Early Enterococcus-associated acute postinfectious glomerulonephritis after kidney transplant

Background

Postinfection as an etiology for glomerulonephritis (GN) is rarely described in post-transplant recipients and may be due to impaired immune response. It is also possible that such cases are not biopsied or not reported. There are rare case reports in the literature [1–3]. We report here a rare first case of Enterococcus-related postinfectious GN in a transplant recipient seen in our center.

Case report

A 64-year-old Caucasian man with past medical history of diabetes and hypertension on hemodialysis for 12 years underwent deceased donor kidney transplant. He underwent induction with Thymoglobulin®, early steroid withdrawal, and was maintained on tacrolimus and mycophenolic acid. Early post-transplant course was uncomplicated and serum creatinine improved to 1.6 mg/dL (141 µmol/L) by Day 8. He was discharged with an indwelling urinary catheter due to urinary retention which was removed 2 days after discharge and had to be reinserted on Day 12 due to retention. On Day 23, he had a fever and was found to have vancomycin-resistant Enterococcus (VRE) urinary tract infection (UTI) which was treated with meropenem for 10 days. On Day 58, he was found to have an elevated creatinine of 2.4 mg/dL (212 µmol/L), and recurrent VRE UTI. Despite treatment of the UTI and clinical improvement, the graft function continued to deteriorate and creatinine peaked at 4.1 mg/dL (362 µmol/L). He had 0.5 g of proteinuria. Urinalysis showed 53 white blood cells (WBCs) and >182 red blood cells (RBCs), with many dysmorphic RBCs, but no evidence of tubular casts. Donor-specific antibodies were negative.

Despite antibiotics, intravenous fluids and reduction of immunosuppression, there was no improvement in renal function, and WBCs and RBCs in urine. A transplant kidney biopsy was performed, which showed hypercellular glomeruli with mesangial and capillary loop infiltration of mononuclear cells as well as neutrophils and eosinophils, along with borderline tubulitis (Banff borderline), acute tubular damage and mild isometric vacuolization of the cytoplasm. Crescents were not noted, but there was a minor duplication of the glomerular basement membranes (Figure 1). C4d and BK virus stains were negative. Electron microscopy showed subendothelial and mesangial electron-dense deposits (Figure 2). Immunofluorescence showed antibodies to IgG (2–3+), IgM (2+), C3 (4+), C1q (2–3+), kappa (2–3+) and lambda (trace to 1+) granular mesangial and capillary loop staining. Antibodies to fibrinogen, IgA and albumin were negative. Complement C3 and C4 were low at 26 mg/dL (reference range 65–180 mg/dL) and 10 mg/dL (reference range 13–52 mg/dL), respectively, with negative ANCA and ANA. The tacrolimus levels were not elevated at any time as to be concerned about toxicity. Cardiac echocardiogram was negative for any vegetation and blood cultures were persistently negative. Findings were most consistent with postinfectious GN in the setting of UTI. He was given methylprednisone 500 mg daily in three doses and did require one session of hemodialysis for volume overload. Oral prednisone was resumed after he had completed his antibiotic therapy, and then discharged on a steroid taper over 10 days. He continued to have repeated episodes of UTI requiring antibiotics with creatinine stabilizing at 2.7 mg/dL (238 µmol/L). Due to persistent urinary retention seen on urodynamics, he underwent ileal loop diversion to the transplanted kidney 2.5 months post-transplant. Following the procedure, his creatinine improved to 1.5 mg/dL (132 µmol/L) and is currently, at 19 months post-transplant, 1.2 (106 µmol/L) with 0.32 g of proteinuria.

Discussion

Postinfectious GN is the most common cause of acute nephritis in children in developing countries. It is more frequent and severe in nontransplant immunocompromised patients such as elderly, HIV, malignancy, alcoholic liver cirrhosis, diabetes or obstructive lung disease. However, despite the high rate of infections in post-transplant patients, it is rarely reported post-transplant. Infections are the most common reason for admissions early post kidney transplant. In one case series, only 3 of 827 patients developed postinfectious GN even though 65% of the patients developed at least one significant bacterial infection during follow-up [1]. Long-term outcome was poor as two of the three patients subsequently lost their graft. The reported 11 cases of transplant biopsy-proven postinfectious secondary GN have been due Staphylococcus aureus sepsis, Staphylococcus urosepsis, post-pharyngitis streptococcal GN, cytomegalovirus infections and Salmonella enteritidis, with variable outcomes.

The pattern of the postinfectious GN seems to be changing in the general population and is no longer associated with only streptococcus or staphylococcal infections. The major pathogenetic mechanism is in situ immune complex formation due to deposition of nephritogenic antigens within the glomerulus. An alternative hypothesis is glomerular trapping of circulating immune complexes. Our case shows the course of VRE-associated postinfectious GN in a transplant patient, which has not been previously described. There is no established therapy for treatment of acute postinfectious GN in these cases aside from aggressive treatment of infection with the appropriate antibiotics to remove the antigenic stimulation. The role of steroids in the treatment of postinfectious GN in these cases is unclear. Five of the reported 11 cases were treated with pulse steroids. Based on limited cases, some authors suggest a short course of pulse steroids while others disagree arguing that it would predispose to further infections. We felt a short steroid
course was beneficial for the anti-inflammatory effect aside from aggressive antibiotic therapy and reversal of any potential inciting events. The long-term prognosis of postinfectious GN is poor even in nontransplanted patients where up to 27% have chronic kidney disease and 10% end up on dialysis at 7.5 years of follow-up. To date, there are only three cases of Enterococcus-related GN in native kidneys, of which two were crescentic GN and one membranous \[4, 5\]. To date, there are no cases of Enterococcus-related GN in transplant recipients.

In conclusion, we report the first case of Enterococcus-related postinfectious GN in a transplant recipient on immunosuppression. These cases are generally associated with extremely poor long-term outcome of graft function. However, with early detection, antibiotics and surgical intervention, sustained long-term renal function is possible.

Conflict of interest statement. The results of this case have not been published previously elsewhere. The authors of this manuscript have no related conflicts of interest to disclose.

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