

39. Hsing AW, McLaughlin JK, Chow WH, Schuman LM, Co Chien HT, et al.: Risk factors for colorectal cancer in a prospective study among U.S. white men. *Int J Cancer* **77**, 549–553, 1998.
40. Knekt P, Jarvinen R, Dich J, and Hakulinen T: Risk of colorectal and other gastro-intestinal cancers after exposure to nitrate, nitrite and N-nitroso compounds: a follow-up study. *Int J Cancer* **80**, 852–856, 1999.
41. Pietinen P, Malila N, Virtanen M, Hartman TJ, Tangrea JA, et al.: Diet and risk of colorectal cancer in a cohort of Finnish men. *Cancer Causes Control* **10**, 387–396, 1999.
42. Kato I, Akhmedkhanov A, Koenig K, Toniolo PG, Shore RE, et al.: Prospective study of diet and female colorectal cancer: the New York University Women's Health Study. *Nutr Cancer* **28**, 276–281, 1997.
43. Slattery ML, Berry TD, Potter J, and Caan B: Diet diversity, diet composition, and risk of colon cancer (United States). *Cancer Causes Control* **8**, 872–882, 1997.
44. Nkondjock A, Shatenstein B, Maisonneuve P, and Ghadirian P: Assessment of risk associated with specific fatty acids and colorectal cancer among French-Canadians in Montreal: a case-control study. *Int J Epidemiol* **32**, 200–209, 2003.
45. Dolecek TA and Granditis G: Dietary polyunsaturated fatty acids and mortality in the Multiple Risk Factor Intervention Trial (MRFIT). *World Rev Nutr Diet* **66**, 205–216, 1991.
46. Schloss I, Kidd MS, Tichelaar HY, Young GO, and O'Keefe SJ: Dietary factors associated with a low risk of colon cancer in coloured west coast fishermen. *S Afr Med J* **87**, 152–158, 1997.
47. Kobayashi M, Tsubono Y, Sasazuki S, Sasaki S, and Tsugane S: Vegetables, fruit and risk of gastric cancer in Japan: a 10-year follow-up of the JPHC Study Cohort I. *Int J Cancer* **102**, 39–44, 2002.
48. Tsugane S, Sasazuki S, Kobayashi M, and Sasaki S: Salt and salted food intake and subsequent risk of gastric cancer among middle-aged Japanese men and women. *Br J Cancer* **90**, 128–134, 2004.
49. Bartsch H, Nair J, and Owen RW: Dietary polyunsaturated fatty acids and cancers of the breast and colorectum: emerging evidence for their role as risk modifiers. *Carcinogenesis* **20**, 2209–2218, 1999.
50. Terry PD, Rohan TE, and Wolk A: Intakes of fish and marine fatty acids and the risks of cancers of the breast and prostate and of other hormone-related cancers: a review of the epidemiologic evidence. *Am J Clin Nutr* **77**, 532–543, 2003.
51. Singh J, Hamid R, and Reddy BS: Dietary fat and colon cancer: modulation of cyclooxygenase-2 by types and amount of dietary fat during the postinitiation stage of colon carcinogenesis. *Cancer Res* **57**, 3465–3470, 1997.
52. Badawi AF, El-Soheily A, Stephen LL, Ghoshal AK, and Archer MC: The effect of dietary n-3 and n-6 polyunsaturated fatty acids on the expression of cyclooxygenase 1 and 2 and levels of p21ras in rat mammary glands. *Carcinogenesis* **19**, 905–910, 1998.
53. Fan YY, Spencer TE, Wang N, Moyer MP, and Chapkin RS: Chemopreventive n-3 fatty acids activate RXRalpha in colonocytes. *Carcinogenesis* **24**, 1541–1548, 2003.
54. Nair J, Vaca CE, Velic I, Mutanen M, Valsta LM, et al.: High dietary omega-6 polyunsaturated fatty acids drastically increase the formation of etheno-DNA base adducts in white blood cells of female subjects. *Cancer Epidemiol Biomarkers Prev* **6**, 597–601, 1997.
55. Hanaoka T, Nair J, Takahashi Y, Sasaki S, Bartsch H, et al.: Urinary level of 1.N(6)-ethenodeoxyadenosine, a marker of oxidative stress, is associated with salt excretion and omega-6 polyunsaturated fatty acid intake in postmenopausal Japanese women. *Int J Cancer* **100**, 71–75, 2002.
56. Dolecek TA: Epidemiological evidence of relationships between dietary polyunsaturated fatty acids and mortality in the multiple risk factor intervention trial. *Proc Soc Exp Biol Med* **200**, 177–182, 1992.
57. Abou-el-Ela SH, Prasse KW, Farrell RL, Carroll RW, Wade AE, et al.: Effects of D,L-2-difluoromethylornithine and indomethacin on mammary tumor promotion in rats fed high n-3 and/or n-6 fat diets. *Cancer Res* **49**, 1434–1440, 1989.
58. Bartram HP, Gostner A, Reddy BS, Rao CV, Scheppach W, et al.: Missing anti-proliferative effect of fish oil on rectal epithelium in healthy volunteers consuming a high-fat diet: potential role of the n-3:n-6 fatty acid ratio. *Eur J Cancer Prev* **4**, 231–237, 1995.
59. Bartram HP, Gostner A, Scheppach W, Reddy BS, Rao CV, et al.: Effects of fish oil on rectal cell proliferation, mucosal fatty acids, and prostaglandin E2 release in healthy subjects. *Gastroenterology* **105**, 1317–1322, 1993.
60. Deschner EE, Lytle JS, Wong G, Ruperto JF, and Newmark HL: The effect of dietary omega-3 fatty acids (fish oil) on azoxymethanol-induced focal areas of dysplasia and colon tumor incidence. *Cancer* **66**, 2350–2356, 1990.
61. Noguchi M, Minami M, Yagasaki R, Kinoshita K, Earashi M, et al.: Chemoprevention of DMBA-induced mammary carcinogenesis in rats by low-dose EPA and DHA. *Br J Cancer* **75**, 348–353, 1997.

ECOLOGICAL STUDY OF SOLAR RADIATION AND CANCER MORTALITY IN JAPAN

Tetsuya Mizoue*

Abstract—Geographic observation of the increased mortality of some cancers at higher latitudes has led to a hypothesis that vitamin D produced after exposure to solar radiation has anti-carcinogenic effects. However, it is unclear whether such association would be observed in countries like Japan, where fish consumption, and therefore dietary vitamin D intake, is high. Pearson correlation coefficients were calculated between averaged annual solar radiation levels for the period from 1961 through 1990 and cancer mortality in the year 2000 in 47 prefectures in Japan, with adjustments for regional per capita income and dietary factors. A moderate, inverse correlation with solar radiation was observed for cancers of the esophagus, stomach, colon, rectum, pancreas, and gallbladder and bile ducts in both sexes (correlation coefficient, ranging from -0.6 to -0.3). The results of this study support the hypothesis that increased exposure to solar radiation reduces the risk of cancers of the digestive organs.

Health Phys. 87(5):532–538; 2004

Key words: cancer; radiation, cosmic; mortality; exposure, population

INTRODUCTION

STUDIES OF the geographical distribution of cancers have shown that certain types of cancer mortality increase with decreasing intensity of solar radiation, including cancers of the colon (Garland and Garland 1980; Gorham et al. 1989; Grant 2002), breast (Gorham et al. 1989; Garland et al. 1990; Grant 2002), prostate (Hanchette and Schwartz 1992; Grant 2002), and ovary (Lefkowitz and Garland 1994; Grant 2002). Since the association is not fully explained by known risk factors, including diet, one hypothesis that has been proposed is that increased exposure to solar radiation helps to prevent cancers through the augmented synthesis of vitamin D (Garland and Garland 1980). This hypothesis is biologically plausible, since vitamin D analogs are reported to affect cell

proliferation and differentiation (Lointier et al. 1987; Brehier and Thomasset 1988).

Previous studies in Japan have detected increased mortality from cancers of certain digestive organs at high latitudes (Kato et al. 1985; Watanabe and Arimoto 1990). These findings suggest the existence of an association between solar radiation and cancer, even in countries like Japan where fish consumption, and therefore dietary vitamin D intake, is high (FAOUN 1996). Geographically, Japan extends across a wide region from north to south, with major cities in its 47 prefectures ranging from 26°N to 43°N latitude. Its islands feature mountain ranges running through their center. Together, these two characteristics result in significant meteorological variations. The present study sought to investigate whether levels of solar radiation at various sites in Japan over the past 30 y are associated with cancers of the digestive organs and sex-hormone-related organs. The potential confounding effect of diet was assessed using data from a national nutritional survey.

MATERIALS AND METHODS

We chose the prefecture ($n = 47$) as the unit of ecological observation. Sex-specific age-standardized mortality rates according to prefecture for the year 2000 (SID 2002) were obtained for cancers of the colon, rectum, prostate, breast (female), and ovary (Appendix A1), for which a relation with solar radiation or latitude has been suggested; and cancers of the esophagus, stomach, pancreas, and gallbladder and bile ducts, for which higher mortality has been reported in northern regions or in regions of Japan with cloudy and snowy winters.

We obtained data of average annual hours of solar radiation received from 1961 to 1990 for the central city of each prefecture (Japan Meteorological Agency 2001), except for two prefectures, for which data on solar radiation was available for another major city (Saitama: Kumagai City, Shiga: Hikone City). The mean average annual solar radiation during 1961–1990 for the prefectures was $3.59 \text{ KWh h d}^{-1}$ (range: 3.29–4.01).

* Department of Preventive Medicine, Faculty of Medical Sciences, Kyushu University, 3-1-1, Maidashi, Higashiku, Fukuoka 812-8582, Japan.

For correspondence or reprints contact: the author at the above address, or email at mizoue@phealth.med.kyushu-u.ac.jp.

(Manuscript received 15 December 2003; revised manuscript received 14 April 2004, accepted 11 July 2004)

0017-9078/04/0

Copyright © 2004 Health Physics Society

We regarded income and nutritional factors as potential confounding factors. Prefectural income per person was obtained from the 1990 Report on Prefectural Economic Calculation (The Cabinet Office 1990). Data on nutrition were obtained from the 1990 National Nutritional Survey (Bureau of Public Health, Ministry of Health and Welfare 1992), in which subjects were selected at random from the general population. Mean intake of food groups and nutrients was available only for 12 geographical districts, each composed of one to six prefectures (Appendix B1). Therefore, dietary intake for a district was assigned to prefectures that compose the district. The dietary factors initially considered included animal protein, total fat, and fish of any kind (excluding processed food) for cancers of the colon, breast, and prostate; and salt for stomach cancer. Soy products in relation to hormone-related cancers and fiber in relation to colon cancer were not considered because recent ecological studies in Japan have not detected significant associations for these factors (Nagata 2000; Nakaji et al. 2003). Among dietary factors considered, intakes of fat and salt, which were inversely associated with solar radiation (correlation coefficient, -0.49 and -0.26 , respectively), were adjusted in the analysis, while intakes of fish and animal protein, which were materially unrelated to solar radiation (correlation coefficient, 0.09 and 0.08 , respectively), were not adjusted.

The Pearson correlation coefficient between solar radiation and cancer mortality was calculated for all prefectures ($n = 47$). Partial correlation coefficient was estimated by including terms of income and fat intake in the analysis of cancers of the colon, rectum, prostate, breast, and ovary; and for income and salt intake in the analysis of stomach cancer. The analysis was then repeated while excluding prefectures with large metropolitan areas (Kanto area: Saitama, Tokyo, Chiba, Kanagawa; Hanshin area: Kyoto, Osaka, Hyogo; Aichi; Fukuoka). This exclusion was made to account for the potential effects of large migration from other prefectures and reduced exposure to sunlight due to the shadows cast by buildings. Low levels of serum 25-hydroxyvitamin D concentrations in elderly persons, especially those who are sun-deprived, have been reported, suggesting a reduced sun exposure in cities (Gloth et al. 1995). Further analysis excluded Okinawa prefecture at 26°N , which is situated 5 degrees latitude south of the prefecture with the second southern-most latitude in Japan. The reasons for excluding this prefecture included unique dietary and disease patterns in Okinawa. For example, so-called westernized foods including bread, beef, and coffee were consumed more heavily, while the frequency of salted food intake was remarkably low in Okinawa (Tsugane et al. 2001). The

importance of such dietary factors is further suggested by the observation that the mortality of stomach cancer is only half of the average mortality in Japan (SID 2002).

RESULTS

In men, a significant inverse association was observed between solar radiation and age-adjusted mortality rates from cancers of the digestive organs: the esophagus, stomach, colon, rectum, pancreas, gallbladder and bile ducts (Table 1). In women, significant associations were also observed for these sites, although the strength of the associations was generally weaker than in men (Table 2). In both sexes, the associations for cancers of the colon, rectum, and stomach remained significant after controlling for income and the above-mentioned dietary factors. In contrast, there was virtually no association between solar radiation and mortality for cancers of the prostate, breast, and ovary. Excluding prefectures with major urban areas and Okinawa prefecture did not materially change the results, except for cancers of the stomach and pancreas in women, for which the correlation coefficients became non-significant after these exclusions.

DISCUSSION

We found a moderate, inverse relation between solar radiation and cancer mortality involving cancers of the digestive organs. The findings for colon cancer are consistent with previous ecological analyses (Garland and Garland 1980; Gorham et al. 1989; Grant 2002). Increased mortality from cancers of the esophagus, stomach, and pancreas in areas of Japan with low solar radiation is also consistent with previous spatial observations, including those in Japan (Kato et al. 1985; Watanabe and Arimoto 1990). The present study also

Table 1. Correlation coefficient (r) between solar radiation (1961–90) and cancer mortality (2000) in men, Japan.

Cancer site	Age-adjusted yearly mortality (min. max) per 100,000 person	r	r^a	
			r^a	r^b
Esophagus	10.4 (6.0, 16.5)	-0.42^c	-0.45^c	
Stomach	39.1 (21.7, 51.5)	-0.50^c	-0.48^c	-0.44^c
Colon	14.4 (10.8, 19.2)	-0.55^c	-0.53^c	-0.41^c
Rectum	9.3 (6.7, 11.7)	-0.54^c	-0.53^c	-0.40^c
Colorectum	23.7 (18.0, 29.6)	-0.63^c	-0.61^c	-0.49^c
Pancreas	12.4 (7.0, 17.2)	-0.51^c	-0.53^c	
Gallbladder and bile duct	8.2 (6.6, 12.6)	-0.31^d	-0.55^c	
Prostate	8.6 (5.9, 11.6)	-0.04	-0.01	-0.07

^a Adjusted for income.

^b Additionally adjusted for fat intake (colon, rectum, prostate) or salt intake (stomach).

^c $p < 0.01$.

^d $p < 0.05$.

Table 2. Correlation coefficient (*r*) between solar radiation (1961–90) and cancer mortality (2000) in women, Japan.

Cancer site	Age-adjusted yearly mortality (min, max) per 100,000 person	<i>r</i>	<i>r</i> ^a	<i>r</i> ^b
Esophagus	1.3 (0.5, 1.8)	-0.45 ^c	-0.41 ^c	
Stomach	15.3 (7.7, 19.2)	-0.37 ^d	-0.32 ^d	-0.35 ^d
Colon	9.5 (7.3, 12.3)	-0.51 ^c	-0.46 ^c	-0.33 ^d
Rectum	4.1 (2.7, 5.3)	-0.47 ^c	-0.47 ^c	-0.39 ^c
Colorectum	13.6 (10.4, 17.6)	-0.58 ^c	-0.54 ^c	-0.42 ^c
Pancreas	7.2 (4.0, 9.9)	-0.32 ^d	-0.31 ^d	
Gallbladder and bile duct	6.3 (4.7, 8.5)	-0.44 ^c	-0.50 ^c	
Breast	10.7 (7.1, 13.4)	-0.20	-0.09	-0.06
Ovary	4.3 (2.1, 5.4)	-0.17	-0.06	-0.04

^a Adjusted for income.^b Additionally adjusted for fat intake (colon, rectum, breast, ovary) or salt intake (stomach).^c *p* < 0.01.^d *p* < 0.05.

suggests that cancers of the gallbladder and bile ducts are associated with levels of solar radiation. In contrast, cancers of the prostate, breast, and ovary did not significantly correlate with solar radiation.

As previously hypothesized, one plausible explanation for the present inverse association is that exposure to solar radiation reduces the risk of cancer through photo-initiation of vitamin D production. Vitamin D is synthesized in skin on exposure to ultraviolet B, then metabolized in the liver and kidney to vitamin 1,25-D₃. The synthesis of 1,25-D₃ from the precursor 25-D₃ also occurs at tissue levels (Schwartz et al. 1998). 1,25-D₃ not only promotes cell differentiation and reduces proliferation (Lointier et al. 1987; Brehier and Thomasset 1988) but also inhibits tumor growth in xenografts (Eisman et al. 1987) and induces apoptosis of various cancer cells, including colon (Vandewalle et al. 1995). Furthermore, there is evidence, although not always consistent, that elevated levels of serum vitamin D analog are associated with reduced risk of cancer or cancer precursors (Garland et al. 1989; Platz et al. 2000). These findings suggest that vitamin D may be involved in several stages of carcinogenic process.

The Japanese consume large amounts of fish, a food source rich in vitamin D. Although fish consumption in the present data set was relatively equal among regional populations subject to different levels of solar radiation intensity, the intake of fatty fish, which are especially rich in vitamin D, tends to be greater in northern regions (Bureau of Public Health, Ministry of Health and Welfare 1992). Consistent with this observation, recent national nutritional survey estimated greater intake of vitamin D for Japan's northern districts (The Study Circle for Health and Nutrition Information 2003); the mean dietary intakes of vitamin D were 11.4 mg and 6.5 mg for the most northern district and the most southern

district, respectively. Without this gradient in dietary intake of vitamin D opposite to that of solar radiation, the effects of the latter may well have been greater. However, without population data showing serum vitamin D levels across the nation, we can neither refute nor adopt the proposed hypothesis involving vitamin D.

Other factors may account for the present association. Mortality rates from suicide, more frequent among those who are clinically depressed, are inversely associated with intensity of solar radiation in Japan (Terao et al. 2002). Depression has been suggested to cause immune suppression (Miller et al. 1993) and has been proposed as a predisposing condition for diseases that develop more easily under decreased surveillance activity of the immune system, including cancer. Several, but not all, epidemiological studies have reported an increased risk of cancer among those who are depressed (Penninx et al. 1998). Thus, it is possible that the exposure to solar radiation may indirectly reduce cancer risks through positive effects on mood that intensify immune surveillance systems. However, since there is no direct evidence linking digestive cancers to immune suppression, further research is required to judge the plausibility of this mechanism as an explanation for the inverse association between solar radiation and digestive cancers.

The present study found no association between solar radiation and sex hormone-related cancers (prostate, breast, and ovary), a finding at odds with previous ecological studies (Gorham et al. 1989; Garland et al. 1990; Hanchette and Schwartz 1992; Lefkowitz and Garland 1994; Grant 2002). Although these cancers have steadily increased in Japan over recent decades, their resulting incidence rates remain at the lowest levels among developed countries (Parkin et al. 2002). Studies have indicated that intake of soy products (rich in isoflavones) and fish (rich in ω -3 fatty acid), staples of the traditional Japanese diet, are associated with reduced risk of cancers of the breast and prostate, respectively (Terry et al. 2001; Yamamoto et al. 2003). High consumption of these foods among the Japanese may mask the effects of solar radiation on these cancers. Alternatively, since hormone-related cancers tend to originate early in life (Swerdlow et al. 2002), an adequate study might require information on lifetime exposure to solar radiation. The increasing gap between cancer incidence and mortality has been observed especially for hormone-related cancers. If people living in northern prefectures have a greater availability of medical care service than those living in southern prefectures, it is possible that an association between solar radiation and these cancers would be masked due to a better prognosis of cancer patients in northern areas. In reality, however, medical care cost tends to be higher in southwest districts than

northeast districts in Japan (MHLW 1999), indicating that such bias is a less likely explanation for the lack of the association for these cancers.

Ecological studies are apt to suffer from the ecological fallacy. We adjusted for fat intake in our analysis for colon and rectal cancers and for salt intake in our analysis for stomach cancer. The results indicate that these dietary factors do not fully explain the inverse relationship with solar radiation. Smoking, a possible risk factor of some digestive cancers (Mizoue et al. 2000), did not show a latitudinal gradient in Japan (Bureau of Public Health, Ministry of Health and Welfare 1992). Internal migration or large population increase in some prefectures may dilute the association between solar radiation and cancer. The amount of time that people spend indoors may affect the result. If a snowy winter confines people inside the home, the difference in exposure to solar radiation across the nation would be enhanced. Women showed a slightly weaker association between solar radiation and cancer mortality than men. One possible reason is that women, who may stay longer hours indoors than men, and whose clothing habits may further restrict exposure of the skin, have a smaller geographical difference in exposure levels to solar radiation than men. We used mean solar radiation levels for the period from 1961 through 1990 and mortality data in 2000. Not knowing the minimum exposure period and latent period for the development of cancer, we cannot assess the validity of these periods. An analysis using appropriate exposure and lag periods should yield a stronger association between solar radiation and cancer risk.

CONCLUSION

The present study adds evidence to the hypothesis that solar radiation reduces mortality from cancers of the digestive organs. The lack of association found in this study between solar radiation and sex-hormone-dependent cancers warrants further investigation.

REFERENCES

- Brehier A, Thomasset M. Human colon cell line HT-29: Characterization of the 1,25-dihydroxyvitamin D₃ receptor and induction of differentiation by the hormone. *J Steroid Biochem* 29:265-270; 1988.
- Bureau of Public Health, Ministry of Health and Welfare. The National Nutrition Survey in Japan 1990. Tokyo: Daiichi-Shuppan; 1992 (in Japanese).
- Eisman JA, Barkla DH, Tutton PJ. Suppression of in vivo growth of human cancer solid tumor xenografts by 1,25-dihydroxyvitamin D₃. *Cancer Res* 47:21-25; 1987.
- Food and Agriculture Organization of the United Nations. FAO yearbook: Fishery statistics. Rome: Food and Agriculture Organization of the United Nations; 1996.
- Garland CF, Garland FC. Do sunlight and vitamin D reduce the likelihood of colon cancer? *Int J Epidemiol* 9:227-231; 1980.
- Garland CF, Comstock GW, Garland FC, Helsing KJ, Shaw EK, Gorham ED. Serum 25-hydroxyvitamin D and colon cancer: Eight-year prospective study. *Lancet* 2:1176-1178; 1989.
- Garland FC, Garland CF, Gorham ED, Young JF. Geographic variation in breast cancer mortality in the United States: a hypothesis involving exposure to solar radiation. *Prev Med* 19:614-622; 1990.
- Gloth FM 3rd, Gundberg CM, Hollis BW, Haddad JG Jr., Tobin JD. Vitamin D deficiency in homebound elderly persons. *JAMA* 274:1683-1686; 1995.
- Gorham ED, Garland CF, Garland FC. Acid haze air pollution and breast and colon cancer mortality in 20 Canadian cities. *Can J Public Health* 80:96-100; 1989.
- Grant WB. An estimate of premature cancer mortality in the U.S. due to inadequate doses of solar ultraviolet-B radiation. *Cancer* 94:1867-1875; 2002.
- Hanchette CL, Schwartz GG. Geographic patterns of prostate cancer mortality. Evidence for a protective effect of ultraviolet radiation. *Cancer* 70:2861-2869; 1992.
- Japan Meteorological Agency. Nihon Kiko Hyo (Japan Meteorological Table). Tokyo: Japan Meteorological Business Support Center; 2001 (in Japanese).
- Kato I, Tajima K, Kuroishi T, Tominaga S. Latitude and pancreatic cancer. *Jpn J Clin Oncol* 15:403-413; 1985.
- Lefkowitz ES, Garland CF. Sunlight, vitamin D, and ovarian cancer mortality rates in US women. *Int J Epidemiol* 23:1133-1136; 1994.
- Lointier P, Wargovich MJ, Saez S, Levin B, Wildrick DM, Boman BM. The role of vitamin D₃ in the proliferation of a human colon cancer cell line in vitro. *Anticancer Res* 7:817-821; 1987.
- Miller AH, Spencer RL, McEwen BS, Stein M. Depression, adrenal steroids, and the immune system. *Ann Med* 25:481-487; 1993.
- Ministry of Health, Labour and Welfare. Health expenditure for the elderly per person according to prefecture 1999. Available at: <http://www.mhlw.go.jp/toukei/saikin/hw/hoken/iryomap/99/12page.html> (in Japanese). Accessed April 2004.
- Mizoue T, Tokui N, Nishisaka K, Nishisaka S, Ogimoto I, Ikeda M, Yoshimura T. Prospective study on the relation of cigarette smoking with cancer of the liver and stomach in their endemic region. *Int J Epidemiol* 29:232-237; 2000.
- Nagata C. Ecological study of the association between soy product intake and mortality from cancer and heart disease in Japan. *Int J Epidemiol* 29:832-836; 2000.
- Nakaji S, Shimoyama T, Wada S, Sugawara K, Tokunaga S, MacAuley D, Baxter D. No preventive effect of dietary fiber against colon cancer in the Japanese population: A cross-sectional analysis. *Nutr Cancer* 45:156-159; 2003.
- Parkin DM, Whelan SL, Ferlay J, Teppo L, Thomas DB. Cancer incidence in five continents, Vol. VIII. Lyon: IARC; IARC Publications No. 155; 2002.
- Penninx BW, Guralnik JM, Pahor M, Ferrucci L, Cerhan JR, Wallace RB, Havlik RJ. Chronically depressed mood and cancer risk in older persons. *J Natl Cancer Inst* 90:1888-1893; 1998.
- Platz EA, Hankinson SE, Hollis BW, Colditz GA, Hunter DJ, Speizer FE, Giovannucci E. Plasma 1,25-dihydroxy- and 25-hydroxyvitamin D and adenomatous polyps of the distal colorectum. *Cancer Epidemiol Biomarkers Prev* 9:1059-1065; 2000.

- Schwartz GG, Whitlatch LW, Chen TC, Lokeshwar BL, Holick MF. Human prostate cells synthesize 1,25-dihydroxyvitamin D3 from 25-hydroxyvitamin D3. *Cancer Epidemiol Biomarkers Prev* 7:391-395; 1998.
- Statistics and Information Department, Minister's Secretariat, Ministry of Health, Labor and Welfare. Age-adjusted death rates by prefecture, Special Report on Vital Statistics 2000. Tokyo: Japan Health and Welfare Statistics Association; 2002 (in Japanese).
- Swerdlow AJ, De Stavola BL, Floderus B, Holm NV, Kaprio J, Verkasalo PK, Mack T. Risk factors for breast cancer at young ages in twins: An international population-based study. *J Natl Cancer Inst* 94:1238-1246; 2002.
- Terao T, Soeda S, Yoshimura R, Nakamura J, Iwata N. Effect of latitude on suicide rates in Japan. [letter] *Lancet* 360:1892; 2002.
- Terry P, Lichtenstein P, Feychting M, Ahlbom A, Wolk A. Fatty fish consumption and risk of prostate cancer. *Lancet* 357:1764-1766; 2001.
- The Cabinet Office. Prefectural economic calculation. Tokyo: The Cabinet Office; 1990 (in Japanese).
- The Study Circle for Health and Nutrition Information. The National Nutrition Survey in Japan 2001. Tokyo: Daiichi-Shuppan; 2003 (in Japanese).
- Tsugane S, Sasaki S, Kobayashi M, Tsubono Y, Sobue T. Dietary habits among the JPHC study participants at baseline survey. Japan Public Health Center-based Prospective Study on Cancer and Cardiovascular Diseases. *J Epidemiol* 11(6 Suppl):S30-S43; 2001.
- Vandewalle B, Wattez N, Lefebvre J. Effects of vitamin D3 derivatives on growth, differentiation and apoptosis in tumoral colonic HT 29 cells: Possible implication of intracellular calcium. *Cancer Lett* 97:99-106; 1995.
- Watanabe S, Arimoto H. Standardized mortality rates of cancer by prefecture in 1979-1981 and 1984-1986 in Japan. *Jpn J Clin Oncol* 20:316-337; 1990.
- Yamamoto S, Sobue T, Kobayashi M, Sasaki S, Tsugane S. Japan Public Health Center-based prospective study on cancer cardiovascular diseases group. Soy, isoflavones, and breast cancer risk in Japan. *J Natl Cancer Inst* 95:906-913; 2003.



APPENDIX B1. Daily intake of selected foods or nutrients by district in Japan.

District number	Prefecture number ^a	Fat (g) ^b	Animal protein (g) ^b	Salt (g) ^b	Fish (g) ^b	Vitamin D (mg) ^c
1	01	56.7	44.2	12.7	50.1	11.4
2	02,03,04,05,06,07	57.1	41.5	13.5	49.6	10.9
3	11,12,13,14	59.5	42.3	12.5	40.7	8.5
4	08,09,10,19,20	55.5	39.9	13.6	40.4	10.2
5	15,16,17,18	56.8	39.5	12.8	45.2	8.4
6	21,22,23,24	55.9	39.8	12.1	40.1	7.8
7	26,27,28	57.5	42.6	11.8	39.3	8.4
8	25,29,30	57.5	43.6	13.4	42.9	9.5
9	31,32,33,34,35	56.6	41.8	12.5	48.0	7.3
10	36,37,38,39	55.6	42.4	12.3	44.0	7.7
11	40,41,42,44	54.7	39.5	11.6	42.6	6.2
12	43,45,46,47	53.5	40.8	13.0	55.8	6.5

^a Refer to the first column of Appendix A1.

^b The National Nutrition Survey in Japan 1990.

^c The National Nutrition Survey in Japan 2000.



Dietary Patterns and Colorectal Adenomas in Japanese Men

The Self-Defense Forces Health Study

Tetsuya Mizoue¹, Taiki Yamaji¹, Shinji Tabata^{1,2}, Keizo Yamaguchi^{1,3}, Eiichi Shimizu³,
Masamichi Mineshita³, Shinsaku Ogawa², and Suminori Kono¹

¹ Department of Preventive Medicine, Faculty of Medical Sciences, Kyushu University, Higashiku, Fukuoka, Japan.

² Self-Defense Forces Fukuoka Hospital, Fukuoka, Japan.

³ Self-Defense Forces Kumamoto Hospital, Kumamoto, Japan.

Received for publication July 7, 2004; accepted for publication September 10, 2004.

The role of dietary patterns in colorectal carcinogenesis remains unclear in Asian populations. Using 1999–2002 data, the authors investigated the association between dietary patterns and colorectal adenomas in 1,341 Japanese men who underwent total colonoscopy. Information about diet was obtained using a 74-item food frequency questionnaire prior to the colonoscopy. Three dietary patterns were generated by factor analysis: 1) a high-dairy, high-fruit and -vegetable, high-starch, low-alcohol pattern; 2) an “animal food” pattern; and 3) a Japanese pattern. Logistic regression analysis was used to estimate the odds ratio of having colorectal adenomas with the adjustment for potential confounding variables including body mass index, smoking, alcohol, and leisure-time physical activities. A significant inverse association was found for the high-dairy, high-fruit and -vegetable, high-starch, low-alcohol pattern; the odds ratios for the second, third, and fourth quartiles were 0.97 (95% confidence interval: 0.70, 1.36), 0.71 (95% confidence interval: 0.50, 1.01), and 0.62 (95% confidence interval: 0.43, 0.90), respectively, compared with the lowest ($p_{\text{trend}} = 0.003$). Similar associations were observed for larger adenomas or for each subsite of the colorectum. The Japanese and “animal food” patterns were not clearly associated with colorectal adenomas. A dietary pattern including greater consumption of dairy products and fruits and vegetables with low alcohol consumption may be associated with decreased risk of colorectal adenomas.

adenoma; cross-sectional studies; diet

Abbreviation: DFSA, high-dairy, high-fruit and -vegetable, high-starch, low-alcohol (dietary pattern).

Colorectal cancer is a major cause of cancer deaths in developed countries. Geographic and time-trend analyses, as well as migrant studies, strongly suggest that environmental factors, especially diet, play an important role in the pathogenesis of colorectal cancer (1–3). However, analytical epidemiologic studies have yielded conflicting findings; for example, a body of evidence suggesting a protective role of vegetables or dietary fiber (4) has been either challenged (5–8) or supported (9, 10) by recent large-scale studies. In Japan, colorectal cancer mortality has markedly increased over the last several decades (11) and is now among the highest levels in the world (12). Time-trend analysis has suggested that decreased consumption of dietary fibers (13)

or grains (14) may account for the increase in mortality. Yet it is largely unknown which lifestyle changes associated with Westernization or modernization have contributed to the rapid increase of colorectal cancer in Japan, or whether the traditional Japanese diet protects against this type of cancer.

Analysis of dietary patterns has recently drawn a great deal of attention as a method of investigating the role of foods or nutrients in studies of chronic diseases. Approaches of this sort, dealing with a combination of several foods, can overcome problems arising from close intercorrelation and potential effect modifications among numerous foods or nutrients (15). Factor-analysis studies of Western populations have suggested that a certain dietary pattern may be

Reprint requests to Dr. Tetsuya Mizoue, Department of Preventive Medicine, Faculty of Medical Sciences, Kyushu University, 3-1-1, Maidashi, Higashiku, Fukuoka 812-8582, Japan (e-mail: mizoue@phealth.med.kyushu-u.ac.jp).

predictive of colorectal cancer risk (16–18). Dietary patterns generated by factor analysis, however, are sample specific and may not be applicable to populations having different dietary cultures. While having adopted a Western-style diet, many Japanese still consume large amounts of traditional foods, including rice, fish, and soybean products (19). Thus, dietary patterns among Japanese may differ considerably from those among Western populations.

The aim of the present study was therefore to investigate dietary patterns in relation to the risk of colorectal adenoma, a precursor of colorectal cancer (20, 21), using data from preretirement check-ups among male Self-Defense Forces officials in Japan.

MATERIALS AND METHODS

Study setting

The data used were derived from the Self-Defense Forces Health Study, a cross-sectional survey of male Self-Defense Forces officials who participated in a preretirement health examination at two hospitals (Fukuoka and Kumamoto) in Japan. The study procedure has been described elsewhere (22, 23). In short, all officials undergo a comprehensive health examination before retirement; total colonoscopy is included as a routine procedure. Study questionnaires about health-related lifestyles were distributed prior to colonoscopy to male examinees on the first day of hospital admission for examination. Research assistants checked the questionnaire for unanswered questions and apparently inconsistent answers and, if necessary, sought clarification from the study subjects.

Results of laboratory tests and colonoscopic findings, including histologies for polyp, were extracted from clinical reports. Written informed consent was obtained from study participants. The study protocol has been approved by the ethics committee of Kyushu University.

Study subjects

The present study used data from April 1999 through March 2002. Among 2,390 male Self-Defense Forces officials who underwent the examination, 2,370 (99 percent) agreed to participate in the present study. After excluding men with histories of cancer, stroke, myocardial infarction, coronary revascularization, inflammatory bowel diseases, colorectal surgery, or diabetes mellitus, we kept 2,141 men in the analysis of dietary patterns. Of these, we excluded men who did not receive colonoscopy ($n = 57$), who underwent partial or unsuccessful colonoscopy ($n = 177$), or who had colorectal polyp removal prior to the examination ($n = 148$). Of the remaining 1,759 subjects who completed total colonoscopy, 764 men were identified as having colorectal polyps including hyperplastic nodules. Of these, 476 men had their polyps histologically confirmed: cancer ($n = 1$), carcinoid ($n = 1$), adenoma ($n = 346$), and other histologies ($n = 128$). Only 29 men had adenomas of 10 mm or larger, and nine had tubulovillous or villous adenomas. The data for the 346 men who had adenoma (case group) and 995 men who were free from any colorectal polyp and cancer

(referent group) were analyzed to assess the association between dietary patterns and colorectal adenomas.

Dietary assessment

Information about diet was collected using a food frequency questionnaire designed to assess the average intakes of 74 food items, food groups, and food preparations over the previous year. The questionnaire was an expanded version of a 45-item food frequency questionnaire that was developed on the basis of a published questionnaire (24) and was validated against the 28-day dietary record (25). The expansion of food items was done with reference to food consumption in the National Nutrition Survey (19) and a dietary questionnaire developed elsewhere in Japan (26). Participants were asked to choose from seven response options for most dietary items, ranging from "never/less than one per month" to "two to three times per day." Different response schemes were used for green tea, coffee, and rice (five options) and for alcoholic beverages (six options). Daily consumers of green tea, coffee, or rice were asked about the number of cups or bowls consumed per day. Current drinkers, defined as those who have consumed alcoholic beverages weekly for at least 1 year in their lifetime and who were drinking at the time of the survey, were asked about the frequency of consumption and the amount of consumption per occasion of five alcoholic beverages, that is, sake (a Japanese wine), shochu (a Japanese distilled beverage), beer, whiskey, and wine. The amount of consumption per occasion was used in the estimation of total ethanol intake from these alcoholic beverages, but only the frequency of consumption for each alcoholic beverage was used in the analysis of dietary patterns.

Grouping of food factors

Before the analysis of dietary patterns, intakes of green tea, coffee, or rice were converted into units of cups or bowls per day, while those of other dietary items were quantified in terms of frequency per week. Five dietary questions that overlapped with or were duplicated by others (collective consumption of cooked vegetables, apples, mandarin oranges, other oranges, watermelons) and three questions about food spreads (butter, margarine, and jam/honey) were not used. Furthermore, some foods or food groups similar in nutritional content or culinary use were combined, leaving 39 food items for the purposes of the present study.

Statistical analysis

Dietary patterns were generated by factor analysis (principal components) using SAS PROC FACTOR statistical software (27). Factor analysis is a technique to reduce a number of variables into fewer independent factors. To make interpretation easier, a linear transformation called a "rotation" is normally performed on the initial factor solution. We used an orthogonal rotation procedure (varimax rotation), which maintains the uncorrelated nature of the factors and tries to get the original variables to load high on one of the factors and low on the rest. When factor scores are used as

independent variables in a subsequent regression analysis, this procedure has the advantage over oblique rotation that the analysis is less subject to problems of collinearity. In determining the number of factors to retain, we consider eigenvalue, the scree test, and interpretability. Eleven factors satisfied the criteria for eigenvalues greater than one, and the scree plot showed small breaks in the eigenvalues after factor 5, suggesting three or four factors to retain. Post-rotated factor loadings revealed that three factors well describe distinctive dietary patterns of the study population.

We thus retained the three dietary patterns and designated them as 1) a high-dairy, high-fruit and -vegetable, high-starch, low-alcohol (DFSAs) pattern; 2) an "animal food" pattern; and 3) a Japanese pattern, according to the food items showing high loading (absolute value) with respect to each dietary pattern. We confirmed that these three dietary factors emerged when all 74 food items in our questionnaire were simply included in factor analysis. A factor score for each dietary pattern was calculated by weighting consumption of each food item by the corresponding factor loading and summing the resulting values. This score ranks individuals in terms of how closely they conform to the dietary pattern.

The potential confounding variables considered were hospital (Fukuoka or Kumamoto), age (treated as a continuous variable), parental history of colorectal cancer (absent or present), occupational rank (three categories), body mass index (<22, 22–23.9, 24–25.9, and ≥ 26 kg/m²), smoking (lifetime nonsmoker, former smoker, and current smoker using <15, 15–24, or ≥ 25 cigarettes/day), and leisure-time physical activity, expressed as the sum of metabolic equivalents for each activity multiplied by the corresponding hours of such activity per week (none, <20, 20–39.9, and ≥ 40 metabolic equivalent-hours). Quartiles of factor scores of each dietary pattern among controls were used for cutoff values. Multiple logistic regression that included terms for the above-mentioned variables was performed to estimate the odds ratio and 95 percent confidence interval of colorectal adenomas according to quartiles of scores for each dietary pattern, taking the lowest quartile group as the referent group. Analyses were repeated for adenomas of 5 mm or larger ($n = 140$) or according to the location of the lesion (proximal colon including the cecum, ascending colon, liver flexure, transverse colon, and splenic flexure; distal colon including the descending colon and sigmoid colon; and the rectum). Logistic regression analysis was performed using SAS PROC LOGISTIC software (27).

RESULTS

Table 1 shows factor loadings, which are equivalent to simple correlations between the food items and the dietary patterns. A positive loading indicates that the food item is positively associated with the dietary pattern, and a negative loading indicates an inverse association with the dietary pattern. The DFSAs dietary pattern was characterized by frequent intake of fermented dairy products, milk, confectionaries, bread, fruits, and vegetables and infrequent intake of shochu, a local alcoholic beverage in the study areas. The "animal food" dietary pattern was characterized by various

TABLE 1. Factor-loading matrix for dietary patterns, Self-Defense Forces Health Study, Japan, 1999–2002*

	DFSAs† dietary pattern	"Animal food" dietary pattern	Japanese dietary pattern
Fermented dairy products	0.61	–	–
Confectionaries	0.55	0.18	–
Canned fruits	0.52	–	–
Bread	0.47	–	–0.39
Fruits (not canned)	0.47	–	0.21
Fruit juices	0.47	–	–
Vegetable juice	0.41	–	0.17
Milk	0.40	–	–
Oil dressing	0.33	0.19	0.26
Soda, cola	0.30	0.20	–0.18
Shochu (alcoholic beverage)	–0.40	0.15	0.24
Red meat	–	0.68	–
Poultry	–	0.63	–
Fried foods	0.25	0.49	0.29
Broiled fish/meat	–	0.48	0.32
Seafood (except fish)	–	0.47	0.18
Processed meat	0.17	0.46	–
Processed fish	–	0.41	0.18
Gyoza‡	–	0.40	–
Liver	–	0.38	–
Eggs	–	0.34	0.22
Noodles	–	0.34	–
Soybean products	–	–	0.64
Cooked vegetables	0.36	0.23	0.56
Seaweed	0.27	–	0.55
Raw vegetables	0.45	–	0.52
Pickles	0.19	–	0.51
Green tea	–	–0.15	0.46
Fish	–	0.27	0.38
Potatoes	0.33	0.24	0.35
Garlic	0.20	–	0.32
Variance explained (%)	8.5	7.9	7.7

* Factor loadings are equivalent to simple correlations between the food items and the dietary patterns. Factor loadings less than ± 0.15 were indicated by a dash; food items with factor loadings less than ± 0.30 for all dietary patterns (rice, mayonnaise, nuts, coffee, wine, beer, whiskey, sake) were omitted.

† DFSAs, high-dairy, high-fruit and -vegetable, high-starch, low-alcohol (dietary pattern).

‡ Dumpling with minced pork and vegetable stuffing.

kinds of animal foods, including red meat, poultry, seafood excluding fish, processed meat and fish products, and fried or broiled foods. The Japanese dietary pattern was characterized by traditional foods in Japan (soybean products, seaweed, pickles, and green tea), vegetables, and fish. The proportion of the total variance explained by the three factors was 24 percent.

TABLE 2. Dietary patterns in relation to potential confounding variables and alcohol intake among referents, Self-Defense Forces Health Study, Japan, 1999–2002

Dietary patterns	Hospital (% Kumamoto)	Age (mean years)	Rank (% highest)	Parental history of colorectal cancer (%)	Body mass index (mean kg/m ²)	Smoking (% current smokers)	Physical activity (median metabolic equivalent- hours)	Alcohol (median ml/day)*
DFSA† dietary pattern								
Quartile 1 (low)	28	52.4	10	6	23.7	45	15	62
Quartile 2, 3	27	52.4	14	4	24.0	38	16	32
Quartile 4 (high)	18	52.4	19	5	23.5	40	16	14
<i>P</i> _{trend} ‡	0.01	0.49	<0.01	0.66	0.46	0.26	0.84	<0.01
"Animal food" dietary pattern								
Quartile 1 (low)	30	52.5	15	4	23.6	40	16	14.5
Quartile 2, 3	24	52.4	14	4	23.8	42	16	34
Quartile 4 (high)	22	52.3	13	6	23.9	37	15	49
<i>P</i> _{trend}	0.05	0.23	0.43	0.52	0.15	0.56	0.87	<0.01
Japanese dietary pattern								
Quartile 1 (low)	21	52.4	17	5	23.7	48	9.5	21
Quartile 2, 3	25	52.3	13	4	23.8	39	16	40
Quartile 4 (high)	29	52.5	14	4	23.9	36	19	38
<i>P</i> _{trend}	0.02	0.58	0.51	0.66	0.51	<0.01	<0.01	<0.01

* Estimated from the consumption of five alcoholic beverages: beer, sake, shochu, wine, and whiskey.

† DFSA, high-dairy, high-fruit and -vegetable, high-starch, low-alcohol (dietary pattern).

‡ Mantel-Haenszel chi-squared test for categorical variables and linear regression analysis for continuous variables, assigning to categories of each dietary pattern their median scores (physical activity and alcohol consumption were log transformed).

Table 2 shows the association of dietary patterns with potential confounding variables and alcohol consumption among men free from colorectal polyp or cancer (referent group). Examinees at the Kumamoto hospital had a higher score for the Japanese dietary pattern but lower scores for the DFSA and "animal food" dietary patterns than those at the Fukuoka hospital. This reflects the geographic characteristics of dietary patterns; the southern parts of Kyushu Island, including Kumamoto, are less urbanized than the northern parts, including Fukuoka. Men with a high score for the DFSA dietary pattern tended to have higher occupational positions and consumed smaller amounts of alcohol. Men with high scores for the "animal food" dietary pattern tended to consume greater amounts of alcohol. Men in the upper quartiles of the Japanese dietary pattern tended to be nonsmokers and engaged in higher levels of leisure-time physical activity, and they consumed greater amounts of alcohol.

As shown in table 3, the DFSA dietary pattern was inversely associated with the risk of colorectal adenomas, showing a 40 percent reduced odds ratio among men in the highest quartile of the dietary pattern compared with those in the lowest. This association was slightly more evident for adenomas with a diameter of 5 mm or larger. No apparent association was observed for either the "animal food" dietary pattern or the Japanese dietary pattern.

The DFSA dietary pattern was inversely associated with adenomas at all subsites of the colorectum (table 4). The

association was slightly stronger for the proximal colon in terms of the odds ratio of 0.5 for the highest quartile of the dietary pattern score and test for the trend association ($P_{\text{trend}} = 0.003$), but the confidence intervals of odds ratios for this site overlapped substantially with those for other sites. The Japanese and "animal food" dietary patterns were not measurably associated with colon adenomas. However, a nonsignificant positive association with rectal adenomas was observed for the Japanese pattern, while a nonsignificant inverse association was found for the "animal food" pattern. The odds ratios for the upper three quartiles combined compared with the lowest were 1.64 (95 percent confidence interval: 0.83, 3.25) and 0.64 (95 percent confidence interval: 0.36, 1.13) for the Japanese pattern and "animal food" pattern, respectively.

DISCUSSION

We investigated the association between major dietary patterns and colorectal adenomas among middle-aged Japanese men. Of the three dietary patterns we identified, the DFSA dietary pattern showed a significant, inverse association with the risk of colorectal adenomas.

Strengths and limitations

Our study had several strengths. Selection bias in terms of study participation was unlikely because of nonselective

TABLE 3. Logistic regression results for the association between dietary patterns and colorectal adenoma, Self-Defense Forces Health Study, Japan, 1999–2002

Dietary pattern	Quartile*							<i>P</i> _{trend}
	1 (low)	2		3		4 (high)		
		Odds ratio†	95% confidence interval	Odds ratio	95% confidence interval	Odds ratio	95% confidence interval	
DFSA‡ dietary pattern								
Adenoma of any size	1.00	0.97	0.70, 1.36	0.71	0.50, 1.01	0.62	0.43, 0.90	0.003
Adenoma of 5 mm or larger	1.00	0.84	0.52, 1.34	0.68	0.41, 1.12	0.59	0.35, 0.996	0.04
"Animal food" dietary pattern								
Adenoma of any size	1.00	0.87	0.61, 1.23	0.91	0.64, 1.28	0.86	0.60, 1.23	0.49
Adenoma of 5 mm or larger	1.00	1.05	0.64, 1.72	0.84	0.50, 1.41	0.98	0.59, 1.63	0.75
Japanese dietary pattern								
Adenoma of any size	1.00	0.96	0.67, 1.38	1.13	0.79, 1.61	1.18	0.83, 1.69	0.26
Adenoma of 5 mm or larger	1.00	1.00	0.59, 1.70	1.11	0.66, 1.86	1.24	0.75, 2.08	0.36

* Among referents.

† Adjusted for hospital, age, parental history of colorectal cancer, occupational rank, body mass index, smoking, and leisure-time physical activity.

‡ DFSA, high-dairy, high-fruit and -vegetable, high-starch, low-alcohol (dietary pattern).

recruitment for the preretirement health examination, which included total colonoscopy as a routine procedure, and high study participation rate. The questionnaire was distributed and collected prior to colonoscopy, and thus recall bias associated

with adenoma status was also unlikely. The control series consisted of only subjects who were confirmed via total colonoscopy to be free from any colorectal polyp and cancer, leading to a more valid assessment compared with studies

TABLE 4. Logistic regression results for the association between dietary patterns and colorectal adenoma according to the location of the lesion, Self-Defense Forces Health Study, Japan, 1999–2002

Dietary pattern	Quartile*							<i>P</i> _{trend}
	1 (low)	2		3		4 (high)		
		Odds ratio†	95% confidence interval	Odds ratio	95% confidence interval	Odds ratio	95% confidence interval	
Colon adenoma (n = 299)								
DFSA‡ dietary pattern	1.00	0.93	0.66, 1.32	0.70	0.48, 1.01	0.59	0.40, 0.87	0.003
"Animal food" dietary pattern	1.00	0.93	0.64, 1.35	0.97	0.67, 1.40	0.95	0.65, 1.38	0.85
Japanese dietary pattern	1.00	0.93	0.64, 1.37	1.09	0.76, 1.59	1.11	0.77, 1.62	0.45
Proximal colon adenoma (n = 158)								
DFSA dietary pattern	1.00	1.00	0.64, 1.54	0.67	0.41, 1.09	0.50	0.30, 0.85	0.003
"Animal food" dietary pattern	1.00	0.84	0.51, 1.39	1.08	0.68, 1.73	0.94	0.57, 1.53	0.95
Japanese dietary pattern	1.00	0.87	0.53, 1.42	0.92	0.56, 1.49	1.08	0.67, 1.74	0.70
Distal colon adenoma (n = 171)								
DFSA dietary pattern	1.00	1.00	0.64, 1.54	0.77	0.48, 1.23	0.68	0.42, 1.11	0.08
"Animal food" dietary pattern	1.00	1.01	0.63, 1.62	0.95	0.59, 1.52	1.01	0.63, 1.62	0.97
Japanese dietary pattern	1.00	1.10	0.67, 1.79	1.43	0.90, 2.28	1.21	0.74, 1.96	0.35
Rectal adenoma (n = 63)								
DFSA dietary pattern	1.00	0.94	0.48, 1.84	0.64	0.30, 1.36	0.71	0.34, 1.48	0.26
"Animal food" dietary pattern	1.00	0.66	0.33, 1.34	0.64	0.31, 1.32	0.62	0.30, 1.28	0.22
Japanese dietary pattern	1.00	1.58	0.71, 3.51	1.56	0.70, 3.47	1.79	0.82, 3.92	0.18

* Among referents.

† Adjusted for hospital, age, parental history of colorectal cancer, occupational rank, body mass index, smoking, and leisure-time physical activity.

‡ DFSA, high-dairy, high-fruit and -vegetable, high-starch, low-alcohol (dietary pattern).

based on partial colonoscopy. We controlled for major known or suspected confounding factors. The uniform background of the study subjects in terms of occupation, sex, and age was also advantageous in maintaining comparability between cases and controls, although this uniformity limits the extent to which we may generalize from the present findings.

The present study also features some limitations. For one, the dietary questionnaire has not been validated. However, the former version, including questions and response options similar to those of the present questionnaire, has been validated against 7-day, year-round dietary records (25). Most nutrients and foods demonstrated fairly good correlation between the dietary record and questionnaire; relatively high correlation coefficients of 0.80, 0.77, 0.58, and 0.58 were observed for bread, fruits, dairy products, and pickled vegetables, respectively. Nondifferential misclassification in our dietary assessment could distort risk estimates toward the null. Such a bias may be minimal for the analysis of the DFSA dietary pattern, composed of food items showing good correlation between the dietary record and questionnaire, but this bias could be the reason for the lack of an apparent association with the "animal food" or Japanese dietary pattern.

Limitations of factor analysis arise from the arbitrary decisions (15) involved in selecting and grouping foods for analysis from the questionnaire, in determining the number of factors to retain, in choosing the method of rotation of the initial factors to increase the interpretability of the dietary pattern, and in labeling dietary patterns according to their factor loadings. Masaki et al. (28) identified four major dietary patterns using baseline data of a cohort of male employees in Tokyo. Similar to our study, their study identified a "Western breakfast" dietary pattern and an "animal" dietary pattern, suggesting the existence of dietary patterns common to the Japanese. Our derived dietary patterns accounted for 24 percent of the total variance, which is comparable with a figure observed in a previous study (17) but less than that reported in a Japanese study (28). Caution needs to be exercised when comparing the variance explained across studies, which is determined by various factors including the number of variables in analysis.

Interpretation of findings

A dietary pattern characterized by frequent intakes of dairy products, confectionaries, and fruits and vegetables, as well as by infrequent consumption of *shochu*, a local alcoholic beverage, was inversely associated with the risk of colorectal adenomas. This dietary pattern seems to consist of relatively healthy selections of foods found in Western countries and includes foods of probably low consumption in Japan. According to the food balance sheet (29), per capita supplies of dairy products in Japan and among developed countries in the year 2001 were 66 kg (181 g/day) and 197 kg (540 g/day), respectively; the corresponding values for fruits were 53 kg (145 g/day) and 83 kg (227 g/day). Although the associations between these foods and colorectal cancer have been inconsistent, the present results are in agreement with the existing body of evidence, including findings from recent studies, indicating that high consumption of dairy products

or calcium (30–34) and high consumption of fruits, fruit juices, or fruit fiber (10, 35–38) are each associated with reduced risk of colorectal cancer or adenoma. A positive association of alcohol consumption and colorectal adenomas or cancer has been reported in many studies, including those in Japan (23, 39). In addition to independent effects, there may be complex interactions among food factors constituting the DFSA dietary pattern. For example, fruit juices may enhance calcium absorption (40), and reduced alcohol intake increases the bioavailability of folate (41). The glycemic effects of a high-starch or a high-sugar diet and their contribution to increased risk of colorectal cancer have been suspected, but epidemiologic findings are inconsistent on this point (42, 43). Our finding of an inverse association between the DFSA dietary pattern and colorectal adenomas provides the following suggestions: that a high-starch diet may inhibit, rather than promote, the formation of colorectal adenomas and that the adverse effects of a high-starch diet, if any, may not be so strong as to negate the protective effects of other foods contributing to this dietary pattern on adenoma risk. The inverse association of this dietary pattern with adenoma risk was somewhat stronger for the proximal colon than for other sites. Random error could be an explanation. Alternatively, the dietary pattern may be more closely involved in the formation of adenomas in the proximal colon.

The Japanese dietary pattern was characterized by high consumption of many plant foods, including traditional Japanese foods (soybean products, seaweed, pickles) and vegetables. A diet rich in various plant foods could potentially reduce cancer risk because of their many biologically active chemicals (44). However, the Japanese dietary pattern was not apparently associated with colonic adenomas. In studies of Western populations, inverse associations between similar dietary patterns (designated "prudent" or "healthy" patterns) and colorectal cancer have been unclear (18) or limited to subgroups (17). The lack of such an association in our study may contradict a body of evidence supporting an inverse association between vegetables and colorectal cancer (4) but agree with results of recent prospective, but not all (10, 37), studies reporting no association between vegetables or fiber and colorectal cancer or adenoma (5–8). As most of the adenomas in the present study were small in size and less malignant in nature, the present finding is in line with a hypothesis that vegetables are inversely associated with the progression of colorectal adenomas to cancer but not with the initial appearance of adenomas (38). We found a nonsignificant positive association between the Japanese dietary pattern and the risk of rectal adenomas. Studies in Japan (45–49) have consistently shown that frequent consumption of preserved foods including pickled vegetables and dried/salted fish, typical of the Japanese diet, is associated with increased risk of colorectal cancer; of these studies, three documented a significant association specifically for the rectum (46–48). These preserved foods contain *N*-nitroso compounds (50), which are potent carcinogens (51). Among other foods characterizing the Japanese pattern, broiled fish is a potential source of exposure to carcinogenic heterocyclic amine (52), although we are not aware of any epidemiologic

findings suggesting a relation between broiled fish and colorectal cancer risk.

Meat, especially red meat, processed meat, or meat broiled at high temperature, has been associated with colorectal cancer (53, 54) or adenoma (55). A study in Japan found a significant positive association between intake of animal protein and the risk of colorectal adenoma (56). However, we find no increase in the risk of colorectal adenomas associated with the "animal food" dietary pattern. Besides possible bias due to misclassification in the dietary assessment (as discussed above), the lack of such an association for colon adenomas in our study may be attributable to the moderate consumption of meat in Japan (mean daily intake of total meat: 96 g for men aged 40–49 years (19)). In addition, poultry has contributed to the healthy or prudent dietary patterns in Western populations (16–18). The diversity of animal food sources may dilute the potential carcinogenic effects of a specific animal food. Furthermore, it is possible that moderate intake of animal foods prevents carcinogenesis because these foods provide nutrients such as methionine and folate, which are beneficial in DNA synthesis and DNA methylation (57). In this context, our finding showing an increased risk of rectal adenomas associated with the lowest quartile of the "animal food" pattern may be of note and is consistent with results of certain studies relating to colorectal cancer in Japan (45, 48, 58).

In conclusion, the present results indicate that a dietary pattern characterized by frequent consumption of dairy products, confectionaries, bread, fruits, and vegetables but low intake of local alcoholic beverages is associated with a reduced risk of colorectal adenomas in Japanese men. Nonsignificant associations for rectal adenomas, based on an analysis including only 63 men with adenomas in the rectum, should set a limit to causal inference. However, since the incidence of rectal cancer in Japan has been high among industrial countries (12), the question as to whether a Japanese-style diet or a diet low in animal foods promotes carcinogenesis of the rectum warrants further investigation.

ACKNOWLEDGMENTS

This work was supported by a grant-in-aid for scientific research on priority areas (12218226) from the Ministry of Education, Culture, Sports, Science, and Technology, Japan, and by a Health and Labor Sciences research grant for research on cancer prevention and health services research from the Ministry of Health, Labor, and Welfare, Japan.

The authors thank the ward nurses of Self-Defense Forces Fukuoka and Kumamoto hospitals for their assistance in conducting the study.

REFERENCES

- Buell P, Dunn JE. Cancer mortality among Japanese Issei and Nisei of California. *Cancer* 1965;18:656–64.
- Tominaga S. Cancer incidence in Japanese in Japan, Hawaii, and western United States. *Natl Cancer Inst Monogr* 1985;69:83–92.
- Shimizu H, Mack TM, Ross RK, et al. Cancer of the gastrointestinal tract among Japanese and white immigrants in Los Angeles County. *J Natl Cancer Inst* 1987;78:223–8.
- Glade MJ. Food, nutrition, and the prevention of cancer: a global perspective. American Institute for Cancer Research/World Cancer Research Fund, American Institute for Cancer Research. *Nutrition* 1999;15:523–6.
- Michels KB, Giovannucci E, Joshipura KJ, et al. Prospective study of fruit and vegetable consumption and incidence of colon and rectal cancers. *J Natl Cancer Inst* 2000;92:1740–52.
- Fuchs CS, Giovannucci EL, Colditz GA, et al. Dietary fiber and the risk of colorectal cancer and adenoma in women. *N Engl J Med* 1999;340:169–76.
- Schatzkin A, Lanza E, Corle D, et al. Lack of effect of a low-fat, high-fiber diet on the recurrence of colorectal adenomas. Polyp Prevention Trial Study Group. *N Engl J Med* 2000;342:1149–55.
- Alberts DS, Martinez ME, Roe DJ, et al. Lack of effect of a high-fiber cereal supplement on the recurrence of colorectal adenomas. Phoenix Colon Cancer Prevention Physicians' Network. *N Engl J Med* 2000;342:1156–62.
- Bingham SA, Day NE, Luben R, et al. Dietary fibre in food and protection against colorectal cancer in the European Prospective Investigation into Cancer and Nutrition (EPIC): an observational study. *Lancet* 2003;361:1496–501.
- Peters U, Sinha R, Chatterjee N, et al. Dietary fibre and colorectal adenoma in a colorectal cancer early detection programme. *Lancet* 2003;361:1491–5.
- Statistics and Information Department, Minister's Secretariat, Ministry of Health, Labor, and Welfare. Age-adjusted death rates by prefecture. In: Special report on vital statistics, 2000. (In Japanese). Tokyo, Japan: Japan Health and Welfare Statistics Association, 2002.
- Parkin DM, Whelan SL, Ferlay J, et al, eds. Cancer incidence in five continents. Vol VIII. Lyon, France: International Agency for Research on Cancer, 2002. (IARC publication no. 155).
- Honda T, Kai I, Ohi G. Fat and dietary fiber intake and colon cancer mortality: a chronological comparison between Japan and the United States. *Nutr Cancer* 1999;33:95–9.
- Kono S, Ahn YO. Vegetables, cereals and colon cancer mortality: long-term trend in Japan. *Eur J Cancer Prev* 2000;9:363–5.
- Jacques PF, Tucker KL. Are dietary patterns useful for understanding the role of diet in chronic disease? *Am J Clin Nutr* 2001;73:1–2.
- Slattery ML, Boucher KM, Caan BJ, et al. Eating patterns and risk of colon cancer. *Am J Epidemiol* 1998;148:4–16.
- Terry P, Hu FB, Hansen H, et al. Prospective study of major dietary patterns and colorectal cancer risk in women. *Am J Epidemiol* 2001;154:1143–9.
- Fung T, Hu FB, Fuchs C, et al. Major dietary patterns and the risk of colorectal cancer in women. *Arch Intern Med* 2003;163:309–14.
- Bureau of Public Health, Ministry of Health and Welfare. The national nutrition survey in Japan, 1996. (In Japanese). Tokyo, Japan: Daiichi Shuppan, 1998.
- Hill MJ, Morson BC, Bussey HJ. Aetiology of adenoma-carcinoma sequence in large bowel. *Lancet* 1978;1:245–7.
- Vogelstein B, Fearon ER, Hamilton SR, et al. Genetic alterations during colorectal-tumor development. *N Engl J Med* 1988;319:525–32.
- Kono S, Handa K, Hayabuchi H, et al. Obesity, weight gain and risk of colon adenomas in Japanese men. *Jpn J Cancer Res* 1999;90:805–11.
- Toyomura K, Yamaguchi K, Kawamoto H, et al. Relation of cigarette smoking and alcohol use to colorectal adenomas by

- subsite: the Self-Defense Forces Health Study. *Cancer Sci* 2004;95:72-6.
24. Shirota T, Yoshizumi F. A study on convenient dietary assessment. (In Japanese). *Nippon Koshu Eisei Zasshi* 1990;37:100-8.
 25. Lee KY, Uchida K, Shirota T, et al. Validity of a self-administered food frequency questionnaire against 7-day dietary records in four seasons. *J Nutr Sci Vitaminol* 2002;48:467-76.
 26. Tokudome S, Ikeda M, Tokudome Y, et al. Development of data-based semi-quantitative food frequency questionnaire for dietary studies in middle-aged Japanese. *Jpn J Clin Oncol* 1998;28:679-87.
 27. SAS Institute, Inc. *SAS/STAT user's guide*, version 6. 4th ed. Vol 2. Cary, NC: SAS Institute, Inc, 1989.
 28. Masaki M, Sugimori H, Nakamura K, et al. Dietary patterns and stomach cancer among middle-aged male workers in Tokyo. *Asian Pac J Cancer Prev* 2003;4:61-6.
 29. FAOSTAT nutritional data. Food balance sheets. Rome, Italy: Food and Agriculture Organization of the United Nations, 2004. (http://www.fao.org/waicent/portal/statistics_en.asp).
 30. Martinez ME, Willett WC. Calcium, vitamin D, and colorectal cancer: a review of the epidemiologic evidence. *Cancer Epidemiol Biomarkers Prev* 1998;7:163-8.
 31. Baron JA, Beach M, Mandel JS, et al. Calcium supplements for the prevention of colorectal adenomas. Calcium Polyp Prevention Study Group. *N Engl J Med* 1999;340:101-7.
 32. Pietinen P, Malila N, Virtanen M, et al. Diet and risk of colorectal cancer in a cohort of Finnish men. *Cancer Causes Control* 1999;10:387-96.
 33. Wu K, Willett WC, Fuchs CS, et al. Calcium intake and risk of colon cancer in women and men. *J Natl Cancer Inst* 2002;94:437-46.
 34. McCullough ML, Robertson AS, Rodriguez C, et al. Calcium, vitamin D, dairy products, and risk of colorectal cancer in the Cancer Prevention Study II Nutrition Cohort (United States). *Cancer Causes Control* 2003;14:1-12.
 35. Sandler RS, Lyles CM, Peipins LA, et al. Diet and risk of colorectal adenomas: macronutrients, cholesterol, and fiber. *J Natl Cancer Inst* 1993;85:884-91.
 36. Platz EA, Giovannucci E, Rimm EB, et al. Dietary fiber and distal colorectal adenoma in men. *Cancer Epidemiol Biomarkers Prev* 1997;6:661-70.
 37. Terry P, Giovannucci E, Michels KB, et al. Fruits, vegetables, dietary fiber, and risk of colorectal cancer. *J Natl Cancer Inst* 2001;93:525-33.
 38. Smith-Warner SA, Elmer PJ, Fosdick L, et al. Fruits, vegetables, and adenomatous polyps: the Minnesota Cancer Prevention Research Unit case-control study. *Am J Epidemiol* 2002;155:1104-13.
 39. Otani T, Iwasaki M, Yamamoto S, et al. Alcohol consumption, smoking, and subsequent risk of colorectal cancer in middle-aged and elderly Japanese men and women: Japan Public Health Center-based Prospective Study. *Cancer Epidemiol Biomarkers Prev* 2003;12:1492-500.
 40. Andon MB, Peacock M, Kanerva RL, et al. Calcium absorption from apple and orange juice fortified with calcium citrate malate (CCM). *J Am Coll Nutr* 1996;15:313-16.
 41. Hillman RS, Steinberg SE. The effects of alcohol on folate metabolism. *Annu Rev Med* 1982;33:345-54.
 42. Slattery ML, Benson J, Berry TD, et al. Dietary sugar and colon cancer. *Cancer Epidemiol Biomarkers Prev* 1997;6:677-85.
 43. Terry PD, Jain M, Miller AB, et al. Glycemic load, carbohydrate intake, and risk of colorectal cancer in women: a prospective study. *J Natl Cancer Inst* 2003;95:914-16.
 44. Steinmetz KA, Potter JD. Vegetables, fruit, and cancer. I. Epidemiology. *Cancer Causes Control* 1991;2:325-57.
 45. Kondo R. Epidemiological study on cancer of the colon and the rectum. (In Japanese). *Nagoya Med J* 1975;97:93-116.
 46. Tajima K, Tominaga S. Dietary habits and gastro-intestinal cancers: a comparative case-control study of stomach and large intestinal cancers in Nagoya, Japan. *Jpn J Cancer Res* 1985;76:705-16.
 47. Hoshiyama Y, Sekine T, Sasaba T. A case-control study of colorectal cancer and its relation to diet, cigarettes, and alcohol consumption in Saitama Prefecture, Japan. *Tohoku J Exp Med* 1993;171:153-65.
 48. Inoue M, Tajima K, Hirose K, et al. Subsite-specific risk factors for colorectal cancer: a hospital-based case-control study in Japan. *Cancer Causes Control* 1995;6:14-22.
 49. Yang CX, Takezaki T, Hirose K, et al. Fish consumption and colorectal cancer: a case-reference study in Japan. *Eur J Cancer Prev* 2003;12:109-15.
 50. Scanlan RA. Formation and occurrence of nitrosamines in food. *Cancer Res* 1983;43(suppl):2435s-40s.
 51. Tricker AR, Preussmann R. Carcinogenic *N*-nitrosamines in the diet: occurrence, formation, mechanisms and carcinogenic potential. *Mutat Res* 1991;259:277-89.
 52. Ohgaki H, Hasegawa H, Kato T, et al. Carcinogenicities in mice and rats of IQ, MeIQ, and MeIQx. Princess Takamatsu Symp 1985;16:97-105.
 53. Sandhu MS, White IR, McPherson K. Systematic review of the prospective cohort studies on meat consumption and colorectal cancer risk: a meta-analytical approach. *Cancer Epidemiol Biomarkers Prev* 2001;10:439-46.
 54. Norat T, Lukanova A, Ferrari P, et al. Meat consumption and colorectal cancer risk: dose-response meta-analysis of epidemiologic studies. *Int J Cancer* 2002;98:241-56.
 55. Yoon H, Benamouzig R, Little J, et al. Systematic review of epidemiological studies on meat, dairy products and egg consumption and risk of colorectal adenomas. *Eur J Cancer Prev* 2000;9:151-64.
 56. Nagata C, Shimizu H, Kametani M, et al. Diet and colorectal adenoma in Japanese males and females. *Dis Colon Rectum* 2001;44:105-11.
 57. Giovannucci E. Epidemiologic studies of folate and colorectal neoplasia: a review. *J Nutr* 2002;132(suppl):2350s-5s.
 58. Kato I, Tominaga S, Matsuura A, et al. A comparative case-control study of colorectal cancer and adenoma. *Jpn J Cancer Res* 1990;81:1101-8.