Small intestinal nodular lymphoid hyperplasia in patients with giardiasis and normal serum immunoglobulins

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Summary

Nodular lymphoid hyperplasia of the upper small intestine was demonstrated in 25 patients with giardiasis. All had normal serum immunoglobulin levels and seven patients initially presented with clinical findings suggestive of an abdominal lymphoma. In only two, however, was the diagnosis of primary jejunal lymphoma confirmed. It is possible that an aetiological relationship exists between recurrent parasitic infestation and nodular lymphoid hyperplasia of the upper small intestine.

The association between nodular lymphoid hyperplasia (NLH) of the upper small intestine and parasitic infestation, particularly with giardiasis, was initially described in patients with humoral immunodeficiency disorders. Subsequently, lymphoid nodules have been described in the terminal ileum and colon of immunologically normal children, but nodular lymphoid hyperplasia of the upper small intestine has only rarely been observed in the absence of humoral immunodeficiency.

In this report we describe a series of 25 patients with giardiasis who had normal serum immunoglobulins and nodular lymphoid hyperplasia of the upper small intestine. It was suspected, on clinical grounds, that seven of these patients had an intestinal lymphoma: in only two, however, was this diagnosis histologically confirmed.

Methods

Patients

Twenty-five patients (six females and 19 males, age range 12 to 42 years, mean 26.5 years) were studied over a three-year period. Patients were divided into two groups according to their clinical presentation.

Group 1 (cases 1–18)

The duration of symptoms in patients in group 1 ranged from three months to 10 years. Patients presented with a history of recurrent diarrhoea or alternating diarrhoea and constipation, anorexia, weight loss, and generalised weakness. Physical examination revealed no significant abnormalities apart from those of mild nutritional deficiencies such as pallor and glossitis. Most of the patients gave a history of recurrent giardiasis with symptom-free intervals after antigiardial therapy.

Group 2 (cases 19–25)

All patients in this group were suspected of having an intestinal lymphoma. The duration of symptoms ranged from six months to 13 years. They gave a history of symptoms similar to those in group 1, but, in addition, complained of marked weight loss, fever, and abdominal pain. On examination they were emaciated, and had pallor and glossitis. Two patients had clubbing and pedal oedema. All patients had palpable abdominal masses suggestive of lymphadenopathy: two had hepatomegaly and one had splenomegaly. Six patients were subjected to exploratory laparotomy. At operation five were found to have no gross abnormality apart from enlarged mesenteric lymph nodes. Small intestinal biopsies were taken. Patient no. 24 in this group presented with a 10-year history of recurrent giardiasis, and a barium meal study done four years before admission was suggestive of nodular lymphoid hyperplasia. An irregular hard mass was palpable in the upper abdomen; at surgery this proved to be a lymphoblastic lymphoma arising from the upper jejunum. The rest of the intestine was studded with multiple nodules, some of which showed malignant change.
Small intestinal nodular lymphoid hyperplasia

Patient no. 25 was admitted with a seven year history of recurrent giardiasis. Barium meal examination showed a markedly disordered small intestinal pattern with numerous rounded filling defects suggestive of nodular lymphoid hyperplasia. Peroral jejunal biopsy revealed a diffuse histiocytic lymphoma. This patient refused surgery and was subsequently lost to follow up.

INVESTIGATIONS

In addition to routine haematological and biochemical investigations the stools were examined for parasites on three separate days. In 22 patients a sample of the jejunal aspirate was examined for parasites and a second sample was cultured quantitatively for aerobic and anaerobic bacteria. Small intestinal absorption was assessed by urinary D-xylose excretion after a 5 g oral dose, the Schilling test, and 72 hours faecal fat excretion on a 75 g fat diet.

Serum (in all patients) and jejunal fluid (in 15 patients) immunoglobulins were measured by single radial immunodiffusion, using WHO reference sera as standards and antisera obtained from Behringwerke. Upper gastrointestinal barium meal studies were performed in all patients. Peroral jejunal biopsy was obtained using a Crosby capsule. In patients in group 2, sera were tested for the presence of free alpha heavy chains by an immunoselection plate technique, using antihuman kappa and antihuman lambda antisera (Behringwerke) in the gel and an antihuman alpha chain specific antiseraum (Behringwerke) in the trough. Operative specimens of lymph node and jejunal biopsies were subjected to histopathological examination. In selected patients, paraffin-embedded sections of jejunal biopsies were stained by an unlabelled antibody enzyme immunoperoxidase technique (using a peroxidase-antiperoxidase soluble immune complex prepared in this laboratory and antisera obtained from Behringwerke) for IgA, IgG, and IgM containing cells.

Nineteen patients were treated with a course of metronidazole (1200 mg/day) for seven to 10 days and stool examination and absorption studies (which were abnormal before treatment) were repeated. All 19 patients who had repeat studies had a minimum of two jejunal biopsies repeated two to three weeks after a course of metronidazole. The patients in whom partial atrophy of the villi and malabsorption persisted after antigiardial treatment were presumed to have an underlying primary tropical malabsorption syndrome and were treated with a long-term course of tetracycline and folic acid and jejunal biopsies repeated three weeks, and, in some, six weeks after starting treatment. Barium meal studies were not repeated after treatment as we did not feel that it would be justified to repeat this investigation within such a short time. Moreover, not all patients had radiographic changes of nodular lymphoid hyperplasia.

Results

Results of relevant investigations are summarised in the Table.

All patients in both groups had *Giardia lamblia* in the stool, and/or jejunal aspirate. Six of 22 patients had a significant bacterial overgrowth with counts >10⁹/ml in the upper jejunum. Twenty patients had malabsorption of one or more test substances. Serum immunoglobulin levels in all patients were within control limits for this population. Jejunal fluid immunoglobulin levels in 15 patients did not differ significantly from controls. Free alpha heavy chains were not detectable in any of the sera tested. Barium series revealed a disordered small intestinal pattern with coarsening and widening of the jejunal loops in 21 patients. In 14 patients, small regular filling defects could be seen in the jejunum (Fig. 1). Jejunal biopsies were independently assessed by two reviewers. Nodular lymphoid hyperplasia was defined as the presence of one or more large aggregates of lymphoid cells in the lamina propria or submucosa. Serial sections through these aggregates in most biopsies revealed a follicular structure with a hyperplastic germinal centre and a surrounding rim of small lymphocytes (Fig. 2). In some sections these follicles were situated deep in the lamina propria or submucosa while in others they protruded into the lumen causing effacement of overlying villi. An increase in the cellularity of the lamina propria and an increase in intraepithelial lymphocytes was present in all biopsies. A consistent finding was the presence of normal numbers of plasma cells in all biopsies. Staining by the unlabelled enzyme antibody technique revealed an abundance of Ig bearing cells in the lamina propria, most of which contained either IgM or IgA in almost equal proportions; there were very few IgG bearing cells. The majority of cells in the lymphoid follicles were found to be non-Ig bearing cells. A few IgA (Fig. 3) and IgM bearing cells, however, were seen in the follicles. Histological examination of enlarged mesenteric lymph nodes in cases 19–25 revealed reactive follicular hyperplasia.

There was no definite correlation between clinical symptoms, absorption parameters, and histological changes in patients in both groups.

RESPONSE TO TREATMENT

Repeat stool examination was negative in all
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Normal values: Vit B₁₂ absorption, >7%; D-xylene excretion, >1.1 g; Faecal fat excretion, <5 g/day; IgG (serum), 700-1500 g/l; IgA (serum), 90-325 g/l; IgM (serum), 40-150 g/l. + Present, - Absent. ND: not done. PVA: partial villous atrophy. NVA: no villous atrophy. NLH: nodular lymphoid hyperplasia.

* Three weeks after treatment. † Six weeks to three months after treatment. ‡ Three to six months after treatment.
patients who were treated with metronidazole. Twelve of the 19 patients improved symptomatically after treatment. In seven patients, bowel symptoms improved but anorexia and weight loss persisted. These seven patients were found to have persistent partial atrophy of the villi and malabsorption, and were presumed to have an underlying primary tropical malabsorption syndrome: they were subsequently treated with tetracycline and folic acid. In 17 of the 19 patients changes of nodular lymphoid hyperplasia could not be detected in two or more jejunal biopsies repeated after treatment.

**Discussion**

Certain features distinguish this group of patients from those with the classical syndrome of nodular lymphoid hyperplasia, which has been described in patients with panhypogammaglobulinaemia or selective IgA deficiency. In this syndrome, jejunal villus architecture is generally normal, bacterial overgrowth in the jejunum is common, and plasma cells are absent or deficient in the lamina propria. Immunofluorescence studies of the jejunal mucosa reveal an almost total absence of Ig bearing cells in the lamina propria. Nodular lymphoid hyperplasia in these patients is believed to represent either an accumulation of plasma cell precursors because of a maturational defect in the B lymphocyte series or a florid local immune response to unknown antigens. Giardiasis has generally been assumed to be secondary to the immunodeficiency state. Although eradication of the parasite results in clinical improvement, a change in the size or number of the nodules has not been observed. None of the patients in our study had a severe humoral immunodeficiency disorder. The majority had partial atrophy of the villi and malabsorption. Plasma cells were present in normal numbers and immunoperoxidase staining of the jejunal mucosa...
plasia may be a patchy lesion and multiple biopsies are often necessary to confirm reversal of lymphoid hyperplasia after treatment. We did not consider it ethical to perform multiple biopsies with a Crosby capsule nor would it have been acceptable to the patients. They did, however, have at least two biopsies performed after treatment with metronidazole and the reversal of nodular lymphoid hyperplasia after anti-giardial therapy appears to suggest a possible causal relationship between giardiasis and nodular lymphoid hyperplasia. It is difficult to define a control population in this area of India where the incidence of parasitic infestation, malnutrition, and primary tropical malabsorption is extremely high. We have, however, analysed data on 464 patients seen during the period of this study in whom information on jejunal histology, absorption status, and stool examination for parasites is available. Of these 464 patients, 67, 77, and 75 had giardiasis, ascariasis, and amoebiasis respectively. Twenty-five of the 67 patients (37.3%) with giardiasis had nodular lymphoid hyperplasia, whereas this was found in only three of 77 (3.8%) with ascariasis and two of 75 (2.6%) with amoebiasis. Of the remaining 245 patients in whom there were no parasites, only two (0.8%) had nodular lymphoid hyperplasia. Thus, of a total number of 32 patients with nodular lymphoid hyperplasia seen during this period, 25 (78.1%) had giardiasis. These data further corroborate our speculation regarding the relationship of giardiasis to nodular lymphoid hyperplasia. It is, however, possible that other organisms, as yet unidentified, may be responsible for the lymphoid hyperplasia.

Wright and Tomkins have demonstrated that patients with giardiasis and malabsorption have increased numbers of intraepithelial lymphocytes in the small intestine which decrease after anti-giardial therapy. They have drawn an analogy to a similar phenomenon in coeliac disease and suggest that a hypersensitivity response to a giardial antigen (similar to that produced by gluten) may be responsible for the mucosal damage and malabsorption. It is possible that the lymphoid hyperplasia in our patients represents a similar hypersensitivity response to chronic antigenic stimulation by *Giardia lamblia*. It is of interest that lymphoid follicles have also been frequently observed in biopsies from children with giardiasis. Owen *et al* in studies of *Giardia muris* infested mice, have also observed large numbers of small lymphocytes in the mucus of the intestinal lumen, some of which were adherent to the *Giardia* trophozoites. They suggested that this represents an immune mediated migration of lymphocytes which may help eliminate the parasite.

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Fig. 3  *Section from jejunal biopsy showing IgA bearing cells in a lymphoid follicle and adjoining lamina propria. Immunoperoxidase, ×250 (original magnification).*

revealed an abundance of IgA and IgM bearing cells in the lamina propria. The incidence of bacterial overgrowth in the jejunum was similar to that of patients with giardiasis without nodular lymphoid hyperplasia, but was higher than that of patients with other causes of diarrhoea (see below). Architectural changes in the villi improved in only 12 of 19 patients but nodular lymphoid hyperplasia apparently disappeared in all but two. Persistence of anorexia, weight loss, and partial atrophy of the villi even after metronidazole therapy in seven of the 19 patients raises the possibility of an underlying primary tropical enteropathy in association with giardiasis. This diagnosis was supported by their subsequent improvement on a regime of tetracycline and folic acid.

It is well-known that nodular lymphoid hyperplasia...
Small intestinal nodular lymphoid hyperplasia

Although all our patients had normal serum immunoglobulins we have not excluded the possibility of a functional antibody deficiency, particularly as some of them were severely malnourished. Although this is unlikely, it is a factor that needs to be further evaluated.

Our finding of a relative increase in IgM bearing cells in the lamina propria confirms the observations of Ridley and Ridley, patients with systemic humoral immunodeficiency have also been shown by Webster et al. to produce IgM locally in the gut.

Two patients in group 2 were found to have a primary upper gastrointestinal lymphoma. Both patients have a long history of recurrent giardiasis and one patient had radiological features suggestive of nodular lymphoid hyperplasia four years before the diagnosis of lymphoma. A similar case of nodular lymphoid hyperplasia and lymphoma in a normo-gammaglobulinaemic patient has been described by Kahn et al. Shaw and Hennigar have described the occurrence of nodular lymphoid hyperplasia in association with lymphoma outside the gastrointestinal tract. The association between nodular lymphoid hyperplasia and jejunal lymphoma has also been described in a patient with adult onset hypogammaglobulinaemia.

The clinical picture in patients in group 2 is similar in many respects to that of patients with alpha heavy chain disease. It is postulated that an unknown chronic antigenic stimulus underlies this disease.

Giardiasis is commonly found in these patients, although no consistent association with any pathogen has been noted. It is interesting that in some cases of alpha heavy chain disease the initial biopsy of the jejunum and/or mesenteric lymph nodes shows a non-specific hyperplastic lymphoid infiltration. Subsequently a plasma cell infiltrate becomes evident. Similar lymphoproliferation is associated with dilantin therapy and coeliac disease where there is a spectrum ranging from a benign non-specific reaction to overt malignancy. Whether a genetic predisposition or an unknown stimulus could trigger the malignant potential of these disorders remains speculative.

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References


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