Case Report

Complement-4 Deficiency in a Child with Systemic Lupus Erythematosus Presenting with Standard Treatment-Resistant Severe Skin Lesion

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The complement system is of great importance in systemic lupus erythematosus. Complete genetically determined deficiencies are with few exceptions reported for the various complement proteins, and most of the deficiency states are rare. Deficiencies of the factors in the classical pathway are also associated with development SLE and SLE-like disorders. Most of the patients with lupus present skin involvement. Approximately, 75–95% of patients with cutaneous lupus erythematosus respond to antimalarial therapy and/or topical glucocorticosteroids. Immunosuppressive agents are usually considered a second-line approach in patients with resistant disease. In this study, we present the clinical features and determine the molecular basis responsible for the complete C4A and C4B deficiencies in a lupus patient presented subacute cutaneous lupus erythematosus and resistance to treatment.

1. Introduction

Lupus erythematosus is a chronic, autoimmune disease resulting from an interaction of genetic, environmental, and hormonal factors and characterized by a spectrum of clinical forms with a variable evaluation from a localized cutaneous form to life-threatening systemic form. Skin involvement occurs in 70–85% of all patients with lupus [1]. Specific skin lesions of cutaneous lupus erythematosus are classified as acute cutaneous lupus erythematosus, subacute cutaneous lupus erythematosus (SCLE), and chronic cutaneous lupus erythematosus, according to the clinical characteristics of the lesions [1].

Subacute cutaneous lupus erythematosus (SCLE) may present with annular or papulosquamous cutaneous lesions that are symmetrically located in the sun-exposed areas of the body [2]. SCLE shows typical serological findings, with anti-Ro (SS-A) antibodies present in up to 100% of patients [3, 4]. However, antinuclear antibodies (ANAs), complement C3/C4 deficiency, rheumatoid factor, circulating immune complexes, lymphopenia, and thrombocytopenia can frequently be detected. Severe neurological or nephrological involvement is rare in SCLE, whereas mild musculoskeletal involvement is commonly observed. It is well known that some patients suffering from CLE develop extracutaneous manifestations during the course of their disease; up to 30% of patients with SCLE show systemic involvement [5]. Most patients with SCLE can be sufficiently treated with photo protection, topical corticosteroids, antimalarials, or a combination of these. When standard therapy fails, second-line agents approach [6].

The complement system is a group of plasma and membrane proteins that are sequentially activated via proteolytic cleavages to defend against microbial infections [7, 8].

The complement system is of great importance in systemic lupus erythematosus (SLE). Complement contributes to inflammation and tissue damage in this disease, but seemingly in paradox, deficiency of some complement proteins dramatically increases the risk for SLE [9]. Deficiency states within the classical pathway are associated with increased risk to develop SLE and SLE-like disease [10].

Genetically determined complement deficiencies are inborn errors and may have an impact on the development...
of the immune system. Classical pathway deficiency has led to different hypothesis to explain roles for complement in SLE pathogenesis [9]. Complement deficiency leads to impaired handling of immune complex and inadequate clearance apoptotic cell debris [11, 12]. Another hypothesis suggests that the complement system has an important role in the development of tolerance against self [13]. Also, complement deficiency results in lack of normal B-cell tolerance and provides production of autoantibodies [9]. In addition, complement components are in some ways also important for regulation of cytokine production, especially Type 1 interferons have been shown to have a central role in the pathogenesis of SLE [14, 15].

While the genetic basis for a majority of SLE cases is polygenic, a homozygous deficiency in one of the early complement components alone can be strong enough to cause the disease, a situation similar to a single gene defect in an autosomal recessive disease [16, 17]. Respectively, 93% and 78% of patients with complete C1q and C4 deficiencies eventually develop SLE or a lupus-like disease [18].

Genetically, the complement C4 gene located in the class III region of the major histocompatibility complex (MHC) on chromosome 6p21.3. About 40 protein variants for complement C4 have been documented [19]. These proteins are segregated into two classes, the acidic C4A and the basic C4B. Each C4 gene either codes for a C4A or a C4B protein [20]. C4A is believed to be important in the clearances of immune complexes, and C4B is more powerful in propagating the complement activation cascades. The complete absence of both C4A and C4B proteins may therefore lead to decreased ability of immune defense against microbes as well as inefficient disposal of immune complexes.

To date, 28 individuals with complete C4 deficiency from 19 families have been reported [21, 22]. Among them, 15 individuals developed SLE, 7 developed lupus-like diseases, and four of the remaining subjects were only afflicted by kidney diseases [23].

In this study, we present the clinical features and determine the molecular basis responsible for the complete C4A and C4B deficiencies in a lupus patient presented with subacute cutaneous lupus erythematosus and progression systemic form and resistance to treatment.

2. Case Report

Seventeen-year-old patient was a male born by parents of non consanguineous marriage. He was diagnosed with SCLE at the age of 15 years old and had local therapy in dermatology clinic before admitting our clinic. Family history for rheumatologic disease was negative. On physical examination, his temperature 38.3, respiratory rate 38/min, pulse rate 100/min, and blood pressure was found as 90/70 mmHg. In auscultation, lungs were clear and the heart sounds were normal. Abdominal palpation was normal. His clinical presentation included severe malar rash with marked photosensitivity and bullous lesions on the tips of fingers and hands (Figure 1). Skin biopsy was performed. There was interface dermatitis with irregularity and loss of basal cells. In the papillary and upper reticular dermis was a perivascular and interstitial lymphocytic infiltrate. There was granular deposition of Ig A, IgM, and C3 at the dermal-epidermal junction. Also he has arthritis which described swelling, tenderness, and pain on all small joints.

Laboratory tests showed an erythrocyte sedimentation rate (ESR) of 100 mm/h, C-reactive protein of <0.3 mg/dL (normal), hemoglobin of 11.1 g/dL, hematocrit of 31%, white blood cell count (WBC) of 4900/mm³ with normal differential count, platelet count of 197000/mm³, serum urea of 25 mg/dL, serum creatinine of 0.8 mg/dL, albumine of 3.5 g/dL, total cholesterol of 106 mg/dL, triglycerides of 57 mg/dL, calcium of 8.7 mg/dL, sodium of 140 mEq/L, potassium of 4.7 mEq/L, chloride of 107 mEq/L, alanine aminotransferase of 23 U/L, and aspartate aminotransferase of
We report a child with SLE and C4 deficiency and with severe skin involvement that proved to be subacute cutaneous lupus erythematosus (SCLE). In our patient, the diagnosis of SLE was based on the presence of constitutional symptoms, malar rash, photosensitivity, oral ulceration, arthritis, anemia, a high titer of ANA, and low C3, C4. The patient resistant to initial therapies including HQK, Prednisolon and CYC. Rituximab is a chimeric monoclonal IgG1 antibody that binds specifically to the CD20 antigen and mediates B-cell lysis. Prior studies have shown that rituximab therapy can be a safe and efficacious addition to therapy with standard immunosuppressive agents in patients with refractory childhood SLE [24–29]. In our case, Rituximab led to a remarkable improvement of skin lesions, resulting in a significant decrease of the SLEDAI from 20 at the beginning to 5 at the end of therapy. In addition, he exhibited a low C4 level during the extended period of observation. Finally, the genetic phenotyping of complement revealed that he was homozygous for C4B deficiency.

SLE is a systemic autoimmune disease characterized by the breakdown of immunotolerance and the production of a wide range of autoantibodies that target multiple tissues and organs [30]. Homozygous complement deficiencies represent rare conditions. These are estimated to be found in less than 1% of SLE patients [31, 32]. Respectively, 93% and 78% of patients with complete C1q and C4 deficiencies eventually develop SLE or a lupus-like disease [22, 33]. In addition, the concordance rates for siblings with homozygous deficiency of C1q or C4 to develop SLE are 90% and 80%, respectively, which are even higher than the rate in monozygotic twins (26–60%) with other genetic defects [17].

C4A deficiency causes autoimmune disease, especially systemic lupus erythematosus [34]. Indeed, approximately 15% of Caucasian SLE patients exhibit homozygous C4A deficiency, whilst more than 50% of Caucasian patients with SLE have heterozygous C4A deficiency [35]. C4B deficiency caused autoimmune-associated diseases, such as systemic lupus erythematosus, or diseases with an autoimmune component, such as autism [36, 37].
Heterozygous and homozygous deficiencies of C4A were present in 40–60% of SLE patients from almost all ethnic groups such as Northern and Central Europeans, Anglo-Saxons, Caucasians in the US, African Americans, Asian Chinese, Koreans, and Japanese [38]. The major causes for C4AQ0 in Caucasian and African SLE patients are the presence of a mono-S RCCX module with a single, short C4B gene and the 2-bp insertion into the sequence for presence of a mono-S RCCX module with a single, short C4B gene, also heterozygous and homozygous deficiencies of C4B in our patient.

In conclusion, we report a case of C4B deficiency accompanied with SLE with therapy-resistant skin involvement. Further studies are needed to clarify the relationship of C4B deficiency and SLE which presented severe skin involvement.

Conflict of Interests

The authors declare no conflict of interests.

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