Antineutrophil cytoplasmic antibodies in sera from colectomised ulcerative colitis patients and its relation to the presence of pouchitis


Abstract

Background—Few studies have evaluated the influence of colectomy on antineutrophil cytoplasmic antibody (ANCA) positivity in ulcerative colitis (UC). In small series of patients it has been suggested that ANCA positivity in UC might be predictive for development of pouchitis after colectomy.

Aims—To assess the prevalence of ANCA in UC patients treated by colectomy and a Brooke’s ileostomy (UC-BI) or ileal pouch anal anastomosis (UC-IPAA), and the relation between the presence of ANCA, the type of surgery, and the presence of pouchitis.

Subjects—63 UC patients treated by colectomy (32 with UC-BI and 31 with UC-IPAA), 54 UC, and 24 controls.

Methods—Samples were obtained at least two years after colectomy. ANCA were detected by indirect immunofluorescent assay.

Results—There were no differences between patients with (36.3%) or without pouchitis (35.0%) and between patients with UC (55%), UC-BI (40.6%), and UC-IPAA (35.4%). However, ANCA prevalence significantly decreases in the whole group of operated patients (38.0%) compared with non-operated UC (p=0.04).

Conclusions—The prevalence of ANCA in operated patients was significantly lower than in non-operated UC, suggesting that it might be related either to the presence of inflamed or diseased tissue. ANCA persistence is not related to the surgical procedure and it should not be used as a marker for predicting the development of pouchitis.

(Gut 1996; 38: 894–898)

Keywords: ulcerative colitis, proctocolectomy, ileal pouch anal anastomosis, antineutrophil cytoplasmic antibodies, pouchitis.

Ulcerative colitis (UC) and associated conditions such as primary sclerosing cholangitis are some of the diseases in which antineutrophil cytoplasmic antibodies (ANCA) have been described.1-3 UC associated ANCA (UC-ANCA) most commonly show an immunofluorescence perinuclear pattern. Although not universally accepted,5 cytoplasmic and mixed ANCA patterns have been described associated to UC.4,6 On the other hand, because the antigenic specificity of UC-ANCA is unknown,1 3 9 10 they are referred to as atypical ANCA or x-ANCA by some authors.1-3 Few studies have specifically evaluated the influence of colectomy on ANCA positivity in UC.7-11-14 As well as only dealing with a small series of patients, none of them includes a large number of cases with Brooke’s ileostomy, which are the only ones that may be considered absolutely free of disease. On the other hand, all these series report a high prevalence of ANCA in patients with pouchitis (80–100%).4 7 11-13 Based on the results of these studies it has been suggested that ANCA assessment in UC may be useful in predicting the development of pouchitis in patients undergoing IPAA.2

The aims of this study were (a) to assess the prevalence of ANCA in UC patients treated with colectomy and either a Brooke’s ileostomy or ileal pouch anal anastomosis and (b) to determine the relation between the presence of ANCA, the type of surgery performed, and the presence of pouchitis.

Methods

PATIENTS AND CONTROLS

Ulcerative colitis patients treated with colectomy

One hundred and forty eight patients with a diagnosis of UC have had a colectomy since 1973 in three of four hospitals taking part in the study. The diagnosis had been established by the Lennard-Jones clinico pathological criteria.15 Patients with a history of indeterminate colitis were excluded from the study. Based on the information available, 92 of 114 patients living in Catalonia (31 895 km²) could be contacted by phone and asked to participate in the study. Sixty three of them agreed to be included. Thirty two had a Brooke’s ileostomy (UC-BI) (15 male, 17 female; 43 years (range: 23–76)) and 31 had an ileal pouch anal anastomosis (UC-ANCA) (18 male, 13 female; 34 years (range: 24–61)). The technique of IPAA included mucosectomy of the anorectal stump in five patients, whereas a staple ileal pouch anal anastomosis was performed in the remaining cases. In all patients a two loop J reservoir was made.

Disease controls

Fifty four UC patients who had not had an operation (35 male, 19 female; 40 years (range:
Antineutrophil antibodies in acute disease, patients with ulceration per low power field were included in the study. The diagnostic criteria were the same as above.

Healthy controls
Twenty-four volunteers, living in the same geographical area (11 male, 13 female; 43 years (range: 23–72)) acted as a healthy control.

CLINICAL ASSESSMENT
In all patients a complete clinical assessment was performed at the time of blood sampling for ANCA detection. In non-operated UC patients this included: activity and extent of the disease, current treatment, time from diagnosis, extraintestinal manifestations, and outcome. The activity of the disease was assessed by means of the Truelove index. Table I shows the clinical characteristics of the non-operated UC patients. The assessment of UC patients treated with a colectomy included: time from diagnosis and surgery, extraintestinal manifestations, previous and current treatment, and postoperative outcome.

Eight patients with UC-BI were previous UC-IPAA that had failed. The primary reason for pouch excision was chronic pouchitis resistant to medical treatment in two patients and fistula and pelvic sepsis in six patients. Four of the second group also had chronic continuous pouchitis. None of these patients had indeterminate or Crohn’s colitis. In UC-IPAA patients, the degree of pouch inflammation was assessed by means of the modified Moskowitz index (Table II). Pouchitis was defined as a score equal or greater than 7. Eleven of 31 UC-IPAA patients fulfilled this criterion at the time of inclusion.

Previous history of confirmed pouchitis was also recorded. Four of 11 patients with pouchitis at the time of inclusion had history of chronic continuous pouchitis resistant to medical treatment. Table III shows clinical data of UC patients treated with a colectomy. Some UC-IPAA patients were taking more than one drug. None of the patients included had primary sclerosing cholangitis.

### Table I: Clinical characteristics of the ulcerative colitis patients

<table>
<thead>
<tr>
<th>Activity</th>
<th>Patients</th>
</tr>
</thead>
<tbody>
<tr>
<td>Activity</td>
<td>No treatment</td>
</tr>
<tr>
<td>Inactive</td>
<td>Salicylates</td>
</tr>
<tr>
<td>Inactive</td>
<td>Corticosteroids</td>
</tr>
<tr>
<td>Mild</td>
<td>Azathioprine</td>
</tr>
<tr>
<td>Moderate</td>
<td>Corticosteroid refractory</td>
</tr>
</tbody>
</table>
| Severe | *First episode. Time elapsed from diagnosis: 4 months (0–25), 19–74), living in Catalonia, attending the ‘Hospital Universitari Germans Trias i Pujol’ were included in the study. The diagnostic criteria were the same as above.

### Table II: Pouchitis disease activity index (PDAI)

<table>
<thead>
<tr>
<th>Criteria</th>
<th>Score</th>
</tr>
</thead>
</table>
| Clinical | Fibrin
| Stool frequency: | Present
| Usual postoperative stool frequency | 1
| 1–2 stools/day>postoperative usual | 2
| 3 or more stools/day>postoperative usual | 3
| Rectal bleeding: | Present
| None or rare | 1
| Present daily | 2
| Fecal urgency or abdominal cramps: | Absent
| None | 0
| Occasional | 1
| Usual | 2
| Fever (temperature >37.8°C): | 0
| Absent | 0
| Present | 1
| Endoscopic inflammation | 1
| Oedema | 1
| Granularity | 1
| Friability | 1
| Loss of vascular pattern | 1
| Mucoseleucate | 1
| Ulceration | 1
| Acute histological inflammation | 1
| Polymorphonuclear leucocyte infiltration: | 1
| Mild | 1
| Moderate+ crypt abscess | 2
| Severe+ crypt abscess | 3
| Ulceration per low power field (mean): | 1
| <25% | 1
| 25–50% | 2
| >50% | 3

### Table III: Clinical characteristics of UC patients treated with a colectomy

<table>
<thead>
<tr>
<th>UC-BI</th>
<th>UC-IPAA pouchitis</th>
<th>UC-IPAA no pouchitis</th>
</tr>
</thead>
<tbody>
<tr>
<td>Years from diagnosis</td>
<td>14 (4–36)</td>
<td>6 (2–34)</td>
</tr>
<tr>
<td>Years from surgery</td>
<td>9 (2–16)</td>
<td>4 (2–9)</td>
</tr>
<tr>
<td>Clinical score*</td>
<td>–</td>
<td>2 (0–4)</td>
</tr>
<tr>
<td>Endoscopic score*</td>
<td>–</td>
<td>4 (3–6)</td>
</tr>
<tr>
<td>Histological score*</td>
<td>–</td>
<td>2 (1–5)</td>
</tr>
<tr>
<td>Global score*</td>
<td>–</td>
<td>8 (7–11)</td>
</tr>
<tr>
<td>Treatment (patients):</td>
<td>–</td>
<td>32</td>
</tr>
<tr>
<td>No treatment</td>
<td>–</td>
<td>2</td>
</tr>
<tr>
<td>Salicylates</td>
<td>–</td>
<td>0</td>
</tr>
<tr>
<td>Local corticosteroids</td>
<td>–</td>
<td>3</td>
</tr>
<tr>
<td>Antibiotics*</td>
<td>–</td>
<td>0</td>
</tr>
<tr>
<td>Pouchitis disease activity index (PDAI)</td>
<td>–</td>
<td>4</td>
</tr>
</tbody>
</table>

*Pouchitis disease activity index; fmetronidazole/tetracyclines.
cytoplasmic staining was observed at a 1:40 dilution. The second cut off dilution was chosen to eliminate unspecified cytoplasmic positivity yielded by some negative sera. All negative sera and those given as positive by only one observer were retested blindly. Serum samples given as positive by only one observer in duplicate assay were considered as negative. Positive ANCA samples were titrated to a dilution of 1/1280.

To differentiate positive ANCA from anti-nuclear antibodies (ANA) that mimic ANCA, all positive sera were tested with HEP-2 cells and rat liver sections, which are highly sensitive for ANA detection. All ANA positive sera with stronger titres than that observed in neutrophil slides were considered as ANCA negative. This occurred in four patients (three non-operated and one operated) and one healthy control.

All assays were performed at the Research Laboratories of the Hospital Universitary ‘Germans Trias i Pujol’.

**Statistical analysis**

Comparison of the frequencies between patients with or without pouchitis were analysed using the two tail χ² and Yates’s correction. The frequencies between UC, UC-IB, and UC-IPAA were compared using a two tail χ² and the comparison between operated and non-operated UC patients was performed using a left sided Fisher exact test. Titres were compared using the Kruskal-Wallis test. The Spearman rank correlation coefficient was used to determine associations between titres and the time elapsed from surgery. The results are expressed by means of median and frequencies with their range and 95% confidence intervals (CI), respectively. The statistical procedures were performed using the programs of the Biomedical Data Processing, BMDP (BMDP, Statistical Software, Los Angeles, California, 1986).

**Ethical considerations**

The study was performed in accordance with the 1975 Declaration of Helsinki ethical guidelines and was approved by the Research and Ethical Committees of the Hospitals.

**Results**

Table IV shows the number of ANCA positive patients and controls in the groups studied.

<table>
<thead>
<tr>
<th>ANCA Positive Patients</th>
<th>ANCA Positive Patients</th>
</tr>
</thead>
<tbody>
<tr>
<td>UC-IB</td>
<td>UC-IPAA</td>
</tr>
<tr>
<td>Ulcerative colitis</td>
<td>30 (55.5%)*</td>
</tr>
<tr>
<td>UC-IB</td>
<td>13 (40.6%)†</td>
</tr>
<tr>
<td>UC-IPAA with pouchitis</td>
<td>4 (36.3%)‡</td>
</tr>
<tr>
<td>UC-IPAA without pouchitis</td>
<td>7 (35.0%)‡</td>
</tr>
<tr>
<td>Healthy controls</td>
<td>0</td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th>UC-IB (total proctocolectomy plus Brooke’s ileostomy)</th>
</tr>
</thead>
<tbody>
<tr>
<td>UC-IPAA (proctocolectomy plus ileal pouch anal anastomosis)</td>
</tr>
</tbody>
</table>

*26 p-ANCA; 1 c-ANCA; 3 mixed pattern; †11 p-ANCA; 2 mixed pattern; ‡3 p-ANCA; 1 mixed pattern; §6 p-ANCA; 1 mixed pattern.

The positivity of ANCA was significantly higher in all groups studied compared with the healthy control group (p<0.0001).

**Relation between ANCA positivity and the presence of pouchitis**

In patients with UC-IPAA, there were no differences in ANCA positivity between those with (36-3% (CI: 10-9 to 69-2)) or without pouchitis (35-0% (CI: 15-5 to 59-2)) (p=0.720).

Two of four (CI: 6-7 to 93-2%) patients with chronic continuous pouchitis and four of eight UC-IPAA failure (CI: 15-7 to 84-3%) were ANCA positive.

As the percentage of ANCA positive patients was the same as those with or without pouchitis, they were considered together and compared with both patients treated with a colectomy and non-operated UC patients to determine the influence of a Brooke’s ileostomy upon ANCA positivity.

**Relation between ANCA positivity and the type of surgery performed**

There were no differences in ANCA positivity between patients with UC-IB (40-6% (CI: 23-7 to 59-3)) and UC-IPAA (35-4% (CI: 19-2 to 54-6)) and non-operated UC patients (55% (CI: 41-4 to 69-1)) (p=0.154).

Because there were no significant differences between both groups of operated patients, they were considered together (both UC-IB and UC-IPAA) and compared with UC non-operated patients to determine if colectomy decreases ANCA positivity. There was a significant decrease in ANCA prevalence in operated patients – 38-0% (CI: 26-1 to 51-2) – compared with non-operated UC (55%) (p=0.044).

The Figure shows the titres of ANCA and UC, UC-IB, and UC-IPAA. There were no differences in ANCA titres between groups...
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(p = 0.755). There was also no correlation between the time elapsed from surgery and the titre of ANCA (r = 0.098).

Discussion

There is no agreement about those immunofluorescence ANCA patterns that should be considered associated with UC. Some laboratories have considered that all UC-ANCA are of the perinuclear type,\(^9\)\(^{19-21}\) whereas others also include those less frequent cases showing either diffuse cytoplasmic or mixed pattern.\(^9\)\(^6\)\(^6\)

We share this second view as, until the specific antigen(s) are identified, it does not seem reasonable to exclude any type of ANCA pattern when an unequivocal immunofluorescence staining is found.

One of the aims of this study was to assess if removal of the colon results in a decline of ANCA in UC patients and if there is a relation to the type of surgery performed. The percentage of ANCA positivity in our UC non-operated patients (55%) is similar to other reports,\(^9\)\(^{22}\)\(^{23}\) and is within the range described in published reports (40–80%).\(^4\)\(^9\)\(^{19-25}\) Whether or not no circulating ANCA persist long term after colectomy in UC patients is not well established. Although most studies have reported the persistence of ANCA after surgery,\(^4\)\(^7\)\(^{11-13}\)\(^{25}\) there are some reports of postoperative ANCA disappearance.\(^14\)\(^{26}\)

However, most studies include a small number of patients and none of them includes a large number of cases with Brooke’s ileostomy.

Although there are no data about the time course of ANCA behaviour after colectomy, we thought that two years might be a reasonable period of time for assessing ANCA clearance, assuming that they are related to the presence of diseased tissue. We have not found a significant difference in ANCA positivity between non-operated UC patients, those with UC-BI, and UC-IBA,\(^6\)\(^{26}\) showing that these antibodies persist in the serum in a high percentage of UC operated patients long term after colectomy. However, we found a significant decrease of ANCA positivity when both groups of operated patients were considered together (that is, increasing twofold the number of operated patients). This finding might disclose the existence of a weak relation between ANCA positivity and inflammation, as has been shown in some studies.\(^6\)\(^{26}\) In fact, our non-operated patients showed a non-significant trend towards a greater percentage of ANCA positivity in those with active versus inactive UC (63% vs 43%; \(p=0.13\)), and in those with extensive disease versus proctitis (63% vs 31%; \(p=0.07\)). An alternative explanation would be that ANCA clearance after colectomy was a very slow process taking more than two years and a prospective and paired assessment before and after surgery in 15 UC patients found a significant decrease in ANCA titre a mean of two years after colectomy.\(^14\) In our study, no differences between groups with respect to ANCA titre, or any relation between the titres of ANCA and the time elapsed from surgery were found. However, as this was not a paired comparison, interindividual differences in the ability of antibody production may have prevented the discovery of such differences.

In this series we found a similar frequency of ANCA positivity between those patients with and without pouchitis. In contrast, several studies have reported a higher prevalence of ANCA in patients with pouchitis.\(^4\)\(^7\)\(^{11-13}\)

However, most of them include a small number of patients with pouchitis or it is not well defined, or both.\(^4\)\(^11-13\) In this regard, to distinguish between relapsing and chronic continuous pouchitis may be relevant. In the largest reported series in which all patients with pouchitis had chronic inflammation resistant or dependent on medical treatment, frequency of ANCA positivity was 100%.\(^7\) The authors suggest that the strong relation between ANCA and pouchitis would be related to a specific disease pattern (chronic continuous pouchitis) rather than to the mere presence of inflammation.\(^7\) In the remaining studies, it is not stated if those patients with pouchitis had relapsing or chronic continuous disease. However, a recent study published as an abstract, showed that those patients without pouch inflammation at inclusion, but with previous episodes of pouchitis, had a 53.3% of ANCA positivity.\(^12\) This figure is similar to the percentage found in our patients with pouchitis, most of them having relapsing disease. On the other hand, in our series two of four chronic resistant pouchitis and four of eight IPAA failures had positive ANCA.

Taken as a whole, all these data suggest that ANCA would not predict the development of relapsing pouchitis, and the strong association between ANCA positivity and chronic continuous pouchitis deserves further confirmation. However, even if a 100% association (sensitivity) could be confirmed, the low specificity of ANCA for chronic continuous pouchitis will make the use of this marker doubtful as a predictor of this condition.

Although all these queries could be only answered by means of a longitudinal study, its practicability seems to be questionable. Such a study would be very long lasting, taking into account that pouchitis may appear as late as 10 years after surgery.\(^27\)\(^{28}\) In the case of chronic continuous pouchitis, an event occurring in only 5% of the total IPAA,\(^28\) the great number of patients to be included would be an additional difficulty.

In conclusion, ANCA persist in a high proportion of UC patients after colectomy and its presence is not related to the type of surgery performed. The lower percentage found in operated patients suggests that ANCA positivity might be related to the presence of either inflamed or diseased tissue. On the other hand, ANCA positivity in UC should not be used as a marker for predicting the development of pouchitis and, at present, it does not seem a good parameter for deciding the type of surgery to be performed in UC patients.

Part of this study has been presented as a poster at the 95th Annual Meeting of the American Gastroenterological Association held in San Diego in May 1995, and published as an abstract in Gastroenterology 1995; 108: A816.


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