Ulceration of the small intestine in children with coeliac disease

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Abstract

Background—Ulceration of the small intestine in children has not been previously described.

Patients—Two children, aged 12 and 18 months, presented with a history of failure to thrive and intractable diarrhoea.

Results and Conclusions—Upper gastrointestinal endoscopy showed multiple ulcers involving the first and second parts of the duodenum. Histology of biopsy specimens taken from these sites confirmed ulceration and showed other features consistent with a diagnosis of coeliac disease. They both showed pronounced clinical improvement and satisfactory linear growth on a gluten free diet. A year later the diagnosis of coeliac disease was confirmed on a biopsy controlled gluten challenge, and repeat endoscopy showed complete resolution of the intestinal ulceration.

Case histories

Case I was a white female child who presented at 18 months of age with a three week history of vomiting, diarrhoea, weight loss, and abdominal distension. She had no previous history of illness. The neonatal period was normal. She had been breast fed until nine months, when solid foods were first introduced. She was the only child of healthy unrelated parents. Her uncle had coeliac disease and her aunt had died of cystic fibrosis. At presentation, she was unwell with pronounced abdominal distension and buttock wasting.

Case II was a one year old male Asian child who presented with a four month history of diarrhoea, recurrent vomiting, and weight loss. He had had no relevant illnesses in the past. The neonatal period was normal. He had been breast fed for the first two weeks of life, and then started on a cows’ milk formula. Commercially available baby foods were introduced at the age of three months. There were no relevant family illnesses. At presentation he was thin and miserable, with abdominal distension and buttock wasting.

Investigation and follow up

The antigliadin, antireticulin, and antiendomysium antibodies were positive in both patients. The serum albumin concentrations were 28 and 37 g/l, IgA concentrations 1·2 and 1·0 g/l, and haemoglobin concentrations 13 and 12 g/dl in cases I and II respectively. Ferritin, red cell folate, clinical chemistry, and IgG and IgM concentrations were all normal and stool microbiology showed no pathogens. Small bowel biopsy specimens were obtained endoscopically, macroscopically there was inflammation of the entire duodenal mucosa with multiple ulcers, mainly involving the first and second parts of the duodenum. Multiple biopsies from the distal part of the duodenum showed mucosal erosions, villous atrophy, crypt hyperplasia, and an increase in the number of intraepithelial lymphocytes; the histological features were suggestive of coeliac disease (Figure).

Both patients were started on a gluten free diet. Their symptoms resolved and they showed significant catch up growth in weight and length. A year later, while on a gluten free diet, upper gastrointestinal endoscopy showed normal duodenal mucosa and biopsy specimens from the distal part of the duodenum were normal in both patients. On gluten challenge, case I developed diarrhoea, anorexia, and weight loss, and the gluten containing diet was discontinued after four weeks when further biopsy specimens were taken; they...
Endoscopic small bowel biopsy specimen showing superficial erosion and loss of villus architecture and accompanying crypt hyperplasia.

showed villous atrophy. Case II was able to tolerate a gluten containing diet and after three months the postchallenge biopsy specimens from the distal part of the duodenum confirmed the diagnosis of coeliac disease. No intestinal ulceration was seen in either patient. Both have remained well with a normal growth rate over a three year follow up period on a gluten free diet.

Discussion
The relation of ulcerative jejunoileitis to enteropathy associated T cell lymphoma in adult patients is well established. This association is so strong that it has been suggested that ‘benign’ ulcerative jejunoileitis usually represents a manifestation of lymphoma with secondary inflammation obscuring the nature of the malignancy. In our patients, ulceration of intestinal flat mucosa occurred in the absence of lymphoma. Ulceration of the small intestine in children with coeliac disease has not been previously reported and this manifestation may now be considered as one of the complications or variants of the presentation of coeliac disease. The relevance of the features seen in these cases is difficult to establish as this is primarily a macroscopic appearance which would not have been identified so readily in the past using the technique of peroral capsule biopsy.

The clinical course of these coeliac children with ulceration of the distal duodenum was different from that described in adult patients. In the adults there is some heterogeneity, but there was a notably poor response to gluten exclusion. It has been suggested that a prolonged alteration in enterocyte turnover could result in local areas of regenerative failure; however, in some adults the ulcers persisted despite the recovery of normal villous architecture. The aetiology of ulceration in the small bowel in coeliac disease remains uncertain, and its importance in relation to other aspects of this disorder remains to be established.