Increased risk of left sided colon cancer in patients with diverticular disease

T Stefánsson, A Ekborn, P Sparén, L Pählman

Abstract
Certain similar epidemiological characteristics suggest a common aetiology for colon cancer and diverticulosis of the colon. The hypothesis that patients with diverticulosis are at increased risk of developing colon cancer was tested in a retrospective, population based, cohort study in Sweden. A total of 7159 patients (2478 men and 4681 women) who had been given a hospital discharge diagnosis of diverticulosis or diverticulitis of the colon between 1965 and 1983 were followed up during 1985 by means of record linkage procedures. After excluding the first 2 years of follow up, there was not a significant increase in risk (SIR) overall for colon cancer (SIR=1.2; 95% confidence intervals (CI) 0.9, 1.6) or for rectal cancer (SIR=1.1; 95% CI 0.7, 1.7). The observed number of right sided colon cancers was as expected (SIR=0.9; 95% CI 0.5, 1.5). In contrast, an increased risk of left sided colon cancer was found both overall (SIR=1.8; 95% CI 1.1, 2.7) and consistently in men and women as well as in different age groups. This risk increased the longer the follow up (p value for trend <0.001). These results do not support the hypothesis of a common aetiology in diverticular disease and colonic cancer but suggest a causal relationship between diverticular disease and cancer of the left colon.

Subjects and methods
COHORT
The Uppsala health care region, covering six counties, is located in central Sweden and had, during the study period, a population of 1.2–1.3 million people. Since there is almost no private inpatient treatment in Sweden, hospital provided medical services are, in effect, population based and referable to the county in which the patient lives. From 1965 until 1983, the Swedish National Board of Health and Welfare received annual reports from all inpatient medical institutions in Sweden and recorded data on individual hospital admissions and discharges in the inpatient register for all inhabitants within the Uppsala health care region. Besides a national registration number, a unique personal identifier assigned to all Swedish citizens, each record contains data on place of residence, hospital department, surgical procedures, and up to eight discharge diagnoses. These diagnoses were coded according to the seventh revision of the International Classification of Diseases until 1968 and according to the eighth revision thereafter. A recent publication estimated that the overall extent of under-reporting to the inpatient register was less than 2%. Severe under-reporting occurred in certain counties during a limited period, representing a few % only of the estimated total number of hospital admissions.16

All patients with records in the inpatient register containing a diagnostic code for diverticulosis (ICD 7 code 572-12 and ICD 8 code 562-10) or diverticulitis (ICD 7 code 572-11 and ICD 8 code 562-11) of the colon were considered for inclusion in the study. The national registration number allowed us to select the first recorded discharge with this diagnosis for each individual. A total of 7630 individuals had been given a discharge diagnosis of diverticulosis or diverticulitis at least once between 1965 and 1983 and were potentially eligible. We excluded 128 of these because they had been entered in the inpatient register with an incomplete or inconsistent national registration number and were not available for follow up. Through the National Census Bureau and the emigration register we were able to confirm that the remaining members of the cohort were alive and living in Sweden on December 31, 1985. A total of 343 cases were not found in any of these registers, probably because they had been entered in the inpatient register with an erroneous national registration number and they were excluded. The number of patients available to follow up was thus 7159–2478 men and 4681 women. At first discharge, 1410 patients were under the age of 60 years, 1665 were 60 to 69 years, and 4084 were 70 years or older.
TABLE I  Observed number of cases (OBS) and risk of developing colorectal cancer expressed as the standard incidence ratio (SIR), with 95% confidence intervals in parentheses, according to the localisation of the cancer and gender. The first 2 years of follow up were excluded.

<table>
<thead>
<tr>
<th>Site</th>
<th>All OBS</th>
<th>Men OBS</th>
<th>Women OBS</th>
<th>All SIR</th>
<th>Men SIR</th>
<th>Women SIR</th>
</tr>
</thead>
<tbody>
<tr>
<td>All colon*</td>
<td>43</td>
<td>1·2 (0·9-1·6)</td>
<td>17</td>
<td>1·6 (0·9-2·5)</td>
<td>26</td>
<td>1·1 (0·7-1·6)</td>
</tr>
<tr>
<td>Right + transverse colon</td>
<td>16</td>
<td>0·1 (0·9-0·5)</td>
<td>6</td>
<td>1·1 (0·4-3·5)</td>
<td>10</td>
<td>0·8 (0·4-1·5)</td>
</tr>
<tr>
<td>Descending + sigmoid colon</td>
<td>21</td>
<td>1·8 (1·1-2·7)</td>
<td>9</td>
<td>2·2 (1·0-4·2)</td>
<td>12</td>
<td>1·6 (0·8-2·7)</td>
</tr>
<tr>
<td>Rectum</td>
<td>22</td>
<td>1·1 (0·7-1·7)</td>
<td>9</td>
<td>1·2 (0·6-2·3)</td>
<td>13</td>
<td>1·1 (0·6-1·8)</td>
</tr>
</tbody>
</table>

* Including multiple and unspecified sites.

FOLLOW UP

Record linkage (based on the national registration number) to the nationwide register of causes of death to obtain information on the date of death among those who had died during 1985. The National Swedish Cancer Registry, founded in 1958, was used to ascertain all incident cancers diagnosed in the cohort from start of follow up until the end of 1985. The time of observation was calculated from the date of the first discharge with diverticulosis or diverticulitis of the colon until the occurrence of a diagnosis of colorectal cancer, death, or the end of the observation period (December 31, 1985). For additional analysis we also used the inpatient register to search for those patients within our cohort who had undergone resection of the sigmoid colon or a left sided hemicolectomy during follow up through 1983. A total of 507 patients out of the 7159 (7·1%) had been operated on. In this category of patients, censoring was done for the occurrence of the cancer at the left colon at the date of surgery.

The expected number of cancers was calculated by multiplying the number of person years for each sex by age specific cancer incidence rates for each 5 year age group and calendar year of observation. These expected rates were derived from the study population, that is the Uppsala health care region. For the main analysis, we used a 2 year latency period between first discharge and calculation of observed and expected numbers of cancers. The reason for this approach was to eliminate or to reduce the possible impact of selection bias. Such a bias would occur in patients in whom the symptoms of the cancer were the reason for admission to hospital and who were given the discharge diagnosis of diverticulosis or diverticulitis.

TABLE II  Observed number of cases (OBS) and risk of developing colorectal cancer expressed as the standard incidence ratio (SIR), according to the localisation of the cancer and age at first discharge. The first 2 years of follow up were excluded. 95% confidence limits in parenthesis.

<table>
<thead>
<tr>
<th>Age at diagnosis of diverticulitis disease</th>
<th>SIR OBS</th>
<th>Men OBS</th>
<th>Women OBS</th>
<th>SIR OBS</th>
<th>Men OBS</th>
<th>Women OBS</th>
</tr>
</thead>
<tbody>
<tr>
<td>&lt;60 y</td>
<td>3</td>
<td>0·8 (0·2-2·3)</td>
<td>10</td>
<td>1·0 (0·5-1·8)</td>
<td>20</td>
<td>1·3 (0·9-1·9)</td>
</tr>
<tr>
<td>60-69 y</td>
<td>1</td>
<td>0·4 (0·1-1·8)</td>
<td>3</td>
<td>0·6 (0·3-3·6)</td>
<td>12</td>
<td>1·7 (0·9-3·0)</td>
</tr>
<tr>
<td>&gt;=70 y</td>
<td>3</td>
<td>1·2 (0·3-3·6)</td>
<td>4</td>
<td>0·7 (0·2-1·8)</td>
<td>8</td>
<td>1·3 (0·7-2·2)</td>
</tr>
</tbody>
</table>

* Including multiple and unspecified sites.

TABLE III  Observed number of cases (OBS) and risk of developing colorectal cancer expressed as the standard incidence ratio (SIR), according to the localisation of the cancer and duration of follow up from the first discharge after diverticulitis disease. 95% confidence limits in parenthesis.

<table>
<thead>
<tr>
<th>Follow up</th>
<th>All OBS</th>
<th>Men OBS</th>
<th>Women OBS</th>
<th>All SIR</th>
<th>Men SIR</th>
<th>Women SIR</th>
</tr>
</thead>
<tbody>
<tr>
<td>&lt;2 y</td>
<td>75</td>
<td>5·8 (4·6-7·3)</td>
<td>18</td>
<td>1·2 (0·7-1·9)</td>
<td>19</td>
<td>1·4 (0·8-2·2)</td>
</tr>
<tr>
<td>2-4 y</td>
<td>20</td>
<td>3·0 (1·9-4·7)</td>
<td>7</td>
<td>0·9 (0·4-1·9)</td>
<td>8</td>
<td>1·1 (0·5-2·2)</td>
</tr>
<tr>
<td>5-9 y</td>
<td>19</td>
<td>2·7 (1·4-4·2)</td>
<td>9</td>
<td>1·1 (0·5-2·1)</td>
<td>8</td>
<td>1·0 (0·5-2·1)</td>
</tr>
<tr>
<td>&gt;=10 y</td>
<td>19</td>
<td>2·7 (1·4-4·2)</td>
<td>9</td>
<td>1·1 (0·5-2·1)</td>
<td>8</td>
<td>1·0 (0·5-2·1)</td>
</tr>
</tbody>
</table>

* Including multiple and unspecified sites.
** Trend tests with the $\chi^2$ method, * after excluding the first 2 years, $\chi^2=13·7$ and $p<0·001$.

STATISTICAL METHODS

The standardised incidence ratio (SIR) was defined as the ratio of observed numbers of cancers to those expected. The 95% confidence interval (CI) of the standardised incidence ratio was then calculated on the assumption that the observed number follows a Poisson distribution. In order to test the existence of a trend the $\chi^2$ method was used.17

Results

The cohort with 7159 patients was followed up for 2-20 years, which generated a total of 42 615 person years at risk, with a mean duration of 5·95 years of follow up.

When the first 2 years after first discharge were included, a total of 118 patients were diagnosed with colon cancer, compared with the expected number of 49·3 for an SIR of 2·4 (95% CI 2·0, 2·9). During the same period there were 41 cases of rectal cancer, compared with the expected number of 26·5 (SIR=1·6; 95% CI 1·1, 2·1). The increased risk of both colon and rectal cancers was almost entirely caused by an excess number of cases during the early follow up period.

When the first 2 years after discharge were excluded, the relative risk of colon cancer was only slightly increased (SIR=1·2 95% CI 0·9, 1·6). There was a significantly increased risk of cancer of the left colon (descending and sigmoid colon) (SIR=1·8; 95% CI 1·1, 2·7) whereas the observed number of cancers in other parts of the colon was close to the expected number (SIR=0·9; 95% CI 0·5, 1·4). A non-significant increase in relative risk was found for rectal cancer (SIR=1·2 95% CI 0·7, 1·7) (Table I).

The relative risk for cancer of the colon and rectum was similar in men and women (Table I), and did not seem to be related to age at first discharge (Table II). The possibility that the excess risk of colon cancer, notably left sided, was unevenly distributed by years of follow up was analysed in Table III. A considerably increased relative risk within 2 years of follow up was found for left sided colon cancer and a more
modest increase for cancer of the right colon and of the rectum. After more than 2 years of observation, the excess risk of cancer of the descending and sigmoid colon persisted throughout the observation period and increased significantly (p for trend <0·001) the longer the follow up (Table III).

Discussion
We found a significant, approximately twofold increase in the relative risk of left sided colon cancer in patients with diverticulosis or diverticulitis. Two years or more after the first hospital discharge, this increased risk occurred consistently in men and women, at different ages, and it persisted more than 10 years after first discharge. In contrast, we found no evidence of increased risk of right sided colon cancer and only a small, non-significant excess of rectal cancers.

The validity of the diagnosis was assessed in the medical records of 537 consecutive patients given a discharge diagnosis of diverticulosis or diverticulitis at the Uppsala University Hospital 1969–1984 (Stefansson, unpublished data). In 486 (91·5%) of the patients the diagnosis had been confirmed by a barium enema, by operation, or both; x ray was negative in 29 (5·%), and the diagnostic method was clinical examination alone in 22 (4%) patients. These data indicate that the specificity of a discharge diagnosis of diverticulosis or diverticulitis is high. It is likely, but unproved, that results from one large hospital are representative for the entire health care region. The strength of the association shown in this study would be marginally underestimated if some 10% of the cohort members did not have diverticulosis or diverticulitis or any other disease which entails an increased risk of left sided colon cancer. Closer surveillance of patients with diverticulosis or diverticulitis in the colon may have lead to increased discovery of cancer of the left colon in our cohort compared with that in the general population. However, if such a bias exists, it should affect the results for rectal cancer most and no excess risk was found for that site. Moreover, closer surveillance might also be protective if premalignant lesions, notably polyps, are detected and removed.

Because the registry started in 1965, a number of patients in the cohort might have had an unnoticed discharge before that date. Thus, this series contains a mixture of incident and prevalent cases of patients with diverticular disease. The prevalent cases are likely to have had their disease longer than incident cases. If patients with diverticular disease have an increased risk for developing left sided colon cancer, the prevalent cases should have higher risk of developing left sided colon cancer than the incident cases. But as the group of prevalent cases is small and the increase in cancer risk is not more than about twice the normal risk, this group will not influence noticeably the results in Table III.

The absence of an increase in the relative risk of cancer in the right colon and rectum 2 years or more after the first discharge contradicts the hypothesis that diverticulosis of the colon and colorectal cancer in general have aetiological factors in common, as proposed in the early 1970s.18 19 This absence of an association also agrees with two previous studies. In one case-control study, 119 patients with colon cancer were compared with hospital controls who had had a barium enema because of gastrointestinal symptoms but in whom no carcinoma was diagnosed. There was no difference in the prevalence of diverticulosis of the colon between cases and controls.20 In another case-control study, 150 patients with diverticulosis of the colon and abdominal symptoms were compared with controls who had abdominal symptoms but no disease.21 22 Two groups underwent colonoscopy and the risk of cancer was found to be the same in both groups.

The significant trend of an increased risk of cancer of the left colon the longer the follow up suggests that the association between diverticular disease and cancer of the left colon may be causal. Different, possible biological explanations of such an association exist. The most common site for diverticular disease is in the sigmoid colon.23 Likewise the concentration of bacteria is high in the left colon and some types are known to produce carcinogens or cocarcinogens through degradation of biliary steroids in the faeces.24 25 Carcinogens may be trapped in the diverticula and thereby establish prolonged contact with the colonic mucosa. According to another suggestion, the DNA in the mucosal cell nuclei becomes more susceptible to damage and malignant transformation when the turnover of cells increases because of chronic inflammation.26 This could be the case in patients with diverticulitis of the sigmoid colon as in patients with ulcerative colitis,27 Crohn’s disease,28 and those patients with gastric helicobacter pylori infection of the stomach.29

We conclude that the increasing relative risk for cancer of the left colon in patients with diverticular disease may reflect a causal relationship. In clinical practice this means that malignant disease should be excluded if a patient who has had diverticular disease has a relapse of symptoms. This excess risk is, however, not great enough to justify any screening procedures for early detection of cancer in patients with known diverticulosis of a history of diverticulitis of the colon.

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