Case report

Juvenile Systemic Lupus Erythematosus associated with Klinefelter's syndrome: A case report

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ABSTRACT

We present the first reported case of juvenile Systemic Lupus Erythematosus with Klinefelter’s syndrome in a 14-year-old Iranian boy who had leg ulcers and arthritis. He had low level of testosterone accompanied with hypergonadotropic hypogonadism. This case emphasizes the importance of two X chromosomes as a risk factor for Systemic Lupus Erythematosus in women and men with Klinefelter’s syndrome (47, XXY).

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Introduction

Klinefelter’s syndrome is the most frequent major abnormality of sexual differentiation in men and affects one in every 500–1000 born males.1 Men with Klinefelter’s syndrome have more than one X chromosome, usually two X chromosomes (47-XXY). The phenotype is characterized by eunuchoid appearance, increased length of legs and arms, scanty facial and body hair, gynecomastia, small and firm testes, and hyperpigmentation of the lower extremities.2 Hormonally, the syndrome is characterized by hypergonadotropic hypogonadism in which testosterone level is usually half of normal.3 Patients with Klinefelter’s syndrome show a higher percent of Systemic Lupus Erythematosus (SLE) and other skin-related autoimmune diseases than the normal population. SLE may be a presenting symptom of Klinefelter’s syndrome and may lead to diagnosis of the disease.4 Here we present a case of juvenile Systemic Lupus Erythematosus with Klinefelter’s syndrome and most probably the first report from Middle East.

Case report

A 14-year-old Iranian boy presented to our department with a 3-week history of arthritis on both of his wrists and ankles, accompanied by myalgia and photosensitivity. His physical examination revealed butterfly rash, palpable and plantar erythema, and some small ulcers on his legs. He had low head hair implantation, and hyperpigmentation on face. Other physical findings were gynecomastia, pectus excavatum, small and firm testicles, and long extremities. The results of his laboratory investigations were as follows: white blood cell count
His renal biopsy showed mesangial glomerulitis. He was treated with oral prednisone, lupus anticoagulant, blood tests were within normal limits or negative, including factors (positive +2), C-reactive protein (positive 17 mg/L), C3 4%, red blood cells count 424 × 10⁹/mm³, platelets count 140 × 10⁹/mm³, hemoglobin 11.3 g/dL, hematocrit 34.1%, erythrocyte sedimentation rate 84 mm/h, aspartate amino transferase 37 U/L, alanine aminotransferase 31 U/L, lactate dehydrogenase 232 U/L, creatinine kinase 67 U/L, aldolase 5.6 U/L, blood urea nitrogen 27 mg/dL, creatinine 0.9 mg/dL, total protein 7.4 g/dL, antinuclear antibody (positive > 1:160 homogeneous pattern), rheumatoid factors (positive +2), C-reactive protein (positive 17 mg/L), C3 54 mg/dL, C4 11 mg/dL, IgG 2210 mg/dL, IgM 220 mg/dL, IgA 191 mg/dL, and IgE 500 mg/dL. His urine analysis revealed 2-glycoprotein, anticardiolipin IgG and IgM (World Health Organization Class II).

Discussion

Klinefelter’s syndrome can present at any age. In adult men, the diagnosis may be made during investigation for sterility, but, at puberty, eunuch body might be the hallmark. Children with Klinefelter’s syndrome might be diagnosed because of learning difficulties or social problems. Klinefelter’s syndrome and SLE occur together more often than would be expected by chance alone. Studies have shown a clear relationship between low levels of testosterone and high prevalence of ulceration in patients with Klinefelter’s syndrome. Treatment with testosterone leads to improvement of leg ulcers in these patients.8 Our patient showed both leg ulcers and low level of testosterone, whose leg ulcers were also reduced during subsequent therapy that included testosterone.

Males with untreated hypogonadism, associated with significant gonadal failure and very low levels of testosterone, have an increased risk of developing rheumatoid/autoimmune diseases. In fact, testicular dysfunction predisposes to the development of rheumatoid/autoimmune diseases. These patients have an increased frequency of antinuclear antibody and anticytokerin antibodies compared with other hypogonadotropic hypogonadlic patients.9 Although, our patient’s sera were not positive for anticytokerin antibodies, it was positive for antinuclear antibody (positive > 1:160 homogeneous pattern). SLE may be a presenting symptom of Klinefelter’s syndrome and may lead to diagnosis of the disease. The first presentation of our patient was arthritis, myalgia, and photosensitivity. However, his follow-up revealed hypergonadotropic hypogonadism and his karyotype confirmed the diagnosis of Klinefelter’s syndrome.

Humoral and cellular immunities are enhanced in Klinefelter’s syndrome, as a result of testosterone deficiency and increased levels of estradiol, which enhance autoantibody production. Treatment with testosterone has also proved to suppress both cellular and humoral immunities in these patients.9,10

Conclusion

The frequency of Klinefelter’s syndrome is increased in men with SLE compared with its prevalence in men without SLE. Therefore Klinefelter’s syndrome and SLE might be associated. Whereas Klinefelter’s syndrome in children is often subclinical, the pediatric rheumatologists treating male children with lupus should be aware of Klinefelter’s syndrome and provide them access to imperative medical management.

References