CASE REPORT

Intestinal goblet cell autoantibody associated enteropathy

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This report describes a case of refractory enteropathy with circulating intestinal goblet cell autoantibodies (IGA). A 19 year old man with hyperthyroidism had suffered from protracted diarrhoea for nearly 10 years. Histological examination showed evidence of collagenous enterocolitis. The diarrhoea did not improve despite fasting under total parenteral nutrition. An immunofluorescence assay demonstrated IGA without anti-enterocyte autoantibodies, the hallmark of autoimmune enteropathy, although other criteria were fulfilled. None of 109 controls, including 55 cases of inflammatory bowel disease and one of lymphocytic colitis, had IGA. This case is considered to be a variant of autoimmune enteropathy, and might be a distinct entity.

Collagenous enterocolitis (CE) is a rare condition characterised by collagenous colitis with involvement of the small intestine. Autoimmune enteropathy (AE) is also a rare disorder defined by the following criteria: protracted diarrhoea, no response to total parenteral nutrition (TPN), predisposition to autoimmune disease, and no immunodeficiency. We report the first case of CE with circulating autoantibodies to mucin in intestinal goblet cells but not autoantibodies to the cytoplasm of enterocytes, which are the hallmark of AE.

CASE REPORT

A 19 year old Japanese man was admitted to our hospital in July 2001 with a longterm history of continuous watery diarrhoea, which had averaged five to six times daily since December 1991 at the age of 10 years. His past medical history included atopic dermatitis and uveitis at the age of 10 and 17 years, respectively. He suffered from hyperthyroidism two years before admission, and has been treated with thiamazole. His mother has systemic lupus erythematosus and Sjögren’s syndrome, and his brother has hypophysial dwarfism and idiopathic leucopenia.

He was 146 cm tall and weighed 32 kg. His bowel movements had increased with distention. Laboratory data on admission showed severe hypokalaemia at a concentration of 1.5 mmol/litre. There were mild liver and pancreatic dysfunction (alanine aminotransferase, 64 U/litre; aspartate aminotransferase, 64 U/litre; serum amylase, 198 U/litre; and trypsin, 675 ng/ml). He had no viral hepatitis B or C infection, but anti-adult T cell leukaemia virus antibody was positive. Quantitative immunoglobulins were normal (IgG, 9340 mg/litre; IgA, 1170 mg/litre; and IgM, 370 mg/litre). All stool cultures were negative. An α1 antitrypsin test showed no evidence of protein losing enteropathy. A 5-h D-xylose (5 g) absorption test was consistent with severe malabsorption, with a five hour urine collection of 0.3 g (normal, > 1.5 g). Because the diarrhoea continued at an average of 2200 g/day over a six week period, despite fasting under TPN, a diagnosis of coeliac disease could not be made.

Abdominal ultrasonography and computed tomography showed no abnormalities. Barium studies of the small intestine and colorectum were normal. Upper and lower endoscopical examinations revealed no gross abnormalities, except for reactive lymphoid hyperplasia in the terminal ileum. However, biopsies of the duodenum and entire colorectum demonstrated subepithelial collagen deposits in the sigmoid colon with a striking depletion of goblet cells (bar, 100 µm).

Figure 1  (A) Subepithelial collagen bands around tubules are seen in the duodenal mucosa with goblet cell depletion and increased lymphoplasmacytic infiltrate (bar, 50 µm; haematoxylin and eosin stain). (B) The Masson-trichrome stain showed subepithelial collagen deposits in the sigmoid colon with a striking depletion of goblet cells (bar, 100 µm).

Abbreviations: AE, autoimmune enteropathy; CE, collagenous enterocolitis; IE, intestinal goblet cell autoantibody associated enteropathy; IGA, intestinal goblet cell autoantibodies; TPN, total parenteral nutrition.
autoantibodies in patients with Crohn’s disease (n = 32), the pancreas at a titre of 1/40. We also tested for the above colonic goblet cells at a titre of 1/80, and in the acinar cells of the diagnosis of CE was made.

The only previously documented case of AE demonstrating IGA was that of Moore and colleagues, who described the case of a 9 year old boy. Two other similar cases were reported, but there was no information regarding IGA or anticolonic goblet cell autoantibodies. The case described by Moore et al has several similarities with our case, apart from the presence of IGA: the boys were both older than patients with AE, and there was protracted diarrhoea, abnormal liver function tests, goblet cell depletion, and no appreciable improvement after administering corticosteroids. Our patient exhibited CE, which was absent in Moore’s patient. CE has only been reported in 17 patients to our knowledge, and all were women, with an average age of 60 (range, 40–80) years. These findings conflict with the findings in our patient. Therefore, the present case should be diagnosed as a variant of AE rather than CE, namely IGA associated enteropathy (IE). IE may account for some patients with refractory enteropathy, such as unclassified sprue, which is poorly responsive to gluten restriction.

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