Timing of infiltration of T lymphocytes induced by gluten into the small intestine in coeliac disease

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SUMMARY Jejunal biopsy was performed on control subjects and patients with treated and untreated coeliac disease. Monoclonal antibodies to T lymphocyte surface markers were used to quantify T cell phenotypic subsets in the jejunal mucosa. The patients with untreated coeliac disease had significantly more of both suppressor/cytotoxic and helper/inducer T cells in the surface epithelium than either the control subjects or the treated patients. Serial jejunal biopsy specimens were taken from five treated coeliac patients for six hours after a gluten challenge. In four of these five infiltration of the surface epithelium by both T cell phenotypes, together with deterioration in the villus architecture, had occurred both within two hours of having started the challenge. This suggests that T lymphocytes may have a role in the pathogenesis of coeliac disease.

Coeliac disease is characterised by a small intestinal enteropathy, resulting in malabsorption. Histological sections of small intestinal mucosa taken from patients with untreated coeliac disease show a variety of abnormalities, including villus atrophy, elongation of the crypts, and abnormally flattened enterocytes with loss of their brush border.† Lymphocytes and plasma cells also infiltrate the mucosa. A gluten free diet resolves these changes.

The mechanism of mucosal damage caused by gluten in susceptible subjects remains unclear. Earlier suggestions that the toxicity of gluten was related to a primary specific small intestinal enzyme deficiency7 8 or to a lectin-like effect of gluten9 have not been substantiated.5 6

A weight of evidence supports the hypothesis that the mucosal damage results from an immunological reaction to gluten. In vitro studies on small intestinal biopsy specimens from patients with coeliac disease showed specific activation to gliadin of both the humoral7 8 and cell mediated9 10 arms of the immune systems within the small intestinal mucosa.

Most small intestinal intraepithelial lymphocytes both in normal subjects and patients with coeliac disease are T cells.11 12 In untreated coeliac disease the number of intraepithelial T cells is increased relative to the number of enterocytes.13 14 This study aimed to quantitate intraepithelial and lamina propria T lymphocyte subsets in the small intestinal mucosa of coeliac and control subjects and to compare the timing of changes in T cell density with that of the morphological damage to the small intestinal mucosa of patients with treated coeliac disease who had been subjected to a gluten challenge.

Patients and methods

Small intestinal biopsy specimens were taken from three subject groups: (i) control subjects (n = 6), who underwent biopsy as part of their diagnostic management and were subsequently found not to have coeliac disease; (ii) patients with treated coeliac disease (n = 5), in whom the diagnosis had previously been established by standard criteria15 and whose biopsy specimens showed either normal histology or a degree of partial villus atrophy; and (iii) patients with untreated coeliac disease (n = 5) whose biopsy specimens showed the typical histological changes of the disease and who subsequently showed improvement both clinically and in their small intestinal morphology after a period of withdrawal from dietary gluten.

Five patients, three males and two females, aged 17–57 years, with coeliac disease, which had been treated by gluten free diet for between eight months and 30 years, were subjected to serial jejunal biopsy. Biopsy specimens were taken prior to and at half hourly intervals up to six hours after a gluten chal-

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length; this consisted of 10 g of gluten suspended in 150 ml of water and taken by mouth. One of these five patients served as a negative control by taking water only on a separate occasion. Another of these five had a further biopsy specimen taken one month after the initial challenge, during which time he consumed a normal diet containing gluten.

All biopsy specimens were obtained using a Quinton hydraulic multiple biopsy instrument, with the capsule positioned under fluoroscopic control in the proximal jejunum and left there for six hours in the patients who had been challenged with gluten. Informed consent was obtained from the patients before the study, which was approved by the ethical committee of St Thomas’s Hospital.

Each biopsy specimen was carefully orientated on silver foil, coated in optimal cutting temperature (OCT) compound (Miles Laboratories), and snap frozen in liquid nitrogen and isopentane. Biopsy specimens were stored under liquid nitrogen until used.

**Morphometry**

Cryostat sections (5 µm) were mounted on glass slides and stained with haematoxylin and cosin. The sections were examined using an eyepiece micrometer and a Leitz Wetzler microscope; five separate measurements each of villus surface-cell height and enterocyte surface-cell height were made and the mean values calculated.

**Lymphocyte Counting**

Cryostat sections (5 µm) mounted on glass slides were stained using an immunoperoxidase method as follows: the sections were air dried, fixed in acetone, and washed first in tap water and then in Tris buffered saline (TBS, pH 7-6). Murine monoclonal antibodies to different human T lymphocyte subsets were diluted 1 in 10 in TBS and applied to the sections for one hour in a moist chamber. The monoclonal antibodies (Becton Dickenson) used with their phenotypic specifications were anti-leu 4 to the pan T cell phenotype, anti-leu 3a to the helper/inducer T cell phenotype, and anti-leu 2a to the suppressor/cytotoxic T cell phenotype.

Sections were then washed in TBS and in a second layer antibody, peroxidase conjugated rabbit anti-mouse immunoglobulin (Dako Ltd), diluted 1 in 50, and applied for 30 minutes. After further washing in TBS the sections were developed using 3, 3-diaminobenzidine as substrate. Cell nuclei were counterstained with Mayer’s haematoxylin. The sections were finally washed, dehydrated in alcohol, cleared in xylene and mounted.

Separate sections from each biopsy specimen were stained with each of the three different monoclonal antibodies so that each T cell phenotypic subset could be counted. In the surface epithelium T lymphocytes were counted per 100 enterocytes in five separate fields using an objective at a magnification of 40 and the mean value taken. Lamina propria T cells were counted per unit area in three different villi, using an eyepiece graticule and the same magnification objective, and again the mean value was calculated. The area of the eyepiece graticule was 1.82 x 10^-2 mm². Several biopsy specimens were found to be either too small or too distorted to permit accurate lymphocyte counting. For this reason it was not always possible to follow the changes in counts in each patient at regular half hourly intervals after the gluten challenge. For comparison, changes in T subsets after gluten challenge were presented as percentage changes from the values in the biopsy specimen taken before challenge (time 0). This was necessary because individual patients had different starting values for the numbers of cells in each T cell subset.

**Results**

**Baseline T Subsets**

Table 1 shows the means of the individual counts of intraepithelial T lymphocytes that stained with the three different monoclonal antibodies and also the Leu 2a (suppressor/cytotoxic): Leu 3a (helper/inducer) ratios for the three subject groups in the first part of the study. In each group Leu 2a T cells greatly outnumbered Leu 3a positive cells. The patients with untreated coeliac disease had significantly more Leu 2a positive (p < 0.0001), Leu 3a positive (p < 0.02), and Leu 4 positive T cells (p < 0.0001) than the control subjects. Counts for the patients with treated coeliac disease were intermediate between the controls and the untreated patients, although they did not differ significantly from the controls.

<table>
<thead>
<tr>
<th></th>
<th>Leu 2a Suppressor/cytotoxic</th>
<th>Leu 3a Helper/inducer</th>
<th>Leu 4 Pan T</th>
<th>Leu 2a:Leu 3a ratio</th>
</tr>
</thead>
<tbody>
<tr>
<td>Controls (n = 6)</td>
<td>22.0 (6.3)</td>
<td>3.2 (1.4)</td>
<td>27.2 (6.8)</td>
<td>6.9:1</td>
</tr>
<tr>
<td>Treated coeliac patients (n = 5)</td>
<td>31.6 (19.9)</td>
<td>3.3 (1.3)</td>
<td>40.1 (19.6)</td>
<td>9.6:1</td>
</tr>
<tr>
<td>Untreated coeliac patients (n = 5)</td>
<td>64.2 (9.9)</td>
<td>5.2 (0.9)</td>
<td>78.0 (10.4)</td>
<td>12.3:1</td>
</tr>
</tbody>
</table>
Coeliac lymphocytic infiltration

Table 2  Small intestinal lamina propria T lymphocyte counts (mean (SD) No/unit area) and suppressor/cytotoxic:helper/inducer cell ratios

<table>
<thead>
<tr>
<th></th>
<th>Leu 2a Suppressor/cytotoxic</th>
<th>Leu 3a Helper/inducer</th>
<th>Leu 4 Pan T</th>
<th>Leu 2a:Leu 3a ratio</th>
</tr>
</thead>
<tbody>
<tr>
<td>Controls (n = 6)</td>
<td>39·5 (17·0)</td>
<td>59·9 (18·1)</td>
<td>101 (31·3)</td>
<td>1:1·5</td>
</tr>
<tr>
<td>Treated coeliac patients (n = 5)</td>
<td>38·2 (11·2)</td>
<td>68·2 (27·6)</td>
<td>110 (33·1)</td>
<td>1:1·8</td>
</tr>
<tr>
<td>Untreated coeliac patients (n = 5)</td>
<td>37·0 (11·1)</td>
<td>57·2 (25·1)</td>
<td>100·1 (37·4)</td>
<td>1:1·5</td>
</tr>
</tbody>
</table>

Figure  Percentage changes in intraepithelial T lymphocyte counts and mucosal morphometry plotted against time in five patients given gluten challenge and one control patient given only water. •—• Suppressor/cytotoxic T cells; □—□ helper/inducer T cells; ○—○ villus height:crypt depth ratio; ■—■ epithelial surface-cell height.
not differ significantly from the control group. The
differences between the three groups in Leu 2a:Leu 3a
ratios were not significant. In the lamina propria there
were slightly more Leu 3a positive cells than Leu 2a
positive cells (table 2), but there were no significant
differences between the three subject groups for any
of the three T lymphocyte subsets nor for the Leu
2a:Leu 3a ratios.

**EFFECT OF GLUTEN CHALLENGE**

Of the five patients with treated coeliac disease who
were given a gluten challenge, four showed a similar
pattern of changes in intraepithelial lymphocyte
counts and morphometry. The figure shows their
results as percentage changes from the values before
challenge (time 0). There were no significant changes
in T lymphocyte counts in the lamina propria in any
of the subjects. Each of these four patients cases (a–d)
showed an increase in both suppressor/cytotoxic and
helper/inducer intraepithelial T cell counts concomi-
tant with a deterioration in mucosal morphometry,
shown by a reduction in villus height: crypt depth
ratio and a fall in enterocyte height, although the lat-
ter actually increased in one patient (c) after an initial
fall. These changes started within two hours of the
gluten challenge in each subject but showed a variable
degree of recovery by six hours. The fifth patient (e),
in whom coeliac disease was diagnosed only eight
months before the study, showed a very different pat-
tern of changes; there was virtually no change in
either intraepithelial suppressor/cytotoxic lympho-
cyte counts nor morphometric variables during the
six hours after the gluten challenge, although there
was an increase in epithelial helper/inducer lym-
phocytes. On rebiopsy, however, after four weeks on a
normal diet containing gluten, he was found to have
similar epithelial infiltration with both T cell subsets
and changes in mucosal morphometry as had
occurred in the other four patients during the first six
hours.

As expected, in the patient who took water only (f)
(figure), there was very little change in either epithelial
lymphocyte counts or mucosal morphometry.

**Discussion**

This study clearly shows considerable differences
between the surface epithelium and lamina propria of
the small intestinal mucosa in terms of the T lym-
phocyte phenotype subpopulations, and emphasises
the importance of studying them separately. In agree-
ment with the findings of previous studies \(^{12,16–18}\) we
found that most intraepithelial T cells, both in control
and coeliac subjects, are of the suppressor/cytotoxic
phenotype, whereas in the lamina propria there was a
slight predominance of helper/inducer T cells. The
ratio of suppressor/cytotoxic:helper/inducer cells did
not differ significantly between the controls and the
treated and untreated coeliac patients in either layer.
This ratio was therefore not measured in the challenge
studies. In addition, there were no significant differ-
ces in the numbers of lamina propria T lym-
phocytes of any of the three phenotypes that were
counted between the control subjects and the patients
with treated and untreated coeliac disease. A gluten
challenge given to patients with treated coeliac disease
produced no significant change in T cell counts in the
lamina propria during the first six hours. Lancaster-
Smith *et al* \(^{19}\) also found no significant difference in
total lamina propria lymphocyte numbers between
control and coeliac patients, although a single gluten
challenge given to treated patients did produce a
small increase in lymphocyte numbers in the lamina
propria 24 to 48 hours later, but which was no longer
detectable at seven days. This increase may have been
accounted for by B rather than T lymphocytes, and
the authors also found a larger increase in lamina pro-
pria plasma cells both at 24–48 hours and seven days.

Finding the correct method of quantitating small
intestinal intraepithelial lymphocytes has proved con-
troversial. Ferguson and Murray \(^{13}\) counted the num-er of intraepithelial lymphocytes per 100 enterocytes
to establish the normal range of values using data
from 160 non-coeliac biopsy specimens. They found
significantly increased intraepithelial lymphocyte
counts in patients with untreated coeliac disease com-
pared with controls. Intraepithelial lymphocyte
counts were also raised above the value for normal
controls in coeliac patients on a gluten free diet,
although these were significantly lower than in the
untreated group. Marsh, \(^{20}\) however, argued that
intraepithelial lymphocyte counts should be related to
a fixed area of underlying muscularis mucosae (areal
density) rather than to the number of enterocytes
when comparing normal and coeliac mucosa, as
coeliac and normal epithelium differ considerably in
respect of their relative surface areas, structure, and
cell sizes. In this study, however, we chose the former
method as, like Corazza *et al* \(^{14}\) and Dobbins, \(^{21}\) we
feel that it is the local density of lymphocytes within
the epithelium that is most relevant when considering
the possible aetiological role of these cells in coeliac
disease. Like us, Jenkins *et al* \(^{18}\) also showed a much
higher relative density of T lymphocytes in untreated
coeliac small intestinal surface epithelium than in nor-
mal surface epithelium. These authors also showed a
change in marker patterns of the intraepithelial lym-
phocytes "which might suggest an altered functional
state rather than just an increase in density because of
the non-specific crowding of cells into a reduced sur-
face epithelial compartment."

Although we studied only a small number of
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patients, in the small intestine of most patients with treated coeliac disease oral gluten challenge caused both damage to the villus architecture, evidenced by a reduction in villus height: crypt depth ratio and enterocyte height, and an increase in density of T lymphocytes in the surface epithelium; both of these events started within two hours of the challenge. The concurrent morphological and lymphocyte density changes suggest that they are interrelated. The delayed response to gluten challenge shown by one of our five patients suggests that there may be a subpopulation of coeliac patients who relapse more slowly after reintroduction of dietary gluten. This is in keeping with the large variation in the time taken to relapse in a study performed by Kumar et al.22

The speed with which changes in morphometry and intraepithelial lymphocyte counts occurred in our study in four of the five patients is somewhat surprising, and is faster than has previously been reported. Anand et al23 in a study of two coeliac subjects, detected the first histological changes three to four hours after gluten challenge and suggested that the timing was in keeping with an Arthus type reaction producing the mucosal damage. Bramble et al24 found changes both in brush border enzymes and villus population counts one to two hours after gluten challenge, but no significant changes in the villus height or crypt depth within 24 hours of the challenge.

Our findings of a rapid increase in density of T lymphocytes in the surface epithelium of the small intestine of treated coeliac patients following gluten challenge, coinciding with the observed damage to the villus architecture, suggest that T lymphocytes have a role in the pathogenesis of the disease, although further work is required to elucidate the exact nature of that role.

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References


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