Bowel movement frequency and risk of colorectal cancer in a large cohort study of Japanese men and women

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The relationship between bowel movement (BM) frequency and the risk of colorectal cancer was examined in a large cohort of 25,731 men and 37,198 women living in 24 communities in Japan. At enrolment, each participant completed a self-administered questionnaire on BM frequency and laxative use. Incidence rate ratios (IRR) with 95% confidence intervals (CI) were estimated using Cox’s proportional-hazard model. During the follow-up period (average length 7.6 years), 649 cases of colorectal cancer, including 257 cases of colorectal cancer, were identified. Among women, subjects who reported a BM every 2–3 days had the lowest risk of developing colorectal cancer (IRR = 0.70, 95% CI = 0.52–0.97) and colon cancer (IRR = 0.70, 95% CI = 0.49–1.00), whereas those reporting a BM every 6 days or less had an increased risk of developing colorectal cancer (IRR = 2.47, 95% CI = 1.01–6.01) and colon cancer (IRR = 2.52, 95% CI = 0.93–6.82) compared with those reporting ≥1 BM per day. A similar, but nonsignificant, association between the frequency of BM and cancer risk was observed in men. There was no association between colorectal or colon cancer risk and laxative use. Regulating BM frequency might therefore have a role in the prevention of colorectal cancer.

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An association between constipation and the risk of colorectal cancer has long been noted. Prolonged intestinal transit time might not only increase the duration of contact between carcinogens in the stools and the gut wall, but could also concentrate carcinogens by increasing colonic water absorption. A meta-analysis of 14 case–control studies that examined the association between constipation or infrequent bowel movements (BM) and colorectal cancer and found a statistically significant 48% increase in the pooled odds ratios for colorectal cancer (Kotake et al, 1995; Le Marchand et al, 1997; Ghadirian et al, 1998; Jacobs and White, 1998; Roberts et al, 2003).

Since bowel habits might be influenced by the presence of colorectal cancer, retrospective studies cannot exclude the effects of the cancer itself, as well as recall bias, on their results. However, few prospective studies have addressed this issue. The only cohort study, which had a 12-year follow-up period involving 84,577 women, of colorectal cancer incidence and BM frequency or laxative use reported negative results (Dukas et al, 2000). The influence of BMs on male colorectal cancer has not been previously studied prospectively.

We conducted a large cohort study to investigate the association between bowel habits, laxative use, susceptibility to diarrhoea and the colorectal cancer risk in Japanese men and women.

MATERIALS AND METHODS

All data were taken from the Japan Collaborative Cohort (JACC) Study, the methods of which have been described in detail elsewhere (Ohno and Tamakoshi, 2001). Briefly, the original study population consisted of 110,792 Japanese adults aged 40–79 years. Enrolment began in 1988 and continued until the end of 1990 in 45 areas across Japan. Most subjects were recruited from the general population or when undergoing routine health checks in the municipalities. Written informed consent for participation was obtained individually from subjects, with the exception of a few
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study areas in which informed consent was provided at the group level after explaining the aims of the study and confidentiality of the data to community leaders. The study protocol was approved by the Ethics Committee of Medical Care and Research of the Fujita Health University School of Medicine, Japan.

Analyses were restricted to data from the 65 184 participants who lived in the 24 study areas in which cancer registries were available. A further 58 subjects with a previous history of colorectal cancer, and 2197 subjects for whom information about bowel habits was not available, were excluded. Therefore, a total of 62 929 individuals (25 731 men and 37 198 women) were involved in this analysis.

All participants completed a self-administered questionnaire on enrolment. This covered demographic characteristics and lifestyle factors such as diet, tobacco smoking, alcohol consumption, physical activity, BM frequency, susceptibility to diarrhoea and laxative use over the past year. The alternative answers provided on the questionnaire for the frequency of BM were: ‘daily’, ‘every 2 – 3 days’, ‘every 4 – 5 days’ and ‘every 6 days or less’. With regard to laxative use, the questionnaire asked only whether the participants used laxatives in the past one year at the time of enrolment; additional data on the type of laxative, the reason for use and the duration of use were not collected. Participants also provided information about susceptibility to diarrhoea by answering ‘yes’, ‘no’ or ‘neutral’ in response to the question: do you often have diarrhoea?

Population registries in the municipalities were used to determine the vital and residential status of subjects. Registration of death is required under the Family Registration Law in Japan, which applies throughout the country. Incidences of cancer were confirmed using records from the population-based cancer registries, which were supplemented by a systematic review of death certificates (Ohno and Tamakoshi, 2001); in some areas, medical records were also reviewed in major local hospitals. The mortality-to-incidence ratio for colorectal cancer was 0.28 in the cohort covered by the cancer registries. This figure is comparable with those calculated in the most accurate population-based cancer registries in Japan (Parkin et al, 2003), which indicates that most cases of colorectal cancer were identified in the study population.

The follow-up period ran from the time of the baseline survey through to the end of 1997 in all but three areas (in which it ran until the end of 1994, 1995 and 1996, respectively). The end point of the study was defined as the incidence of colorectal cancer (10th Revision of the International Classification of Diseases, ICD-10: C18 – C20) or colon cancer (ICD-10: C18). The risk of rectal cancer was not analysed separately because of the relatively small number of cases observed. Subjects who moved out of the study area or died from causes other than colorectal cancer were treated as censored cases. During the study period, only 3.3% (2071) of the participants were lost from the follow-up as a result of a change of residence.

All analyses were carried out by sex using the SAS statistical package release 8.2 (SAS Inc., Cary, NC, USA). Differences in baseline characteristics between categories of BM frequency were tested using the chi-squared ($\chi^2$) test or one-way analysis of variance (ANOVA). The follow-up period for each participant was calculated as the time between completing the questionnaire and either the diagnosis of colon or rectal cancer, death, moving out of the study area or the end of the study – whichever occurred first.

The incidence rate ratios (IRR) and 95% confidence intervals (CI) for colorectal and colon cancer were estimated, by sex, using Cox’s proportional-hazard model according to the levels of BM frequency, laxative use and susceptibility to diarrhoea. The categories of ‘every day’ for BM, ‘nonuse’ for laxative use and ‘no’ or ‘neutral’ for susceptibility to diarrhoea were used as reference groups.

Analyses were adjusted for the following potential confounding factors: age (continuous variable); body mass index (BMI) calculated as weight (kg) [height (m)]$^2$ and categorised as $\geq 25$ kg m$^{-2}$ or $<25$ kg m$^{-2}$; intake frequency of green leafy vegetables (‘daily’ or ‘not daily’); intake frequency of alcohol (‘$\geq$ 5 days per week’ or ‘$<5$ days per week’); current smoking status (‘smoker’ or ‘nonsmoker’); time spent walking per day (‘$\leq$ 30 min or $>30$ min’); history of colorectal cancer in parents or siblings (‘yes’ or ‘no’); and age at leaving full-time education (‘$\geq$ 20 years’ or ‘$<20$ years’). For each covariate, missing values were treated as an additional category and were included in the model. To determine the influence of symptoms of colorectal cancer on bowel habits, analyses were repeated excluding the first 3 years of follow-up. In all cases, two-sided $P$-values $<0.05$ were considered to be statistically significant.

RESULTS

Within the study group, 1.1% of men and 4.0% of women reported infrequent BMs (every 4 days or less). The use of laxatives was more common among women (14.7%) than men (6.9%), whereas men were more likely to report frequent diarrhoea (20.3%) than women (9.7%).

Table 1 shows the baseline characteristics of the study population by BM frequency. Regardless of sex, individuals who reported infrequent BMs – compared with those who reported BMs daily or every 2 – 3 days – had a lower average BMI, were less likely to spend $>30$ min walking per day and were more likely to use laxatives.

A significant difference in the intake frequency of green leafy vegetables and in smoking status across the BM groups was observed only among women: those who reported BMs daily or every 2 – 3 days were more likely to consume green leafy vegetables daily and less likely to be smokers. In addition, women who reported BMs every 2 – 3 days were, on average, younger than those in the other BM groups. Alcohol consumption did not differ between BM groups in women.

Among men, the number that reported daily alcohol intake increased linearly with BM frequency. Male subjects who reported BMs every 2 – 3 days had the lowest rate of frequent diarrhoea, whereas the number of women who reported frequent diarrhoea decreased linearly with BM frequency.

During the follow-up period (average length 7.6 years, standard deviation 1.9), a total of 649 cases of colorectal cancer were identified (379 in men and 270 in women), which included 429 cases of colon cancer (225 in men and 204 in women).

Age-adjusted IRRs were calculated for colorectal and colon cancer according to BM frequency (not shown). Regardless of sex, the ratios were $<1.00$ for subjects who reported BMs every 2 – 3 days relative to those who reported daily BMs: the IRRs for colorectal cancer were 0.74 in men (95% CI $= 0.51$ – 1.09) and 0.71 in women (95% CI $= 0.52$ – 0.97), whereas the IRRs for colon cancer were lower in men (0.45; 95% CI $= 0.25$ – 0.82) and the same in women (0.71; 95% CI $= 0.49$ – 1.00). In contrast, the age-adjusted IRRs for subjects who reported highly infrequent BMs (every 6 days or less) relative to those with daily BMs were $>1.00$: the IRRs for colorectal cancer were 1.14 in men (95% CI $= 1.06$ – 1.10) and 2.53 in women (95% CI $= 1.04$ – 6.15), whereas the IRRs for colon cancer were 1.78 in men (95% CI $= 0.25$ – 12.7) and 2.59 in women (95% CI $= 0.96$ – 6.98).

Adjustment for potential confounding factors [as discussed above] had no significant effects on the IRRs (Table 2). Furthermore, even after excluding the first 3 years of follow-up, there was a lower risk of colorectal or colon cancer in women who reported BMs every 2 – 3 days relative to those who reported daily BMs: the multivariate-adjusted IRRs were 0.64 for colorectal cancer (95% CI $= 0.43$ – 0.96) and 0.68 for colon cancer (95% CI $= 0.43$ – 1.05). Increased risks of colorectal and colon cancers were also observed in association with highly infrequent BMs.
although they were not statistically significant.

Table 3 shows the associations between laxative use, susceptibility to diarrhoea and colorectal or colon cancer risk. There were weak nonsignificant positive associations between laxative use and cancer risk in both men and women, but no association between cancer risk and frequent diarrhoea.

DISCUSSION

This is the first prospective study, to our knowledge, that has reported a significant association between BM frequency and colorectal cancer risk. Infrequent BMs were associated with a significantly increased risk of colorectal cancer and a marginally increased risk of colon cancer in women. A similar, but
nonsignificant, association was found in men. These results were not altered by adjusting for potential confounding factors or excluding the first 3 years of follow-up from the analysis, which indicated that the effects of the cancers themselves on bowel habits were not responsible for the associations.

These results support the findings of recent case-control studies and of the meta-analysis carried out by Sonnenberg and Müller (1993), which reported a significantly increased risk of colorectal cancer in association with constipation or infrequent BMs. However, the findings from the Nurses’ Health Study in the United States – only one published prospective data on the association between BM frequency and female colorectal cancer risk (Dukas et al, 2000) – did not support an association between infrequent BMs and the risk of colorectal cancer. One possible reason for the discrepancy between these results and those of the present study is that different criteria were used to define ‘infrequent BM’. The Nurses’ Health Study defined this as an average frequency of ‘every third day or less’. However, in the present study, a significantly increased risk of colorectal and colon cancer was found only in subjects who reported BMs every 6 days or less relative to those reporting daily BMs. Therefore, we suggest that only highly infrequent BMs elevate the risk of colorectal cancer.

Daily BMs were found to increase the risk of colorectal and colon cancer compared with BMs every 2–3 days, in both men and women. This observation is in line with the results of a previous case-control study carried out in Japan (Kato et al, 1993). However, the Nurses’ Health Study (Dukas et al, 2000) found no difference in colorectal cancer incidence between subjects who reported \( \geq 2 \) BMs per day and those who reported BMs once per day (multivariate-adjusted IRR = 0.89, 95% CI = 0.65–1.20). Unfortunately, limitations of the questionnaire used in the present study precluded us from determining the risk associated with \( \geq 2 \) BMs per day. On the basis of the combined findings of these studies, we speculate that subgroups that have highly frequent BMs might be at an increased risk of colorectal cancer. Experimental studies have reported elevated levels of prostaglandin \( E_2 \) (PGE\(_2\)) in the gastrointestinal tract in many diarrhoeal states (Burakoff and Percy, 1992), and increased levels of PGE\(_2\) might be associated with carcinogenesis in the large intestine (Reddy et al, 1993). We did not observe a significant association between self-reported susceptibility to diarrhoea and colorectal cancer risk, and the results of previous epidemiological studies (case-control studies only) were inconsistent (Dales et al, 1979; Kune et al, 1987).

Similar to ‘constipation’, the definition of ‘diarrhoea’ is equivocal. Some case-control studies have suggested that ‘soft’ or ‘loose’ faeces might increase the risk of colorectal cancer (Kato et al, 1993; Inoue et al, 1995). To have a conclusion, additional data on factors such as faecal consistency should be collected and analysed together with data on susceptibility to diarrhoea and BM frequency.

A weak nonsignificant positive association was found between laxative use and the risk of colorectal cancer in both men and women. Previously, the meta-analysis of Sonnenberg and Müller (1993) revealed a significant 46% increase in the risk of colorectal cancer associated with the use of laxatives. On the other hand, recent case-control studies (Jacobs and White, 1998; Nascimbeni et al, 2002; Roberts et al, 2003) and a prospective study (Dukas et al, 2000) found no relationship between these factors – although Dukas et al suggested that some types of laxative might influence intestinal pH and the metabolism of intestinal flora, thereby modifying colorectal cancer risk. The effects of laxative type were not investigated in the present study because of limitations of the questionnaire. Further prospective studies investigating the types of laxative and duration of use will be necessary to resolve this question.

The risk of rectal cancer was not analysed independently because of the small number of cases in the study group. Larger-scale prospective studies will be necessary to reveal the effects of bowel habits on the development of cancers of the large intestine at specific sites.

There were some limitations to the scope of the present study. For example, although the main risk factors for colorectal cancer were adjusted for in the analysis, other factors such as aspirin use and hormone replacement therapy in women might have confounded the results. Also, bowel habits were evaluated only through a self-reported questionnaire that was administered once at the baseline; the reproducibility and validity of the responses of subjects were therefore not confirmed.

In conclusion, this study shows that highly infrequent BMs can increase the risk of colorectal cancer in both men and women. Highly frequent BMs may also enhance this risk. Further

### Table 3: IRR for colorectal and colon cancer according to laxative use and susceptibility to diarrhoea

|                | Colorectal cancer | Colon cancer |
|----------------|-------------------|--------------|
|                | Observed          | Multivariate- |                |
|                | person-years      | adjusted* IRR| 95% CI\(^b\)   |
|                | No. of cases      |              |                |
| Men Laxative use | No                | 155 068      | 292            | 1.00 | 1.00 |
|                | Yes               | 100 015      | 33             | 1.28 | 0.89 – 1.86 |
| Susceptibility to diarrhoea | Normal | 143 808      | 285            | 1.00 | 1.00 |
|                | Having frequent diarrhoea | 35 775 | 68             | 1.08 | 0.82 – 1.41 |
| Women Laxative use | No                | 206 189      | 183            | 1.00 | 1.00 |
|                | Yes               | 33 097       | 41             | 1.20 | 0.85 – 1.69 |
| Susceptibility to diarrhoea | Normal | 230 880      | 224            | 1.00 | 1.00 |
|                | Having frequent diarrhoea | 23 417 | 26             | 1.18 | 0.79 – 1.78 |

IRR = incidence rate ratios; BMI = body mass index. *Adjusted for age, BMI, intake frequency of green leafy vegetables, daily alcohol drinking, current smoking status, time spent for walking per day, family history of colorectal cancer and education. \(^b\)CI: confidence interval.
prospective studies are needed to confirm our findings and to clarify the risk associated with BMs for colorectal cancer by subsite.

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REFERENCES

Burakoff R, Percy WH (1992) Studies in vivo and in vitro on effects of PGE2 on colonic motility in rabbits. Am J Physiol 262: G23 – G29

Dales LG, Friedman GD, Ury HK, Grossman S, Williams SR (1979) A case – control study of relationships of diet and other traits to colorectal cancer in American blacks. Am J Epidemiol 109: 132 – 144

Dukas L, Willett WC, Colditz GA, Fuchs CS, Rosner B, Giovannucci EL (2000) Prospective study of bowel movement, laxative use, and risk of colorectal cancer among women. Am J Epidemiol 151: 958 – 964

Ghadriyan P, Maisonneuve P, Perret C, Lacroix A, Boyle P (1998) Epidemiology of socioeconomic characteristics, lifestyle, medical history, and colon cancer: a case – control study among French Canadians in Montreal. Cancer Detect Prev 22: 396 – 404

Inoue M, Tajima K, Hirose K, Hamajima N, Takezaki T, Hirai T, Kato T, Ohno Y (1995) Subsite-specific risk factors for colorectal cancer: a hospital-based case–control study in Japan. Cancer Causes Control 6: 14 – 22

Jacobs EJ, White E (1998) Constipation, laxative use, and colon cancer among middle-aged adults. Epidemiology 9: 385 – 391

Kato I, Tominaga S, Matsuura A, Yoshi Y, Shirai M, Kobayashi S (1993) Case–control study of bowel habits and colorectal adenoma and cancer. J Epidemiol 3: 1 – 5

Kotake K, Koyama Y, Nasu J, Fukutomi T, Yamaguchi N (1995) Relation of family history of cancer and environmental factors to the risk of colorectal cancer: a case–control study. Jpn J Clin Oncol 25: 195 – 202

Kune GA, Kune S, Watson LF (1987) The Melbourne Colorectal Cancer Study. Characterization of patients with a family history of colorectal cancer. Dis Colon Rectum 25: 600 – 606

Le M Marchand L, Wilkens LR, Kolonel LN, Hankin JH, Lyu LC (1997) Associations of sedentary lifestyle, obesity, smoking, alcohol use, and diabetes with the risk of colorectal cancer. Cancer Res 57: 4787 – 4794

Nascimento MC, Donato F, Girard M, Mariani P, Villanacci V, Salerni B (2003) Constipation, anorectal laxatives, melanosin coli, and colon cancer: a risk assessment using aberrant crypt foci. Cancer Epidemiol Biomarkers Prev 11: 753 – 757

Ohno Y, Tamakoshi A (2001) Japan collaborative cohort study for evaluation of cancer risk sponsored by monbusho (JACC study). J Epidemiol 11: 144 – 150

Parkin DM, Whelan SL, Ferlay J, Teppo L, Thomas DB (2003) Cancer Incidence in Five Continents, Vol . 8 Lyon: IARC Press

Reddy BS, Rao CV, Rivenson A, Kelloff G (1993) Inhibitory effect of aspirin on azoxymethane-induced colon carcinogenesis in F344 rats. Carcinogenesis 14: 1493 – 1497

Roberts MC, Millikan RC, Galanko JA, Martin C, Sandler RS (2003) Constipation, laxative use, and colon cancer in a North Carolina population. Am J Gastroenterol 98: 857 – 864 doi;10.1016/S0002-9270(03)00505-9

Sonnemberg A, Muller AD (1993) Constipation and cathartics as risk factors of colorectal cancer: a meta-analysis. Pharmacology 47(Suppl 1): 224 – 233