Mucosal antibodies in inflammatory bowel disease are directed against intestinal bacteria

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Abstract
In contrast with normal subjects where IgA is the main immunoglobulin in the intestine, patients with active inflammatory bowel disease (IBD) produce high concentrations of IgG from intestinal lymphocytes, but the antigens at which these antibodies are directed are unknown. To investigate the specificities of these antibodies mucosal immunoglobulins were isolated from washings taken at endoscopy from 21 control patients with irritable bowel syndrome, 10 control patients with intestinal inflammation due to infection or ischaemia, and 51 patients with IBD: 24 Crohn's disease (CD, 15 active, nine quiescent), 27 ulcerative colitis (UC, 20 active, seven inactive). Total mucosal IgG was much higher (p<0.001) in active UC (median 512 μg/ml) and active CD (256 μg/ml) than in irritable bowel syndrome controls (1.43 μg/ml), but not significantly different from controls with non-IBD intestinal inflammation (224 μg/ml). Mucosal IgG bound to proteins of a range of non-pathogenic commensal faecal bacteria in active CD; this was higher than in UC (p<0.01); and both were significantly greater than controls with non-IBD intestinal inflammation (CD p<0.001, UC p<0.01) or IBS (p<0.001 CD and UC). The mucosal IgG binding was shown on western blots and by enzyme linked immunosorbent assay (ELISA) to be principally directed against the bacterial cytoplasmic rather than the membrane proteins. Total mucosal IgA concentrations did not differ between IBD and controls, but the IgA titres against faecal bacteria were lower in UC than controls (p<0.01). These experiments show that there is an exaggerated mucosal immune response particularly in active CD but also in UC directed against cytoplasmic proteins of bacteria within the intestinal lumen; this implies that in relapse of IBD there is a breakdown of tolerance to the normal commensal flora of the gut.

(Gut 1996; 38: 365–375)

Keywords: inflammatory bowel disease, mucosal antibodies, intestinal bacteria.

The aetiologies of ulcerative colitis (UC) and Crohn’s disease (CD) are unclear. There are a number of suggestions of possible environmental or genetic causes including persistent infections, vasculitis, and immunological abnormalities (including autoimmunity). It is thought that CD and UC have different aetiologies on the basis of differences in location, the macroscopic and microscopic appearance of the disorders, and differences in mucosal immune activation (resulting in different cytokine and immunoglobulin isotype profiles). Despite this the two disorders may have very similar presentations and precipitants of clinical relapse.12

We have previously shown that during relapse of CD intestinal permeability is increased3–5 allowing access of luminal constituents to the mucosa. The acute inflammatory response is, however, quantitatively an order of magnitude greater than in other conditions with similarly increased intestinal permeability.9 This is probably caused by increased intestinal immune responsiveness caused by the underlying disease. The importance of the interaction between luminal antigens and the mucosal immune system as the mechanism of relapse in inflammatory bowel disease (IBD) is suggested by the following findings.

(1) Known causes of relapse (intestinal infections, non-steroidal anti-inflammatory drugs, and stress) all increase permeability across the epithelial cell layer or have the potential to do so.6 Moreover, normal intestinal permeability in patients with CD predicts a long remission, whereas increased permeability heralds relapse.47

(2) Active CD can be treated by faecal stream diversion,8 and recurrence of distal disease is largely avoided if an ileostomy is performed at the time of right hemicolectomy.9

(3) In strains of mice that have been genetically engineered to ‘knockout’ the T lymphocyte receptor,10 or either of the cytokines interleukin (IL) 211 or IL10,12 intestinal inflammation develops with some parallels to the clinical presentation and pathological findings in human IBD. Significantly, where these mice were bred in a germ free environment they did not get intestinal inflammation. This shows that in these animal models of IBD the primary genetic abnormalities (which may be quite different from those in human UC or CD) each cause an increased mucosal immune response to the commensal intestinal bacteria, leading to active disease. In some human patients with active IBD antibiotic treatment may be effective.13 Clinical studies of bowel decontamination regimens in humans as a method of reducing disease activity have been inconclusive, however, possibly owing to difficulties in actually achieving decontamination or due to the use of non-absorbed antibiotics, which are directly toxic to the intestinal mucosa.
In this paper we have directly tested the hypothesis that there is an abnormally increased mucosal immune response against non-pathogenic commensal intestinal bacteria in active CD and UC. Previous studies of circulating antibodies in serum samples of IBD patients initially proposed that there were high titres against enterobacteria (cross reactive with colonic mucins), but serum titres were subsequently shown not to be significantly different from control patients. Similarly the proposal of enterobacterial antigen specific activation of peripheral blood lymphocytes was not confirmed. However circulating immunoglobulin specificities may be unrepresentative of locally produced mucosal immunoglobulins. IgA is the main immunoglobulin produced in the intestinal mucosa of normal subjects, but in patients with active UC or CD both immunohistochemistry and cultured mucosal lymphocytes show that IgG production is dramatically increased (IgG1 and IgG3, in UC; IgG1, IgG2, and IgG3, in CD). The antigens against which these mucosal immunoglobulins are directed have received comparatively little attention, except that monoclonal antibodies generated from mesenteric lymph nodes of IBD patients showed a higher frequency of binding to commensal intestinal bacteria in CD than in UC although control data could not be obtained.

We report the isolation of immunoglobulins directly from the intestinal mucosa of patients with IBD and control subjects with and without intestinal inflammation and characterisation of their antigen specificity against non-pathogenic commensal intestinal bacteria.

Methods

Patients

Patients were studied while undergoing colonoscopy or flexible sigmoidoscopy to investigate lower abdominal pain and changed bowel habit, or to define the extent or activity of known IBD. The diagnoses of UC (n=27) and CD (n=24) were made on standard criteria, with serial biopsies providing histopathological confirmation of disease in each case and histological UC disease activity. Crohn's disease was defined as quiescent with a Harvey Bradshaw index of 0-1 and normal serum C reactive protein. There were two groups of control patients. The first was a group of 21 patients suspected of having a functional disorder; no biochemical, haematological or microbiological abnormalities were found, colon and terminal ileal histology was normal, and a final diagnosis of irritable bowel syndrome was made in each case. The second control group was a series of 10 patients with intestinal inflammation; seven patients had infectious gastroenteritis, two patients had diverticulitis, and one had ischaemic colitis. The Table shows patient details including activity, duration, and extent of disease, and current treatment.

Patients were prepared for colonoscopy by two oral doses of sodium picosulphate (10 mg) taken 48 and 24 hours before the procedure and were sedated immediately before starting the examination by intravenous administration of 10-15 mg diazepam and 75 mg pethidine. Patients for flexible sigmoidoscopy were unprepared apart from a phosphate enema 90 minutes beforehand, and they were not sedated. A sample of serum was taken from each patient at the time of colonoscopy. Informed consent was obtained from all patients, and the study was approved by the King's Healthcare Trust Ethics Committee.

Isolation of intestinal mucosal immunoglobulin

The colonic mucosa was washed under direct vision with 100 ml 0-9% (w/v) saline injected through the flush channel of an Olympus CF200HL videocolonoscope or flexible sigmoidoscope, with the patient positioned to ensure that the relevant colonic segment was in a dependent position. Great care was taken to aspirate any fluid encountered in the colon during insertion of the colonoscope, and washings were only taken during withdrawal of the instrument so virtually all the wash fluid was aspirated (median 88 ml, range 72-96 ml). The wash fluid was aspirated after two minutes through the suction channel, and immediately aliquoted into 10 ml 1 mM-phenylmethylsulphonylfluoride and 0·2 ml 500 mM EDTA (pH 8·0) to inhibit proteolysis. Two biopsy specimens were subsequently taken from each

<table>
<thead>
<tr>
<th>Group</th>
<th>Age Mean (SD)</th>
<th>Duration of disease (mean and range)</th>
<th>Treatment (details of patients)</th>
<th>Disease distribution</th>
</tr>
</thead>
<tbody>
<tr>
<td>Control subjects (Irritable bowel syndrome) n=21; 16 colonoscopy, 5 flexible sigmoidoscopy</td>
<td>52 (19) 26-80</td>
<td>5 days-13 days</td>
<td>None apart from 1 patient receiving cefuroxime and metronidazole IV</td>
<td>Salmonella colitis (5) Shigella (1) Campylobacter (1) Ischaemic colitis (1) Diverticulitis (2) Ileocolonic (17) Colonic (7)</td>
</tr>
<tr>
<td>Control subjects with intestinal inflammation all flexible sigmoidoscopy</td>
<td>34 (13) 17-62</td>
<td>None</td>
<td>None apart from 1 patient receiving cefuroxime and metronidazole IV</td>
<td>Salmonella colitis (5) Shigella (1) Campylobacter (1) Ischaemic colitis (1) Diverticulitis (2) Ileocolonic (17) Colonic (7)</td>
</tr>
<tr>
<td>Crohn's disease (n=24, colonoscopy 15 active, 9 quiescent)</td>
<td>36 (15) 21-67</td>
<td>4 years (0-16 years)</td>
<td>None</td>
<td>Remission (7) Left-sided (14) Pancolitis (6)</td>
</tr>
<tr>
<td>Ulcerative colitis (n=27; 21 colonoscopy 6 flexible sigmoidoscopy; 20 active 7 inactive)</td>
<td>37 (15) 17-80</td>
<td>7 years (0-32 years)</td>
<td>None</td>
<td>Remission (7) Left-sided (14) Pancolitis (6)</td>
</tr>
</tbody>
</table>
washing site to assess independently histo-
logical involvement (CD) or disease activity
(UC).

Precipitation of mucosal immunoglobulin from
endoscopic washings
To precipitate immunoglobulins 25 ml of 50% (w/v) polyethylene glycol (PEG) 4000 were
added to 100 ml of mucosal washings and
centrifuged at 6000 g for 10 minutes at 4°C. The
pellet containing immunoglobulin was
resuspended in 2 ml of 15% (w/v) polyethylene
 glycol 4000, which was resuspended by
centrifugation. The precipitate was dried, and
resuspended in 1 ml 10 mM-TRIS Cl, 50 mM NaCl
pH 7.4. This was confirmed routinely to contain undegraded immuno-
globulins by western blot analysis, which were
quantified and used in experiments to deter-
ine antibacterial membrane titres.

Preparation of bacterial proteins
Bacterial isolates of non-pathogenic Escherichia
coli (K12; human faeces; O Rough H48; from
Professor P J F Henderson, University of Cam-
bidge), Klebsiella aerogenes (KCH908; human
faeces; type 11; King's College Hospital),
Bacteroides fragilis (KCH727; human
faeces; King's College Hospital), Enterobacter
casais (KCH806; human faeces; King's
College Hospital), Staphylococcus epidermidis
(KCH1004; skin swab; King's College Hospital),
Haemophilus influenzae (KCH633; blood
culture; King's College Hospital), and
Clostridium perfringens (KCH 1001; human
faeces; type A; King's College Hospital) were
grown in liquid culture using aerobic or anaer-
obic medium from a single colony picked from
selective agar plates. Bacteria were disrupted
by sonication at mid-log phase (Soniprep 150,
four bursts 25 μm peak to peak) in 50 ml of
10 mM-TRIS Cl, 1 mM-EDTA, 50 mM NaCl
pH 7-4; the cytoplasmic fraction was then
obtained as the supernatant after centrifuga-
tion (47 000 g, 20 min, 4°C), the pellet was
then washed as previously described35 and
recovered by recentrifugation (47 000 g, 20
min, 4°C) to obtain the membrane fraction.

Kaback vesicles were prepared from E.coli as
previously described.34

ELISA quantification of mucosal
immunoglobulins and antibacterial
immunoglobulins
To quantify mucosal immunoglobulins and antibacterial immunoglobulins microtitre
plates (96 wells, Maxisorb, Nunc) were coated
with 50 μl of monoclonal antihuman IgG
and IgA (Sigma) at a dilution of 1:10 000 and
1:1000 respectively in 15 mM-Na2CO3,
35 mM-NaHCO3, 0-02% sodium azide, pH
9-6) or 50 μl of a combined preparation of
cytoplasmic and membrane proteins at a con-
centration of 10 μg/ml (5 μg/ml each) in the
same carbonate/bicarbonate coating buffer by
incubation at 4°C overnight. The plates were
washed three times with phosphate buffered
saline containing 0-05% (w/v) TWEEN 2035
before blocking with 10% (w/v) bovine serum
albumin in phosphate buffered saline for two
hours at 25°C and washed a further three
times in phosphate buffered saline-TWEEN.
Samples of mucosal immunoglobulins were
serially diluted in phosphate buffered saline-
TWEEN, and 50 μl of each was loaded in
duplicate to the plates. A set of standards were
made for every plate by serial dilutions of a
immunoglobulin calibrant (Sheffield Protein
Reference Unit) or a standard serum calibrant
of known high titre to each bacterial pro-
tein preparation. The standard serum calibrant
for each bacterial preparation was obtained by
screening serum samples from a set of entirely
healthy volunteers; those chosen for their high
titre were E.coli man aged 26; K.aerogenes man
aged 26; B.fragilis woman aged 28; Ent. faecalis
man aged 35; S.epidermidis man aged 58;
H.influenzae man aged 23; G.perfringens woman
aged 62; each serum was frozen in aliquots and
thawed only once before use. After incubation
for two hours at 25°C the plates were again
washed in phosphate buffered saline-TWEEN,
and 50 μl of 1 μg/ml biotinylated antihuman
IgG (Vector Laboratories, BA 3080) or bio-
tinylated antihuman IgA (Vector Laboratories,
BA 3030) was added to each well incubated
for a further two hours at 25°C. The plates were
again washed three times with phosphate buffered saline-TWEEN, and each
were incubated with avidin-horseradish
peroxidase complex for 30 minutes at 25°C
(Vectastain Elite ABC, Vector Laboratories).
Unbound reagent was washed off with
phosphate buffered saline-TWEEN, and the
enzyme was detected by the addition of 50 μl
4 mg/ml o-phenylenediamine dihydrochloride
in 50 mM phosphate citrate buffer (pH 5-0),
0:03% (w/v) sodium perborate. This reaction
was stopped after five minutes by the addition
of 50 μl 3 M sulphuric acid, and the absorb-
ance in each well was read at 495 nm using a
Titertek Multiscan MCC/340 plate reader.

Each sample and standard was performed in
triplets, and mean results of IgG and IgA
were expressed in μg/ml. Bacterial protein
binding was based on a standard curve drawn
for each plate using a serum calibrant (assigned
as 100 ELISA units) known to bind well
to each protein preparation. After subtraction
of blanks (where no serum or mucosal
immunoglobulin was added) each sample was
read at the mid-linear portion of the curve, and
mean results (of triplets) were expressed in
ELISA units/μg total IgG or IgA.

Binding of bacterial cytoplasmic or mem-
brane proteins was measured as described
above, except that the microtitre plates were
coated with either cytoplasmic or membrane
proteins at a concentration of 10 μg/ml in car-
bonate/bicarbonate buffer overnight. The
swiss sandwich assay for immunoglobulins was also
adapted to measure IgG isotypes by coating
the plates in carbonate/bicarbonate buffer
overnight with specific IgG capture antibody
(all mouse antihuman monoclonal antibodies
were from Sigma (I-9388, I-9513, I-7260
and I-9888) and were used at the following

Extensive preliminary experiments were performed to optimise these ELISA methods. Coating of the solid phase was most satisfactory in freshly prepared carbonate/bicarbonate buffer at 4°C overnight, and the concentration of each protein used to coat the plates was adjusted to obtain a steep reference curve (approximately 400 ng of antihuman IgG, 350 ng of antihuman IgA, and 550 ng of bacterial protein were bound during the coating procedure). A small amount of lipopolysaccharide (approximately 0.08 mEU) also bound to the wells with the bacterial membranes, but not the bacterial cytoplasmic protein preparations. In the case of bacterial proteins, the metabolically labelled proteins were used in control experiments to confirm that dissociation did not occur during the protocol. The incubation times after coating were set at two hours to allow the binding reactions to reach equilibrium (after about 30 minutes), and the concentration of secondary antibody in the detection system was shown not to be limiting for each assay. We also confirmed that there was no significant cross reaction of the different antibodies used in the sandwich ELISAs.

**Protein assays**

Total proteins were assayed using the bicinchoninic acid (BCA) method (Pierce). Albumin measurements were made by the Technicon RA radioimmunoassay method. To estimate the fraction of IgG derived from serum leakage, IgG and albumin were measured in both serum and unprocessed mucosal washing samples and the following equation was used:

\[
\text{IgG (washings)} = \frac{\text{albumin (washings)}}{\text{IgG (serum)}}
\]

**Western blots**

SDS poly-(12.5%) acrylamide gels were run as described previously using a mini-V.8.10 electrophoresis system (BRL). Molecular weight markers (rabbit muscle myosin, 205,000; E. coli β-galactosidase, 116,000; rabbit muscle phosphorylase B, 97,400; bovine serum albumin, 66,000; ovalbumin, 45,000; bovine erythrocyte carbonic anhydrase, 29,000; egg white lysozyme, 14,300; Sigma) were run in a separate lane at the edge of the gel. Protein samples (200 μg) from E. coli membrane or cytoplasmic proteins were prepared as described previously and were run in a single large well on each gel to achieve uniform separation of the proteins (Fig 1). After electrophoresis, the gels were either stained with Comassie Blue R or transferred to Hybond C (Amersham) using the electroblot stack assembly (150 V, 1 hour) with 25 mM-TRIS HCl, 192 mM glycine, 10% (v/v) methanol (pH 8.3) transfer buffer. The blots were dried and a strip at each end was removed (Fig 1) and stained with Amido black to ensure clean separation of the mucosal proteins and uniform transfer. Markers were also visualised in a separate narrow well on the edge of the gel by Amido black staining. Between the end strips the blots of mucosal proteins were sectioned into 2 mm strips and processed further as described below.

Each strip of the mucosal protein blot was washed in phosphate buffered saline containing 0.05%-TWEEN 20, and blocked by incubation in 10% bovine serum albumin at room temperature for one hour. They were then washed again in phosphate buffered saline-TWEEN and incubated with different samples of serum (500 μl) or mucosal immunoglobulin (300 μl) preparations at a concentration of 150 μg/ml or 10 μg/ml IgG respectively in separate test tubes for two hours. The strips were individually washed three times with 5 ml of phosphate buffered saline-TWEEN in each tube and then incubated with biotinylated

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**Figure 1:** Method of assessment of mucosal and serum IgG binding to bacterial cytoplasmic or membrane proteins. Bacterial cytoplasmic or membrane proteins (200 μg) were run in a single large well by 12.5% SDS polyacrylamide gel electrophoresis. Molecular weight markers were run in a separate lane at the side of the gel and stained separately. The separated bacterial proteins were transferred to nitrocellulose, which was cut into strips, with each edge being stained for the bacterial protein bands with Amido black to ensure uniform transfer. Each intermediate strip was then incubated with either serum or mucosal immunoglobulin from an individual patient; after washing, specifically bound antibodies were detected using secondary biotin labelled antihuman IgG. Binding of secondary antibody independently of the mucosal or serum IgG was shown using 'blank' strips where no serum or mucosal immunoglobulin had been added.
antihuman IgG (Vector Laboratories, BA3080) or antihuman IgA (Vector Laboratories, BA3030) at dilutions of 1:1000 and 1:1500 respectively. After washing three more times in 5 ml phosphate buffered saline-TWEEN, bound immunoglobulin was detected on each strip by incubation with avidin-horseradish peroxidase complex for 30 minutes at 25°C (Vectastain Elite ABC, Vector Laboratories); unbound enzyme was washed off with phosphate buffered saline-TWEEN and the enzyme was detected using diaminobenzidine as a substrate.

Statistical methods

Statistical analysis was performed using the Wilcoxon signed rank test for unpaired samples using the SPSS package35 on the Digital Equipment Corporation VAX cluster (VAX 8800) of the computing service of the University of London.

Results

Immunoglobulins isolated from intestinal washings in control subjects and in IBD

Immunoglobulins isolated from colonscopic washings by polyethylene glycol precipitation were quantified by sandwich ELISAs. Proteolytic degradation was routinely excluded by detecting intact α (IgA) or γ (IgG) heavy chains of M, 55 000 and 50 000 respectively in western blots (not shown). In control subjects with irritable bowel syndrome most of the immunoglobulin was IgA (median concentration 240 µg/ml), with much lower amounts of IgG (median 1·43 µg/ml; Fig 2). In patients with active CD and UC total IgG was significantly increased compared with irritable bowel control subjects (medians CD=256 µg/ml, UC=512 µg/ml; p<0·001 for both). IgG concentration were similarly increased, however, in patients with intestinal inflammation due to gastrointestinal infections or ischaemia (median 224 µg/ml), suggesting that the increase in IgG was not specific to IBD. Patients with inactive UC or in quiescent Crohn’s had lower IgG concentrations compared with active disease in each case (medians UC 27 µg/ml p<0·05; CD 2 µg/ml p<0·005). Concentrations of total IgA (Fig 2) or IgM (not shown) did not differ significantly between either control or IBD groups.

The IgG subtypes in washings from patients with IBD were predominantly IgG1, and IgG2, (medians 479 µg/ml and 51 µg/ml respectively) in active UC and IgG1, IgG2, and IgG3, (121 µg/ml, 185 µg/ml, and 36 µg/ml) in CD.

Albumin estimation of serum leakage

To estimate the contribution of serum leakage to proteins isolated from the colonic mucosal washings, albumin was measured in the crude mucosal extracts (before polyethylene glycol precipitation) and compared with serum albumin in 16 control patients with irritable bowel syndrome, 11 patients with active CD, and 13 with active UC. Assuming that the same proportion of serum IgG leaks into the washings as serum albumin, the fraction of IgG in the washings derived from serum was then calculated from the ratios of albumin and IgG in washings and serum. The proportion of IgG derived from serum in CD patients was between 3·2–18% (median 8·6%), UC 0·13–16·2% (median 7·5%), and controls 0·01–8% (median 2%). Thus most of the immunoglobulin in the washings was produced in the intestinal mucosa, and serum leakage contributed only a small fraction of the preparations.

Mucosal antibody binding to intestinal and non-intestinal bacteria proteins in IBD

Mucosal IgG was found to bind to proteins from non-pathogenic commensal strains of E coli, B fragilis, and C perfringens in patients with active CD and UC (Fig 3). In these experiments we corrected for different yields of mucosal immunoglobulins by expressing ELISA signals (binding) in relation to the total concentration of mucosal IgG or IgA. In each case, the titres for active CD patients were significantly higher than those for active ulcerative colitis (p<0·01). The mucosal IgG titres from control patients with irritable bowel syndrome against these commensal bacteria were at the lower limit of detection of the assay system. Moreover IgG titres to the bacterial protein preparations from patients with non-IBD inflammation were also significantly lower than active CD (p<0·001) or active UC (p<0·01; Fig 3). Very similar results were obtained using proteins from non-pathogenic strains of K aerogenes (non-IBD inflamed controls median 0·9 ELISA units/µg IgG, range 0·17–17·8, n=10; active CD median 259·4, range 68·3–398·6, p<0·001, n=13; active UC median 7·5 range 0·39–22·7, p<0·05, n=14).
and *Ent. faecalis* (non-IBD inflamed controls median 0.42 ELISA units/μg IgG, range 0.11–3.9, n=10; active CD median 146.0, range 12.7–661.7, p<0.01, n=12; active UC median 4.12 range 1.19–52.36, p<0.01, n=12). However, binding of intestinal bacterial proteins by mucosal immunoglobulins from patients with quiescent CD or inactive UC was not significantly greater than the non-IBD inflamed control group, except for quiescent CD, which showed significantly increased binding to *B. fragilis* (p<0.05). In six of the patients with left sided UC washings were also taken from the macroscopically and histologically inactive area in the ascending colon; in each case the titres from the inactive area (for example, for *E. coli* median 0.7 ELISA units/μg IgG; range 0.08–6) were lower than those from the active region (median 29.5, range 7–68; p<0.001). There were no significant differences in serum titres to any of these bacterial preparations between controls, UC or CD groups (shown for *E. coli*, Fig 3B).

In contrast with the results with intestinal bacteria, measurements of mucosal immunoglobulin titres in IBD patients and controls using the non-intestinal bacteria *S. epidermidis* (Fig 4), *S. aureus*, and *H. influenzae* (not shown) showed very little measurable mucosal IgG directed against these organisms, whereas there were strong serum IgG titres. This shows that the mucosal immunoglobulin response against bacterial antigens is directed specifically against the colonic flora. Furthermore as the serum IgG titres against non-intestinal bacteria did not appear in the mucosa this also suggests that direct leakage from the serum does not contribute significantly to the IgG mucosal titres.

There were strong mucosal IgA titres in both groups of control patients against proteins from commensal intestinal bacteria (Fig 5). However IgA mucosal titres were significantly lower in patients with active UC compared with controls against *E. coli* (p<0.005), *B. fragilis* (p<0.005), and *C. perfringens* (p<0.01), although some control patients had IgA titres against commensal bacteria at or below the concentrations seen in UC. In no case were there significantly reduced mucosal IgA titres in active CD compared with the irritable bowel syndrome control group. Only very weak IgA titres were obtained to proteins from non-intestinal organisms *S. epidermidis*, *S. aureus*
Mucosal antibody binding to cytoplasmic and membrane proteins of E coli

To investigate further the bacterial proteins bound by mucosal immunoglobulins in patients with IBD, we carried out the following experiments.

1. ELISA assays were performed to assess mucosal immunoglobulin binding to the individual bacterial membrane or cytoplasmic protein fractions.

2. Western blots were run in which cytoplasmic or membrane proteins were loaded en bloc from a single large well, and the nitrocellulose was cut into strips after electrophoretic transfer, thus there was an identical amount of each E coli cytoplasmic or membrane protein band on every strip, which could be used to detect binding of immunoglobulins in individual mucosal or serum patient samples.

Both these techniques gave similar results. The ELISA assays showed that there was significantly greater binding of mucosal IgG in active CD to the cytoplasmic proteins than to the membrane fraction (p<0.001; Fig 6A), however low concentrations of serum immunoglobulin bound to both bacterial cytoplasmic and membrane protein fractions (Fig 6B). Mucosal IgA bound to both
membrane and cytoplasmic proteins in controls, yet membrane binding was significantly decreased (p<0.01) in patients with UC (Fig 6C). Very similar results were obtained for membrane and cytoplasmic proteins prepared from B fragilis and C perfringens as those shown in Fig 6 for E coli.

On western blots mucosal IgG samples from patients with CD clearly bound over 12 distinct cytoplasmic protein bands but very weak binding was seen in mucosal IgG samples of patients with UC and no bands were seen in inflamed controls (even though the same concentration of total mucosal IgG (10 μg/ml) was used in each case; Fig 7A). A single prominent band at M, 59 000 on the E coli cytoplasmic proteins in controls and UC patients was not due to binding of mucosal immunoglobulin, as it was also seen in blank strips to which no mucosal immunoglobulin had been added (Fig 7A), showing that the secondary anti-human IgG used to detect bound immunoglobulins crossreacts with this E coli cytoplasmic protein. In contrast, mucosal IgG from active CD, UC or control patients did not bind to membrane proteins either at a total mucosal IgG concentration of 10 μg/ml (Fig 7B) or at concentrations of up to 150 μg/ml even when the mucosal antibody preparations were precleared with lipopolysaccharide (not shown) despite the fact that satisfactory protein separation was shown by staining the end strips of the blot for protein (Fig 7B). Indeed, serum IgG from controls, CD, and UC patients (incubated with the strips at the higher concentration of 150 μg/ml to visualise the bound immunoglobulin clearly as the signals were very weak at 10 μg/ml) bound both E coli membrane and cytoplasmic proteins from the same set of strips as in Fig 7A and 7B suggesting that the failure of mucosal IgG binding was due to the absence or blocking of specific immunoglobulins (Fig 7C). Similar results were also obtained on western blots of cytoplasmic and membrane proteins prepared from B fragilis or C perfringens and cytoplasmic and inner membrane proteins of Kaback vesicles from E coli (not shown).

Discussion
In this paper we present and validate a new technique that permits the quantification and assessment of antigen specificity of degraded mucosal immunoglobulins. In keeping with immunohistochemical studies we found greatly increased IgG concentrations from patients with IBD, which relates to disease activity. IgG recovery from patients with IBD after total intestinal lavage (with a non-absorbable polyethylene glycol solution) also showed a correlation with disease activity.\(^{37}\) We have confirmed the different subclasses of IgG secreted in CD (IgG1, IgG2, and IgG3) and UC (IgG, and IgG2).\(^{25}\) Most importantly we have shown that the intestinal IgG is specifically directed against cytoplasmic proteins from commensal bacteria in active CD and UC, implying that these may be of major importance in the relapse of the two diseases. There is increased binding of mucosal IgG in active CD to proteins of both Gram positive and Gram negative non-pathogenic intestinal
Mucosal antibodies in inflammatory bowel disease are directed against intestinal bacteria

**Figure 7: Binding of IgG from mucosal and serum immunoglobulins to E coli cytoplasmic or membrane proteins.** A western blot of E coli cytoplasmic or membrane proteins was processed as shown in Figure 1, with each strip representing binding of a different patient mucosal or serum immunoglobulin sample. Binding is shown (A) by mucosal IgG (10 µg/ml) to E coli cytoplasmic proteins; (B) by mucosal IgG (10 µg/ml) to E coli membrane proteins; and (C) by control serum IgG (150 µg/ml) to E coli cytoplasmic and membrane proteins to strips cut from the same blots shown for (A) and (B). UC and CD serum samples gave similar results to those shown for (C). The biotinylated (anti-human IgG) secondary antibody also bound to a single E coli cytoplasmic protein of Mr, 59 000 (shown on the ‘blank’ strips where no mucosal immunoglobulin sample was added), showing the band at Mr, 59 000 in control and UC samples in (A) and (C) is due to crossreaction of this detecting antibody and not to mucosal or serum IgG binding.

bacteria compared with active UC or inflamed (non-inflammatory bowel disease) controls. Mucosal IgG from active UC patients also shows increased binding compared with non-IBD inflamed controls although there is some overlap between the two ranges.

A significant advantage of the endoscopic mucosal wash technique over total intestinal lavage is that mucosal immunoglobulins can be obtained and isolated from defined regions of the intestinal tract. As the epithelial carrier for IgA (secretory component) does not translocate IgG, however, the appearance of IgG in endoscopic mucosal washings, or in ‘total intestinal lavage’ fluid, may reflect either local mucosal production of immunoglobulin or serum leakage. Although this effect would reduce our IgG yields from controls with irritable bowel syndrome, the bacterial protein binding ELISA results are corrected for IgG (and IgA) yields in each sample. It nevertheless remains possible that some of the immunoglobulin that binds to enterobacterial proteins is derived from the serum by leakage rather than being produced locally within the mucosa, however the following three findings argue against this being a significant effect. Firstly, there is higher binding of Crohn’s and UC mucosal IgG than serum immunoglobulin to commensal bacterial proteins when both are corrected for immunoglobulin concentration. Secondly, mucosal immunoglobulins did not show significant binding to non-intestinal commensal bacterial proteins, whereas there was strong binding by serum IgG. Thirdly, mucosal IgG bound to cytoplasmic but not membrane proteins of E coli, and serum bound to both protein fractions. We have found that isolation of immunoglobulin from biopsy or surgical material results in most of the immunoglobulin being derived from IgG being derived from IgG.

It has been well established that in CD there are activated T cells within the lamina propria. The increased production of IgG specific for commensal bacterial cytoplasmic proteins in active CD may reflect Th helper T cell help from these activated cells. Thus the recognition of bacterial peptides by this population of lymphocytes via their T cell receptors would result in profound immune activation from any damage to epithelial integrity. This may also explain the increased response seen in active CD compared with active UC.

The mechanism that causes mucosal IgG production against bacterial cytoplasmic but not membrane proteins in active CD and UC remains unclear. Serum immunoglobulins bind well both to membrane and cytoplasmic proteins, reflecting the systemic immune activation required to combat E coli bacteraemias. It is possible that mucosal immunoglobulins do not bind membrane proteins because they also contain antibodies to other cell wall components present in the membrane preparations (for example, lipopolysaccharide), which mask the protein binding, indeed some of the binding to the bacterial membrane fractions seen in ELISAs may reflect binding to lipopolysaccharide. Clearing with lipopolysaccharide or using inner membrane protein preparations, however, did not result in protein binding on the immunoblots. Alternatively, the response against membrane proteins may be inhibited. Knockout mice in which either the T cell receptor or IL2 genes have been disabled get intestinal inflammation unless they are bred in a germ free environment. This has been interpreted as inactivating a CD4+ IL2 producing population of αβ T cells, which are capable of inhibiting the mucosal B cell.
responses. As T cell clones can be rendered anergic (and hence inhibitory) in culture by repeated stimulation with antigen; this mechanism may apply mainly to membrane proteins as they are comparatively protected from proteolytic digestion by bound lipids. A further possibility is that antigen organisation within the bacterial membrane may directly influence membrane mediated T cell tolerance.

Although the range of immunoglobulin recovery in the washings may conceal small differences we found no significant change in total IgA concentrations in the various patient groups. We did observe low IgA titres to membrane preparations of colonic flora in UC compared with controls. Others have shown low concentrations of IgA, or of secretory component in some patients with UC, and decreased spontaneous IgA secretion, with an increased proportion of IgA, and monomeric IgA secretion in isolated intestinal mononuclear cells from patients with IBD. It has been suggested that in IBD, there may be a functional deficiency in IgA (owing to decreased dimeric IgA and IgA2) resulting in increased exposure of the colonic mucosa to bacterial proteins. Our data support this, although the importance of this finding in relation to pathogenesis is uncertain as the lowest control values overlap with the IgA titres found in UC.

In summary, there is a prominent immune response in the intestinal mucosa of CD directed against the normal colonic flora. There is also an IgG mediated immune response against normal colonic flora in the mucosa of patients with UC, but this is smaller than that in CD, suggesting that IgG produced in the former is also directed against other antigens, such as epithelial cell proteins. We suggest that relapse in CD results from an exaggerated immune response to luminal bacterial contents. While the mechanisms underlying this remain unclear, many of the known precipitants of relapse (for example, infections, non-steroidal anti-inflammatory drugs, and stress) have the potential to increase intestinal permeability, resulting in exposure of sensitised intestinal mucosa to luminal microbial antigens.

We are most grateful to Professor Alan McGregor for his support and encouragement during the course of this work. We would like to thank Drs A Marker and P O'Donnell for the histological assessments and Sister S Rolyard and the staff of the day surgery unit at King's College Hospital for assistance during the collection of patient samples. We are also grateful to Drs W Forgaerts and M H Jones for help with immunological techniques. Dr R Sherwood and Mr J Keating helped with the microalbumin estimations. We are also grateful to Dr K Chester and Professor D Vergani for helpful discussion. This work was supported by an MRC Clinician Scientist Fellowship to AJSM.

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