**Impact of the increasing use of immunosuppressants in Crohn’s disease on the need for intestinal surgery**

J Cosnes, I Nion-Larmurier, L Beaugerie, P Afchain, E Tret, J-P Gendre


**Background/Aim:** Immunosuppressants are now used much earlier in the course of Crohn’s disease; however their effect on the natural history of the disease, especially on the need for surgery, is not known. The aim of this study was to assess the evolution of the need for surgery in Crohn’s disease during the last 25 years.

**Patients and Methods:** The medical charts of 2573 patients were reviewed retrospectively. The use of immunosuppressants (azathioprine or methotrexate), the need for intestinal resection, and the occurrence of intestinal complications were assessed using Kaplan-Meier analysis in five consecutive cohorts of patients defined by the date of diagnosis of Crohn’s disease (1978–82; 1983–87; 1988–92; 1993–97; 1998–2002).

**Results:** In 565 patients seen in the authors’ unit within the first three months after diagnosis, characteristics of Crohn’s disease at diagnosis did not differ from one cohort to another. The five year cumulative probability to receive immunosuppressants increased from 0 in the 1978–82 cohort to 0.13, 0.25, 0.25, and 0.56 in the 1983–87, 1988–92, 1993–97, and 1998–2002 cohorts, respectively (p<0.001). Concomitantly, the cumulative risk of intestinal resection remained unchanged (from 0.35 to 0.34 at five years; p = 0.81). The cumulative risk of developing a strictureing or a penetrating intestinal complication remained also unchanged. Similar results were obtained in the 2008 patients seen during the same period who were referred to us more than three months after diagnosis.

**Conclusion:** Although immunosuppressants have been used more frequently over the last 25 years, there was no significant decrease of the need for surgery, or of intestinal complications of Crohn’s disease.

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The treatment strategy for Crohn’s disease (CD) is evolving.1 Steroids and 5-aminosalicylates are still commonly used; however, during recent years there have been some significant changes. Firstly, the remarkable efficacy and the overall good tolerance of immunosuppressants were recognised2,3 and when compared with the morbidity of untreated disease,4 led to their wider greater use. Some authors even proposed using immunosuppressants from the very beginning of the disease, with the objective to modify its natural history,5 and this strategy has been shown to improve disease course and to avoid steroids in children with moderate to severe disease.6-8 Concomitantly, anti-tumour necrosis factor therapy became available in the late 1990s. Infliximab was shown to be able to induce clinical remission in patients unresponsive to standard treatment,9 to clear anatomic damage, and, when used as maintenance treatment, to maintain clinical remission10 and fistula closure.11

However, the impact of an increasing use of immunosuppressants and novel therapies on the natural history of CD remains poorly evaluated. An important and unbiased criterion for assessing the overall severity of CD is the need for surgery: for decades there has been a consensus to limit surgery to complications and refractory intractable forms.

The aim of our study, performed in a large series of patients with CD, was to evaluate the effect upon the need for surgery of an increasing use of immunosuppressants over the last 25 years.

**Patients and Methods**

**Patients**

Patients with CD who were seen consecutively in our unit between January 1978 and December 2002 were included retrospectively. In January 2003, our unit moved from Rothschild Hospital to St-Antoine Hospital and recruitment for this study was interrupted. Diagnosis of CD was based on Lennard-Jones criteria.12 Two groups of patients were analysed. The first group consisted of 565 patients who were seen in our unit early in the course of their disease—within the first three months following diagnosis of CD. The time of diagnosis was defined as the date of first detection of unequivocal inflammatory abnormalities of the intestine, as assessed from radiological, endoscopic, or peroperative observations. Patients in the second group (n = 2008) were seen more than three months after diagnosis, being referred to us for various reasons. Within each group, patients were divided into five consecutive five year cohorts, according to the date of diagnosis: cohorts 1978–82, 1983–87, 1988–92, 1993–97, and 1998–2002.

**Study design**

The study analysed the evolution of surgical requirements over 25 years by comparing the need for excisional surgery in relation to the use of immunosuppressants among the five chronologic cohorts. Only excisional intestinal surgery was taken into account. Appendectomy, stricturoplasty, bypass, and elective surgical treatment of abscess were not considered. The time to first intestinal resection was the main outcome criteria. In addition were noted the time to first large intestinal resection—either alone or cumulative, defined by a post-surgical index equal or superior to 20.13— and the time to establishment of a definitive stoma. The post-surgical handicap index has been developed to predict the functional consequences of intestinal resection for CD. It is calculated from operative records, taking into account the location and extent of intestinal resection. An index score equal or superior to 20 has a high predictive value of

**Abbreviations:** CD, Crohn’s disease; HR, hazard ratio.
diarrhoea following intestinal resection. Intestinal complications of CD were defined according to the Vienna classification: intestinal strictures as the occurrence of constant luminal narrowing demonstrated by radiological, endoscopic, or surgical examination combined with pre-stenotic dilatation and/or obstructive signs or symptoms but without evidence of penetrating disease. Perforations were intra-abdominal fistulas, inflammatory masses, and/or abscesses. First morphological demonstration of narrowing or penetrating complication was used to date the occurrence of the complication.

**Treatment of Crohn’s disease**

Our treatment policy has been described elsewhere. Flare up episodes were treated with mesalamine (3–4 g daily) or prednisolone (1 mg/kg per day, progressively tapered after four weeks), according to their clinical severity. When steroid unresponsive or intolerant to azathioprine. Its dosage was increased to 2.5–3 mg/kg per day. Intramuscular methotrexate (20–25 mg weekly) was used in patients operated on, should be interpreted in relation to different durations of follow up. Azathioprine was maintained for a prolonged period in most cases but had to be stopped within a five year cumulative probability of needing immunosuppressants or surgery after diagnosis. Survival curves were compared by means of a two sided log rank test. A Cox proportional hazards regression model with a backward variable elimination procedure was used to assess the strength of the associations while controlling for possible confounding variables. All baseline variables suspected to be possible predictors of intestinal surgery (age, sex, initial disease location (upper GI tract, jejunal, ileal, colonic, rectal, or anoperineal lesions), smoking status, appendectomy, familial history, ethnicity (White, North-African non-Jewish ancestry), geographic origin outside Paris area, and calendar period of diagnosis (before or after 30 June 1990), were entered into the model. Results of analysis are presented as hazard ratios (HRs) with 95% confidence intervals. Calculations were performed using GB-STAT statistical software (Silver Spring, MD, USA).

**RESULTS**

The characteristics of CD at diagnosis in the five cohorts of group 1 are given in table 1. Patients were very similar at diagnosis from one cohort to another, with a predominance of females, a mean age about 30 years, a large proportion of smokers (1/2 of the patients), and a similar disease location. Table 2 gives the cumulative characteristics of the disease at the end of 2003. No attempt was made to contact the patients at that time and 31% of them had been lost to follow up. Because patients from the oldest cohorts had a longer disease duration, they developed more stricturing or penetrating complications and were classified so according to Vienna classification. Similarly, the respective proportions of patients needing steroids or immunosuppressants, and operated on, should be interpreted in relation to different durations of follow up. Azathioprine was maintained for a prolonged period in most cases but had to be stopped within the first month because of adverse events in 16 patients (11%). It was switched to methotrexate in eight of those latter patients.

**Changes in the use of immunosuppressants over 25 years in group 1 patients**

Figure 1 shows the cumulative use of immunosuppressants in the five cohorts. Data of patients for whom immunosuppressants had to be stopped early are included. As expected, immunosuppressants have been used more and more early over the last 25 years, with a five year cumulative probability

**Table 1 Characteristics at diagnosis of the five consecutive chronological cohorts of patients seen in our unit within the three months following diagnosis (group 1)**

<table>
<thead>
<tr>
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</tr>
</thead>
<tbody>
<tr>
<td>Ileocolonic</td>
<td>13 (38)</td>
<td>19 (41)</td>
<td>41 (40)</td>
<td>60 (34)</td>
<td>72 (35)</td>
</tr>
<tr>
<td>Small bowel</td>
<td>11 (32)</td>
<td>16 (35)</td>
<td>32 (31)</td>
<td>59 (34)</td>
<td>70 (34)</td>
</tr>
<tr>
<td>Colon</td>
<td>7 (21)</td>
<td>10 (22)</td>
<td>27 (26)</td>
<td>52 (30)</td>
<td>60 (29)</td>
</tr>
<tr>
<td>Other</td>
<td>3 (9)</td>
<td>1 (2)</td>
<td>2 (2)</td>
<td>5 (2)</td>
<td>5 (2)</td>
</tr>
<tr>
<td>Appendectomy</td>
<td>13 (38)</td>
<td>20 (43)</td>
<td>35 (34)</td>
<td>49 (28)</td>
<td>63 (30)</td>
</tr>
<tr>
<td>Familial history</td>
<td>4 (12)</td>
<td>13 (28)</td>
<td>10 (10)</td>
<td>20 (17)</td>
<td>33 (16)</td>
</tr>
<tr>
<td>Smokers</td>
<td>18 (53)</td>
<td>31 (67)</td>
<td>51 (50)</td>
<td>86 (49)</td>
<td>100 (48)</td>
</tr>
<tr>
<td>Caucasian</td>
<td>30 (88)</td>
<td>40 (87)</td>
<td>93 (91)</td>
<td>133 (87)</td>
<td>167 (81)</td>
</tr>
<tr>
<td>North-African non-Jewish ancestry</td>
<td>2 (6)</td>
<td>6 (13)</td>
<td>8 (8)</td>
<td>16 (9)</td>
<td>26 (13)</td>
</tr>
<tr>
<td>Living outside Paris area</td>
<td>13 (38)</td>
<td>19 (41)</td>
<td>26 (25)</td>
<td>57 (32)</td>
<td>65 (31)</td>
</tr>
</tbody>
</table>

Numbers in parentheses are percentages. Intergroup comparisons showed no significant differences.
of prescription of zero in the 1978–82 cohort to 0.56 (95% CI 0.31 to 0.78) in the 1998–2002 cohort.

Cumulative need for excisional surgery in the five group 1 cohorts

One hundred and ninety patients (34%) were operated on at least once. Excisional surgery was performed in 41 cases before first admission in our unit and thereafter in 149 cases. Figure 2 gives the cumulative need for first excisional surgery. The curves were superimposed, with no significant difference from one curve to another (log rank, p = 0.81). Excluding the 80 patients who had had surgery within the first three months following diagnosis yielded the same result (log rank, p = 0.49), although immunosuppressants were used in that group much earlier over the years (log rank, p = 0.0001). In the whole group of 565 patients, Cox analysis confirmed that the year of diagnosis had no significant effect upon the need for surgery. Factors associated with surgery were ileal involvement (HR 2.78; 95% CI 2.19 to 3.51) and absence of rectal involvement (HR 0.34; 95% CI 0.27 to 0.43).

In the cohorts 1978–82, 1983–87, 1988–92, 1993–97, and 1998–2002, the five year cumulative probabilities of having a large intestinal resection, defined by a post-surgical handicap index >20, were 0.29 (0.15–0.50), 0.20 (0.10–0.36), 0.24 (0.16–0.35), 0.13 (0.07–0.22), and 0.17 (0.04–0.49), respectively. The curves were not significantly different according to log rank test (p = 0.23). The five year cumulative probability of having a definitive stoma varied not significantly between 0 and 0.03 from one cohort to another (p = 0.33).

Indications for first intestinal resection in group 1 patients

Table 3 gives the indications for the first intestinal surgery in the five cohorts. The proportion of patients being operated on for medical failure, stricture, and perforation, respectively, did not change significantly between the five cohorts, although in the most recent cohort there was a clear reduction of operations for medical failure (13% vs 22–38% in the other cohorts). Kaplan-Meier analysis of the cumulative probability of intestinal stricture and perforation did not show significant differences between cohorts. In the cohorts 1978–82, 1983–87, 1988–92, 1993–97, and 1998–2002, the five year cumulative probabilities of intestinal stricture were 0.23 (0.11–0.43), 0.14 (0.06–0.29), 0.19 (0.12–0.30), 0.17 (0.11–0.26), and 0.10 (0.02–0.42), respectively (log rank

Table 2 Cumulative characteristics of the five consecutive chronologic cohorts of patients seen in our unit within the three months following diagnosis (group 1)

<table>
<thead>
<tr>
<th>Cohort</th>
<th>Patients (n)</th>
<th>Deceased</th>
<th>Lost to follow up</th>
<th>Median disease duration, months (interquartile range)</th>
<th>Vienna classification</th>
<th>Structuring</th>
<th>Penetrating</th>
<th>Medical treatment</th>
<th>IS, immunosuppressants</th>
</tr>
</thead>
<tbody>
<tr>
<td>1978–82</td>
<td>34</td>
<td>4</td>
<td>14 (41)</td>
<td>188 (23–256)</td>
<td>Inflammatory 15 (44)</td>
<td>9 (26)</td>
<td>10 (29)</td>
<td>No steroids, no IS 6 (18)</td>
<td>Early withdrawal of IS 0</td>
</tr>
<tr>
<td>1983–87</td>
<td>46</td>
<td>2</td>
<td>20 (43)</td>
<td>181 (64–207)</td>
<td>Structuring 4 (9)</td>
<td>25 (54)</td>
<td>10 (29)</td>
<td>Steroids, no IS 22 (65)</td>
<td>IS 6 (18)</td>
</tr>
<tr>
<td>1988–92</td>
<td>102</td>
<td>3</td>
<td>42 (41)</td>
<td>123 (43–144)</td>
<td>Penetrating 17 (17)</td>
<td>43 (42)</td>
<td>25 (54)</td>
<td>Steroids, no IS 29 (63)</td>
<td>IS 10 (22)</td>
</tr>
<tr>
<td>1993–97</td>
<td>176</td>
<td>3</td>
<td>70 (40)</td>
<td>59 (19–101)</td>
<td>Inflammatory 51 (50)</td>
<td>51 (50)</td>
<td>31 (30)</td>
<td>No steroids, no IS 80 (45)</td>
<td>IS 31 (30)</td>
</tr>
<tr>
<td>1998–2003</td>
<td>207</td>
<td>0</td>
<td>76 (40)</td>
<td>66 (32–144)</td>
<td>Structuring 22 (13)</td>
<td>59 (34)</td>
<td>39 (22)</td>
<td>Early withdrawal of IS 0</td>
<td></td>
</tr>
<tr>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td>Medical treatment 6 (25)</td>
<td>66 (32)</td>
<td>66 (32)</td>
<td>IS 6 (18)</td>
<td>Early withdrawal of IS 0</td>
</tr>
</tbody>
</table>

Table 3 Indications for the first intestinal resection in the five consecutive chronologic cohorts of patients seen in our unit within the three months following diagnosis (group 1)

<table>
<thead>
<tr>
<th>Cohort</th>
<th>Resections (n)</th>
<th>Diagnosis</th>
<th>Stricture</th>
<th>Perforation</th>
<th>Failure of medical treatment</th>
</tr>
</thead>
<tbody>
<tr>
<td>1978–82</td>
<td>16</td>
<td>1 (6)</td>
<td>6 (38)</td>
<td>3 (19)</td>
<td>6 (38)</td>
</tr>
<tr>
<td>1983–87</td>
<td>23</td>
<td>0 (0)</td>
<td>3 (13)</td>
<td>1 (5)</td>
<td>5 (22)</td>
</tr>
<tr>
<td>1988–92</td>
<td>49</td>
<td>0 (0)</td>
<td>11 (22)</td>
<td>21 (43)</td>
<td>16 (33)</td>
</tr>
<tr>
<td>1993–97</td>
<td>56</td>
<td>1 (2)</td>
<td>8 (14)</td>
<td>23 (45)</td>
<td>19 (34)</td>
</tr>
<tr>
<td>1998–2003</td>
<td>46</td>
<td>1 (2)</td>
<td>7 (15)</td>
<td>31 (67)</td>
<td>6 (13)</td>
</tr>
</tbody>
</table>

Numbers in parentheses are percentages.

*Other included mesenteric infarction, haematochezia, and small bowel adenocarcinoma, respectively.
Evolution of the need for surgery 1978–2003

In the total cohort of 2573 patients from groups 1 and 2, 1070 underwent 1426 intestinal resections from January 1978 to December 2003 (22 928 patient years). Two hundred and seventeen resections (15%) were performed within the first three months following diagnosis. Except for the year 1978 (46 patients only), the percentage of patients who were operated on during the first three months remained less than 5% (fig 3). After the first three months, the operative rate (number of operations performed per year) fluctuated within a narrow range (3.3–7.5%), without any significant change over 26 years (fig 3).

DISCUSSION

This study shows that although immunosuppressants were initiated much earlier during the course of CD, the need for intestinal resection remained stable over 25 years. The percentage of patients requiring intestinal surgery each year remained equal. The probability of having a definitive stoma appeared also to be unaffected from 1978 to 2002. However, large intestinal resections became more unusual.

This study has some limitations. Firstly, the retrospective nature of the study may have led to bias in the interpretation of the data—however, it was necessary to obtain an observation period long enough to ascertain the long term effect on surgery of changes in the medical strategy of CD. In addition, intestinal resection can be considered as an unbiased and solid criterion, even retrospectively, as it is performed only when necessary. Besides, during a period of 20–25 years, many factors other than the treatment strategy may have influenced the indications for surgery. However, all patients seen from the beginning were followed up in the same unit by the same small group of physicians, who used homogeneous guidelines and took collegial decisions. Moreover, comparison of cohorts at inclusion showed that compared with the first group of patients, a higher proportion of referred patients had had surgery before admission in our unit (p<0.001). Otherwise, results observed were similar to those of group 1 regarding an increased use of immunosuppressants but a stable need for excisional surgery over the years (table 4). The risk of having a definitive stoma remained also stable. However, in that group, the risk of having a large intestinal resection decreased significantly with time.

Table 4: Five year cumulative rates (95% CI) of therapeutic intervention in the patients referred to our unit more than three months after diagnosis of Crohn’s disease (group 2, n = 2008)

<table>
<thead>
<tr>
<th>Cohort</th>
<th>Patients (n)</th>
<th>IS therapy</th>
<th>Cumulative probability (95% CI)</th>
<th>Patients at risk (n)</th>
<th>Intestinal resection</th>
<th>Cumulative probability (95% CI)</th>
<th>Patients at risk (n)</th>
<th>Large intestinal resection</th>
<th>Cumulative probability (95% CI)</th>
<th>Patients at risk (n)</th>
<th>Definitive stoma</th>
<th>Cumulative probability (95% CI)</th>
<th>Patients at risk (n)</th>
</tr>
</thead>
<tbody>
<tr>
<td>1978–82</td>
<td>218</td>
<td></td>
<td>0.04 (0.02–0.08)</td>
<td>195</td>
<td>0.36 (0.29–0.48)</td>
<td>0.29 (0.23–0.35)</td>
<td>145</td>
<td>0.01 (0.00–0.03)</td>
<td>202</td>
<td>0.02</td>
<td>0.01 (0.00–0.04)</td>
<td>387</td>
<td></td>
</tr>
<tr>
<td>1983–87</td>
<td>342</td>
<td></td>
<td>0.14 (0.10–0.18)</td>
<td>278</td>
<td>0.30 (0.25–0.35)</td>
<td>0.22 (0.18–0.27)</td>
<td>250</td>
<td>0.02 (0.01–0.04)</td>
<td>315</td>
<td>0.02</td>
<td>0.01 (0.01–0.04)</td>
<td>387</td>
<td></td>
</tr>
<tr>
<td>1988–92</td>
<td>486</td>
<td></td>
<td>0.27 (0.23–0.32)</td>
<td>296</td>
<td>0.32 (0.28–0.37)</td>
<td>0.19 (0.16–0.23)</td>
<td>321</td>
<td>0.02 (0.01–0.04)</td>
<td>387</td>
<td>0.02</td>
<td>0.01 (0.01–0.04)</td>
<td>355</td>
<td></td>
</tr>
<tr>
<td>1993–97</td>
<td>563</td>
<td></td>
<td>0.45 (0.40–0.50)</td>
<td>203</td>
<td>0.31 (0.27–0.36)</td>
<td>0.15 (0.11–0.19)</td>
<td>310</td>
<td>0.02 (0.01–0.04)</td>
<td>387</td>
<td>0.02</td>
<td>0.01 (0.01–0.04)</td>
<td>335</td>
<td></td>
</tr>
<tr>
<td>1998–2002</td>
<td>399</td>
<td></td>
<td>0.63 (0.49–0.76)</td>
<td>17</td>
<td>0.36 (0.22–0.53)</td>
<td>0.12 (0.05–0.28)</td>
<td>27</td>
<td></td>
<td></td>
<td></td>
<td></td>
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<td></td>
</tr>
</tbody>
</table>

Log rank p value

- IS therapy: <0.0001
- Intestinal resection: <0.0001
- Large intestinal resection: <0.0001
- Definitive stoma: 0.72
they were very similar regarding demographic characteristics and disease location. In particular there is no reason to believe that CD became more severe with time while other disease characteristics did not change. Secondly, it should be noted that a relatively large proportion of patients were lost to follow up. We made no attempt to contact the patients or physicians to update the data. We do not believe this may have minimised the need for surgery of the oldest cohorts because patients who are lost to follow up are usually those doing well and not requiring further surgery. The cumulative probability of surgery in our patients was very similar to that reported in two unbiased and complete series of the literature, the NCCDS17 and the Copenhagen County cohort study.14 Finally, our unit is a tertiary referral centre and referral bias is unavoidable. To limit this bias, we restricted the analysis to patients seen during the first three months of the disease course. This precaution was not sufficient to eliminate such a referral bias because an important proportion of these patients came to surgery during that period. However, when we excluded these latter patients, analyses gave similar results and, in particular, the discrepancy between an increased use of immunosuppressants and a stable need for surgery remained unchanged. These results were confirmed in a second large group of patients.

The occurrence of strictureing and perforation complications was the main reason for excisional surgery. The frequency of these complications did not change significantly from one cohort to another. This is a disappointing result because it could be expected that immunosuppressants could have an anatomic effect and prevent these complications. Indeed, D’Haens et al reported that in 74% of patients with colonic or ileocolonic disease who were clinically responders to azathioprine, endoscopic lesions had healed completely or nearly completely after a mean of two years of treatment. Histologically there was disappearance of the inflammatory infiltrate, with only a degree of architectural disturbance remaining.19 In the present study there was, over the most recent years, a clear decrease of intestinal resection for medical failure, which represented only 13% of surgical indications in the 1998–2002 cohort. This result may be related to immunosuppressants, but also to infliximab even though it was used sparingly in very few patients. In any case, the absence of a decrease in the need for surgery over the last 25 years questions the efficiency of our medical strategy. In fact, there is the possibility that in this study immunosuppressants had no impact on complications and the need for surgery because they were given too late during the course of the disease. Supporting this hypothesis, a large majority of patients came to surgery while not having received immunosuppressants, or were operated on early (within the first three months), while the mean time of response to purine analogues is three months.20 By contrast, if we make the hypothesis that immunosuppressants are able to change the natural history of CD, nearly half the patients who were operated on more than three months after diagnosis could have avoided surgery. Thus we can extrapolate that in our series immunosuppressants were given too late and too scarcely to have a significant impact on the need for surgery. Such an assertion is not in accordance with the anatomic effect observed by D’Haens et al.,19 that immunosuppressants, even when given early, may have no preventive effect on the occurrence of strictureting and penetrating complications.

An interesting and more encouraging result of our study was the decrease of the probability of having a large intestinal resection over the last 25 years in the group of referred patients. A similar trend, although not significant, was observed in patients who were seen early after diagnosis and were, for the most part, operated on in our surgical unit. The reason for such a decrease, from 29% to 12% five years after diagnosis, may be linked to a greater use of immunosuppressants, but may also be a change in the surgical strategy favouring segmental and limited resections in the most recent years.

In summary, this study shows that immunosuppressants have been used increasingly over the years. However, this evolving therapeutic strategy was not associated with a decrease in the need for surgery or in a decrease of the occurrence of intestinal complications. This result does not question the efficacy of immunosuppressants for achieving and maintaining remission,21 sparing steroids,22 and improving quality of life, but it does questions the timing of starting immunosuppressants in patients with moderate to severe CD.

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REFERENCES
Efficacy and strategy of pneumatic dilatation in achalasia

We read with interest the article by Eckardt et al regarding the long term results of pneumatic dilatation in achalasia (Gut 2004;53:629–33). Fifty four patients were followed up for a median of 14 years after a single pneumatic dilatation using the Browne-McHardy dilator. Five and 10 year remission rates were 40% and 36%, respectively, and repeated dilatations only mildly improved the clinical response. Most of the relapses occurred within one year of dilatation. Patients with post-dilatation lower oesophageal sphincter pressures of <10 mm Hg had a significantly better outcome. The authors suggest that failure to respond to the first dilatation should lead to consideration of alternative therapy.

We disagree with this conclusion and we would like to bring to your attention a recent prospective study on the long term effects of pneumatic dilatation in 11 patients with achalasia.1 A different approach was taken—that is, treatment consisted of one or more pneumatic dilations under conscious sedation in order to achieve stable clinical remission, defined as persisting one year after dilatation. To this end, close follow up was performed in the first year after dilatation (scheduled assessments at three and 12 months). Thereafter, clinical and manometric assessments were performed yearly for six years. The clinical score was according to Eckardt et al. Five patients needed one (30 mm diameter Rigiflex dilator) and six needed two (30 and 35 mm diameter) dilatations. No complications occurred. All patients remained in clinical remission and their lower oesophageal sphincter pressure decreased to <10 mm Hg and remained unchanged over time.

There are similarities in the results of the two studies. One of the outcome of our 11 patients was comparable with that of the eight patients of Eckardt et al with a lower oesophageal sphincter pressure of <10 mm Hg who had a remission rate of 73% at 1 year; and (2) the observation that the six patients in our series who needed a second dilatation all relapsed within one year of the first dilatation agrees with the data by Eckardt et al, showing that most relapses occur in 12 months. However, our dilatations were more successful and, importantly, a second dilatation led to a sustained remission in all patients. We do not know the reasons for this difference but we believe it may be at least partly related to our use of the non-compliant Rigiflex dilator, which is currently considered the best choice, although there are no adequately powered comparisons with the Browne-McHardy dilator in the literature. Similarly to our result, a recent paper has shown very good efficacy of a second dilatation with the Rigiflex dilator in patients who had relapsed. Another possible reason is the use of conscious sedation during the procedure which allowed us to complete all dilatations; Eckardt et al, who used topical anaesthesia only, had to prematurely terminate 17% of the procedures.

In conclusion, our published experience and our current clinical practice, involving treatment and follow up of 10–15 new achalasia patients each year, suggest that performance of one or two dilatations until stable clinical remission is a valuable strategy, and that pneumatic dilatation under conscious sedation with the Rigiflex dilator is an effective long term treatment in most patients with achalasia.

References

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Conflicts of interest: None declared.

Authors’ reply

Penagini and Cantu should be congratulated for the remarkable results they were able to obtain in 11 patients with achalasia treated by pneumatic dilatation. To my knowledge, not a single study has so far produced similar results. A review of prospective studies in patients undergoing pneumatic dilatation with the Rigiflex dilator indicated that approximately 80% will have a good or excellent short term response. However, if such patients are observed for prolonged periods, the results obtained do not differ significantly from those observed following treatment with the older balloons. In a recent study, in which 56 patients were treated with the Rigiflex dilator and observed for more than 10 years, the long term success rate was 55%. Thus it is my impression that differences in treatment results are not so much related to differences in technique and operator experience but rather to the number of patients investigated, duration of follow up, and finally the quality of the study design. It is hoped that carefully designed randomised studies, which are now in progress, will tell us whether we should continue to offer pneumatic dilatation to the great majority of patients with achalasia or whether we should advise them to undergo surgery instead.

References


injected live LAB was 5

hypothesised that toxicity may also differ

of the gut flora, and the role of epithelial cells

local and innate immunity, the general status

this study confirms the importance of the

inflammation, attachment, and allergy models. Even if

of the TNBS-induced colitis model to evaluate the

Moreover, our study raised two important

mucosal tissue, the cause of the

biotherapeutic agents: present knowledge and

Inhibition of leukotriene synthesis markedly

disease causation as cationic

mutations in anionic trypsins in pancreatic juice. We investigated

whether mutations in the anionic trypsino-

gene, 17

inflammatory bowel disease.

Owing to the extremely high sequence

homology between PRSS1 and PRSS2, a nested polymerase chain reaction (PCR) was used to

clearly identifying the presence of

mutations in the anionic trypsinogen

found any of the reported or any novel

sequencing approach. However, we did not

were PCR amplified, purified, and sequenced

flanking region in an attempt to look for any

whether mutations in the anionic trypsino-

trypsins in pancreatic juice. We investigated

PRSS2, and PRSS3 are functional genes

and hence classified in the exons 2 and 3 only. The

mutations in the anionic trypsinogen

gene are not associated with
tropical calcific pancreatitis

Pancreatitis is considered to be an autodigestive
disease due to premature activation of

tryptsinogen inside the pancreas. Its genetic basis has recently been established with the

identification of causal mutations in cationic

trypsinogen gene (PRSS1) in patients with

hereditary1 and non-hereditary pancreatitis.7

Mutations in other genes such as SPINK1
(encoding pancreatic secretory trypsin inhibi-

and cystic fibrosis transmembrane conductance regulator (CFTR))7,8 genes have also been associated with the disease.

Tropical calcific pancreatitis is a type of

idiopathic pancreatitis, reported particularly in the tropics. Recently, we and others

have shown absence of PRSS1 mutations

but significant prevalence of the N345 mutation

in the SPINK1 gene in these patients.4-6

However, our study raised two important

questions: firstly, the exact role of SPINK1

mutations in disease causation as cationic

tryptsinogen is normal with an intact auto-

lysis site; and secondly, the cause of the
disease in the remaining patients negative for both

PRSS1 and SPINK1 mutations.

Of the nine members of the human

trypsinogen gene family, only PRSS1, PRSS2, and PRSS3 are functional genes

coding for cationic, anionic, and meso-try-

psinogens. These genes have similar isoforms, respectively. The anionic form accounts for about one third of the total

trypsins in pancreatic juice. We investigated

whether mutations in the anionic trypsino-

gene may contribute to the pathogenesis of tropical calcific pancreatitis. Mutations

reported to date in the PRSS1 gene, 17

are clustered in exons 2 and 3 only. The

remaining three are in the promoter region

but reported in isolated patients. Hence we

initially screened exons 2 and 3 of the anionic

trypsinogen gene in 68 well characterised

Indian patients with tropical calcific pancreatitis.4-6 Subsequently, we also sequenced

the promoter, complete coding region, and the

flanking region in an attempt to look for any

novel mutation.

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TNBS-induced colitis model to evaluate the anti-

inflammatory properties of lactic acid bacteria. Dig Dis


Lactobacillus bulgaricus and Streptococcus thermophilus in mice. Eur Cytokine News 1991;2:299–303.

3. Rachmilewitz D, Kataoka K, Kamali F, et al. Toll-like receptor 9 signaling mediates the anti-


Figure 1 Lactic acid bacteria (LAB) protection on macroscopic damages induced by 2,4,6-

trinitrobenzene sulphonic acid (TNBS): the impact of the oral or intraperitoneal route of LAB

administration on reduction of TNBS-induced colitis in mice. Results are expressed as per cent

reduction of mean macroscopic inflammation of mice treated with LAB, in relation to the mean

score of non-treated mice. Colitis index was assessed 48 hours after TNBS administration. Each bar

represents an independent experiment of control (n = 10) and LAB treated mice (n = 10).

OR: p<0.05,

**p<0.01,**

* p<0.001, ** p<0.001, significantly different compared with the corresponding TNBS control group (Mann-

Whitney U test). CFU, colony forming unit; d, day.

<table>
<thead>
<tr>
<th></th>
<th>Oral route: 1 x 10^8 cfu/d – 5 consecutive days prior to TNBS</th>
<th>Intraperitoneal route: 5 x 10^7 cfu – 1 day prior to TNBS</th>
<th>1 x 10^9 cfu – 2 h prior to TNBS</th>
<th>5 x 10^9 cfu/d – 6 and 1 day prior to TNBS</th>
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Acknowledgements

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Mutations in anionic trypsino-
gene are not associated with
tropical calcific pancreatitis

As emphasised by Gosh and colleagues
(Gut 2004;53:620–2), approaches involving
and homozygous states with a mutant allele frequency of 0.58 (9 AA, 20 GG, and 39 AG) and was comparable with 0.61 in 50 controls (7 AA, 18 GG, and 25 AG) analysed.

Our results thus exclude any association of mutations in the anionic trypsinogen gene in tropical calcific pancreatitis and suggest a role for other genetic or non-genetic factors in the pathogenesis of the disease. Screening of genes such as CFTB may explain the disease in the remaining patients. It also affirms the importance of the N345 mutation in SPINK1 as the major genetic factor for this type of pancreatitis.

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doi: 10.1136/gut.2004.055335

*M M Idris and S Bhaskar contributed equally to this work.

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References


Elevated plasma protein C levels correlate with the presence of fatty liver (NASH and NAFLD)

The clinical implications of non-alcoholic fatty liver disease (NAFLD) are derived mostly from its common occurrence in the general population and the potential of the condition to progress to fibrosis and cirrhosis. Markers that help in making an early diagnosis and treatment are warranted. Protein C is a vitamin K dependent glycoprotein that functions as a circulating anticoagulant through proteolytic cleavage and inactivation of the coagulation factors Va and VIII. Whether or not protein C levels increase in patients with NAFLD has not been assessed.

We measured protein C levels in 44 patients (28 men and 16 women; mean ages 45 (11) and 49 (12) years, respectively; 15 patients with fatty liver (FL), 15 with non-alcoholic steatohepatitis (NASH), and 14 with chronic viral hepatitis B+C (CH). All were diagnosed by histology and liver technetium scan or ultrasound. Ten healthy subjects were used as controls. Obesity, hyperlipidaemia, and diabetes were present in 60%, 73%, and 23% of cases, respectively; 64% of patients had elevated liver enzyme tests (alanine aminotransferase 45 (21) IU/l in FL and 43 (18) IU/l in NASH). Mean protein C levels were significantly increased in patients with NAFLD (n = 30) compared with those with chronic viral hepatitis (140 (36) % vs 101 (24); p < 0.0009 and healthy individuals (140 (36) % vs 120 (12); p < 0.04).

No significant difference in protein C levels was noted between simple fatty liver and NASH (149 (34) % and 130 (37) %, respectively; p = 0.07). A significant correlation was found between protein C and extent of fatty infiltration (r = 0.63; p < 0.001) (fig 1), insulin resistance index (r = 0.3; p < 0.01), and triglyceride levels (r = 0.45, p < 0.001). Protein C could discriminate correctly between NAFLD and chronic viral hepatitis in 82% of cases. No significant association was found between protein C and aminotransferase levels.

In conclusion, protein C was elevated in patients with NAFLD. The underlying mechanism remains unknown. Agewall et al suggested an increase in hepatic synthesis of protein C due to increased hepatic insulin resistance. Increased levels have been reported in patients with diabetes, hypertriglyceridaemia, and nephrotic syndrome, with the use of anabolic steroids, oral contraceptives, and alcohol, and with increasing age. Diabetes and hypertriglyceridaemia are predisposing conditions to fatty liver and were present in 23% and 73% of cases, respectively. The remaining conditions were excluded by clinical and biochemical findings. Although more studies are needed, these preliminary findings suggest that elevated protein C levels together with elevated liver enzymes may be used as markers for NAFLD and may obviate the need for liver biopsy.

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doi: 10.1136/gut.2004.060251

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References


Coexistent chronic idiopathic intestinal pseudo obstruction and inflammatory bowel disease

Chronic idiopathic intestinal pseudo obstruction (CIIP) is a severe condition presenting with abdominal pain and dysmotility. Inflammatory or degenerative changes of the autonomic nervous system or of the muscles of the bowel have been observed in CIIP. As patients with inflammatory bowel disease (IBD) may show clinical and histological signs of autonomic neuropathy and dysmotility, the aim of this study was to examine whether there is an association between CIIP and IBD.

Six patients at our hospital presenting with signs and symptoms of intestinal dysmotility were diagnosed with CIIP based on increasing features, antroduodenojejunal manometry, and full thickness biopsies (table 1). Patient No 1 had an acute erosive colitis some years previously with bloody diarrhoea and an enhanced sedimentation rate, which was treated with steroids, and patient No 2 had relapsing proctitis treated with 5-aminosalicylic acid (5-ASA). Patient No 3 was
totally and patient No 4 partially colectomised because of slow transit constipation. Patient No 6 was proctocolectomised due to refractory colitis. The patients were further investigated with magnetic resonance (MR) enterography\(^a\) and video capsule enteroscopy to establish whether there were any signs of IBD. If these examinations showed any pathology, push enteroscopy and ileocolonoscopy were also performed. All biopsies collected over the years were re-evaluated.

MR enterography did not reveal any pathological changes in any of the subjects. In three patients (Nos 1, 3, and 4), video capsule enteroscopy revealed Crohn-like ulcerations/erosions in the stomach and small intestine. Further examination of patient No 1 by push enteroscopy confirmed the erosions in the stomach and one third of the proximal small intestine. In patient No 3, capsule enteroscopy showed aphthous ulcers typical of Crohn’s disease throughout the distal jejunum and ileum (fig 1A). Ileocolonoscopy showed the same picture in the ileum and ileocecal anastomosis.

Histopathological examination of the full-thickness biopsies from patient Nos 1 and 2 showed visceral degenerative neuropathy, combined with vacuolisation of the interstitial cells of Cajal (ICC)s. In patient No 3, lymphoctic ganglioneuromatosis was found in both neural plexa of the resected colon and ileum (fig 1B), with signs of neurone degeneration, and 50% and 80% reduction of ICCs in the perimyenteric ICC-plexus and deep muscular plexus of the circular muscle layer, respectively. Patient No 4 had a normal biopsy, and patient No 5 was not biopsied. Examination of mucosal biopsies from patient No 1 revealed focal active inflammation in the duodenum and caecum, and chronic inflammation in the rectum; patient No 5 had multifocal mild antral cryptitis, and both patients were diagnosed with suspected Crohn’s disease. Colon biopsies from patient No 6 revealed epithelioid cell granulomas and multinucleated giant cells, as well as multifocal transmural lymphoid hyperplasia consistent with Crohn’s disease.

In three patients (Nos 1, 3, and 4), dysmotility preceded the mucosal changes. In patient Nos 2 and 5, these two entities occurred simultaneously, while in patient No 6, dysmotility developed after proctocolectomy. Ganglioneuritis in patient No 3 could have been caused by Crohn’s disease before other symptoms of the disease developed. Treatment with 5-ASA has reduced her abdominal pain. The normal histology of the sigmoidum in patient No 4 does not exclude the possibility of ganglioneuritis in other parts of the bowel due to the known patchy involvement of the gut in Crohn’s disease.

The present observations indicate that apart from inflammation, even purely degenerative neuronal and ICCs changes seen in CIP can occur in patients who also have IBD/IBD-like condition. At present, it is not known whether the observed abnormalities are part of IBD or independent of each other. This small patient sample prevents us from drawing any definite conclusion regarding this question. Further observations are needed to establish whether or not this connection is causal.

### Table 1

<table>
<thead>
<tr>
<th>Patient No:</th>
<th>Debut age/CIIP</th>
<th>Main symptoms</th>
<th>Clinical diagnosis</th>
<th>Endoscopic pathology</th>
<th>Histopathology</th>
<th>Antroduodenal manometry</th>
</tr>
</thead>
<tbody>
<tr>
<td>1 23/F</td>
<td>16/22</td>
<td>Pain, bloody diarrhoea</td>
<td>Crohn’s disease, CIIP</td>
<td>Small and large bowel</td>
<td>Degenerative neuropathy</td>
<td>Suspected Crohn’s disease</td>
</tr>
<tr>
<td>2 26/F</td>
<td>15/25</td>
<td>Pain, vomiting</td>
<td>Prolitis, CIIP</td>
<td>Rectum</td>
<td>Degenerative neuropathy</td>
<td>Abnormal</td>
</tr>
<tr>
<td>3 35/F</td>
<td>26/35</td>
<td>Constipation, dyspepsia</td>
<td>Crohn’s disease, CIIP</td>
<td>Small and large bowel</td>
<td>Ganglioneuritis</td>
<td>Normal</td>
</tr>
<tr>
<td>4 44/F</td>
<td>35/39</td>
<td>Constipation, pain</td>
<td>Suspected Crohn’s disease, CIIP</td>
<td>Small bowel</td>
<td>Normal</td>
<td>Abnormal</td>
</tr>
<tr>
<td>5 55/M</td>
<td>39/41</td>
<td>GORD, later pain and diarrhoea</td>
<td>Suspected Crohn’s disease, CIIP</td>
<td>Large bowel</td>
<td>Suspected Crohn’s disease</td>
<td>Ganglioneuritis, Crohn’s disease</td>
</tr>
<tr>
<td>6 67/M</td>
<td>61/64</td>
<td>Pain, weight loss</td>
<td>Crohn’s disease, CIIP</td>
<td>Normal</td>
<td>Normal</td>
<td>Abnormal</td>
</tr>
</tbody>
</table>

CIIP, Chronic idiopathic intestinal pseudo-obstruction; GORD, gastro-oesophageal reflux disease.

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**References**


We read the article by Allen and colleagues (Gut 2004;53:1566–70) with interest and would like to report a case of probable cannabinoid hyperemesis seen in a district general hospital in the UK. A 21 year old chef was admitted to our hospital on seven occasions over a two year period (April 2001 to December 2002) with profuse vomiting. Apart from a history of migraine as a child, he was fit and well. He smoked cannabis. Physical examination was unremarkable. The observation that the patient wanted to take regular baths because he had found that bathing eased the sickness was documented in the nursing notes but its significance was not appreciated. Investigations during attacks disclosed neutrophilia but blood urea, electrolytes, liver biochemistry, and serum amylase were normal. Abdominal x ray was also normal. Upper gastrointestinal endoscopy showed grade i oesophagitis and gastritis. Gastric biopsies were histologically normal. An abdominal ultrasound scan and small bowel barium follow through examination were normal. Additional normal or negative investigations included: autoantibodies and immunoglobulins, C reactive protein, and urinary porphyrin screen. Computed tomography scan of the brain was also normal.

During his last admission, the patient’s girlfriend showed us an article published in an Australian newsletter which she had obtained via the internet, in which Dr JH Allen had raised the possibility of a link between recurrent vomiting and cannabis abuse. With the aid of the internet we traced an Australian newsletter which she had included: autoantibodies and immunoglobulins, C reactive protein, and urinary porphyrin screen. Computed tomography scan of the brain was also normal.

We hypothesised that as a result of chronic inflammatory conditions in the large intestine, increased colonic iNOS activity may produce an excess of NO, nitrogen oxides, and nitrite, which in turn react with NOC present in the colon to produce relatively higher levels of NOC. Increased formation of NOC in IBD patients may thus contribute to the relatively high incidence of colorectal cancer associated with this disease.

A recent population-based case control study showed that in cases with a history of IBD, increased exposure to drinking water nitrate was associated with an increased risk of colon cancer whereas no such association was found in the overall population. We hypothesised that as a result of chronic inflammatory conditions in the large intestine, increased colonic iNOS activity may produce an excess of NO, nitrogen oxides, and nitrite, which in turn react with NOC present in the colon to produce relatively higher levels of NOC. Increased formation of NOC in IBD patients may thus contribute to the relatively high incidence of colorectal cancer associated with this disease.

Most research on endogenous NOC exposure has focused predominantly on the intragastric formation of these compounds in relation to the gastric cancer risk. However, we now report that faecal NOC levels in IBD patients are considerably higher than those we reported previously in gastric juice (0.25 (0.3) ng/g), which indicates that NOC exposure may be even more relevant in colon carcinogenesis. Based on these results, we conclude that the colon of IBD patients is exposed to relatively high concentrations of this carcinogenic compound, probably as a direct consequence of continuous NO production by the inflammatory process. As this exposure may strongly contribute to the increased colon cancer risk associated with IBD, dietary recommendations for IBD patients, avoiding high NOC intake, seem warranted.

**Table 1 Faecal N-nitrosodimethylamine (NDMA) concentrations in patients with inflammatory bowel disease (IBD) and in healthy controls**

<table>
<thead>
<tr>
<th></th>
<th>Controls (n = 17)</th>
<th>All IBD cases (n = 17)</th>
<th>Non-hospitalised cases (n = 10)</th>
<th>Hospitalised cases (n = 7)</th>
</tr>
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<tbody>
<tr>
<td>NDMA positive</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>NDMA (ng/g)</td>
<td>35</td>
<td>41</td>
<td>56</td>
<td>25</td>
</tr>
<tr>
<td>NDMA positive</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>NDMA (ng/g)*</td>
<td>1.4</td>
<td>10.9†</td>
<td>14.3†</td>
<td>2.4‡</td>
</tr>
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</table>

*Average concentration of NDMA positive samples. tp<0.05, t†p<0.01: significantly higher compared with the control group (Mann-Whitney U test). tp<0.05: significantly lower compared with non-hospitalised cases (Mann-Whitney U test).

Conflict of interest: None declared.

**References**


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Conflict of interest: None declared.

**Inflammatory bowel disease stimulates formation of carcinogenic N-nitroso compounds**

In patients with inflammatory bowel disease (IBD) the clinical presentation of colorectal cancer is observed. Although severe inflammatory conditions per se represent a risk factor for neoplasia, we would like to draw attention to the possible role of increased activity of inducible nitric oxide synthase (iNOS), as found in IBD patients, in the endogenous formation of carcinogenic N-nitroso compounds (NOC). In healthy individuals, relatively small amounts of NOC are formed by the interaction between NOC precursors (NOCP), present in dietary items such as meat and fish, and nitrosating agents derived from dietary nitrate. It has been proposed that endogenous formation of NOC may explain the link between meat consumption and colon cancer risk found in epidemiological studies. We hypothesised that as a result of chronic inflammatory conditions in the large intestine, increased colonic iNOS activity may produce an excess of NO, nitrogen oxides, and nitrite, which in turn react with NOC present in the colon to produce relatively higher levels of NOC. Increased formation of NOC in IBD patients may thus contribute to the relatively high incidence of colorectal cancer associated with this disease.

A recent population-based case control study showed that in cases with a history of IBD, increased exposure to drinking water nitrate was associated with an increased risk of colon cancer whereas no such association was found in the overall population. This clearly indicates that the risk of colon cancer in IBD patients is not only determined by the disease itself but dietary factors known to influence the endogenous formation of NOC are also associated with an increased risk in these patients. Although both the increased formation of NOC found in mice with chemically induced colitis and increased levels of NO and nitrite found in the colonic lumen of patients with ulcerative colitis support this hypothesis, the relatively small amounts of NOC levels have never been investigated in IBD patients.

Therefore, we collected faecal samples from 17 patients diagnosed with ulcerative colitis and 17 healthy controls, and determined levels of N-nitrosodimethylamine (NDMA), a predominant carcinogenic NOC, using gas chromatography-mass spectrometry, as previously described. The study was approved by the medical ethics committee of the Maasland Hospital, Sittard, the Netherlands, and all patients gave their consent. In 41% of patients, we found levels of NDMA above the detection limit of 1 ng/g, compared with 35% of controls. Comparison of concentrations in NDMA positive samples showed that the average concentration in patients was significantly higher than that in the control group (table 1). When IBD patients were subdivided into hospitalised and non-hospitalised cases, the difference between the non-hospitalised group and controls was even more pronounced, whereas NDMA concentrations in hospitalised patients and controls were similar. All hospitalised patients received only liquid nutrition (Nutrison; Nutricia, UK) without additional intake of NOC rich dietary foods, these results confirm that the combination of high dietary NOC intake and inflammation may present a risk factor.

Most research on endogenous NOC exposure has focused predominantly on the intragastric formation of these compounds in relation to the gastric cancer risk. However, we now report that faecal NOC levels in IBD patients are considerably higher than those we reported previously in gastric juice (0.25 (0.3) ng/g), which indicates that NOC exposure may be even more relevant in colon carcinogenesis. Based on these results, we conclude that the colon of IBD patients is exposed to relatively high concentrations of this carcinogenic compound, probably as a direct consequence of continuous NO production by the inflammatory process. As this exposure may strongly contribute to the increased colon cancer risk associated with IBD, dietary recommendations for IBD patients, avoiding high NOC intake, seem warranted.
Hepatocellular carcinoma occurring after successful treatment of childhood cancer with high dose chemotherapy and radiation

Hepatocellular carcinoma (HCC) is one of the world’s most common malignancies and accounts for more than 90% of all primary liver cancers. A number of different risk factors have been identified for the development of HCC. Hepatitis B carrier state, environmental toxins, chronic hepatitis C virus infection, hereditary haemochromatosis, and liver cirrhosis of almost any cause are well known risk factors for HCC. In addition, environmental toxins such as aflatoxins and contaminated drinking water may contribute to the pathogenesis of HCC, especially in Asia and underdeveloped countries. Finally, a number of HCC cases have occurred after the use of thorotrast for diagnostic procedures, and survivors of the atomic bomb of Hiroshima were also at higher risk for HCC development,

indicating that radiation might also induce the development of HCC. Herein we describe a rare case of HCC occurring in a patient 17 years after successful treatment of peripheral neuroectodermal tumour (PNET).

A 32 year old female presented with pain in the right upper quadrant of her abdomen. Seventeen years prior to presentation in our hospital this patient was treated for a PNET with a combination of high dose chemotherapy (vincristine, adriablastin, ifosfamide, and actinomycin D) and surgical removal of the tumour. There were no signs of any recurrence of the tumour observed on her last check up 12 month earlier. Physical examination of the patient in our clinic showed typical signs of late radiation damage (erythema of the skin and an underdeveloped right breast) (fig 1). A firm 3–5 cm mass was palpable at the lower edge of the liver. Laboratory tests showed elevated α-fetoprotein (41881 μg/l). Hepatitis serology was negative and there was no evidence of any other liver disease. Magnetic resonance imaging revealed multiple intrahepatic masses up to 6.5 cm. A biopsy from the hepatic tumour was taken and confirmed the clinical diagnosis of HCC. The patient died three months after the diagnosis was made.

To the best of our knowledge, secondary HCC following high dose chemotherapy has never been described and therefore we searched the German Childhood Cancer Registry, which started to register all cases of malignancies in children (<15 years) in 1980. This database also collects data from secondary malignancies following chemotherapy. In this database we were able to detect a total of four more cases of secondary HCC, which are summarised in table 1. Interestingly one patient was hepatitis B surface antigen positive.

Radiotherapy has been shown to be associated with an increased risk of solid tumours 10–15 years after treatment and later.

There is one report in the literature of a radiation induced hepatoma in a patient with a non malignant haemangioendothelioma, which occurred 20 years after radiation of the liver with 28.5 Gy. To date, the molecular mechanism of hepatocarcinogenesis is not completely understood. The main causative agents—hepatitis B virus, hepatitis C virus, and aflatoxin B1—have been identified, which together are responsible for approximately 80% of all HCC in humans. This series of cases clearly supports the notion that secondary HCC can follow not only radiation therapy of children but also high dose chemotherapy, and may prompt careful follow up examinations of the liver in patients with a possible risk for the development of HCC.

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<table>
<thead>
<tr>
<th>First malignancy</th>
<th>Age (y)</th>
<th>Treatment</th>
<th>Age when HCC was diagnosed (y)</th>
<th>Time from first to second malignancy (y)</th>
</tr>
</thead>
<tbody>
<tr>
<td>ALL</td>
<td>4</td>
<td>CTx*</td>
<td>20</td>
<td>15</td>
</tr>
<tr>
<td>Osteosarcoma</td>
<td>4 na</td>
<td>RTx</td>
<td>18</td>
<td>9</td>
</tr>
<tr>
<td>ALL</td>
<td>4 na</td>
<td></td>
<td>10</td>
<td>5</td>
</tr>
<tr>
<td>PNET</td>
<td>15</td>
<td>CTx, RTx</td>
<td>33</td>
<td>18</td>
</tr>
<tr>
<td>Teratoma</td>
<td>2 na</td>
<td></td>
<td>19</td>
<td>16</td>
</tr>
</tbody>
</table>

ALL, acute lymphocytic leukaemia; PNET, peripheral neuroectodermal tumour; CTx, chemotherapy; RTx, radiation therapy.

This patient was hepatitis B surface antigen positive.

References


Biologics in inflammatory disease: infliximab associated risk of lymphoma development

In their excellent overview of currently available biologic compounds that are in use or under investigation for Crohn’s disease (CD), Sandborn and Faubion (Gut 2004;53:1366–73) reconfirm the unique standing of infliximab. They also note the ongoing discussion concerning the increased occurrence of lymphoproliferative disorders in patients who received infliximab.

Recently, we followed a 61 year old patient with a 31 year history of relapsing CD. Initial treatment was with steroids but after 10 years with infliximab. Relapse occurred two decades after hepatic irradiation.

...
However, tumour cells were negative for CD30 and ALK protein. Molecular analysis demonstrated a monoclonal immunoglobulin heavy chain rearrangement in the absence of a T cell receptor rearrangement, confirming the diagnosis. The tumour was neither responsive to CHOP-Rituximab nor to the ensuing second and third line chemotherapies. When the patient presented for fourth line chemotherapy, spontaneous partial remission was seen, persisting now for 10 months up to the last clinical follow up.

As mentioned by Sandborn and Faubion, the 33 published cases (1) of lymphomas following infliximab therapy raise the question of a contributory role of infliximab in the propagation of lymphoproliferative disorders. We now add a novel case of a metachronous duplex non-Hodgkin lymphoma of initially T and then B cell phenotype. Imbalanced function of T lymphocytes may have acted as a key feature in this patient as the development of CD and the EBV related B cell non-Hodgkin lymphoma of 6.6/100 000 treated patients was estimated in 2002, which approximately 500 000 patients have been treated.

Table 1 Patients with infliximab therapy and development of lymphoma

<table>
<thead>
<tr>
<th>Case</th>
<th>Age (y), sex, disease</th>
<th>Dose</th>
<th>No of infusions</th>
<th>Lymphoma</th>
<th>EBV</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>77 M, NR</td>
<td>NR</td>
<td>10</td>
<td>Burkitt lymphoma</td>
<td>NR</td>
</tr>
<tr>
<td>2</td>
<td>NR</td>
<td>NR</td>
<td>10</td>
<td>Hodgkin lymphoma</td>
<td>NR</td>
</tr>
<tr>
<td>3</td>
<td>43 F, NR</td>
<td>NR</td>
<td>10</td>
<td>Hodgkin lymphoma</td>
<td>NR</td>
</tr>
<tr>
<td>4</td>
<td>34 M, NR</td>
<td>NR</td>
<td>10</td>
<td>DLBCL</td>
<td>Positive</td>
</tr>
<tr>
<td>5</td>
<td>70 M, NR</td>
<td>NR</td>
<td>10</td>
<td>DLBCL</td>
<td>Positive</td>
</tr>
<tr>
<td>6</td>
<td>29 M, CD 5 mg/kg, 3</td>
<td>NR</td>
<td>10</td>
<td>Hodgkin lymphoma</td>
<td>NR</td>
</tr>
<tr>
<td>7</td>
<td>68 F, NR</td>
<td>NR</td>
<td>10</td>
<td>B cell NHL</td>
<td>NR</td>
</tr>
<tr>
<td>8</td>
<td>62 M, NR</td>
<td>NR</td>
<td>10</td>
<td>DLBCL</td>
<td>NR</td>
</tr>
<tr>
<td>9</td>
<td>73 M, NR</td>
<td>NR, multiple</td>
<td>10</td>
<td>Manet cell lymphoma</td>
<td>NR</td>
</tr>
<tr>
<td>10</td>
<td>74 F, RA 10 mg/kg, 8</td>
<td>NR</td>
<td>10</td>
<td>B cell NHL</td>
<td>NR</td>
</tr>
<tr>
<td>11</td>
<td>48 M, RA 10 mg/kg, 2</td>
<td>NR</td>
<td>10</td>
<td>B cell NHL</td>
<td>NR</td>
</tr>
<tr>
<td>12</td>
<td>59 F, RA 3 mg/kg, 5</td>
<td>NR</td>
<td>10</td>
<td>Multiple myeloma</td>
<td>NR</td>
</tr>
<tr>
<td>13</td>
<td>61 F, RA 1 mg/kg, 1</td>
<td>NR</td>
<td>10</td>
<td>Hodgkin lymphoma</td>
<td>NR</td>
</tr>
<tr>
<td>14</td>
<td>36 M, CD, HIV 10 mg/kg</td>
<td>NR</td>
<td>10</td>
<td>B cell NHL</td>
<td>NR</td>
</tr>
<tr>
<td>15</td>
<td>62 M, CD 10 mg/kg, 1</td>
<td>NR</td>
<td>10</td>
<td>Intravascular B-NHL</td>
<td>NR</td>
</tr>
<tr>
<td>16</td>
<td>48 F, DM 5 mg/kg, 3</td>
<td>NR</td>
<td>10</td>
<td>DLBCL</td>
<td>NR</td>
</tr>
<tr>
<td>17</td>
<td>47 M, 6 mg/kg, 3</td>
<td>NR</td>
<td>10</td>
<td>CD30+ T-cell lymphoma</td>
<td>Negative</td>
</tr>
</tbody>
</table>

Psoriasis
18 70 M, CD 5 mg/kg, 3 Folicular lymphoma | NR |
19 51 M, CD 5 mg/kg, 4 Hodgkin lymphoma | NR |
20 25 M, CD 5 mg/kg, 1 NK cell lymphoma | NR |
21 79 M, CD 5 mg/kg, 1 B cell NHL | NR |
22 24 F, CD 5 mg/kg, 5 Multiple myeloma | NR |
23 NR M, RA 5 mg/kg, 2 Mixed cell NHL | NR |
24 NR M, RA 5 mg/kg, 2 Mixed cell NHL | NR |
25 NR M, RA 5 mg/kg, 2 B cell NHL | NR |
26 NR M, RA 5 mg/kg, 2 B cell NHL | NR |
27 NR M, RA 5 mg/kg, 2 B cell NHL | NR |
28 NR M, RA 5 mg/kg, 2 B cell NHL | NR |
29 NR M, RA 5 mg/kg, 2 B cell NHL | NR |
30 NR M, RA 5 mg/kg, 2 B cell NHL | NR |
31 NR M, RA 5 mg/kg, 2 B cell NHL | NR |
32 NR M, CD 5 mg/kg, 1 NK cell lymphoma | NR |
33 NR M, CD 5 mg/kg, 1 NK cell lymphoma | NR |
34 61 M, CD 10 mg/kg, 3 Metachronous lymphoma (ALCL, DLBCL) | Positive |


Genotypes 677TT and 677CT+1298AC of methylenetetrahydrofolate reductase are associated with the severity of ulcerative colitis in central China

Increased blood levels of homocysteine have been found to be associated with inflammatory bowel disease (IBD) in several studies. The main genetic determinant associated with elevated plasma levels of homocysteine (t-Hcy) is the MTHFR 677C→T gene polymorphism of methylenetetrahydrofolate reductase, a critical enzyme involved in the remethylation pathway of homocysteine. An association of the MTHFR 677T allele with IBD has been reported in Northern Europe but not in three series from Italy and France. Double heterozygosity MTHFR 677CT+1298AC also produces reduced enzyme activity and increased t-Hcys, but its association with IBD has never been studied. Similarly, the association of IBD with transcobalamin (TCN1 776C→G), a genetic determinant that influences transcobalamin levels and t-Hcys, is not known. Transcobalamin is the protein that promotes intestinal transcytosis and cell delivery of vitamin B12, the cofactor of the methionine synthase dependent remethylation pathway.

In this study, we have evaluated the association of ulcerative colitis (UC) with MTHFR 677C→T, MTHFR 1298A→C, and TCN1 776C→G in a series of 72 patients from central China who gave informed consent. This series was compared with 111 age and sex matched controls. The research protocol was approved by the local appointed committee. Extraction of DNA and determination of polymorphisms were performed as described previously by us. A continuity corrected χ² test and an ANOVA test were used, respectively, to assess differences in categorical and continuous variables between groups. Odds ratios of independent categorical variables.
that differed significantly between patients and controls were determined by logistic regression analysis. A p value <0.05 was considered to indicate statistical significance.

The main clinical characteristics are summarised in table 1. Most of the cases were recently diagnosed. None had any thrombotic manifestations. TCNI 776G allele frequency was approximately 1.5-fold higher compared with Caucasians, and we failed to find any association with the risk of UC or severity of disease. MTHFR 677T allele frequency in our control group was close to that reported in the US, whereas in China the frequency of the MTHFR 677T allele was approximately 1.5-fold higher compared with the coast of West Africa.

In conclusion, our study showed that the genotypes of MTHFR, associated with a decrease in enzyme activity, seemed to be more significantly associated with extension of disease than with the primary risk, at least in central China.

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References

CORRECTION

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The original article by Cosnes et al (Impact of the increasing use of immunosuppressants in Crohn’s disease on the need for intestinal surgery. Gut 2005;54:237–41), published in the February 2005 issue was incomplete. figure 2 was missing from the proof. A corrected version of the pdf can be viewed at http://gut.bmjournals.com/cgi/data/54/2/237/DC1/1, and the missing figure can be seen here.