INTRODUCTION

Tuberous sclerosis complex (TSC), also known as Bourneville disease or Bourneville–Pringle disease, is an autosomal dominant genetic disorder with various clinical manifestations that affects the brain, skin, kidneys, heart, and other organs [1]. The association of STB with psoriasis is very rare and suggests a common pathophysiology. We report such a rare case of tuberous sclerosis in a 45-year-old male with psoriasis.

CASE REPORT

A 45-year-old male with no significant pathological antecedent presented himself to the urology department with low back pain. The patient had had asymptomatic cutaneous lesions since the age of six years with extension and increase in number and size, as well as similar cases in the family (father, four brothers, granddaughter, nephews, and nieces), with no notion of consanguinity in the parents, and pruriginous, erythematous, squamous lesions evolving by pushed remission since the age of ten years. The patient did not report epileptic seizures, psychomotor disorders, or other possible associated signs. A dermatological examination revealed multiple angiofibromas (Fig. 1a) symmetrically distributed over the centrofacial areas, fibrous cephalic plaques (Fig. 1b) sitting at the level of the forehead and scalp, a shagreen plaque with a 10-cm long axis (Fig. 1c) sitting at the left axillary level, multiple skin tags around the neck, and periungual fibromas (Figs. 1d and 1e). Erythematous squamous plaques sitting at the knees and elbows were bilateral and symmetrical with a methodical scratching of positive pitcher; the body surface was 4% (Fig. 2). The patient did not show any symptoms of cardiovascular, endocrine, respiratory, immune, or musculoskeletal disorders. Ultrasonography and abdominal CT revealed a multicystic kidney without pylocelical dilation with pylocelictic stones. The diagnosis of STB was retained and was associated with psoriasis. The patient benefitted from a double “J” probe uplift in the urology department and application of a topical corticosteroid on the psoriasis plaque with an improvement.
DISCUSSION

Tuberous sclerosis complex (TSC) is one of a group of related disorders known as neurocutaneous syndromes or phakomatoses with an incidence rate of approx. 1 in 5000–10,000 live births [2,3]. It is an autosomal dominant disease with high penetrance caused by genetic mutations in one of the TSC1 or TSC2 genes [2], which results in the overactivation of mammalian target of rapamycin complex 1 (mTOR), a key intracellular regulator of cell growth and proliferation, leading to hamartomatous lesions in several organs [3,4].

TSC is a multisystem disorder with various clinical manifestations. The wide spectrum of clinical features results from the formation of hamartomas in various organs. Hamartomas are frequently present in the skin, brain, kidneys, heart, and, less frequently, in the lungs, retina, gingiva, bones, and gastrointestinal tract [5]. The diagnosis is based on the association of major criteria and minor criteria (Table 1) [6]. The diagnosis is made when two major criteria, or one major and two minor, are fulfilled. The diagnosis and management of TSC is often challenging. The treatment involves addressing the symptoms caused by the hamartomas. Inhibitors of the mTOR pathway, such as rapamycin, have an immunosuppressive and antiproliferative action. This drug is effective in reducing the volume of the tumors.

On the other hand, psoriasis is a chronic autoimmune inflammatory skin disorder, following the proliferation and abnormal differentiation of keratinocytes, which are under the influence of several factors whose genetic component remains the most likely [7], including mutation of the genes of the chemokine, including MCP1, CCR2, and CCR5. The PSORS2 locus is a gene that is the active transcription factor in inflammation and immunity located in the region of the gene encoding the RAPTOR protein, a protein associated with mTOR regulation. Thrombin/threonine protein kinase (mTOR) regulates growth and cell proliferation in response to environmental stimuli. It is overexpressed preferentially at the psoriatic level of lesional and nonlesional skin [8].

In psoriasis, dysregulation of cytokines and growth factors may lead to the activation of the mTOR signaling system, which initiates the proliferation of keratinocytes and synovial cells responsible for psoriatic arthritis. For the first time, we explored how a dual kinase inhibitor of mTOR signal proteins may be an equally effective...
therapeutic agent for psoriasis. The association with TSC in our patient suggests the genetic predisposition of psoriasis vulgaris [9]. It blocks IL-2-induced LT proliferation via mTOR inhibition.

CONCLUSION

To our knowledge, this is the first report that describes the association of psoriasis with STB in a patient. These inflammatory diseases are chronic with mainly cutaneous manifestations and share a hereditary character and a common pathophysiology, but with different mutations.

Consent

The examination of the patient was conducted according to the principles of the Declaration of Helsinki.

REFERENCES


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