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Basal serum cortisol and adrenocorticotrophic hormone levels in patients with atopic dermatitis

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Background: Atopic dermatitis (AD) is an inflammatory skin disease with eczematous pruritic lesions. Topical corticosteroids are the most widely used and the mainstay of treatment for AD. There are some studies that percutaneous systemic absorption of topical steroids may occur and lead to suppression of hypothalamic-pituitary-adrenal axis (HPAA). However, almost in all of these studies, "basic" HPAA function (before application of topical steroids) was not evaluated.

Aim: The aim of this study was to investigate basal serum cortisol, adrenocorticotrophic hormone (ACTH) and IgE levels in patients with AD and their correlation with disease severity.

Materials & Methods: Levels of basal serum cortisol, ACTH and IgE were assessed by ELISA in 31 patients with AD and 31 control subjects. Clinical severity of AD was evaluated by the SCORAD (SCORing Atopic Dermatitis) index.

Results: Data analysis showed no statistical difference for basal serum cortisol and ACTH levels between two groups. The serum IgE level was significantly higher in AD group ($P=0.02$). The SCORAD index was correlated with serum IgE level but not with the basal serum cortisol level and ACTH level.

Conclusion: Basal serum cortisol and ACTH levels are normal in AD patients. Serum IgE level is significantly higher in AD patients and correlated with disease severity.

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Depression in patients with psoriasis: Effects of biologics

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Psoriasis is a common chronic immune-mediated inflammatory disorder affecting 2-3% of the population. Psoriasis is associated with a variety of psychological difficulties, including poor self-esteem, social stigmatization, anxiety, depression and suicidal ideation. Psoriasis is associated with substantial impairment of health-related quality of life (HRQOL). Some studies estimated that around one-quarter of patients in dermatology practices have psychiatric disorders. Significantly elevated concentrations of TNF- α have been observed in the plasma of patients with psoriasis and in patients with major depression. TNF α has been implicated in the pathogenesis of both psoriasis and depression. In humans, administration of TNF α blockers such as etanercept (Enbrel), infliximab (Remicade) and adalimumab (Humira) and IL-12/23 inhibitors have been found to improve depression that accompany psoriasis.

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