



Plasmacytosis and Nephrotic Syndrome Revealing Sjögren's Syndrome

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Sjögren's syndrome is an autoimmune disease characterized by the lymphocytic infiltration of exocrine glands resulting in dry eyes and mouth but can also cause multi-organ damage.¹ Renal involvement is rare during primary Sjögren's syndrome, occurring in 3 to 5% of patients, and even rarely resulting in end-stage renal disease (5%).²⁻⁵

We herein report the case of a 43-year-old female patient admitted to our hospital with sudden onset dyspnea, abdominal pain, diffuse edema, weight gain of 7 kg, dyspepsia and polyarthralgia. The patient had been recently diagnosed with systemic erythematosus lupus based on polyarthralgia, normocytic anemia and positivity of antinuclear antibody and anti-Sjögren's syndrome type A (anti-SSA) antibody. The laboratory investigations showed an acute kidney injury with nephrotic syndrome, anti-SSA and anti-Sjögren's syndrome type B (anti-SSB) antibody positivity, low complement component 4 levels and hypergammaglobulinemia. Cryoglobulinemia was excluded. A percutaneous kidney biopsy was performed the day after her admission. The patient was rapidly treated with high-doses of intravenous corticosteroids, but her clinical condition and renal failure worsened with an estimated glomerular filtration rate of 19 mL/min. The kidney biopsy revealed acute tubulointerstitial nephritis with

massive lymphoplasmacytic infiltrate positive for anti-cluster of differentiation 20 antibody and no expression of anti-immunoglobulin G4 antibody (Figure 1a, b). Because of the important lymphocytic infiltrate, a bone marrow biopsy was performed to exclude lymphoma and showed a rich marrow cellularity with polyclonal plasmacytosis (Figure 1c, d). Positron emission tomography (PET)-fluorodeoxyglucose (FDG) scan did not demonstrate any enlarged lymph nodes or hepatosplenomegaly in favor of lymphoma or Castleman disease. Schirmer's test came back positive and a minor salivary gland biopsy depicted lymphocytic sialadenitis with a Chisholm and Mason score of 4 (Figure 1e, f). Electromyography results showed inflammatory polyradiculoneuritis in C6 and C7. A gastric emptying scan was consistent with gastroparesis. Moreover, immunohistochemistry results of the kidney biopsy also came back negative for human herpesvirus 8 (HHV8) antibody. The absence of the typical histological signs, as well as HHV8 negativity and normal PET-FDG ruled out Castleman disease. Our patient presented primary Sjögren's syndrome with complex systemic manifestations, including renal, hematological (lymphoproliferative syndrome), neurological (polyradiculoneuritis and gastroparesis) and articular disease. The patient was treated with

Received: July 27, 2020 **Accepted:** September 16, 2020 **Published online:** December 10, 2020

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Citation:

La C, Soyfoo M. Plasmacytosis and Nephrotic Syndrome Revealing Sjögren's Syndrome. Arch Rheumatol 2021;36(x):i-iii.

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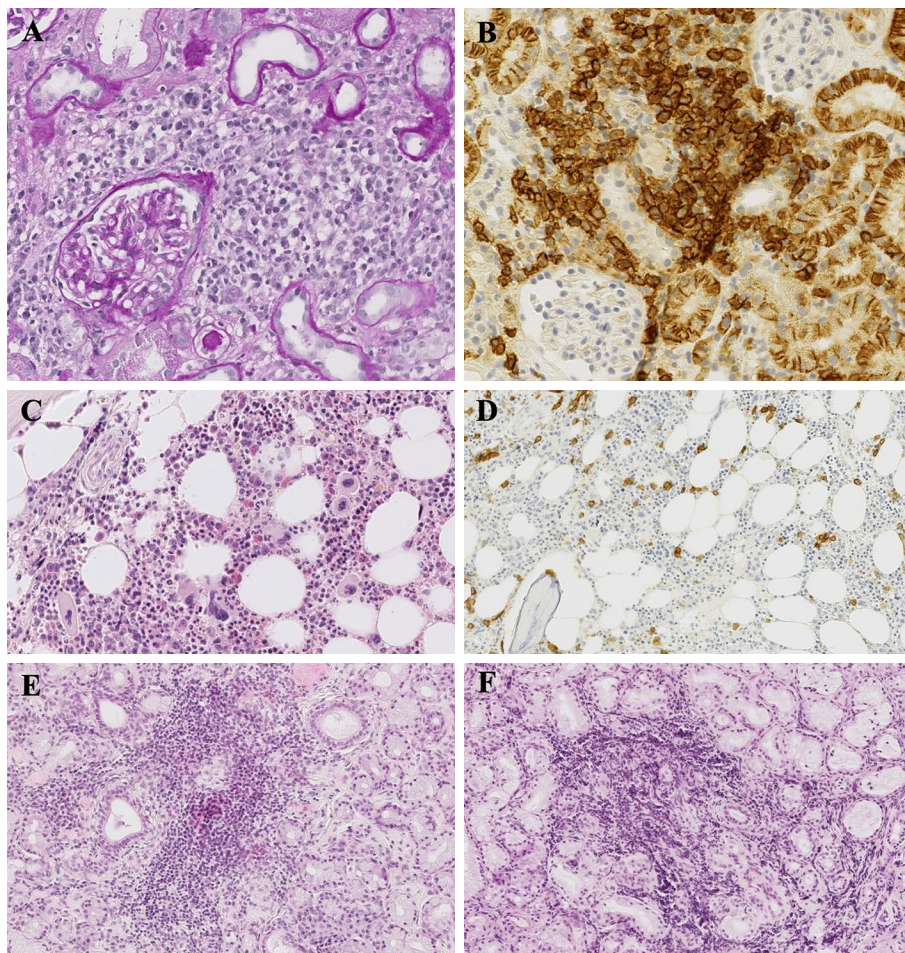


Figure 1. (a) Histopathologic images of kidney biopsy show a massive lymphoplasmacytic infiltrate disrupting renal parenchyma and renal tubules but sparing glomeruli. (b) Immunostaining of CD 138 in kidney biopsy confirms plasmacytic nature of tubulointerstitial infiltrate. (c) Histopathologic pictures of bone marrow biopsy show a rich cellularity, with all three lineages represented. (d) Immunostaining of CD138 in bone marrow biopsy indicates a mild plasmacytosis without monoclonality. (e, f) Salivary gland biopsy showed a lymphocytic sialadenitis with a Chisholm and Mason score of 4.

CD: Cluster of differentiation.

rituximab and oral steroids with good clinical and biological improvement. Cyclophosphamide and steroids were started a year later for a renal flare but suspended because of a major hematological toxicity and the patient had to undergo chronic hemodialysis. Finally, she received a renal transplant from a non-heart-beating-donor five years after initial diagnosis and is currently treated with tacrolimus and mycophenolate. A written informed consent was obtained from the patient.

In conclusion, renal involvement is rare in primary Sjögren's syndrome. Tubulointerstitial nephritis represents the largest majority of cases,

while glomerular lesions are rather associated with cryoglobulinemia.^{6,7} The prognosis is usually favorable, but the patients should be carefully screened if renal involvement is suspected, because of the risk of developing chronic kidney disease. Furthermore, a renal biopsy should be discussed to exclude other causes of renal insufficiency and to guide the therapeutic decision.

Declaration of conflicting interests

The authors declared no conflicts of interest with respect to the authorship and/or publication of this article.

Funding

The authors received no financial support for the research and/or authorship of this article.

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