tertiary care hospital at Thiruvananthapuram. Informed consent was obtained. They were divided into two groups namely the tacrolimus group (14) and corticosteroids group (14). Counseling was provided to both groups and the Dermatology Life Quality Index (DLQI) scale was used to score. The collected data has been analyzed and presented. Results: A total of 28 patients with Atopic Dermatitis fulfilling the study criteria were included for analysis. The predominance of atopic dermatitis is seen in females than in males and in the age group of 21 - 30 years. Using paired t-test, the statistical analysis clearly depicts that there is a clear shift from the severe effect on the quality of life to a milder effect in both groups separately as well as combined with significant p-value < 0.001. However, there is no significant change between the groups. Conclusions: From this study, it is concluded that there is betterment in the quality of life of patients with atopic dermatitis both pre and post counseling. Pharmacist's intervention in the form of counseling is hence an integral factor in improving the quality of life of patients with atopic dermatitis.

INTRODUCTION

Atopic dermatitis is an itchy, chronic, or chronic relapsing, inflammatory skin condition. The rash is characterized by itchy papules [occasional vesicles in infants] which become excoriated and lichenified and typically have a flexural distribution. The eruption is frequently associated with other atopic conditions in the individual or other family members [1-3].

Etiological factors mainly include genetic influence. Various studies have demonstrated this in detail. Examples include susceptible loci like 1g21^[4], 3p24-22^[5], etc. Abnormal epidermal barrier function is a well-recognized clinical association of atopic eczema and until recently it wasn't clear whether the dry skin was only a consequence of the underlying immune inflammatory response genetic defects in filaggrin gene has been associated with early onset of atopic dermatitis, a worse prognosis, and persistence of disease into adulthood [6-8]. Variations in the SPINK5 gene have also been associated with atopic dermatitis [9-11]. Cord blood IgE is high in babies whose mothers are atopic or have high IgE, whereas paternal atopy or raised IgE are not associated with raised cord blood IgE indicating the risk is higher in maternal atopy [12-13]. Exposure to alcohol, cigarettes and other pollutants has attracted attention, but none has emerged as a major factor. The hygiene hypothesis is another factor [14, 15]. There are differences in the prevalence of allergic diseases among people living in an urban and rural set up [16-19]. Biopsy of developing lesions of atopic dermatitis reveals that Tcells infiltrate the skin in the early disease process [20, 21]. Deficiencies of certain immunoglobulins have also been associated with the development of atopic dermatitis. Altered sensations associated with the neuropeptides can result in intense itching and aggravates this condition. Clinical features in adult phase are similar to in later childhood with lichenification of flexures and hands, nipples, lips [22]. Diagnostic criteria include the Hanifin and Rajka criteria [23]. Diagnosis is usually aided by estimation of total serum IgE, specific radioallergosorbent tests [RASTs], prick tests and patch tests most important [24, 25]. Treatment usually involves the reduction of trigger factors like irritants, habitual scratching, excessive heat and sweating, food allergens, airborne allergens, stress, etc., Topical therapy [bathing and emollients] [26, 27], topical corticosteroids, topical calcineurin inhibitors like tacrolimus, pimecrolimus [28]. Oral therapy includes Antihistamines like H1 receptor antagonists like promethazine, trimeprazine [29], antibiotics to relieve from exudation and pustule formation. Other approaches include interferon gamma, intravenous

immunoglobulins, montelukast. Phototherapy includes UVB, narrowband UVB, medium and high dose UVA1 and PUVA [30-34]. Desensitization [allergen specific immunotherapy] is also advised but has limited part in management [35].

Quality of life (QOL) is perceived as the quality of an individual's daily life, that is, an assessment of their well-being or lack thereof. QOL is a broad concept that includes such things as the standard of living, community, and family life [36]. Health-related QOL (HRQOL) assesses qualities directly related to the disease as well as those that are independent of the disease but may be affected by it. The literature on QOL has increased substantially over the past decade, whereas before, medical concerns were mainly centered on issues.

Dermatology Life Quality Index [DLQI] is used here. This present study aims to assess the effect of counseling on patients with AD and the objective is to evaluate the QOL of the patients both pre and post counseling.

MATERIALS AND METHODS

- 2.1 Data Source. All relevant information regarding the study was collected from case records and direct interview with the patients. Data from patients or caretaker was collected by using a specially designed proforma. Quality of life of the patients was assessed by using DLQI scoring. The study was approved by the Research Ethics Committee of Cosmopolitan Hospital, Thiruvananthapuram.
- 2.2. Study Population: Patients were taken from the Dermatology department of Cosmopolitan Hospital. An age group of 16-70 years was recruited. Informed consent was obtained. The study was conducted for a period of 2 months.
- 2.3. Assessment of the effect of counseling: Patients with atopic dermatitis, who had received prescriptions for atopic dermatitis drugs including tacrolimus and corticosteroids. Counseling was provided to the patients regarding the disease. Assessment of the quality of life of patients before and after counseling was done using DLQI scores.
- 2.4 Statistical Analysis. Effect of counseling in both groups of Tacrolimus and Corticosteroids was done using the paired t-test.

RESULTS

In this study, a total of 28 subjects participated of which 14 subjects were allotted to the Tacrolimus group and 14 subjects to the Corticosteroids group. 75% (21) of the subjects were females and 25% (7) were males.

In the tacrolimus group, 11(78.6%) subjects were females and 3(21.4) subjects were males. In the corticosteroids group, 10(71.4%) subjects were females and 4 (28.6%) subjects were males.

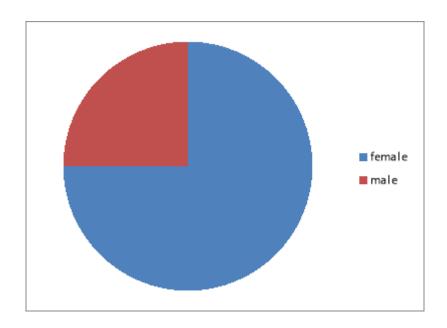


Figure 1: Distribution of subject

In this study, subjects of the age range of 16-70 years were included and then categorized into 6 groups. It was observed graphically that a number of subjects were in the age group of [21 -30] years (28.6%) and least in the age group of [41 -50] years.

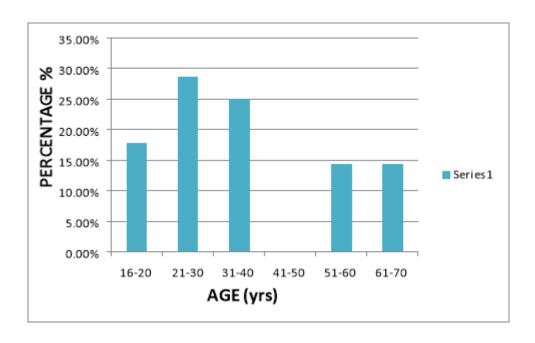


Figure 2: Distribution of subjects by age

The plan of the study was to compare the quality of life of patients. Here the patients taking tacrolimus and corticosteroid were combined to form a single group and the quality of life before the counseling and after the counseling was evaluated and compared using the t-test.

EFFECT OF COUNSELLING ON QUALITY OF LIFE

Table 1: Effect of counseling on quality of life

| GROUP | BEFORE COUNSELLING | AFTER COUNSELLING | PERCENTAGE CHANGE | t- TEST | P- VALUE |
|-----------------|-----------------------|----------------------|----------------------|------------|-------------|
| TACROLIMUS | 2.92 | 1.07 | 64.72 | 13 | P<0.001% |
| CORTICOSTEROIDS | 2.85 | 1.35 | 54.3 | 10.81 | P<0.001% |
| TOTAL | 2.89 | 1.21 | 58.131 | 16.21 | P<0.001% |

From the table, there is evidence that through paired t-test, there is a significant change in the quality of life due to counseling in the tacrolimus group(p<0.001). Before counseling, QOL of subjects in the Tacrolimus group was severely affected (2.92). Post counseling, significant improvement in QOL was observed (1.07) which indicates a clear shift from severe to mild effects. Similarly, a shift from severe effect (2.85) to mild effect (1.35) on the QOL was observed in the Corticosteroid group.

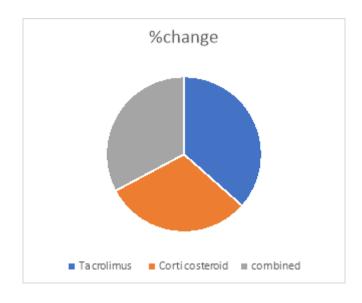


Figure 3: Percent change observed

It is evident from the study that the quality of life of patients can be increased from severe to mild. In this study, a positive outcome that the quality of life of the patient will increase on both group and in the combined group is observed. However, there is no significant change between the groups.

DISCUSSION

In the study done by Mozaffari H et al ^[37], 86 patients with AD who were referred to an immunology clinic and 98 patients [> 4 years old] attending a general clinic acting as controls [without any chronic or severe disease] participated in this survey. Physician filled the Children's Dermatology Life Quality Index [CDLQI] questionnaire for individuals more than 16 years via face-to-face interview. There were significant differences between the mean of CDLQI score and DLQI score in case and control groups (p<0.001). For children and adults with AD, the mean score of each question was significantly higher than in the control group (p<0.001).

In the cross-sectional study of 132 outpatients with AD by Maksimovic N et al ^[38], to assess the QOL, Short Form 36 (SF-36), Dermatology Life Quality Index (DLQI) and Children's Dermatology Life Quality Index (CDLQI) were administered. Increasing disease severity was associated with greater impairment in QOL in both children and adults. Our study found the influence of stressful life events on the emotional role of AD patients. These results demonstrate that AD influences health-related QOL, especially in children.

According to the study done by Agner T et al ^[39], a total of 416 patients with hand eczema from 10 European patch test clinics participated in the study. Data on QOL were obtained from a self-administered questionnaire using DLQI. No significant difference was found between males and females with respect to QOL (DLQI median values and 25/75 percentiles for males and females being 7.0 [3-13] and 8.0 [3-14] respectively), although males were more severely affected than females [p<0.025]. A significant positive correlation was found for hand eczema severity and age (p<0.001), while no significant correlation was found for QOL and age. QOL was found increasingly reduced when sick leave was getting higher [p<0.001]. A statistically significant correlation between QOL [as measured by DLQI] and hand eczema severity as measured by HECSI was found [p<0.001]. No significant difference in QOL was found between males and females, in spite of significantly more severe eczema in males, indicating that QOL I female patients is more easily affected.

Holm et al ^[40] measured HRQOL in patients with AD to analyze discriminant, divergent, and convergent validity by examining the association between various QOL methods and to examine the association between disease severity assessed by an objective Severity Scoring of Atopic Dermatitis (SCORAD) and QOL. HRQOL was assessed at two visits at a 6-monthly interval in 101 patients with Dermatology Life Quality Index (DLQI) or Children's DLQI (CDLQI)], one generic instrument (SF-36), and three visual analog scales of severity and pruritus. Objective SCORAD was used to measure disease severity. Thirty-five children aged 3–14 years were included. Results showed that patients with AD had significantly lower QOL than healthy controls and the general population. DLQI/CDLQI, pruritus, and patient and investigator overall assessment of eczema severity was significantly (p < 0.0001) and positively correlated with SCORAD, while the generic questionnaire showed only poor correlation. The authors concluded that AD has an impact on HRQOL.

Another study that included 239 AD patients aged 4–70 years showed that patients with AD had inferior scores on vitality, social functioning, and mental health subscales compared with individuals in the general population [40]

An international study performed in the Czech Republic, Singapore, Brazil, the Netherlands, and South Korea on QOL and family QOL in children with AD found a similar impact of the disease as rated by parents of 419 children under the age of 4 years in all countries [41]

CONCLUSION

With increasing worldwide prevalence, atopic dermatitis has major social and financial implications for individuals, health-care providers, and society as a whole. The good news is that even if symptoms of AD can be uncomfortable and at times difficult to control, the disease, in general, can be successfully managed and in some cases even prevented [42]. Individuals affected by AD can lead almost normal lives. From this study, it is concluded that there is betterment in the quality of life of patients both pre and post counseling. Hence pharmacists play a very crucial role in improving the quality of life of patients with atopic dermatitis through proper counseling and hence this indicates that counseling is an integral factor in improving QOL. However, larger sample size and a longer duration of study are necessary for better and reliable results.

REFERENCES

- 1. Darsow U, Lubbe J.Taieb A, et al, for the European Task Force on Atopic Dermatitis. Position paper on the diagnosis and treatment of atopic dermatitis, J. Eur Acad Dermatol Venereol 2005, 19; 286 95
- 2. Williams H. Atopic dermatitis N Engl J Med 2005; 352-2314-24
- 3. Brown.S.Reynolds NJ. Atopic and nonatopic eczema. BJ 2006; 352; 584-8
- 4. Cookson WO, Ubhi B, Lawrence R et al. Genetic linkage of childhood atopic dermatitis to psoriasis susceptibility loci. *Nat Genet* 2001; 27- 372-3
- 5. Bradley M, Soderhall C, Luthman H et al. Susceptibility loci for atopic dermatitis on chromosomes 3, 13, 15, 17 and 18 in a Swedish population. *Hum Mol Genet* 2002; 11-1539-48
- 6. Palmer CN, Iwine AD, Terron-Kwiatkowski A et al. Common loss-of-function variants of the epidermal barrier protein filaggrin are a major predisposing factor for atopic dermatitis. *Nat Genet* 2006; 38- 441-6
- 7. Stemmler S, Parwez Q, Petrasch- Parwez E et al. Two common loss-of-function mutations within the filaggrin gene predispose for early-onset of atopic dermatitis. *J Invest Dermatol* 2007; 127-722-4
- 8. Barker JN, Palmer CN, Zhao Y et al. Null mutations in the filaggrin gene [FLG] determine major susceptibility to early-onset atopic dermatitis that persists into adulthood. *J Invest Dermatol* 2007; 127- 564-7
- 9. Walley AJ, Chavanas S, Moffat MF et al. Gene polymorphism in Netherton and common atopic disease. *Nat Genet* 2001; 29- 175-8
- 10. Kato A, Fukai K, Oiso N et al. Association of SPINK5 gene polymorphisms with atopic dermatitis in the Japanese population. *Br J Dermatol* 2003; 148-665-9
- 11. Nishio Y, Noguchi E, Shibasaki M et al. Association between polymorphisms in the SPINK5 gene and atopic dermatitis in the Japanese. *Genes Immun* 2003; 4-515-7.
- 12. Dold S, Wjst M, von Mutius E et al. Genetic risk for asthma, allergic rhinitis, and atopic dermatitis. *Arch Dis Child* 1992;67- 1018-22
- 13. Ruiz RG, Kemeny DM, Prince JF. Higher risk of infantile atopic dermatitis from maternal atopy than from paternal atopy. *Clin Exp Allergy* 1992; 22-762-6.
- 14. Strachan DP. Family size, infection, and atopy; the first decade of 'hygiene hypothesis'. *Thorax* 2000; 53 [Suppl. 1]; S2 10
- 15. Strachnan DP. Hay fever, hygiene, and household size. BMJ 1989; 299; 1259-60
- 16. Von Ehrenstein OS, von Mtius E, Iili S et al. Reduced risk of hay fever and asthma among children of farmers. *Clin Exp Allergy* 2000;30-187-93
- 17. Braun- Fahrlander C. The role of farm environment and animal contact for the development of asthma and allergies. *Clin Exp Allergy* 2000;31- 1799-803

- 18. Braun Fahrlander C, Riedler J, Herz U et al. Environmental exposure to endotoxin and its relation to asthma in school-age children. *N Engl J Med* 2002; 347 869-77
- 19. Riedler J, Eder W, Oberfield G, et al. Austrian children living on a farm have less hay fever, asthma, and allergic sensitization. Clin Exp Allergy 2000; 30 194 200
- 20. Grewe M, Walther S, Gyufko K et a. Analysis of the cytokine pattern expressed in situ in inhalant allergen patch test reactions of atopic dermatitis patients. *J Invest Dermatol* 1995; 105-407-10.
- 21. Thepen T, Langeveld- Wildschut EG, Bihari IC, et al. Biphasic response against aeroallergen in atopic dermatitis showing a switch from initial TH2 response to a TH1 response in situ; an immunocytochemical study. *J Allergy Clin Immunol* 1996; 97-828-37
- 22. Bannister MJ, Freeman S. Adult-onset atopic dermatitis. Australias J Dermatol 2005; 41-225-8
- 23. Hanifin JM, Rajka RG. Diagnostic features of atopic dermatitis. *Acta Derm Venereol [Stockh]* 1980; 92[Suppl, 144]; 44-7
- 24. Cronin E, McFadden JP. Patients with atopic eczema do become sensitized to contact allergens. Contact Dermatitis 1993; 28- 225-8
- 25. Lever R, Forsyth A. Allergic contact dermatitis in atopic dermatitis. *Acta Derm Venereol Suppl [Stockh]* 1992; 76- 95-8
- 26. Holden CA, English J, Hoare C et al. Advised best practice for the use of emollients in eczema and other dry conditions. *J Dermatol Treat* 2002; 13-103-6
- 27. Chamlin SL, Frieden IJ, Fowler A et al. Ceramide-dominant, barrier repair lipids improve childhood atopic dermatitis. *Arch Dermatol* 2001; 137-1110-2
- 28. Ashcroft DM, Dimmock P, Garside R et al. Efficacy and tolerability of topical pimecrolimus and tacrolimus in the treatment of atopic dermatitis; a meta-analysis of randomized controlled trials. *BMJ* 2005; 330-516-22
- 29. Klein PA, Clark RA. An evidence-based review of the efficacy of antihistamines in relieving pruritis in atopic dermatitis. *Arch Dermatol* 1999; 135; 1522-5
- 30. Kruttmann J. Phototherapy for atopic dermatitis. Clin Exp Dermatol 2000; 25; 552-8
- 31. Reynolds NJ, Franklin V, Gray JC et al. Narrowband ultraviolet B and broadband ultraviolet A phototherapy in adult atopic eczema; a randomized controlled trial. *Lancet* 2001; 357-2012-6
- 32. Tzaneva S, Seeber A, Schwaiger M et al. High dose versus medium-dose UVA1 phototherapy for patients with severe generalized atopic dermatitis. *J Am Acad Dermatol* 2001; 45; 503-7
- 33. Der-Petrossian M, Seeber A, Honigsmann H, Tanew A. Half side comparison study on the efficacy of 8-methoxypsoralen bath- PUVA versus narrowband ultraviolet B phototherapy in patients with severe chronic atopic dermatitis. *Br J Dermatol* 2000; 142-39-43
- 34. Clayton TH, Clark SM, Turner D, Goulden V. The treatment of severe atopic dermatitis in childhood with narrowband ultraviolet B phototherapy. *Clin Exp Dermatol* 2007; 32- 28-33
- 35. Glover MT, Atherton DJ. A double-blind controlled trial of hyposensitization to *Dermatophagoides* pteronyssinus in children with atopic eczema. Clin Exp Allergy 1992; 22- 440-6
- 36. Jenney MEM, Lane RL, Lurie N: Developing a measure of health outcomes in survivors of childhood cancer: a review of the issues. *Med Pediatr Oncol* 1995; 24: 145–153.
- 37. Mozaffari H et al. Quality of life in atopic dermatitis patients. *J Microbiol Immunol Infect*, 2007 Jun; 40(3): 260-4
- 38. Maksimovic N et al. Health-related quality of life in patients with atopic dermatitis. J Dermatol.2012 Jan;39(1):42-7
- 39. Agner T et al. Hand eczema severity and quality of life: a cross-sectional, multicentre study of hand eczema patients. Contact dermatitis. 2008 Jul;59(1):43-7
- 40. Kiebert G, Sorensen SV, Revicki D, Fagan SC, Doyle JJ, Cohen J, Fivenson D: Atopic dermatitis is associated with a decrement in health-related quality of life. *Int J Dermatol* 2002; 41:151–158.
- 41. Chernyshov PV, Jirakova A, Ho RC, Moed H, Caldeira AP, Alvarenga TM, Park CW, Hercogova J: An international multicenter study on quality of life and family quality of life in children with atopic dermatitis. *Indian J Dermatol Venereol Leprol* 2013; 79: 52–58.
- 42. Spieldenner J, Belli D, Dupont C, Haschke F, Iskedjian M, Nevot Falcó S, Szajewska H, von Berg A: Partially hydrolyzed 100% whey-based infant formula and the prevention of atopic dermatitis: comparative pharmacoeconomic analyses. *Ann Nutr Metab* 2011; 1: 44–52