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Krishna Thapa
Department of Dermatology,
Universal College of Medical
Sciences, Nepal

Determine of S100A8 level in patients of chronic spontaneous urticaria

Krishna Thapa

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Abstract

Introduction: Chronic spontaneous urticaria (CSU) is characterized by the spontaneous appearance of transient itchy wheals, angioedema or both, for at least 6 weeks. The present study was conducted to determine level of S100A8 in patients of chronic spontaneous urticaria.

Materials & Methods: The present study was conducted on 82 cases (Group I) of chronic spontaneous urticaria of both genders. Equal number of healthy subjects (Group II) was also taken. CSU was diagnosed according to the EAACI/GA2 LEN/ EDF/WAO guidelines. Plasma levels of S100A8 were assessed using human enzyme linked immunosorbent assay (ELISA) kit.

Results: There were 34 male and 48 female in group I and 42 males and 40 female in group II. Mean age in group I was 27.1 years and in group II was 29.2 years. Basophil count was 0.21% in group I and 0.45% in group II. Eosinophil count was 1.62 % in group I and 2.15% in group II. The mean S100A8 level in group I was 2415.6 pg/ml and in group II was 1380.9 pg/ml.

Conclusion: Authors found that Plasma S100A8 level was higher in CSU patients. It may serve as biomarkers of CSU.

Keywords: Basophil, Chronic spontaneous urticaria, S100A8

Introduction

Chronic spontaneous urticaria (CSU) is characterized by the spontaneous appearance of transient itchy wheals, angioedema or both, for at least 6 weeks. While it affects 0.5%–1% of the population, it not only has negative visual impact on patients, but also decreases the quality of life ^[1]. CSU is an inflammatory disease, probably caused by an interactive combination of immune, genetic, and environmental factors, including infections. Various changes in levels of immune-inflammatory, coagulation/fibrinolytic, hormonal, and metabolic markers have been reported in CSU patients ^[2].

Urticaria presents with wheals (hives), angioedema, or both, and has a lifetime prevalence of about 9%. The appearance of pruritic, erythematous dermal swellings that blanch with pressure, indicating the presence of vasodilation and superficial dermal edema, is characteristic of wheals ^[3]. Angioedema is caused by similar pathologic alterations that occur in the reticular dermis and subcutaneous tissue, with poorly defined swelling and burning. One-third of patients present with both hives and angioedema, 30% to 40% present with isolated hives, and 10% to 20% with isolated angioedema ^[4].

S100 family consists of a serial of EF-hand calcium (Ca²⁺)-binding proteins, with more than 20 distinguished proteins. It is reported that S100A8, S100A9, and S100A12 play important roles in the pathogenesis of immunological disorders in the human body. They are involved in the development of autoimmune-associated diseases, such as psoriasis, rheumatoid arthritis, and systemic lupus erythematosus ^[5]. The present study was conducted to determine level of S100A8 in patients of chronic spontaneous urticaria.

Materials & Methods

The present study was conducted in the department of Dermatology. It comprised of 82 cases (Group I) of chronic spontaneous urticaria of both genders. Equal number of healthy subjects (Group II) was also taken. All were informed regarding the study and written consent was obtained. Ethical clearance was taken before starting the study.

General information such as name, age, etc. was recorded. In all cases, the clinical and

Corresponding Author:
Krishna Thapa
Department of Dermatology,
Universal College of Medical
Sciences, Nepal

dermoscopic characteristics were recorded. CSU was diagnosed according to the EAACI/GA2 LEN/ EDF/WAO guidelines. Urticaria Activity Score (UAS) was used to assess the activity of urticaria, including the wheal numbers and pruritus. Plasma levels of S100A8 were assessed using human enzyme linked immunosorbent assay (ELISA) kit. Results thus obtained were subjected to statistical analysis. P value less than 0.05 was considered significant.

Results

Table I: Assessment of parameters

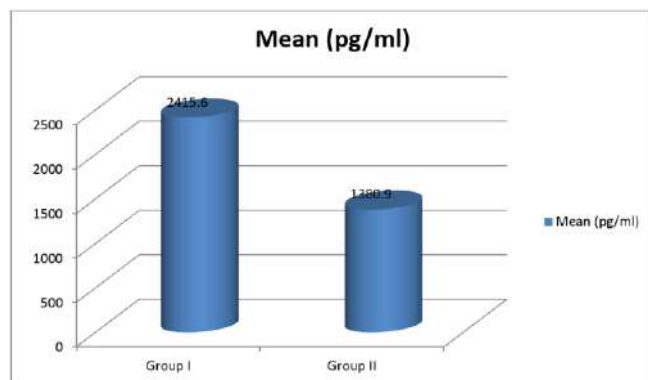
Variables	Group I (Cases)	Group II (Control)
Male	34	42
Female	48	40
Mean age (Years)	27.1	29.2
Basophil %	0.21	0.45
Eosinophil %	1.62	2.15

Table I shows that there were 34 male and 48 female in group I and 42 males and 40 female in group II. Mean age in group I was 27.1 years and in group II was 29.2 years. Basophil count was 0.21% in group I and 0.45% in group II. Eosinophil count was 1.62 % in group I and 2.15% in group II.

Table II: Assessment of S100A8 in both groups

Variables	Mean (pg/ml)	P value
Group I	2415.6	0.05
Group II	1380.9	

Table II, graph I shows that mean S100A8 level in group I was 2415.6 pg/ml and in group II was 1380.9 pg/ml.



Graph I: Assessment of S100A8 in both groups

Discussion

Urticaria may be classified into acute or chronic. Acute urticaria (AU) is defined by the occurrence of spontaneous wheals or angioedema for <6 weeks. In acute cases, it is important to exclude anaphylaxis in the presence of respiratory, gastrointestinal, or neurologic symptoms or hemodynamic instability [6]. Eliciting factors have been found in <50% of cases, with upper respiratory infections being the most common trigger (40%), followed by drug reactions (9.2%) and suspected food intolerance (0.9%). Among infectious agents, upper respiratory tract agents, Mycoplasma pneumonia, and parasitic infections have been commonly reported in children, while viral hepatitis and infectious mononucleosis are important culprits in adults [7]. The present study was conducted to determine level of

S100A8 in patients of chronic spontaneous urticaria.

In this study, there were 34 male and 48 female in group I and 42 males and 40 female in group II. Mean age in group I was 27.1 years and in group II was 29.2 years. Basophil count was 0.21% in group I and 0.45% in group II. Eosinophil count was 1.62 % in group I and 2.15% in group II.

Zhou *et al.* [8] assessed the levels of plasma S100A8, S100A9, and S100A12 were measured in 51 CSU patients and 20 healthy controls using enzyme linked immunosorbent assay kits. The values in the patient group and that of the healthy controls were statistically compared. The plasma levels of S100A8, S100A9, and S100A12 were significantly higher in CSU patients than those in controls. Interestingly, the level of S100A12 was significantly correlated with S100A8 and S100A9 in CSU patients ($P < 0.05$ and $P < 0.001$, respectively). In addition, S100A8, S100A9, and S100A12 were all significantly inversely correlated with blood basophil percentage.

We observed that mean S100A8 level in group I was 2415.6 pg/ml and in group II was 1380.9 pg/ml. About one-third to one-half of patients with CSU show a positive response against their own serum (positive autologous serum skin test [ASST]). IgG antibodies to the high-affinity IgE receptor FcεR1a, or less commonly IgG antibodies to IgE, have been documented. There seems to be an increased risk for thyroid disorders (hypothyroidism more often than hyperthyroidism), diabetes mellitus type I, systemic lupus erythematosus, and rheumatoid arthritis in patients with CaU. Although these autoantibodies are of academic interest, as some studies report a more intense refractory course, their clinical relevance remains unclear [9].

The exact pathogenic role of S100A8, S100A9, and S100A12 in the development of CSU is still unknown. Three S100 proteins are involved in the pathogenesis of CSU in several ways. Firstly, these three S100 proteins bind to and activate responses by two widely expressed but divergent receptors, namely, toll-like receptor 4 (TLR4) and the receptor for advanced glycation end-products (RAGE). The engagement of these two receptors by S100 proteins is linked to an array of signaling pathways, notably NF-κB and mitogen-activated protein kinases; the induction of p38 signaling was known to trigger the release of proinflammatory cytokines including interleukin (IL)-6, tumor necrosis factor (TNF)-α, and IL-1β through the action of NF-κB [10].

Conclusion

Authors found that Plasma S100A8 level was higher in CSU patients. It may serve as biomarkers of CSU.

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