

## Relapse of nephrotic syndrome after a bee sting

Sir,

Relapses in patients with minimal change disease (MCD) have been rarely reported following exposure of inhaled allergens, foods, insect stings, and vaccination.<sup>[1]</sup> Herein, we report a case with relapse after a bee sting while in complete remission.

A 28-year-old male presented with sudden onset generalized edema. One week before the admission he had a bee sting on his forearm. He had a known medical history of MCD, but was in complete remission for 6 years. Seven days after the bee sting the patient had marked edema on the face, and legs. On admission the patient was afebrile, with eyelid and 3+ pretibial edema and a normal blood pressure of 110/60 mmHg. Laboratory examinations revealed proteinuria (2708 mg/day), normal renal function (creatinine 0.8 mg/dl), total serum protein 5.1 g/dl, serum albumin 2.8 g/dl, total cholesterol 315 mg/dl, triglycerides 68 mg/dl, LDL cholesterol 228 mg/dl, and white blood cell 19,600/mm<sup>3</sup>. We accepted these findings and symptoms as a relapse of disease, and methylprednisolone treatment was introduced at the dose of 1 mg/kg/day. One week later his proteinuria resolved to 356 mg/day and clinically improvement was observed. The dosage of corticosteroid was tapered to 4 mg/day over the next 4 weeks, and there was no relapse during 1-year follow-up.

Bee stings usually cause minor local allergic reactions. But, systemic complications such as glomerulonephritis (GN), interstitial nephritis, acute renal failure, myocarditis, centrilobular necrosis of liver, Guillain-Barre syndrome, and vasculitis can be seen.<sup>[2-4]</sup> GN is rare and there are little known data about histological findings, long-term follow-up, incidence, and response to therapy of GN after an insect sting. Cuoghi *et al.*, in their series, include 180 children with nephrotic syndrome (NS); found that three children had recurrent NS triggered by the

insect sting and the remission was achieved with steroid therapy in all.<sup>[3]</sup> Although spontaneous remission may occur in some cases, the most reported cases required corticosteroid therapy for remission.<sup>[2-4]</sup> Similarly, oral steroid treatment-induced prompt remission in our case.

Recent data suggest that atopic disorders are common in patients with MCD despite of little evidence that they have a direct pathogenic role in this disorder. Many patients with MCD have increased serum immunoglobulin (Ig) E and interleukin (IL)-13 levels. IL-13 has the ability to cause switch from IgM to IgE in B cells and induce CD80 expression by podocytes.<sup>[1]</sup> It may be responsible for developing proteinuria and increased IgE levels. Reiser *et al.*, showed that induction of CD80 by podocytes results in proteinuria in rat with glomerular epithelial cell foot-process fusion.<sup>[5]</sup> Moreover, urinary CD80 levels increased in patients with MCD during relapse and return to normal after remission.<sup>[1]</sup> Consequently, recent studies suggest that IL-13 may mediate proteinuria in patients with MCD because of its ability to directly induce CD80 expression on the podocyte.

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