Immunohistochemical findings in jejunal specimens from patients with IgA deficiency

T Klemola, E Savilahit, A Arato, T Örmälä, J Partanen, C Eland, S Koskimies

Abstract

Jejunal biopsy specimens from 25 patients with IgA deficiency (IgAd) were studied immunohistochemically to find markers of inflammation. Five of the 25 patients had coeliac disease (CD): they were on a gluten free diet and had normal jejunal morphology. Only two of 15 specimens from control subjects had CD25+ cells in the surface epithelium, while this was seen in 19 out of 20 specimens from IgAd patients (p<0.0001). A significant increase of CD25+ cells was also noted in the lamina propria of IgAd patients. The median percentage of crypt cells in mitosis (Ki67+ cells) was higher in the specimens from IgAd patients (26%) than in those from controls (13%, p<0.001). The densities of γδ T cell receptor positive cells in the surface epithelium and lamina propria did not differ in the specimens from IgAd patients and those of controls nor was the expression of HLA class II antigens augmented in the surface epithelium. These findings were similar for the IgAd patients whether or not the patient had DQB 0201 allele, a genetic marker which is strongly associated with CD. The inadequacy of the local immunoglobulins in patients with IgAd may lead to increased T cell activation, which is accompanied by the appearance of intraepithelial CD25+ cells and with an increase in the mitotic rate in the crypts.

Keywords: IgA deficiency, coeliac disease, jejunal mucosa, HLA class II antigens, γδ T cell receptor

Mucosal IgM antibodies of patients with IgA deficiency (IgAd) may be less efficient than IgA antibodies, resulting in a longer excretion of vaccine viruses than seen in normal individuals.1 IgAd patients with a history of frequent respiratory tract infections were seen to respond to oral cholera vaccination with significantly higher intestinal cholera toxin-specific IgG and IgM antibody responses than healthy IgAd individuals and controls.2 The increased intestinal IgG response may maintain inflammatory and tissue damaging processes in the intestine of IgAd patients by complement activation. Gut permeability to foreign proteins is increased in IgAd patients. They often have high titres of milk antibodies,3 4 and excessive uptake of dietary antigens into the circulation has been reported.5 5 We studied jejunal biopsy specimens from patients with IgAd to find if the inadequacy of mucosal antibodies in these patients causes changes in the expression of T cell activation markers in jejunal mucosa.

Asymptomatic coeliac disease (CD) is often found among patients with IgA deficiency (IgAd) – the frequency of CD in these patients is about 10%.6 A strong association has been found between IgAd and the HLA DQB 0201 gene.7 HLA associated susceptibility to develop CD is primarily conferred by HLA DQ (α1*0501, β1*0201) heterodimer coded by the DQA 0501 and DQB 0201 genes.8 9 An increase in the density of lymphocytes bearing γδ T cell receptor (γδTCR+), αβ T cell receptor (αβTCR+), and enhanced expression of HLA class II antigens DR and DP in the jejunal epithelium are typical of CD.10−18 The intestinal specimens of some family members of CD patients may also display similar changes.17 19 Among them, the changes are associated with the occurrence of genetic markers of CD.17 19

Jejunal biopsy specimens from patients with IgAd were studied to test if the inflammation typical of CD was present in the jejunal mucosa of patients with IgAd but normal jejunal mucosa. We wished to see if the intestine of those patients with IgAd who had inherited the CD marker HLA DQB 0201 differed from that IgAd patients without that gene.

Methods

PATIENTS

Jejunal biopsy was performed on all patients with IgAd because of the increased risk of CD. In this study jejunal biopsy specimens from 25 patients with IgAd (10 girls and 15 boys, mean age 9-8 years) were studied. Twenty of these patients had a normal jejunal structure while eating normal, gluten containing diet but five patients had CD. This had been diagnosed before the study and these patients were on a gluten free diet. Their jejunal morphology was normal at the time the jejunal biopsy was taken for this study.

Control group specimens were made up as follows: 14 specimens from patients with normal serum IgA and normal intestine (six girls and nine boys, mean age 8-6 years); and 11 from patients with CD and a normal serum IgA value (six girls and five boys, mean age 11-5 years). All CD patients were on a gluten free diet and the morphology of their jejunal specimens was normal.

HLA TYPING

DQB1 alleles were determined using oligonucleotide hybridisation techniques according to the 11th HLA Workshop.20

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Frequencies of DQB 0201 and DQB 0501 positive IgAd patients and controls

<table>
<thead>
<tr>
<th>Allele</th>
<th>IgAd patients (n=15)</th>
<th>Controls (n=95)</th>
<th>p (Fisher)</th>
</tr>
</thead>
<tbody>
<tr>
<td>DQB 0201</td>
<td>8/15 (53%)</td>
<td>23/95 (24%)</td>
<td>0.17</td>
</tr>
<tr>
<td>DQB 0501</td>
<td>8/15 (53%)</td>
<td>30.95 (32%)</td>
<td>0.12</td>
</tr>
<tr>
<td>DQB 0201, or 0501</td>
<td>15/15 (100%)</td>
<td>51/95 (53%)</td>
<td>0.03</td>
</tr>
</tbody>
</table>

IMMUNOHISTOCHEMISTRY

Processing the intestinal specimens

The biopsy specimens were immediately embedded in OCT compound and stored at −70°C. Serial cryostat sections were cut at 5 μm, fixed in acetone for 10 minutes, then in chloroform for 30 minutes, and washed three times in Tris buffer, pH 7.4.

Immunohistochemical staining

The buffer was removed and the sections were covered with the diluted monoclonal antibody (mAb) in Tris buffer for one hour. Endogenous peroxidase was blocked by incubation in 0.5% peroxidase for 30 minutes. A Vectastain Elite AB kit (PK-6102, Vectro Laboratories, Burlingham, CA, USA) was used to stain mAb. Interleukin 2 receptor (CD25) was stained by triple incubation of the monoclonal antibody and alkaline phosphatase-antialkaline phosphatase antiserum. The phosphatase was visualised with a fast red reagent. Positively stained cells were counted with a light microscope at 900X magnification.

MAb, intestinal specimens

Monoclonal antibody TCR6 (T cell Sciences, Cambridge, MA, USA) that recognises a constant region of the δ chain of T cell receptor and all γδ T cells, and antibody βF1 (T cell Sciences) which react with nearly all αβ T cell receptor (αβTCR) molecules were used at a dilution of 1:100. Also used were mAbs against the following lymphocyte surface antigens: CD3 (Anti-Leu4, Becton Dickinson, Mountain View, CA, USA), CD4 (anti-T4, Coulter Immunology Hialeah FL), and CD8 (OKT8, Ortho Diagnostic System, Raritan, NJ, USA), CD25 (anti-CD25, Becton-Dickinson). Monoclonal mouse anti-human antibody against proliferating cells (DAKO-PC, clone Ki67) was used to detect dividing cells (Ki67⁺). Monoclonal antibodies to constant fragments of HLA class II antigens, HLA-DR, and HLA-DP were products of Becton-Dickinson and anti-HLA-DQ mAb was a product of Serotec Ltd, Oxford, England.

STATISTICAL ANALYSIS

The densities of cell staining in the jejunal specimens from patients and controls were compared using the two tailed Mann-Whitney U test. Epithelial staining with HLA class II antibodies was graded from 1 to 6 according to the intensity (0 to 3) and cellular distribution (0 to 3, apical membranes – all cell membranes) and the area of staining (parts of the villi and crypts). Fisher’s exact test of probability was used to determine the differences in HLA-DR and HLA-DP expression of epithelial cells. The statistical significance of differences in the DQB frequencies was evaluated by Fisher’s exact test. To avoid comparisons between populations of unequal sizes, the actual calculations were carried out between the patient group (n=19) and a control subgroup of 30 individuals which was obtained after randomisation of the original control group consisting of 95 unselected Finnish individuals.

ETHICS

The study protocol was reviewed and approved by the Ethical Committee of the Children’s Hospital, University of Helsinki.

Results

DQB ALLELES IN PATIENTS WITH IGAD

DQB alleles could be determined in 19 patients with IgAd. Four of these had CD and they all had the DQB 0201 allele. The remaining 15 had IgAd only. Fifty three per cent (8/15) of them had the DQB 0201 allele. The frequency of DQB 0201 in the unselected Finnish control population was lower (24%; 23/95), but the difference was not statistically significant. In addition, DQB 0501 was found in 53% (8/15) of the patients with IgAd only – but again comparison with the control population (32%; 30/95) showed that this was not significant.

All patients with IgAd (19/19) were positive for DQB 0201 or DQB 0501 compared with only 54% (51/95) in the controls (p=0.003 after randomisation). As both the DQB alleles 0201 and 0501 carry ‘non-Asp’ at position 57, all patients were ‘non-Asp’ carriers, while 76% (72/95) of controls were ‘non-Asp’ carriers. The difference in the frequency of ‘non-Asp’ carriers, however, is statistically non-significant (p=0.18 after randomisation).

STAINING WITH HLA CLASS II ANTIBODIES

Epithelial staining with anti-HLA-DR and

![Figure 1: Epithelial staining with anti-HLA-DR antibodies in controls, patients with IgAd having DQB 0201, without it and those in whom the gene analysis was not done (ND). All patients with IgA deficiency (IgAd) and celiac disease (CD) had DQB 0201. Epithelial staining with class II antibodies was graded from 1 to 6 according to the intensity and cellular distribution and area of staining.](http://gut.bmj.com/Downloadedfrom)
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INTERLEUKIN 2 RECEPTOR (CD25) BEARING CELLS IN THE LAMINA PROPRIA AND IN THE SURFACE EPITHELIUM

Patients with IgAd had more CD25+ cells in the lamina propria (p<0.002) (Fig 3) and in the surface epithelium of jejunal mucosa (p<0.0001) (Fig 3) than the controls. The increase in CD25+ cells was not associated with the occurrence of the DQB 0201 gene. IgAd patients with or without this gene had similar densities of these cells in the epithelium and lamina propria (data not shown). Nor was the increase associated with CD – the increase was present only in IgAd coeliac and not in coeliac with normal IgA level (Fig 3).

KI67+ CELLS IN THE CRYPT EPITHELIUM

The percentage of dividing, Ki67+ cells was increased in the jejunal crypt epithelium in specimens from IgAd patients (p<0.001 vs controls) (Fig 4). This increase was not associated with the occurrence of CD associated DQB genes in IgAd patients with normal jejunum (data not shown). Specimens from CD patients on a gluten free diet had the same percentage of Ki67+ cells in the crypts as the controls, while IgAd patients with CD had similar percentages as IgAd patients with a normal jejunum.

Discussion

Secretion of IgA is the most important defence mechanism of the mucosal surfaces against microbial infections and the penetration of foreign proteins. In patients with selective IgA deficiency, this secretion is severely diminished or lacking and is replaced by an excessive

![Figure 2: Densities of γδ T cell receptor (TCR) positive (A) and αβ TCR positive (B) cells in the surface epithelium of jejunal biopsy specimens from patients with IgA deficiency (IgAd) and controls expressed as cells/mm. Medians are represented by solid lines (B). For the densities of γδ TCR positive cells the patients with IgAd are divided on the basis of the occurrence of DQB 0201 gene as in Figure 1.](image)

![Figure 3: Interleukin 2 receptor bearing (CD25+) cells in the surface epithelium (cells/mm²) (A) and in the lamina propria (cells/mm²) (B) of jejunal mucosa in IgAd patients and controls. Symbols as in Figure 2.](image)

anti-HLA-DP was similar in patients with IgAd without CD and controls (Fig 1). Staining in specimens from IgAd patients with the DQB 0201 gene did not differ from that in IgAd patients without the DQB 0201 gene. The expression of HLA-DP was augmented in the jejunal epithelium of the patients with CD and IgAd (p<0.05) and in the CD patients with normal IgA (p<0.005) (Fig 1). Positive staining of epithelial cells with DQ antibody was present in five specimens from IgAd patients and in two control specimens but the difference was not statistically significant. The densities of HLA-DR, HLA-DP, and HLA-DQ-positive cells in the lamina propria did not differ in IgAd patients and controls (data not shown).

T CELLS, γδ TCR+ AND αβ TCR+ CELLS

Densities of CD3+, CD4+, and CD8+ cells in the surface epithelium and lamina propria did not differ in the patient groups and controls (data not shown). Similar densities of αβ TCR+ and γδ TCR+ cells were present in the surface epithelium (Fig 2) and lamina propria (data not shown) of jejunal biopsy specimens from IgAd patients without CD and controls. Those with or without the DQB 0201 gene had similar densities of γδ TCR+ cells in the epithelium (Fig 2A). IgAd patients with CD on a gluten free diet had significantly higher densities (p<0.0001) of γδ TCR+ cells in the jejunal surface epithelium (Fig 2B).
 secretion of IgM and the presence of an increased density of IgM secreting cells in the jejenum. Intestinal diseases are common among IgAd patients and some of them also have minimal morphological changes in the jejunal mucosa. In addition, the density of intraepithelial lymphocytes is increased in some patients with IgA. In our study, it was found that the density of activated CD25+ cells in increased in both the lamina propria and in the surface epithelium in the jejunal mucosa of IgAd patients. Raised serum concentrations of soluble CD8, neopterin, and B2 microglobulin have been reported in patients with common variable immunodeficiency and in IgAd patients. It has been suggested that these findings reflect enhanced T cell mediated immunity in these patients. Our results also support the theory that enhanced T cell activation is present in the intestine of IgAd patients. The T cell activation may be due to increased antigenic stimulation caused by ineffective antigen exclusion or to a chronic infection in the intestinal mucosa of IgAd patients.

In fetal cultured intestine, T cell activation generates total or partial villous atrophy and a profound crypt cell hypertrophy. The changes are associated with an increase of dividing Ki67+ cells in the cryptal epithelium. These changes are mediated by lamina propria T cells. The activated T cells secrete interleukin 2 and interferon γ, but whether these or other cytokines generate the proliferation of crypt cells is still unknown. Activated T cells have been found in the lamina propria of jejunal specimens from CD patients. These cells were present in a higher density in both the lamina propria and epithelium of patients with IgAd compared with controls, indicating T cell activation in the intestine of patients with IgAd. The increase in percentage of dividing Ki67+ cells in the crypt epithelium of IgAd patients may be due to this T cell activation in the lamina propria.

Nilssen et al found an increased density of γδTCR+ intraepithelial lymphocytes in the jejunal mucosa in a patient with IgAd and overt partial villous atrophy, and it was suggested that this may reflect a compensatory mechanism of surface protection in patients with IgAd and a low frequency of infection. In our study a constant increase in γδTCR+ intraepithelial lymphocytes was found only in the jejum of patients with IgAd and CD during gluten free diet when their jejunal morphology was normal. In contrast, all IgAd patients with normal jejunal eating a normal, gluten containing diet had a density of γδTCR+ cells similar to immunologically normal controls.

Increased density of γδTCR+ cells and CD25+ in the jejunal epithelium, and augmented epithelial expression of HLA-DR and HLA-DP antigens were found in a patient with normal jejunal mucosa, who later developed typical CD. These findings also occurred together with CD marker DQ genes in relatives of CD patients. These patients are thought to show latent CD and are at an appreciably increased risk of developing CD.
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