A Case of IgG4-related Mesenteritis and Ankylosing Spondylitis

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ABSTRACT

Immunoglobulin G4-related disease (IgG4-RD) is a systemic fibroinflammatory disease of unknown etiology and pathogenesis, which can affect all organs. The most commonly involved organs are the pancreas, hepatobiliary system, salivary glands, orbits and lymph nodes. Rarely, the thyroid, pituitary, aorta, lung and kidney may also be involved; mesenteric involvement is rare. The association of IgG4-RD with some rheumatological diseases is observed; while there are data in the literature about rheumatoid arthritis and Sjögren syndrome, but association with ankylosing spondylitis is rare. Anti-tumor necrosis factor alpha (anti-TNFα) treatment has been tried in refractory IgG4-RD cases; but this case is of great interest in terms of progressive presentation under treatment with etanercept, an anti-TNFα agent, which is also used for the treatment. But, IgG4-related mesenteritis with ankylosing spondylitis receiving anti-TNFα therapy, which has also been tried in the treatment of IgG4 disease, has not been reported in the literature.

Key Words: IgG4-related disease, Mesenteritis, Ankylosing spondylitis.


INTRODUCTION

Immunoglobulin G4-related disease (IgG4-RD) is a chronic inflammatory syndrome with an imprecisely elucidated pathogenesis. It is characterised by the presence of tumoral mass lesions, storiform fibrosis, obliterative phlebitis, and IgG4 producing plasma cells in the affected organs.¹ The most commonly involved organs are the pancreas, hepatobiliary system, salivary glands, orbits and lymph nodes. Mesenteric involvement is rare.²

There are data in the literature on association of this syndrome with rheumatoid arthritis (RA) and Sjögren syndrome (SS), but association with ankylosing spondylitis (AS) is rare.¹² We, herein, present a patient receiving anti-TNFα for AS, who progressed under treatment and was diagnosed with IgG4-related sclerosing mesenteritis.

CASE REPORT

A 38-year male was diagnosed with HLA B27-positive AS, according to the Modified New York criteria in 2015; and anti-TNF-α agent, etanercept, was started three years earlier, as there was no response to sulfasalazine 2×2 and non-steroidal anti-inflammatory therapy. Grade I hydronephrosis was observed in the right kidney during 15th month of etanercept treatment on ultrasonography obtained due to swelling in both testicles and abdominal pain. Abdominal computed tomography (CT) revealed a mass of 4.5 cm in diameter, with hyperdense soft tissue density, in the medial aspect of the ascending colon (Figure 1). The patient’s ESR and CRP levels were 85 mm/1st h and 6.7 mg/dl, respectively. Anti-TNFα treatment was discontinued and the patient was operated with a preliminary diagnosis of colon malignancy. Intraoperatively, a hard, fixed mass of 4×5 cm was observed to extend behind the cecum to the mesentery. Gonadal structures entered the mass, with which the ureter was closely related. The mass involved the ileocolic mesentery in full-thickness and extended to the inferior vena cava. Complete resection of the mass along with right hemicolectomy was performed. Pathological examination revealed lymphoid follicles, increased plasma cells, obliterative phlebitis and multinuclear giant cells, accompanied by pronounced fibrosis in the mesenteric fat tissue. CD20, CD3, and Ki-67 were positive in scattered cells on immunohistochemistry. IgG4 positive cells were on average 70/high power field (HPF). The IgG4 (+)/IgG (+) cell ratio was 64%, and vascular structures were CD34 positive on immunostaining.
Pathological diagnosis was IgG4-related sclerosing mesenteritis (Figures 2 and 3). It was found that plasma IgG4 levels were increased to 457 mg/dL and IgE, and eosinophil levels were normal. Postoperatively, obtained PET-CT was normal. The patient was started on 0.6 mg/kg methylprednisolone, which was gradually decreased and stopped on an average of three months. Control CTs of the patient revealed no disease recurrence. IgG4 level six months after the treatment was 190 mg/dL, and the patient remained well on follow-up for two years.

**DISCUSSION**

IgG4-RD is a chronic and autoimmune disease associated with increased IgG4 levels in multiple organs and/or tissues. It can affect many organs. Rate of mesenteritis in IgG4-RD varies between 1.6-7% in the literature.\(^\text{1,2}\) The disease may occur incidentally with radiological and pathological findings without clinical symptoms. In a large series of 235 cases, 71 (30%) patients were asymptomatic and 96 (41%) patients were detected with symptoms related to the affected organ.\(^\text{1}\) In the present case, it was diagnosed with a presentation with findings related to ureter pressure. The average age of onset of the disease is in the sixth or seventh decade, and it is more common in men. In some studies, it was seen in all age groups, including children.\(^\text{4}\)

Serum IgG4 level, the most sensitive and specific test for IgG4-RD, is increased in most cases. Serum IgG4 levels >135 mg/dl is important in diagnosis. However, in 30% of patients, serum IgG4 level may be normal.\(^\text{1,2}\) In this patient, serum IgG4 level was high and returned to normal after treatment. Allergic symptoms, and increased serum IgE levels in the blood and peripheral eosinophilia may also be seen in IgG4-RD.\(^\text{5}\) Eosinophil and IgE levels were normal in this patient.

Glucocorticoids are the primary treatment option and the response is generally good. The recommended treatment includes prednisolone at a dose of 0.6 mg/kg/day for two to four weeks and a dose reduction afterwards to 2.5-5 mg/day for maintenance. It is recommended that maintenance therapy should be continued for at least three months to prevent relapse.\(^\text{2,6}\) In the present case, steroid treatment was continued for three months after surgery. No recurrence was detected at two years follow-up after treatment discontinuation. Immunosuppressive therapy, such as azathioprine or mycophenolate mofetil, may be considered in steroid-resistant cases. In patients with recurrent or refractory disease, rituximab or bortezomib treatments are also recommended. Anti-TNF\(\alpha\) treatments have been tried in refractory cases.\(^\text{2,7}\) This case is of great interest in terms of progressive presentation under treatment with etanercept, an anti-TNF\(\alpha\) agent, which is also used in the treatment.

The association of IgG4-RD with other rheumatological diseases is reported. In the cohort of Fernández-Codina et al. including 54 patients, four had RA, one had antineutrophil cytoplasmic antibodies (ANCA)-negative systemic vasculitis, and one had AS.\(^\text{8}\) In the French cohort, 25 patients had RA and one patient had AS.\(^\text{9}\) In the cohort of 235 cases, RA, antiphospholipid antibody syndrome, and ANCA-related vasculitis were detected in one case each, respectively.\(^\text{3}\) This study supports that association with AS is rare. In the literature, no case was observed to progress under anti-TNF\(\alpha\) treatment.

The association of IgG4-RD with AS is exceedingly rare in the litera-
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IgG4-RD has not been observed in any AS patients under biological treatment. The authors presented this case because it is of interest that the patient presented with progressive disease under anti-TNFα treatment, which is also used in the treatment of the disease.

ETHICAL APPROVAL:
The study was approved by the Ethics Committee of University of Health Sciences, Umraniye Training and Research Hospital, Istanbul, Turkey.

PATIENT’S CONSENT:
Written informed consent form was obtained from the patient for the publication of the case details.

CONFLICT OF INTEREST:
The authors declared no conflict of interest.

AUTHORS’ CONTRIBUTION:
OP, SOS, EEY: Conception or the design of the manuscript, acquisition, analysis and interpretation of the data.
All authors have participated in drafting the manuscript; whereas, Ozlem Pehlivan and Sibel Ocak Serin revised it critically.

REFERENCES