Vitiligo, morphea and Sjögren’s syndrome: a case study

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Introduction

Sjögren’s syndrome, vitiligo, and morphea are three distinct autoimmune diseases of unknown etiology. The existence of autoantibodies support an autoimmune hypothesis of these diseases.1,2,3

Case Report

A 56-year-old woman was admitted to the rheumatology clinic with symptoms of burning and the feeling of grit or sand in her eyes. She also had dryness of the mouth, giving rise to difficulties with speech and swallowing food. The presence of anti-Ro/SSA and anti-La/SSB autoantibodies, and grade 4 chronic lymphocytic sialadenitis on biopsy of the salivary gland, together with the clinical findings revealed Sjögren’s syndrome. The patient was referred to the dermatology clinic because of widespread depigmentation in the middle of her back (Figure 1). Following the dermatological examination, the patient was considered to have vitiligo. With the help of Wood’s light examination, the diagnosis was confirmed. Another remarkable dermatological finding was violaceous-colored, slightly depressed and atrophic oval plaques of variable size (10x10 cm to 7x8 cm) on the abdominal and gluteal regions (Figure 2). Histopathological examination of these lesions demonstrated morphea. The patient was unaware of these dermatological conditions.

In her past history, the patient had received radioactive iodine therapy for an adenoma of the thyroid gland eight years ago. In the family history, her daughter had been diagnosed with rheumatoid arthritis. Systemic examination of the patient was unremarkable. There was no systemic involvement associated with the morphea and Sjögren’s syndrome. Laboratory findings for a complete blood cell count, vitamin B12 levels, biochemical analysis, and urinalysis were normal. However, on admission abnormal laboratory findings included: erythrocyte sedimentation rate, 30 mm/hour (normal range, 0-20 mm/hr); C-reactive protein, 3 mg/L (0-5 mg/L); rheumatoid factor (RF), 604 IU/ml (0-15 IU/mL); and anti-nuclear antibody (ANA), 1/320 granular pattern. While anti-Ro/SSA and anti-La/SSB antibodies were positive, anti-double-stranded DNA (anti-dsDNA), and anti-centromere antibody were negative. A biopsy of the salivary gland revealed grade 4 chronic lymphocytic sialadenitis based on the Chilson classification. Schirmer’s tear test was used for the evaluation of tear secretion by the lacrimal glands. There was wetting of less than 5 mm per 5 min. This demonstrated a strong indication of diminished secretion.

Autoimmune etiology is thought to play a role in these diseases. Sjögren’s syndrome is an autoimmune disorder with a higher female incidence, distinctive human leukocyte antigen (HLA) associations, familial clustering with other autoimmune processes, and the presence of autoantibodies, such as anti-Ro/SSA and anti-La/SSB antibodies.4 The association of vitiligo with autoimmune diseases has suggested an immunological basis for vitiligo. Involvement of humoral immunity was first demonstrated by the finding of circulating antibodies directed to melanocytes. The vitiligo antibodies are attracted predominantly to various melanocyte antigens, including tyrosinase, tyrosinase-related protein 1, and tyrosinase-related protein 2. Recently, autoantibodies to a transcription factor called SOX10 have been found in vitiligo related to polyglandular dysfunction.1 Some studies have indicated an association of autoimmune diseases with morphea. In addition to positive titers of ANA, anti-dsDNA, and anti-scleroderma (ScI) 70 antibodies, increased RF levels suggest that an immunological process occurring in morphea.6,7

Although Sjögren’s syndrome, vitiligo, and morphea are autoimmune diseases, they are seen together rarely. A few studies have report...

Figure 1. Vitiligo localized in the lumbar region.

Figure 2. Morphea localized in the gluteal region.
ed an association between morphea and vitiligo to date. There is one reported case of alopecia areata, lichen planus, and onychodystrophy in a patient with vitiligo and morphea. Among other reported cases are Hashimoto thyroiditis with vitiligo and morphea, as well as alopecia areata and ulcerative colitis with vitiligo and scleroderma. There is no data in the literature to evaluate the coincidental occurrence of these diseases; the associations between vitiligo, morphea, and other autoimmune diseases were revealed by case studies.

To our knowledge, there is no reported case of an association between vitiligo, morphea, and Sjögren’s syndrome. The existence of an autoimmune disease makes people prone to other autoimmune diseases. Because of this condition, the patients with an autoimmune disease should be followed carefully.

References