PRURITOCEPTIVE AND PSYCHOGENIC PRURITUS IN LICHEN SIMPLEX CHRONICUS

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ABSTRACT

Pruritus is a diagnostic hallmark for Lichen Simplex Chronicus (LSC). It elicits a scratch response, initiating the itch-scratch cycle, which in turn aggravates the inflammatory response and exacerbates the disease severity. Treating this symptom can be challenging The purpose of this study is to distinguish which type of pruritus has a leading role in the pathogenesis of LSC by studying changes skin barrier functions and evaluating patients psychopathology. We compare the results from the tests of skin barrier and evaluation of psychiatric scales HAD-A and HAD-D observing 56 adults with diagnosis LSC.

Keywords: lichen simplex chronicus, skin barrier evaluation, anxiety

INTRODUCTION

Lichen Simplex Chronicus (LSC) is a common skin disorder characterized by lichenification of skin as a result of excessive scratching (1). The dominant symptom of dermatosis is pruritus. (2) The pathogenesis of this skin disease is not well distinguished. Disorders of skin barrier are described as a trigger or enhancer of the pathological symptoms of LSC (3). There is an altered skin barrier with varying combination of allergens, irritants and skin pathogens which result in a changed immunoregulatory process. (4) On the other hand psychological factors have an important role in the pathogenesis of LSC. Recent evidence has indicated that psychological stress is associated with exacerbation of different skin condi-

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Received: March 18, 2016 **Accepted**: June 3, 2016 tions, including LSC (5). The itch has been classified into four different clinical categories. These include neurogenic, psychogenic, neuropathic, and pruritoceptive (6). These categories were developed based on anatomical, pathophysiological and psychological factors. The itch can be triggered by emotional stress or other underling pruritogenic stimuli. Some authors considered LSC to be a psychosomatic illness (7). This study explores the possible correlation or predomination of pathogenic factors and underlying the type of pruritus in LSC patients

MATERIALS AND METHODS Settings and sample

The study was conducted among 56 non-hospitalized patients 35 females (62.5%) and 21 males (37.5%); the mean age 49.46 years; (range 29-64 years) who visited the dermatology unit in the St. Anna Medical Center between January 2013 and January 2015. The patients have the following inclusion criteria: one or more lichen plaques, highly pruritic, accumulation of normal skin lines, excoriations.

Measures

We used instrumental methods, including, measuring hydration(H) and transepidermal water

loss (TEWL) in healthy and damaged skin to evaluate the functioning of stratum corneum (SC) The 14item Hospital Anxiety and Depression scale HAD-A and HAD-D was used to measure the levels of anxiety and depression. Personality questionnaires were given to patients during consultation and instruc24 months-mean 27g.m2.h (range 20g.m2.h - 32g. m2.h). The results obtained from measuring H show similar correlations. The pathological changes were more visible in patients with a duration of LSH of 19-24 months - mean 23.15 and more than 24 months, mean 18.75 (p=0.008) (Table 1).

Distribution of patients (N%) according to the duration of the disease (months)	TEWL healthy skin g.m2.h	TEWL dam- aged skin g.m2.h	р	H healthy skin	H damaged skin	Р
4-6 N=16.66%	5-15 (10.57)	15-20 (16.42)		20-40 (34.25)	18-35 (28.12)	
7-12 N=19.04%	10-17 (13)	18-23 (20.75)		26-45 (34.4)	18-35 (27.1)	
13-18 N=21.42%	5-18 (12.71)	10-28 (22.85)	0.048	32-40 (37.71)	15-33 (25.28)	0.008
19-24 N=30.95%	8-24 (15.23)	12-38 (27.30)		22-40 (32.15)	10-31 (23.15)	
More than 24 N=11.90%	15-21 (17.8)	20-32 (27)		20-40 (28)	15-22 (18.75)	

Table 1. Distribution according to duration

tions on how to fill them in. The patients were asked to fill-in the questionnaires at home. Completed questionnaires were returned at the next visit.

The statistical analysis was performed with SPSS v.21.0 for Windows. Hypotheses were tested using χ^2 -criteria (for the descriptive profile data). Construct validity was tested by factor analysis. Results with p<0.001 were interpreted as statistically significant.

The most distinct trend for correlation was determined comparing TEWL and H in pathological skin and severity of the disease. In a group of patients stage II TEWL had a mean value of 31.22g.m2.h, while in a group stage I TEWL had a mean value of 16.23g.m2.h. (p=0.009). In a group of patients stage III H had a s mean value of 20.25, while in a group stage I H had a mean value of 28.46 (p<0.003) (Table 2).

Distribution of patients (N%) according to the stage of the disease	TEWL healthy skin g.m2.h	TEWL dam- aged skin g.m2.h	р	H healthy skin	H damaged skin	р
Stage I N=30.95%	5-18 (9.76)	10-23 (16.23)		37.38 (26-45)	28.46 (15-35)	
Stage II N=40.47%	10-17 (14.9)	18-32 (24.1)	0.009	33.17 (20-45)	25.88 (15-31)	0.003
Stage III N=28.57%	12-24 (16.66)	22-38 (31.22)		30.16 (22-40)	20.25 (10-30)	

Table 2. Distribution according to the severity of the disease

RESULTS AND DISCUSSION

We compared the TEWL dividing patients according to the duration of the disease. The pathological changes were more visible in patients with duration of LSC of 19-24 months - mean 27.30g. m2.h (range 12g.m2.h - 38g.m2.h) and more than Bouwstra and associates (2006) reported a correlation between the clinical characteristics of skin diseases and the changes in the barrier function (8). Other authors found no statistically significant relationship between the disturbed homeostasis of the skin barrier and the expression of the disease symptoms (9). The results from our study show the presence of such dependence in dividing patients by disease severity (Table 3).

tween 0-7; 17.98% have mild anxiety- range 8-10; 3.68% have moderate anxiety range 11-14 and just

TEWL and H	Stage of LSC					
	Stage I	Stage II	Stage III	Stage IV	р	
Normal	0%	0%	0%	0%		
Mild changes	59.3%	37.0%	3.7%	0%	< 0.001	
Moderate changes	45.5%	54.5%	64.7%	0%		
Severe changes	0%	35.3%	21.8%	0%		

Table 3. The correlation between levels of disturbance in the skin barrier and severity of LSC

In 1985 Werner and associates published data about increased TEWL in healthy and pathological skin in a study of patients with atopic dermatitis. As a result, the authors presume primary defect in the epidermal barrier (10). Our results showed that from all patients included in the study only 30.95% had pathology in the skin barrier (increased TEWL and decreased H) when measuring a healthy skin stretch. Disturbed pathological function of skin barrier (increased TEWL and reduced H) when measuring o damaged skin was reported in 85.71%. These results questioned the existence of a primary defect in the epidermal barrier and its leading role in unlocking LSC. The correlation between the degree of damage of the skin barrier and the severity of the disease give a reason to assume that the itch aggravates the disease and leads to a dysfunction of the barrier homeostasis.

Results from the 14item Hospital Anxiety and Depression scale HAD-A and HAD-D show that in 77.29% of patients the rate of anxiety is normal be-

anxiety



Fig. 1. Anxiety distribution

1.05% have severe anxiety range 15-21 (Figure 1). The data from the depression evaluation show that most of the patients 94.73% do not have depression range 0-7; 4.21% have mild depression and 1.05% have moderate depression (Figure 2).



Fig. 2. Depression distribution

With regard to the duration of the disease the range was from 57.14% (6 to 8 months) to 87.75% (18-24 months). Mild anxiety was marked the most significant in patients with disease duration of 24-30 months-33.34%, and at least 4.28% in LSC with duration from 6 to 8 months. Moderate anxiety was most significant in the group of patients with duration 6-8 months - 28.57%. Regarding the severity of disease results show no correlation between the levels of anxiety and the severity of LSC. The patients with severe anxiety were in a group of stage II of the disease and patients in a group of stage III show normal levels of anxiety. So the results obtained from the comparison the level of anxiety and disease characteristics show

little correlation with the duration and absence of correlation with the severity of LSC (Table 4).

Interesting results were obtained when comparing the anxiety level and evaluating the functioning

Anxiety		Stage of the disease					
	Stage I	Stage II	Stage III	Stage IV	p-		
normal	12 (33.3%)	18 (50%)	6 (16.7%)	0 (0%)			
mild	7 (53.8%)	3 (23%)	3 (23.1%)	0 (0%)	0.002		
moderate	2 (50.0%)	1 (25%)	1 (25.0%)	0 (0%)	0.092		
severe	0 (0%)	0 (0%)	2 (100%)	0 (0%)			

Table 4. Distribution of anxiety according severity of LSC

LSC is a skin disease included in a group of medical conditions due to psychological factors (11,12). Simonetta and associates (2015) published data reaffirming that psychogenic factors play an important role in the development of LSC (13). Liao and associates (2014) in their retrospective study found that patients with marked anxiety are at 1:41 higher risk of developing LSC compared to the healthy population (14). In our study the results from comparing the level of anxiety with the duration of LSC support the theory of the psychogenic origin of itch in the pathogenesis of the disease (15). The lack of correlation between the severity of LSC and the levels of anxiety support the data export by Perez et al (2009) that in some patients the psychogenic factor is a leading cause of the disease (16).



Figure 3. The correlation between the functioning of the skin barrier and the anxiety levels

of the skin barrier. Results show that patients with mild and moderate anxiety have light changes in the functioning of the skin barrier (19.64%). The patients with the lowest H and the highest TEWL have normal results from the HAD-A and HAD-D scales (21.13%) (p=0.001) (Figure 3).

CONCLUSION

Disorders of the skin barrier are described as a trigger of pruritoceptive pruritus while psychosomatic disorders are associated with psychogenic pruritus in patients with LSC. The absence of correlation between changes in H and TEWL and psychosomatics means that in patients suffering from LSC there are two independent pathological ways of development of the illness.

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