Acute granulomatous interstitial nephritis and ulcerative colitis: a case report and literature review

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ABSTRACT

Tubulo-interstitial nephritis (TIN) in patients affected by inflammatory bowel disease, both ulcerative colitis and Crohn’s disease, is usually considered as drug-associated to aminosalicylate. We report a rare case of granulomatous active tubulo-interstitial nephritis in a young patient with a recent diagnosis of ulcerative colitis naïve to aminosalicylate treatment. The patient has been successfully treated with steroids administration. Our purpose is to sensitize that TIN should always to be considered in differential diagnosis an extra-intestinal manifestation of bowel disease.

Introduction

Extra-intestinal manifestations are rather common in inflammatory bowel disease (both Crohn’s disease and ulcerative colitis) with a reported prevalence that ranges from 6% to 46% of patients.¹ Renal and urological involvement has a prevalence of 4-23% and is mainly represented by nephrolithiasis, urinary fistulas and malignancies in particular renal cell carcinoma commonly diagnosed at a younger age compared to the general population.²-⁴ Parenchymal renal involvement is a rare but serious extra-intestinal manifestation that can affect both glomerular and tubulo-interstitial compartment.³ The pathogenesis of kidney damage in inflammatory bowel disease (IBD), as in other extra-intestinal manifestation, has not been completely understood; the main hypotheses consider that it is related to intestinal disease activity and dependent on the same immunological mechanism or, alternatively, as a result of systemic autoimmune deregulation.⁵ Moreover drug toxicity can be considered as a relevant cause of kidney damage in IBD patients.⁶ The most common glomerulonephritis are immunoglobulin (Ig)A nephropathy, membranous glomerulonephritis, focal segmental glomerulosclerosis and amyloidosis, whereas the most common types of tubulointerstitial nephritis are drug-related or not drug-related.³

Case Report

A 37-year-old man with a past history of severe obesity [body mass index (BMI) >35] and mild hypertension, successfully treated with β-blocker, a family history positive for ulcerous colitis (father) and kidney disease (cousin with end stage kidney disease for unknown nephropathy), with a recent histological diagnosis of active ulcerative colitis established after 4 months of changing bowel habits, abdominal pain and mild fever.

Patient had been referred to nephrology clinic by
a gastroenterologist after the finding of progressive worsening in renal function (serum creatinine baseline 1 year earlier was 0.8 mg/dL, 6 months after 1.6 mg/dL increased to 2.2 mg/dL at the moment of the evaluation, estimated glomerular filtration rate (eGFR) CKD-EPI 47 mL/min/1.73 m²) associated with micro-hematuria and sub-nephrotic proteinuria (0.4 gr/day). No 5-acetylsalycilic acid (ASA) was administered in order to avoid confounding factors on renal picture. Azathioprine (50 mg twice daily) was prescribed to control intestinal symptoms.

Physical examination at the admission revealed a blood pressure of 110/60 mmHg, heart rate of 76 bpm, weight of 93 kg (BMI 29 kg/m²). Blood tests shown a serum creatinine of 2.27 mg/dL, eGFR CKD-EPI 45 mL/min/1.73 m², anemia (hemoglobin 12 g/dL), white blood cells count and C-reactive protein were normal, serum electrolytes were in optimal range, parathyroid hormone 51 pg/mL. Urinary abnormalities have been confirmed, urine culture was negative.

Immunological tests such as antinuclear antibodies, anti-neutrophil cytoplasmic antibodies, anti-double stranded DNA antibodies, C3/C4 fragment of complement system, and rheumatoid factor were normal.

Abdomen ultrasound showed kidneys of normal size and echogenicity, without structural abnormalities. A kidney biopsy was performed.

In kidney biopsy, light microscopy demonstrated intense and diffuse interstitial infiltration by inflammatory cells including lymphocytes and monocytes, leukocyte infiltration of the tubular epithelium called tubulitis and a small amount of tubular necrosis. There were also a lot of interstitial granulomata with epithelioid histiocytes and a few giant cells without caseous necrosis. The glomeruli were normal. Immunofluorescence was negative for all serum tested, a nonspecific granular positivity for C3 was present in some tubular basement membranes. Electron microscopic showed the presence of interstitial inflammatory infiltration of lympho-monocytes also present in peritubular capillaries. The diagnosis was acute granulomatous interstitial nephritis (Figure 1).

Figure 1. Light microscopic of the kidney biopsy showing interstitial infiltration by inflammatory cells, a huge number of granulomata with epithelioid histiocytes and a few giant cells without caseous necrosis. A) Hematoxylin and eosin (H&E) 10×; B) H&E 20×; C) Masson Trichrome 4×; D) PAS 40×.
The diagnosis of extra-intestinal manifestation of ulcerative colitis has been placed once other causes of granulomatous interstitial nephritis were excluded: quantiferon test for tuberculosis was negative; angiotensin converting enzyme serum levels, high-resolution computed tomography of the lung and whole body Fluorine-18 fluorodeoxyglucose positron emission tomography were performed in order to exclude sarcoidosis.

Patient has been treated with high-dose intravenous methylprednisolone bolus (250 mg ×3) gradually tapered to oral 32 mg of prednisone daily, associated with azathioprine for the ulcerative colitis treatment (50 mg/daily).

Blood tests performed after 1 month of therapy shown significant improvement in renal function, serum creatinine was 1.69 mg/dL (eGFR CKD-EPI 62 mL/min/1.73 m²) and complete regression of intestinal symptoms; blood pressure is well controlled without anti-hypertensive drugs.

Discussion and Conclusions

Acute granulomatous interstitial nephritis is considered an uncommon pathological finding in renal biopsy (<1% of native renal biopsies) that has been associated with various etiologies, of which drug hypersensitivities and sarcoidosis encompass the majority of cases.

Ulcerative colitis is a chronic inflammatory disease with localization in the colon and rectum. The exact pathogenesis is not fully clarified but involves a dysregulation of the gastrointestinal immune system in patients with a genetic predisposition. Extraintestinal manifestations are common (up to 40% of patients) and they are associated with autoimmune mechanism.

In our case we observed a progressive worsening of renal function in a patient with a recent finding of ulcerative colitis without an ongoing treatment with mesalazine or other nephrotoxic drugs. Since our patient had not used any drugs, we could postulate that the main pathogenetic mechanism of renal injury was the disease activity of ulcerative colitis.

Parenchymal renal disease in IBD has been described but granulomatous interstitial nephritis is considered extremely rare.

In current literature many case reports describe renal involvement in patients affected by ulcerative colitis but systematic epidemiological studies are lacking. Ambroz et al. published in 2013 the most comprehensive case series on histopathological findings in IBD patients: they retrospectively review all native kidney biopsy examined and of 33,713 patients, 83 where affected by inflammatory bowel disease; the most frequent diagnosis was IgA nephropathy (24%), followed by interstitial nephritis (19%), nephrosclero-

sis (12%), acute tubular injury (8%) and proliferative glomerulonephritis (7%) equally distributed between Crohn’s disease and ulcerative colitis. The interstitial nephritis cases where classified as acute, granulomatous and chronic; all granulomatous cases where related to exposure to 5-ASA.

In literature most cases of tubulo-interstitial nephritis (TIN) are linked to 5-aminosalicylate exposure, nevertheless is not always possible to discern if kidney damage originated from extra-renal manifestations of bowel disease or drug-induced toxicity. Lately some case reports described patients with TIN not related to drugs exposure or as primary manifestation of bowel disease. Moreover, recent studies showed that tubular damage is tensely related to disease activity rather than drugs and that TIN biopsy-proven in patients with IBD and absent 5-ASA exposure, progressed independently of the intestinal disease activity.

Recent promising studies have reported as the expression of some protein biomarkers, such as neutrophil gelatinase-associated lipocalin, a molecule with multiple activities, including anti-inflammatory and immunomodulatory effects, can provide information able to early detect IBD activity such as possible renal damage.

In conclusion, granulomatous interstitial nephritis is a rare extraintestinal manifestation of ulcerative colitis, which is potentially due to systemic immune dysregulation and T-cell activation that it could be treated successfully with immunosuppression. We suggest that kidney function should be monitored and a renal biopsy should be considered for those patients with a persistent increase in serum creatinine.

References