Excessive alcohol consumption favours high risk polyp or colorectal cancer occurrence among patients with adenomas: a case control study

M Bardou, S Montembault, V Giraud, A Balian, E Borotto, C Houdayer, F Capron, J-C Chaput, S Naveau

Background and aims: Excessive alcohol consumption is a risk factor for developing colorectal adenomas. This study aimed to investigate the influence of excessive alcohol consumption on the occurrence of high risk polyps (adenoma ≥10 mm, villous component, high grade dysplasia) or colorectal cancer among patients with at least one colonic adenoma.

Patients and methods: Three groups of patients with at least one colorectal adenoma were included in a case control study: 401 heavy drinkers (group HD, mean daily alcohol intake 117 (SD 4) g/day for a mean duration of 22 (SD 0.6) years), aged 57 (0.5) years (78% men); 152 patients suffering from irritable bowel syndrome (IBS), aged 61 (0.9) years (57% male); and 108 patients with a family history (FH) of colorectal adenoma or cancer, aged 55 (1) years (64% male). Exclusion criteria were: anaemia, haematochezia, personal history of colorectal adenoma or cancer, and for groups HD and IBS a family history of colorectal adenoma and/or cancer. Relative risks were estimated by the odds ratio (OR) using a logistic regression model and were expressed with 95% confidence interval (CI).

Results: After age and sex adjustment, the likelihood of having an adenoma ≥10 mm was higher in group HD than in the IBS group (OR 1.8, 95% CI (1.2–2.7)) and the likelihood of having high risk adenomas or cancer was higher in group HD compared with the IBS group (OR 1.6, 95% CI (1.2–2.1)) and the FH group although this was not significant (OR 1.6, 95% CI (0.97–2.6) (p=0.081); 90% CI (1.03–2.4)). After age and sex adjustment, the likelihood of having an adenoma with high grade dysplasia or cancer was higher in group HD than in the IBS group (OR 1.7, 95% CI (1.02–2.8)) or group FH, although this was not significant (OR 3.7, 95% CI (0.98–15) (p=0.076); 90% CI (1.10–12.47)).

Conclusion: In patients with at least one colorectal adenoma, excessive alcohol consumption increases the likelihood of developing high risk adenomas or colorectal cancer.

Material and methods

Patients

All patients older than 40 years who were referred to our unit between January 1987 and December 1996 for excessive alcohol consumption (heavy drinkers (group HD) >50 g of alcohol/day for the year before admission) underwent screening colonoscopy if they had no contraindications. Two other groups of patients who drank less than 30 g/day were included during the same period: patients who were undergoing colonoscopy because of a family history (group FH) of colorectal cancer.

Abbreviations: IBS, irritable bowel syndrome; OR, odds ratio.
In the second study, cases (n=172) were patients with at least one high risk colorectal polyp (>10 mm and/or with a villous component and/or with high grade dysplasia) or a histologically proved colorectal cancer, and controls (n=489) were those with adenomas <10 mm and without high grade dysplasia, a villous component, or colorectal cancer.

In the third study, cases (n=37) were patients with high grade dysplasia or colorectal cancer, and controls (n=624) were those with adenomatous polyps, irrespective of size, and no high grade dysplasia or colorectal cancer.

All colonoscopies had to reach at least the caecum and were performed by a senior endoscopist. In all patients with at least one adenoma of the colon, the location and number of adenomas was recorded as was the location of high risk adenomas or cancer.

Histopathological diagnosis was performed in the anatomo-pathology unit of our hospital and the degree of dysplasia was classified according to the World Health Organisation classification system.

### Statistical analysis

Univariate analysis studies were performed using the \( \chi^2 \) method for dichotomous variables and the Student’s \( t \) test for continuous variables.

Relative risks of adenomas \( \geq 10 \) mm, high risk adenoma or cancer, and high grade dysplasia or cancer, according to study group, were estimated by odds ratios (ORs) adjusted for age and sex using a stepwise logistic regression. In this model, if the value of the probability level is less than some predefined alpha level, say 0.1, then the variable is said to be statistically significant. The statistical significance of each studied variable was tested using the maximum likelihood method.

### RESULTS

#### Characteristics of the adenomas

Adenomas \( \geq 10 \) mm were found in 122 patients: 84 of these 122 patients had adenomas \( \geq 10 \) mm without villous contingent or high grade dysplasia, nine patients in the IBS group, 16 in the FH group, and 59 in the HD group. Adenomas with a villous component were found in 57 patients (13, 5, and 39 patients in the IBS, FH, and HD groups, respectively). Adenomas with high grade dysplasia were found in 28 patients (three, two, and 23 patients in the IBS, FH, and HD groups, respectively). High risk adenoma (\( \geq 10 \) mm or with a villous contingent or with severe dysplasia) were found in 172 patients (25, 23, and 124 patients in the IBS, FH, and HD groups, respectively). Colorectal cancer was found in 12 patients (three, none, and nine patients in the IBS, FH, and HD groups, respectively). Most (68.64%) of the high risk colorectal adenomas were located in the distal colon (left colon, sigmoid, and rectum). A total of 90% of patients had a single high risk colorectal adenoma while 3% had three or more high risk adenomas.

### Table 1

Age and sex characteristics of patients with at least one colorectal adenoma in the irritable bowel syndrome (IBS), family history (FH), and heavy drinkers (HD) groups

<table>
<thead>
<tr>
<th></th>
<th>IBS group</th>
<th>FH group</th>
<th>HD group</th>
<th>( p ) Value*</th>
</tr>
</thead>
<tbody>
<tr>
<td>( n )</td>
<td>152</td>
<td>108</td>
<td>401</td>
<td></td>
</tr>
<tr>
<td>Age (years)</td>
<td>61 (0.9)</td>
<td>55 (1)</td>
<td>57 (0.5)</td>
<td>&lt;0.05</td>
</tr>
<tr>
<td>Male/Female ratio</td>
<td>87/65 (57%)</td>
<td>69/39 (64%)</td>
<td>313/88 (78%)</td>
<td>&lt;0.001</td>
</tr>
</tbody>
</table>

*\( \chi^2 \) test or F test from ANOVA.
Table 2: Group distribution between cases and controls for adenomas >10 mm

<table>
<thead>
<tr>
<th>Patient groups</th>
<th>Cases</th>
<th>Controls</th>
<th>p Value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Age [y] [mean (SD)]</td>
<td>61 (1)</td>
<td>57 (0.5)</td>
<td>&lt;0.05</td>
</tr>
<tr>
<td>Male/female ratio</td>
<td>61/16 (73%)</td>
<td>343/146 (70%)</td>
<td>NS</td>
</tr>
<tr>
<td>IBS</td>
<td>9 (11%)</td>
<td>127 (26%)</td>
<td></td>
</tr>
<tr>
<td>FH</td>
<td>16 (19%)</td>
<td>85 (17%)</td>
<td>&lt;0.01</td>
</tr>
<tr>
<td>HD</td>
<td>59 (70%)</td>
<td>277 (57%)</td>
<td></td>
</tr>
</tbody>
</table>

The p value is given for the comparison by global χ² method of the relative frequency in the three groups of patients (irritable bowel syndrome (IBS), family history (FH), and heavy drinkers (HD) groups) of cases and controls. The number of patients (cases plus controls=671) is less than the total number of patients (601) because cases with adenomas >10 mm and high grade dysplasia or a villous component were excluded from this part of the study.

Table 3: Group distribution between cases and controls for high risk adenomas or colorectal cancer

<table>
<thead>
<tr>
<th>Patient groups</th>
<th>Cases</th>
<th>Controls</th>
<th>p Value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Age [mean (SD)]</td>
<td>59 (0.9)</td>
<td>57 (0.5)</td>
<td>0.049</td>
</tr>
<tr>
<td>Male/female ratio</td>
<td>126/46 (73%)</td>
<td>343/146 (70%)</td>
<td>NS</td>
</tr>
<tr>
<td>IBS</td>
<td>25 (15%)</td>
<td>127 (26%)</td>
<td></td>
</tr>
<tr>
<td>FH</td>
<td>23 (13%)</td>
<td>85 (17%)</td>
<td>0.00096</td>
</tr>
<tr>
<td>HD</td>
<td>124 (72%)</td>
<td>277 (57%)</td>
<td></td>
</tr>
</tbody>
</table>

The p value is given for the comparison by global χ² method of the relative frequency in the three groups of patients (irritable bowel syndrome (IBS), family history (FH), and heavy drinkers (HD) groups) of cases (patients with at least one high risk adenoma or a cancer) and controls (patients with at least one colorectal adenoma but without high risk adenomas or cancer).

Results of analysis

In the first case control study, we examined the relative frequency in the three risk groups (IBS, FH, HD) of patients with adenomas ≥10 mm but without high grade dysplasia or a villous contingent (cases) and patients without adenomas ≥10 mm and without high grade dysplasia or a villous contingent (controls). This relative frequency was significantly different (p<0.01) between cases (n=84) and patients with adenomas <10 mm (controls, n=489). Heavy drinking was more frequent (70%) in cases than in controls (57%) (table 2).

After age and sex adjustment, the likelihood of having an adenoma ≥10 mm was higher in group HD than in group IBS (OR 1.6 (95% CI 1.07–2.2); p<0.05). After age and sex adjustment, the likelihood of having an adenoma ≥10 mm was not significantly different between groups HD and FH (OR 1.14 (95% CI 0.7–2.0); NS). In the second case control study, we examined the relative frequency in the three risk groups of cases with a high risk adenoma (adenomas ≥10 mm or with high grade dysplasia or a villous component) or colorectal cancer, and of controls. As in the first part of our study, the relative frequency in the three groups of patients was significantly (p<0.001) different between patients with high risk adenomas or colorectal cancer (cases, n=172) and those without high risk adenomas or cancer (controls, n=489). Heavy drinkers were more frequent (72%) in cases than in controls (57%) (table 3).

After age and sex adjustment, the likelihood of having a high risk adenoma or colorectal cancer was higher in group HD than in group IBS (OR 1.6 (95% CI 1.2–2.1); p<0.001) and was also higher, although not significantly at the 5% level, in group HD than in group FH (OR 1.4 (95% CI 0.97–2.6) (p=0.081), 90% CI (1.02–2.3)).

In the second case control study, we examined the relative frequency in the three risk groups of patients with high grade dysplasia or colorectal cancer among alcoholic patients with endoscopically proved adenomas compared with two other populations, one with a risk higher than in the general population (patients with a family history of colorectal tumours) and one with a risk thought to be at least equal to the general population (those suffering from IBS). They concluded that alcohol may be involved in the development of colorectal cancer in patients with IBS.

Influence of cirrhosis

In the HD group, we evaluated the influence of cirrhosis in the three case control studies and found that cirrhotic patients were not significantly more numerous between cases and controls in any of the three studies performed (table 5).

DISCUSSION

We have shown a relationship between alcoholism and the risk of high risk adenomas or colorectal cancer. To our knowledge, this is the first study that has investigated high risk adenomas or cancer among alcoholic patients with endoscopically proved adenomas compared with two other populations, one with a risk higher than in the general population (patients with a family history of colorectal tumours) and one with a risk thought to be at least equal to the general population (those suffering from IBS). The association between colorectal cancer and alcohol consumption is controversial. A meta analysis of the relation between alcohol and colorectal cancer is controversial. Heavy drinking was more frequent (81%) in cases with high grade dysplasia adenomas or colorectal cancer than in controls (59%) (table 4). After age and sex adjustment, the likelihood of having a high grade dysplasia adenoma or a colorectal cancer was higher in group HD than in group IBS (OR 1.7 (95% CI 1.02–2.8); p<0.05) and, although without being statistically significant at the 5% level, than in group FH (OR 3.7 (95% CI 0.98–15) (p=0.076), 90% CI (1.10–12.47)).

The p value is given for the comparison by global χ² method of the relative frequency in the three groups of patients (irritable bowel syndrome (IBS), family history (FH), and heavy drinkers (HD) groups) of cases (patients with at least one high risk adenoma or a colorectal adenoma but without high grade dysplasia or cancer).
promotion of growth of the adenoma, and that alcohol acts as an early promoter of colorectal carcinogenesis but plays no role in malignant changes.

The mechanism by which alcohol might promote colorectal cancer is still controversial. Alcohol in itself would seem to have no direct carcinogenic effect. Its metabolite acetaldehyde has been proved to be cytotoxic. A direct action of acetaldehyde on the faecal microflora has been suggested. Seitz et al have demonstrated in an animal model that the intestinal flora may be involved in acetaldehyde production as germ free rats had significantly lower acetaldehyde concentrations in the rectum (84 (11) v 234 (33) nmol/g/colon; p<0.01) and in the caecum (59 (13) v 121 (33) nmol/g/colon; p<0.05) compared with conventional animals, and that alcohol feeding induced rectal but not caecal hyperregeneration. Acetaldehyde can react with various intracellular and extracellular proteins to form both stable and unstable condensation products. Several proteins involved in these reactions have been shown to form cross links with acetaldehyde, including albumin, tubulin, haemoglobin, collagen, and cytochrome P-450III A. Acetaldehyde-protein adduct formation may lead to cellular damage or dysfunction through alteration of biological protein function or adduct triggered immunological effects. Holstege et al have hypothesised that only on their extracellular location, acetaldehyde modified epitopes could serve as neoantigens and thereby elicit humoral or cellular immune responses. Ki-ras mutations are thought to be early events in the carcinogenic process leading to colon tumours. It has been reported that alcohol may be involved in the disease pathway of colon tumours represented by specific Ki-ras mutations.

Ethanol may increase the risk of adenomatous polyps by interfering with dietary habits, and the relation between ethanol and adenomatous polyps could be the result of confounding by unspecified dietary factors such as folate or vitamin C as alcoholic patients often have a low fruit and vegetable diet. Folate, which is plentiful in vegetables and fruits, vitamin C as alcoholic patients often have a low fruit and vegetable diet. Folate, which is plentiful in vegetables and fruits, also found that the risk of colorectal cancer in patients within the highest quartile for folate intake was lower than patients suffering from IBS. This is in agreement with Faivre and colleagues who found that the proportion of large adenomas in men which could be attributed to combined alcohol and tobacco consumption was 78%.

We did not study the role of cholesterol but it needs to be discussed as we have previously observed that serum cholesterol concentration was lower in patients with adenomas than in patients without adenomas. In the same study, we showed that alcoholism was independently associated with the risk of having colorectal adenomas.

The ORs after adjustment for age and sex were in agreement with the findings that alcoholism may be involved in the severity of colonic adenomas as the relative risk of heavy drinkers having a large adenoma, a high risk adenoma or cancer, or a high grade dysplasia or cancer was significantly increased compared with IBS patients (ORs 1.8, 1.6, and 1.7, respectively). Comparison between the HD and FH groups showed that there was a trend towards an increased risk for high risk adenomas or cancer and for high grade dysplasia or cancer in group HD compared with group FH, but this was not statistically significant at the 5% level.

In conclusion, in our study alcoholism was a risk factor for the development of high risk adenomas or colorectal cancer. This suggest that alcoholic patients may benefit from a screening policy of colonic adenomatous polyps when their liver function allows.

**Table 5 Relation between cirrhosis and: adenomas >10 mm; high risk adenomas or cancer; and high grade dysplasia or cancer among alcoholic patients**

<table>
<thead>
<tr>
<th>Cirrhosis</th>
<th>Adenomas &gt;10 mm (n=59)</th>
<th>Adenomas &lt;10 mm (n=276)</th>
<th>p Value</th>
<th>NS</th>
</tr>
</thead>
<tbody>
<tr>
<td>cirrhosis</td>
<td>High risk adenoma or cancer (n=124)</td>
<td>No high risk adenoma or cancer (n=277)</td>
<td>106 (38%)</td>
<td>NS</td>
</tr>
<tr>
<td>cirrhosis</td>
<td>High grade dysplasia or cancer (n=30)</td>
<td>No high grade dysplasia or cancer (n=371)</td>
<td>143 (39%)</td>
<td>NS</td>
</tr>
</tbody>
</table>

Per cent values represent the number of patients with proven cirrhosis (cirrhosis was assessed either on a liver biopsy or using clinical and biological signs) among the total number of patients in each subgroup of the heavy drinkers.

**Authors’ affiliations**

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**REFERENCES**


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GUT 2002 50: 38-42
doi: 10.1136/gut.50.1.38

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