Detection of Malondialdehyde DNA Adducts in Human Colorectal Mucosa: Relationship with Diet and the Presence of Adenomas

Chiara Leuratti, Mark A. Watson, Elliot J. Deag, Ailsa Welch, Rajinder Singh, Elke Gottschalg, Lawrence J. Marnett, Wendy Atkin, Nicholas E. Day, David E. G. Shuker, and Sheila A. Bingham

Medical Research Council Toxicology Unit, University of Leicester, Leicester LE1 9HN, United Kingdom; Norfolk and Norwich Health Care Trust, NR4 7UA, United Kingdom; [M. A. W.]; Strangeways Research Laboratory, European Prospective Investigation on Cancer (EPIC), University of Cambridge, Cambridge, CB1 4RN, United Kingdom [A. W.]; Department of Biochemistry, Vanderbilt University, Nashville, Tennessee 37232 [L. J. M.]; Imperial Cancer Research Fund Colorectal Cancer Unit, St. Mark’s Hospital, Middlesex, HA1 3UJ United Kingdom [W. A.]; and Medical Research Council Dunn Human Nutrition Centre, Cambridge, CB2 2X4 United Kingdom [S. A. B.

Abstract
Colorectal biopsies from normal mucosa of participants in the United Kingdom Flexible Sigmoidoscopy Trial and European Prospective Investigation on Cancer (EPIC; n = 162) were analyzed for the presence of malondialdehyde-deoxyguanosine (M₁-dG), a DNA adduct derived from lipid peroxidation. The aim was to investigate whether dietary factors can modulate M₁-dG levels and whether M₁-dG in normal mucosa is a risk factor for colorectal adenomas. Samples were analyzed using a sensitive immunoslot blot assay. This study has shown for the first time that M₁-dG is present in human colorectal tissue. M₁-dG levels ranged from undetectable (n = 13) to 12.23 per 10⁷ total bases. Mean levels were 4.3 ± 3 and 4.6 ± 2.9 per 10⁷ total bases in men and women, respectively. In men, there were positive associations of adduct levels with height and age, and inverse associations with body mass index. Legumes, fruit, salad, and whole meal bread were inversely associated with M₁-dG adducts, whereas consumption of offal, white meat, beer, and alcohol were positively associated with elevated levels. In women, there was an inverse association of the adduct with the ratio of polyunsaturated:saturated fatty acids (P = 0.019) and a weak positive correlation with saturated fat (P < 0.061). When levels of adducts were compared in individuals with and without adenomas, there was a trend for higher levels in individuals presenting with adenomas especially in the highest category of M₁-dG adducts (P < 0.005).

Introduction
CRC is the second most common cause of death from malignant diseases in Western Europe and the United States, both in men and in women. Approximately 94% of CRC is sporadic in nature (1), and it is estimated that 75–80% might be attributable to environmental causes (2, 3). CRC develops through a multistep sequence of dysplastic morphological change, most commonly including an adenoma, characterized by an accumulation of genetic defects (4, 5). The risk of developing cancer rises with the number and size of adenomas and villous histology.

Diet is regarded as the most important environmental influence on CRC (6, 7). Increased risk of CRC has been associated with high intake of red meat and total fat (6–9). High consumption of vegetables and fruit has been shown to reduce the risk (reviewed in Refs. 6 and 7). In addition, Lee et al. (10) and Slattery et al. (11) reported that individuals who maintained high levels of physical activity throughout their lives were at a lower risk for developing colon cancer.

Dietary fat, lipid peroxidation, and arachidonic acid metabolism have all been implicated in colorectal carcinogenesis (9, 12, 13). Lipid peroxidation is initiated by free-radical attack of membrane lipids, generating large amounts of reactive products, which have been implicated in tumor initiation and promotion. Because modification of DNA is believed to be an important early step in carcinogenesis, endogenous DNA adducts derived from oxidative stress, lipid peroxidation, or other endogenous processes have been proposed as contributors to the etiology of human cancer (14).

MDA is a major genotoxic carboxyl compound generated by lipid peroxidation (14, 15). It is also a by-product of the arachidonic acid metabolism in the synthesis of prostaglandins (16). Both of these endogenous processes are modulated by dietary factors. For example, lipid peroxidation is stimulated by the presence of high levels of ω-6 PUFAs and is inhibited by dietary antioxidants. Increased levels of MDA, together with increased levels of PUFAs and prostaglandins, e.g., PGE₂, have been reported in tumor tissues of CRC patients as compared with their normal mucosa (13).

MDA is mutagenic in bacterial and mammalian systems (17). It reacts with DNA to form adducts with deoxyguanosine, deoxyadenosine and deoxyctydine (18). The adduct formed on reaction with deoxyguanosine, the highly fluorescent cyclic pyrimidopurinone, M₁-dG, was originally detected in liver

4 The abbreviations used are: CRC, colorectal cancer; MDA, malondialdehyde; PUFA, polyunsaturated fatty acid; M₁-dG, 1,2-malondialdehyde-deoxyguanosine; EPIC, European Prospective Investigation on Cancer; CT-DNA, calf thymus DNA; HPLC, high-pressure liquid chromatography; BMI, body mass index; ISB, immunoslot blot; PTM, PBS-Tween 20 + fat-free milk powder.

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2 To whom requests for reprints should be addressed, at MRC Toxicology Unit, University of Leicester, Leicester LE1 9HN, United Kingdom. Phone: 44-116 2525172; Fax: 44-116-2525616; Email: chiara_leuratti@hotmail.com
3 Present address: Chemistry Department, The Open University, Milton Keynes, MK7 6AA United Kingdom.

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DNA from healthy individuals, at levels of 5–11 adducts per 107 total bases (19). Since then, M1-dG has been detected in several human tissues at levels ranging from 0.02 to 21 per 107 normal bases (20–24). Very interestingly, it has been shown that M1-dG can also be formed after direct oxidative attack on DNA (25). M1-dG is a premutagenic lesion and can induce guanine-thymine transversions and guanine-adenine transitions in DNA (26, 27). The adduct is repaired by bacterial and mammalian nucleotide excision repair and by bacterial mismatch repair pathways (15).

Fang et al. (28) reported that volunteers on a diet containing high amounts of PUFAs had higher levels of M1-dG in leukocyte DNA than individuals on a diet rich in monounsaturated fatty acids. The difference between adduct levels after the two diets was greater in women than in men. These results suggested a role of diet in modulating adduct levels. Results from our group have also suggested a dietary influence on M1-dG levels in human leukocyte DNA (29).

The United Kingdom Flexible Sigmoidoscopy Screening Trial is a multicenter randomized controlled trial designed to investigate the efficacy and effectiveness of a single flexible sigmoidoscopy, with removal of all polyps observed during screening, in decreasing morbidity and mortality from CRC (30). The Norfolk branch of the study is unique because one-third of the 3000 participants also independently participated in the EPIC study. As part of EPIC, dietary, health, and lifestyle data from all of the participants had been collected prospectively through detailed dietary diaries and health checks (31). During flexisigmoidoscopy, colorectal biopsies were taken from the flat mucosa in previous EPIC participants with adenomas (83 cases) and polyp free controls (79 controls) were included in the study. Only patients with histologically proven adenomas (83 cases) and polyp free controls (79) were found to have one or more polyps (adenomatous or villous), metaplastic polyps) and to have completed a detailed diary for 7 days of all food and drink consumed (32). Food diaries were coded and analyzed for 7 days of all food and drink consumed (32). Food diaries from our group have also suggested a dietary influence on M1-dG levels in human leukocyte DNA (29).

Materials and Methods

Materials. CT-DNA, proteinase K, RNase A (from bovine pancreas), RNase T1, and propidium iodide were purchased from Sigma Chemical Co. Ltd. (Dorset, United Kingdom). PBS tablets (Dulbecco A) were purchased from Oxoid Ltd. (Hampshire, England). All other reagents and solvents of analytical or HPLC grade were obtained from either BDH or Fisher Scientific Ltd. (Loughborough, Leicestershire, United Kingdom).

Human Study

Patients. Ethical approval for this study was obtained from the Norwich District Ethics Committee in 1997. Patients (2999) from general practices were screened in Norfolk as part of the United Kingdom Flexible Sigmoidoscopy Study. Among the EPIC patients, at the first health check from 1992–1997, 144 were found to have one or more polyps (adenomatous or metaplastic polyps) and to have completed a detailed diary for 7 days of all food and drink consumed (32). Food diaries from 144 controls, age-, sex-, and general practitioner-matched, were also available. The food diaries were coded and analyzed for nutrients and foods using methods described elsewhere (33). Only patients with histologically proven adenomas (83 cases) and polyp free controls (79 controls) were included in the study. Information on diet, smoking status, weight, height, and BMI [weight (kg)/height2 (m2)] was obtained at the initial EPIC health check.

Biopsies and Polyp Identification. During flexible sigmoidoscopy screening, colorectal biopsies were taken from the normal mucosa of the posterior wall of the rectum, 3 cm above the dentate line. When polyps were present, they were removed at the same time. Biopsies were transferred to cryovials, immersed in liquid nitrogen immediately and stored at −80°C. Patients with multiple or large (>1-m) adenomas or with >20% villous histology were considered high risk.

DNA Extraction. Biopsies were defrosted and washed in PBS twice to minimize bacterial contamination.

For DNA extraction, biopsies were homogenized using glass mortars and PTFE pestles (Polytron or Fisher). DNA was extracted using Qiagen genomic DNA extraction kit (Qiagen Ltd., Crawley, Sussex, United Kingdom) and was digested using proteinase K (160 units), RNase A (400 units), and RNase T1 (400 units). DNA was dissolved in ultrapure water. DNA purity was assessed by 260/280 nm ratio using a GeneQuant II RNA/DNA calculator (Pharmacia Biotech) and by reverse-phase-HPLC after digestion to deoxynucleotides (34).

ISB

Standard MDA-modified CT-DNA was prepared as described previously (34). The amount of M1-dG was measured by HPLC-fluorescence using a calibration curve obtained with synthetic M1-dG (35).

The ISB assay was performed as described previously (34), with several modifications. MDA-modified CT-DNA was diluted with control CT-DNA (both at a concentration of 0.1 μg/μl) to obtain decreasing amounts (5 fmol to 0.2 fmol) of M1-dG adduct and to generate standard curves for the ISB. Standard CT-DNA and colorectal DNA (3.5 μg) were dissolved in 10 molar dipotassium hydrogen P2, (pH 7; 100 μl) and PBS (150 μl). DNA samples were sonicated for 20 min in a water bath sonicator, heat-denatured for 5 min in a boiling water bath, cooled on ice for 10 min and mixed with an equal volume of 2 m ammonium acetate. Resulting single-stranded DNA (143 μl containing 1 μg DNA/sample, in triplicate) was loaded onto nitrocellulose (NC) filters (0.1 μm, BA79; Schleicher & Schuell, Dassel, Germany) using a Minifold II, 72-well slot-blot microfiltration apparatus (Schleicher & Schuell). The slots were rinsed with 200 μl of 1 m ammonium acetate. The filters were subsequently removed from the support and baked at 80°C for 90 min to immobilize the DNA. Filters were then bathed in 100 μl of PTF [PBS-Tween 20 (0.1% Tween) + 0.5% fat-free milk powder] for 1 h at room temperature to inhibit unspecific antibody binding. After two 5-min washes with PBS-Tween 20 (0.1% Tween), the filters were bathed in 40 μl of PTF containing the anti-M1-dG monoclonal antibody D10A1 (36), diluted 1:48,000. Filters were incubated for 2 h at room temperature, followed by overnight incubation at 4°C. After one 1-min and two 5-min washes with PBS-Tween, the filters were incubated for 2 h at room temperature with horseradish peroxidase-conjugated secondary antibody (goat antimouse IgG: Dako A/S, Glostrup, Denmark), diluted 1:4,000 in 32 ml of PTF. After additional washes in PBS-Tween, the filters were incubated for 5 min with the chemiluminescent reagent consisting of 4 ml of luminol enhancer solution plus 4 ml of stable peroxidase buffer (SuperSignal; West Dura, Pierce, Rockford, IL). An image of the filters was acquired using a Fluor-S Multimager (Bio-Rad, Hercules, CA) and the following setting: filter: chemiluminescence; integration: manual (time ranging from 5 to 20 min depending on the intensity of the signal); light source: Chemi: no light; scan width: 80 mm; high sensitivity. The intensity of chemiluminescent signal for each band was determined using the image analysis software. Adduct level in each sample was

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determined from the calibration line generated by MDA-modified DNA (diluted with control DNA) containing known amounts of M$_1$-dG.

**Propidium Iodide Staining for Quantitation of DNA Bound to the Nitrocellulose Filter**

The nitrocellulose filters were washed overnight in PBS and incubated with propidium iodide (250 μg) in 50 ml of PBS for 3 h at room temperature and in the dark. The filters were then washed with PBS for 90 min. An image of the filter was captured using the Bio-Rad Fluor-S Multimager with the following settings: filter: 520 nm; integration: automatic; light source: Epi UV light; high resolution. The intensity of fluorescent signal for each band, which was proportional to the amount of DNA bound to the filter (data not shown), was determined using the image analysis software. The level of adducts in each sample was corrected for the amount of DNA bound to the filter (37). M$_1$-dG levels are given as adducts per $10^7$ total normal bases.

**Statistical Methods**

Results for DNA adducts were subjected to Kruskal-Wallis ANOVA followed by Mann-Whitney test. The difference between distributions of cases and controls was evaluated using the χ² test and Fisher’s exact test. Minitab (for Windows) was used in the analysis. Associations of food and nutrients with M$_1$-dG were evaluated by Pearson’s correlation analysis. Statistical packages SAS version 6.12 and STATA version 6 were used.

**Results**

**Subject Characteristics.** Dietary data and colorectal biopsies from normal mucosa were available from 100 men and 62 women participating in both the United Kingdom Flexiscope Sigmoidoscopy Screening Trial and the EPIC study. Of the 162 participants, 83 were cases with adenomas and 79 were polyps. Twenty-nine of the cases, 11 women and 18 men, had high-risk adenomas. Average age for men and women was 58 ± 3 years. The mean values for weight, height, and BMI in men and women are summarized in Table 1. There were no significant differences between cases and controls. Women were shorter but had a higher BMI than men. Of all participants, 9.5% were current smokers, 10 and 9% in men and women, respectively.

**DNA Adducts.** DNA samples were analyzed using a previously developed ISB assay, which has a limit of detection of ~0.2 adducts per $10^7$ total bases (34). This assay, which uses a specific monoclonal antibody against M$_1$-dG (36), requires only 1 μg of DNA/sample (in triplicate). DNA extraction from colorectal biopsies using a Qiagen kit, as described in “Materials and Methods,” gave a yield that varied between 5 to ~200 μg of DNA. Enough DNA for ISB analysis was therefore obtained from all biopsy samples. In the present work, adduct levels measured by ISB in each sample, were corrected for the amount of DNA bound to the filter, as determined by propidium iodide staining. M$_1$-dG adducts were detected in 92% of the samples analyzed. Thirteen samples had adduct levels below the limit of detection of the assay.

Substantial interindividual variation was observed, with adduct levels ranging from undetectable to 12.23 M$_1$-dG per $10^7$ total bases (Fig. 1). The mean adduct levels were 4.30 ± 3.0 and 4.60 ± 2.9 per $10^7$ bases in men and women, respectively.

<table>
<thead>
<tr>
<th>Findings</th>
<th>All (162)</th>
<th>Polyps (83)</th>
<th>C (79)</th>
<th>Correlation ( \rho ) with M$_1$-dG</th>
<th>Correlation ( \rho ) with M$_1$-dG</th>
</tr>
</thead>
<tbody>
<tr>
<td>Age (y)</td>
<td>57.7 ± 3.9</td>
<td>58.4 ± 3.1</td>
<td>57.7 ± 3.9</td>
<td>0.197 (P = 0.02)</td>
<td>0.189 (P = 0.02)</td>
</tr>
<tr>
<td>Height (cm)</td>
<td>171.7 ± 8.8</td>
<td>169.5 ± 9.2</td>
<td>175.1 ± 6.4</td>
<td>0.278 (P = 0.003)</td>
<td>0.278 (P = 0.005)</td>
</tr>
<tr>
<td>Weight (kg)</td>
<td>76.6 ± 10.9</td>
<td>78.4 ± 9.4</td>
<td>74.3 ± 31.2</td>
<td>0.012 (P = 0.000)</td>
<td>0.012 (P = 0.005)</td>
</tr>
<tr>
<td>BMI (kg/m$^2$)</td>
<td>26.1 ± 5.3</td>
<td>25.9 ± 2.4</td>
<td>25.3 ± 7.5</td>
<td>-0.056 (P = 0.03)</td>
<td>-0.056 (P = 0.02)</td>
</tr>
</tbody>
</table>

Values are means ± SD (range)

Table 1: Anthropometric characteristics and MDA-adduct levels (M$_1$-dG per $10^7$ total normal bases) for all of the participants, men and women in cases (Polyps) and controls (C).
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Fig. 1. Distribution plot of M₁-dG levels in normal colorectal mucosa of cases and controls (men and women). M₁-dG levels ranged from undetectable (n = 13) to 12.23/10⁷ total bases (average: 4.42 ± 2.95/10⁷ bases). *, the difference between the distribution of cases and controls is marginally significant (P = 0.065, by χ² test).

There was no significant difference between the sexes. A significant positive association between adduct levels and age was observed in men (P < 0.01), whereas women showed only a weak, not-significant correlation (P = 0.5). A negative correlation between adducts and BMI was evident in men (P < 0.02), whereas no correlation was observed in women (Table 1). In men, M₁-dG levels were also significantly correlated with height (P < 0.05). No correlation with smoking status was observed either in men or in women.

DNA Adducts and Diet. Information from the 7-day dietary diaries was computerized and converted into estimates of intake for a series of more than 30 food items and nutrients. Significant associations between various food components and adduct levels were observed, with marked differences between men and women (Table 2).

In men, M₁-dG levels were inversely associated (P ≤ 0.05) with the reported consumption of legumes, nuts, and whole meal bread. Cereals, fruit, vegetables, salads, and raw tomatoes showed inverse association with adduct levels, but these were not significant (data not shown). There were significant correlations with adduct levels and beer, offal, white meat, and alcoholic drinks.

In women, there were no significant associations between adduct levels and food consumption.

Intakes of nutrients were also related to adduct levels. Table 3 shows associations for those nutrients that would be expected to relate to MDA adducts, e.g., fat, fatty acids, and antioxidant vitamins. All of the associations with adduct levels failed to reach significance, although there was a weak positive association with saturated fatty acids in women (P = 0.061). The ratio of monounsaturated:saturated fatty acids was inversely correlated with M₁-dG levels in women, although the correlation was only slightly significant (P = 0.099). In addition, the ratio of polyunsaturated:saturated fatty acids was inversely associated with adduct levels in women (P = 0.019). No association between fatty acids and M₁-dG was observed in men.

DNA Adducts and Adenomatous Polyps. Table 1 also shows mean adduct levels in participants found to have adenomas, compared with adenoma-free individuals. Mean differences were not significant, although Fig. 1 showed that cases tended to have higher levels. Twelve individuals with polyps had adducts levels above 9 per 10⁷ bases, compared with 1 polyposis-free individual. χ² testing of differences in adduct levels between the distributions (Fig. 1) in cases and controls failed to reach statistical significance (P < 0.065). However, Fisher’s exact test, not taking into account that the analysis was data driven, showed that there was a large excess of cases (12) compared with controls (1) in this category (P < 0.005). Five of the 12 samples with M₁-dG levels above 9 per 10⁷ bases were from patients presenting with high-risk adenomas.

In females, a trend in relation to the severity of the adenomas was observed (Fig. 2). M₁-dG levels (per 10⁷ total bases) increased from 4.06 ± 2.0 in controls to 4.76 ± 3.6 in low-risk and to 5.40 ± 2.6 in high-risk women. The increase from controls to high-risk women was marginally significant (P = 0.09) by Kruskal-Wallis ANOVA followed by Mann-Whitney U test. There were no differences in men.

Discussion

Colorectal biopsies from normal mucosa of 162 participants in both the United Kingdom Flexiscope Sigmoidoscopy Screening Trial and the EPIC study were analyzed for the presence of M₁-dG, the major DNA adduct formed by reaction of MDA with DNA. The objectives of the study were to investigate whether this adduct can be regarded as a biomarker of specific dietary intake and whether it can be considered a risk factor for the presence of premalignant lesions in humans.

Samples were analyzed using a quick and sensitive ISB assay, which allows a high throughput of samples in large population-based studies, using only small amounts of DNA.

The present study has shown for the first time that M₁-dG is present in the normal colorectal mucosa in humans. The levels measured had an average value, considering all participants, of 4.42 ± 2.95 per 10⁷ total bases. These levels are lower than those detected in our laboratory in human gastric biopsy DNA (23). It is likely that this difference in M₁-dG levels is attributable to tissue-specific differences in adduct formation and removal.

In the present study, there was substantial interindividual variation in adduct levels. Levels in women were slightly, but not significantly, higher than in men. Fang et al. (28) reported higher levels of adducts in WBCs of women as compared with men. Similar findings were reported in the same leukocyte samples for etheno adducts, another product of lipid peroxidation (38). Similar to our findings for colorectal biopsies, no sex differences in M₁-dG levels were reported for gastric tissue (23).

In general, associations between adduct levels and anthropometrically. There was no significant difference between the sexes. A significant positive association between adduct levels and age was observed in men (P < 0.01), whereas women showed only a weak, not-significant correlation (P = 0.5). A negative correlation between adducts and BMI was evident in men (P < 0.02), whereas no correlation was observed in women (Table 1). In men, M₁-dG levels were also significantly correlated with height (P < 0.05). No correlation with smoking status was observed either in men or in women.

### Table 2
Associations of selected food groups with M₁-dG adducts in male and female participants

<table>
<thead>
<tr>
<th>Food Group</th>
<th>All</th>
<th>Males</th>
<th>Females</th>
</tr>
</thead>
<tbody>
<tr>
<td>Whole meal bread</td>
<td>-0.1562a</td>
<td>-0.1778b</td>
<td>-0.1009b</td>
</tr>
<tr>
<td>Legumes</td>
<td>-0.1523b</td>
<td>-0.1850b</td>
<td>-0.0730b</td>
</tr>
<tr>
<td>Nuts</td>
<td>-0.1370</td>
<td>-0.2006b</td>
<td>0.0128</td>
</tr>
<tr>
<td>White meat</td>
<td>0.0693</td>
<td>0.2052b</td>
<td>-0.2087</td>
</tr>
<tr>
<td>Offals</td>
<td>0.0973</td>
<td>0.2769b</td>
<td>0.0191</td>
</tr>
<tr>
<td>Beers</td>
<td>0.1418</td>
<td>0.2110b</td>
<td>-0.1169</td>
</tr>
<tr>
<td>All alcohol</td>
<td>0.1318</td>
<td>0.1969b</td>
<td>-0.0695</td>
</tr>
</tbody>
</table>

* Pearson’s correlation test was used in the analyses.  
1 P ≤ 0.05.  
2 Offal, kidney and liver.  
3 P ≤ 0.005.
M1-dG levels in normal colorectal mucosa were modulated by age in men but not in women. No relationship with smoking was observed in women. Adduct levels increased with mass is positively correlated with CRC risk (6, 7). No correlation test was used in the analyses.

One objective of our study was to investigate whether M1-dG levels in normal colorectal mucosa were modulated by dietary and life-style habits. The hypothesis was that diets rich in PUFAs, which are precursors of MDA, or poor in fruits and vegetables, salads, and raw tomatoes being protective. High consumption of some of these food items has been associated with protection against colon and rectal cancer. Antioxidants present in fruits and vegetables could decrease formation of M1-dG by inhibiting either lipid peroxidation or direct DNA oxidation, which are believed to give rise to M1-dG through the formation of base propenal intermediates (25). Alternatively, micronutrients and bioactive compounds, such as phenols and flavonoids, could stimulate detoxification pathways and DNA repair mechanisms (6, 7).

High intake of beer, alcohol, and white meat was positively correlated with adduct levels in men. Epidemiological studies have reported either an increased risk or no association between beer/alcohol consumption and CRC (7). In studies of diet and colorectal adenomas, alcohol is usually considered to be a potential confounder, although there is little clear evidence of effects in the literature (39–41).

There were no associations between adduct levels and the recorded consumption of about 30 groups of food in women, possibly because of the smaller number of samples analyzed and, therefore, less statistical power. Differences between the sexes in leukocyte adduct levels in response to diet have been reported in a smaller previous study, in which mainly younger sex differences in leukocyte adduct levels have been investigated (28). In the present study, all of the women were ages 55–65 years, and colonic adduct levels were measured.

Here, there was a weak positive association between colorectal biopsy M1-dG levels and saturated fatty acid intake in women and an inverse association with the standard dietary ratio of PUFAs:saturated fat. The effect of saturated fat on M1-dG adduct levels has not been investigated elsewhere, although the previous study had shown higher levels in leukocytes with monounsaturated fatty acids, polyunsaturated fat:monounsaturated fat:polyunsaturated fat:monounsaturated fat:

\[
\text{Saturated fat} \quad 0.1224 \\
\text{Monounsaturated fat} \quad 0.0473 \\
\text{Polyunsaturated fat} \quad -0.0869 \\
\text{M:P ratio} \quad -0.1173 \\
\text{P:S ratio} \quad -0.1436 (P < 0.07) \\
\text{Vitamin C} \quad -0.0364 \\
\text{Vitamin E} \quad -0.0954 \\
\text{Alcohol} \quad 0.1117 \\
\text{Fiber} \quad -0.1126 \\
\text{Fat} \quad -0.0048 \\
\text{Iron} \quad 0.0405 \\
\text{Carotene equivalents} \quad -0.0735
\]

Table 3. Associations of selected nutrient intake with M1-dG levels

<table>
<thead>
<tr>
<th>Nutrients</th>
<th>All</th>
<th>Males</th>
<th>Females</th>
</tr>
</thead>
<tbody>
<tr>
<td>Saturated fat</td>
<td>0.1224</td>
<td>0.1131</td>
<td>0.2390 (P = 0.061)</td>
</tr>
<tr>
<td>Monounsaturated fat</td>
<td>0.0473</td>
<td>0.0096</td>
<td>0.1529</td>
</tr>
<tr>
<td>Polyunsaturated fat</td>
<td>-0.0869</td>
<td>-0.0828</td>
<td>-0.0643</td>
</tr>
<tr>
<td>M:P ratio</td>
<td>-0.1173</td>
<td>-0.0711</td>
<td>-0.2143 (P = 0.09)</td>
</tr>
<tr>
<td>P:S ratio</td>
<td>-0.1436 (P &lt; 0.07)</td>
<td>-0.0438</td>
<td>-0.2970 (P &lt; 0.02)</td>
</tr>
<tr>
<td>Vitamin C</td>
<td>-0.0364</td>
<td>-0.1542</td>
<td>0.1468</td>
</tr>
<tr>
<td>Vitamin E</td>
<td>-0.0954</td>
<td>-0.0954</td>
<td>0.1387</td>
</tr>
<tr>
<td>Alcohol</td>
<td>0.1117</td>
<td>0.1629</td>
<td>0.0103</td>
</tr>
<tr>
<td>Fiber</td>
<td>-0.1126</td>
<td>-0.1703</td>
<td>0.0067</td>
</tr>
<tr>
<td>Fat</td>
<td>-0.0048</td>
<td>-0.0527</td>
<td>0.1631</td>
</tr>
<tr>
<td>Iron</td>
<td>0.0405</td>
<td>0.0308</td>
<td>0.0906</td>
</tr>
<tr>
<td>Carotene equivalents</td>
<td>-0.0735</td>
<td>-0.0672</td>
<td>-0.0887</td>
</tr>
</tbody>
</table>

*Pearson’s correlation test was used in the analyses.
*M:P, monounsaturated fat:polyunsaturated fat; P:M, polyunsaturated fat:monounsaturated fat; P:S ratio, the difference between controls and high-risk women is marginally significant (P = 0.09) by Kruskal-Wallis ANOVA followed by Mann-Whitney U test.

Fig. 2. M1-dG levels in low-risk (LR) adenomas, high-risk (HR) adenomas, and controls (C) for all of the participants, men and women. Values are means ± SD. *The difference between controls and high-risk women is marginally significant (P = 0.09) by Kruskal-Wallis ANOVA followed by Mann-Whitney U test.

[p. 271]
PUFAs could, therefore, have had a confounding effect in the present analysis. However, no associations with fatty fish consumption (the major source of ω-3 fatty acids) were shown.

The other objective of our study was to analyze M₁-dG levels in relation to the presence or absence of adenomatous polyps, to examine whether adducts could be related to disease outcome. A trend toward higher levels of adducts in cases than in controls was observed for both men and women, although the difference failed to reach significance. The difference in adduct levels between women presenting with high risk adenomas and polyp-free controls was statistically significant, although only marginally (Fig. 2). This is probably attributable to the wide interindividual variation in adduct levels and consequent high SDs from the adduct means. In addition, the number of women in the high-risk group was quite small (n = 11).

At present, the relationship between MDA-DNA damage and the risk for adenomatous polyps remains unclear. What is probably very important is the location of M₁-dG in DNA, i.e., which genes are modified, and the rate to which the damage is converted into mutations. DNA repair rate and fidelity are also extremely important. Moreover, other endogenous as well as exogenous DNA damage has been reported in human colorectal tissue (44–47). A previous study (48) has demonstrated increased levels of bulky and/or aromatic DNA adducts associated with increased CRC risk. It is likely that the contribution of DNA damage derived by both exogenous and endogenous sources, together with host susceptibility factors such as polymorphisms in relevant genes (49), could be of importance in colorectal carcinogenesis.

In conclusion, M₁-dG was detected in the normal colorectal mucosa of participants in the United Kingdom Flexiscope Sigmoidoscopy Screening Trial and the EPIC study. Food items that, in epidemiological studies, were associated with modulation of CRC risk, modulated M₁-dG levels in men. Saturated fat was correlated to increased adduct levels in women. Cases showed higher adduct levels than did controls, although the difference was not statistically significant. Results from the present study raise the possibility that M₁-dG plays a role in human colorectal carcinogenesis, certainly in combination with other genetic and environmental factors.

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Chiara Leuratti, Mark A. Watson, Eliot J. Deag, et al.


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