

A Multimineral Natural Product from Red Marine Algae Reduces Colon Polyp Formation in C57BL/6 Mice

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The goal of this study was to determine if a multimineral natural product derived from red marine algae could reduce colon polyp formation in mice on a high-fat diet. C57BL/6 mice were maintained for up to 18 mo either on a high-fat “Western-style” diet or on a low-fat diet (AIN 76A), with or without the multimineral-supplement. To summarize, colon polyps were detected in 22 of 70 mice (31%) on the high-fat diet but in only 2 of 70 mice (3%) receiving the mineral-supplemented high-fat diet ($P < 0.0001$). Colon polyps were detected in 16 of 70 mice (23%) in the low-fat group; not significantly different from high-fat group but significantly higher than the high-fat-supplemented group ($P = 0.0006$). This was in spite of the fact that the calcium level in the low-fat diet was comparable to the level of calcium in the high-fat diet containing the multimineral-product. Supplementation of the low-fat diet reduced the incidence to 8 of 70 mice (11% incidence). Taken together, these findings demonstrate that a multimineral natural product can protect mice on a high-fat diet against adenomatous polyp formation in the colon. These data suggest that increased calcium alone is insufficient to explain the lower incidence of colon polyps.

INTRODUCTION

Past epidemiological (1–6) and interventional (7,8) studies in human subjects have demonstrated the capacity of calcium to reduce colon polyp formation and reduce colon cancer risk. *In vitro* studies have demonstrated the growth-regulating activity of calcium and have provided insights into growth-suppressing mechanisms (9–12). Although calcium has demonstrable chemopreventive activity against colon polyp formation, protection provided by calcium alone can be described as modest. It is estimated that under conditions of optimal use, a reduction in polyp incidence of 20%–22% might be achieved (13). Additional interventions are needed.

In a series of studies, Newmark and colleagues (14–16) demonstrated that mice fed a high-fat Western-style diet (HFWD) developed colon polyps at a higher rate than mice fed a standard low-fat rodent chow diet. The HFWD, along with its high content of saturated fat and carbohydrate, has several additional features that may contribute to the formation of colon polyps. These include a reduction in folic acid, a reduction in methyl group donors, a lower content of fiber and a reduction in calcium. Replacement studies demonstrated that supplementation with calcium reduced polyp formation (though not to background levels) (16). Replacing the missing folic acid, fiber, or methyl group donors had little effect. In the present study we have assessed colon polyp formation

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in C57BL/6 mice fed a high-fat diet in the absence and presence of a multimineral-containing natural product derived from the skeletal remains of the red marine algae, *Lithothamnion calcareum*. The multimineral-containing product dramatically reduced polyp formation in the high-fat-fed mice (i.e., from 31% incidence to 3% incidence). In contrast, mice fed a low-fat diet without the multimineral-containing product but with a comparable level of calcium to that provided by the multimineral product had a colon polyp incidence of 23%. These data confirm and extend a previous small scale study in which the multi-mineral product protected mice from colonic polyps on a high-fat diet. In the pilot study, colon polyps were seen in 4 of 20 mice (20% incidence) on the HFWD mice vs. 0% incidence in mice on the supplemented HFWD (17).

MATERIALS AND METHODS

Multimineral Natural Product

The multimineral natural product used in this study is obtained from the skeletal remains of the red marine algae, *Lithothamnion calcareum* (Pallas), also known as *Phymatolithon calcareum* (Pallas) (18). The algae thrive in the cold Atlantic waters off the southwest coast of Ireland and northwest coast of Iceland. Minerals from sea water are accumulated in the algae fronds over the lifespan of the organism. Eventually, the mineralized fronds break off of the living organism and fall to the ocean floor, from where they are harvested. The mineralized fronds are separated from extraneous materials, sterilized, dried, and milled under ISO and HACCP certification. The final product contains approximately 12% calcium (wt/wt), 1% magnesium, and measurable levels of 72 other trace minerals—essentially all of the minerals that the algae are able to accumulate from the sea water. The product is sold as a food supplement under the name Aquamin[®] (GRAS 000028) and is used in various products for human consumption in Europe, Asia, Australia, and North America (Marigot Ltd, Cork, Ireland). The mineral composition of the algae product can be found in Supplement Table 1.

Diet Groups

Inbred C57BL/6 mice were obtained from Charles River, Portage, MI at 3 wk of age. The animals were started within 1 wk of arrival on either a standard low-fat rodent chow diet (AIN 76A), to serve as a control or a HFWD, prepared according to the formulation of the Newmark stress diet (14). The diet is a modification of AIN 76A and is designed to mimic the diet consumed by many individuals in Western society (19–21). The HFWD contains 20 g% fat from corn oil as compared to 5 g% in AIN 76A (Supplement Table 2). The percentage of calories from fat in this diet is 37.8% as compared to 11.5% in the AIN 76A diet. Although sucrose is reduced in the HFWD relative to the AIN 76A control diet, the overall calories provided in the HFWD is 4764 kcal/kg vs. 3902 kcal/kg in AIN 76A. In addition to its high fat content, the HFWD has additional modifications. Methionine is replaced with cysteine, amounts of fiber, folic acid, and choline are reduced, and the calcium level is reduced from 5.25 gm/kg to 0.41 gm/kg. Both the HFWD and AIN 76A contain a mix of essential trace elements including potassium, magnesium, manganese, chromium, copper, iron, selenium, and zinc. For half of the animals on the HFWD, the diet was supplemented with the multimineral natural product at 62 g/1.062 kg, resulting in a calcium concentration comparable to that in AIN 76A. The concentration of calcium is 1.34 mg/kcal in low-fat (AIN 76A) diet, 3.24 mg/kcal in supplemented AIN 76A, 0.086 mg/kcal in the HFWD and 1.64 mg/kcal in the HFWD supplemented with the multimineral product. Diets were formulated and provided by Research Diets Incorporated (New Brunswick, NJ). The multimineral-rich supplement was compounded into the two diets, which were fed to the animals as solid food pellets. The composition of each diet is presented in Supplement Table 2.

Treatment Protocol and Necropsy

Separate cohorts of male and female C57BL/6 mice were housed 5 mice to a cage, and maintained for 5, 12, or 18 mo on their respective diet as summarized in the flow chart (Fig. 1). For the 5- and 12-mo periods, there were 10 male and 10 female mice

TABLE 1
Distribution of polyps within the colon

Location	Number of polyps			
	Low-fat diet (AIN 76A)		High-fat Western diet	
	Unsupplemented	Supplemented	Unsupplemented	Supplemented
Cecum	15	6	17	1
proximal colon	1	1	4	1
Distal colon	0	1	1	0

Distribution of polyps within the colon was based on the examination under a stereoscopic dissecting microscope. This is based on combined data for polyps at all-time points and from both genders. There were a total 35 males and 35 females present in each diet group including all-time points.

TABLE 2
Histological classification of lesions within the colon

Lesion type	Histological classification			
	Low-fat diet (AIN 76A)		High-fat Western diet	
	Unsupplemented	Supplemented	Unsupplemented	Supplemented
Hyperplasia	2 (1F/1M)	2 (2F/0M)	5 (4F/1M)	1 (1F/0M)
Adenoma	8 (8F/0M)	3 (1F/2M)	9 (9F/0M)	0
Adenocarcinoma	6 (3F/3M)	3 (2F/1M)	4 (4F/0M)	0

Histological classification was done according to the internationally standardized guidelines for the assessment of rodent tumors (23). Values represent total number by category. Values in parenthesis reflect females and males. The five macroscopic lesions that were not confirmed histologically were all very tiny lesions. Three of these were in high-fat males, one was in a high-fat female, and the fifth was in a high-fat female with the mineral supplement.

per diet group. For the 18-mo period, there were 15 mice of each sex per diet group. Food was provided ad libitum. Animals were monitored at 2-day intervals throughout the maintenance phase and were weighed every 2 wk. Mice on the HFWD diet gained more weight than did animals on the low-fat diet, but the multimineral supplement had no effect on average weight gain for either male or female mice in either diet. Between 0–5 months, there were no premature deaths. Between 6–12 months, 5 animals died and between 13 and 18 months, a total of 25 mice died or were euthanized. The majority of these deaths occurred in months 16 and 17. Of the animals that died prematurely, 2 were on unsupplemented AIN 76A, 3 were on mineral-supplemented AIN 76A, 17 were on the unsupplemented HFWD, and 8 were on the mineral-supplemented HFWD. A complete autopsy was done on all animals that died prematurely. The most common cause of premature death was euthanasia for ulcerative dermatitis, a condition that is well-known to occur in C57BL/6 mice in long-term studies (22). Liver tumors were observed in 3 mice and mouse urologic syndrome was observed in 4 male

mice. In the remaining animals, there, was no obvious abnormality. Some of these mice died spontaneously without any cause and tissue could not be collected because of autolysis. Animals that died prematurely were included in their respective groupings.

Whether the animals died prematurely or were euthanized at the respective 3 time-points, they were autopsied as follow: After the abdominal cavity was opened, the gastrointestinal tract was removed from the stomach to the rectum and flushed with saline. The entire length was opened longitudinally, fixed in 2% formalin, stained with methylene blue and examined under a stereoscopic dissecting microscope. Visible raised tumors were identified in this manner and counted. Each observable mass was placed in 2% buffered formalin and prepared for histology. Tissue was stained with hematoxylin and eosin and examined at the light microscopic level by a board-certified veterinary pathologist (I.B.). Lesions were classified according to the recently revised, standardized guidelines established by the International Harmonization of Nomenclature and Diagnostic Criteria for Lesions in Rats and Mice project (23). This project represents consensus criteria for histopathological lesions in rodents as established by the North American, European, British, and Japanese Societies of Toxicologic Pathology. Proliferative lesions were classified as hyperplasias, adenomas, or adenocarcinomas.

Blood was obtained at the time of necropsy from each animal. Serum calcium levels were determined using the Vet-Test Dry Chemistry Analyzer (Idexx Laboratories, Westbrook, ME). Calcium levels in long bones were determined in parallel. Long bones (1 femora and 1 tibia from each animal) were carefully separated from the surrounding connective tissue. Bones from all animals in each group were “pooled” to obtain a sufficient quantity of bone, and the bone tissue converted to ash. Calcium levels were then determined by atomic absorption spectroscopy (Advanced Laboratories, Inc., Salt lake City, UT). All of the procedures involving animals were reviewed and approved by the University Committee on Use and Care of Animals at the University of Michigan.

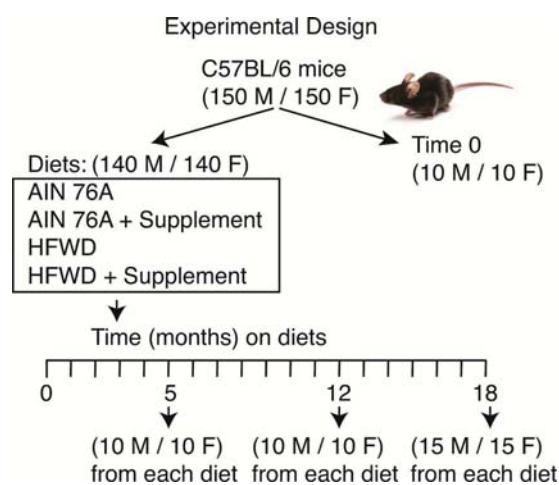


FIG. 1. Experimental design. HFWD = high-fat Western diet; M = male; F = female (Color figure available online).

Statistical Evaluation

195 Differences in polyp incidence were assessed for statistical significance using the Fisher exact test (2-tailed). Differences were considered significant at the $P < 0.05$ level.

RESULTS

Colon Polyp Incidence in Mice with or Without the Multimineral Natural Product as a Dietary Supplement

200 In this study, 20 animals per group (10 males and 10 females) were sacrificed at 5 mo and at 12 mo, and 30 animals per group (15 males and 15 females) were sacrificed at 18 mo. The incidence of colon polyp formation by time, gender, and diet group is shown in Fig. 2. A total of 22 out of 70 mice on the high-fat diet without the mineral supplement had at least 1 visible polyp at necropsy (31% incidence). In contrast, only 2 out of 70 mice on the high-fat diet with the supplement had

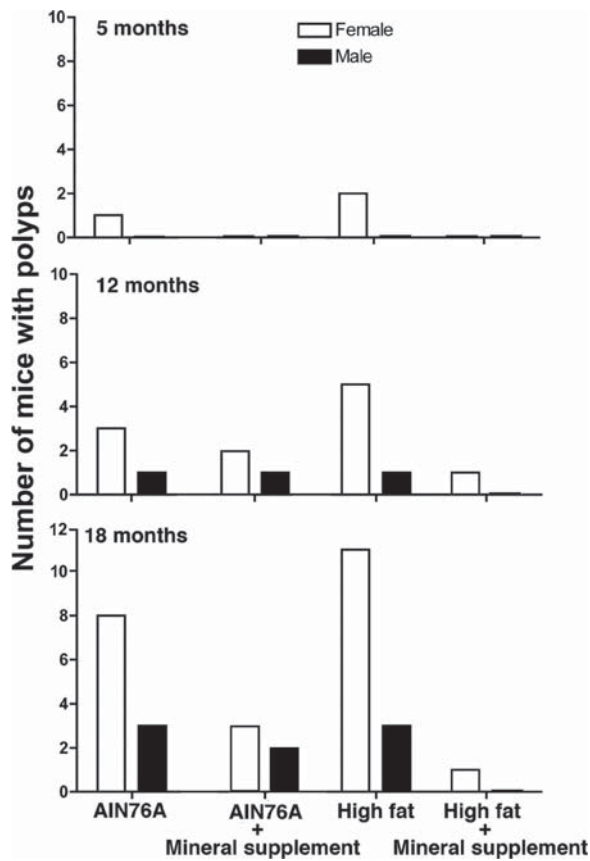


FIG. 2. Colon polyp formation in mice; Effects of time, gender, and presence or absence of the multimineral natural product. Data are based on stereomicroscopic identification. At 5 and 12 mo, there were 10 males and 10 females per group. At 18 mo, the total number of animals per group was 15 males and 15 females. When findings from the 3 time-points were analyzed together (Fisher Exact Test), polyp formation was greater in female mice than in males ($P < 0.0001$) and the inclusion of the multimineral supplement in the high-fat Western diet (HFWD) lowered polyp formation ($P < 0.0001$) relative to the HFWD alone and relative to the AIN 76A diet ($P = 0.0006$).

a visible polyp (3% incidence) ($P < 0.0001$ by Fisher exact test). A control group of mice on the (low-fat) AIN 76A diet had detectable polyps in 16 of 70 animals (23% incidence). The number of animals with polyps in the AIN 76A group was not statistically different from the number of mice on the HFWD alone but was statistically higher than the number of mice with polyps on the HFWD with supplementation ($P = 0.0006$ by Fisher exact test). This was in spite of the fact that mice in this group received a comparable amount of dietary calcium to animals in the high-fat group with the mineral supplement. When the multimineral product was included in the diet of mice on the low-fat diet, polyps were seen in 8 of 70 mice (11% incidence). When data from the high-fat and low-fat diets were combined, overall polyp incidence was reduced from 27% in mice on unsupplemented diets to 7% in mice on supplemented diets ($P < 0.0001$ by Fisher exact test).

Overall, the incidence of polyp formation was strongly weighted toward females. Of the total number of animals with polyps (48 in all), 37 were females and 11 were males ($P < 0.0001$ by Fisher exact test). Not surprisingly, the incidence of polyp formation increased with age. A total of 3 polyps were observed at 5 mo, 14 polyps at 12 mo, and 31 polyps at 18 mo. Among animals on the high-fat diet for 18 mo (15 males and 15 females), there were a total of 14 mice with polyps (47%) whereas in mice on the HFWD with the mineral supplementation (15 males and 15 females), the incidence dropped to 3% ($P = 0.0002$ by fisher exact test). Thus, the most significant protection was observed at the longest time-point.

Table 1 shows the location of the polyps within the colon. Virtually all of the polyps developed in the cecum or at the ileocecal junction (39 total) while the remaining polyps were scattered throughout the proximal colon (7 polyps) and distal colon (2 polyps). There were no significant differences between females and males in polyp location. Likewise, no diet-specific effect on polyp location was seen. The cecum (especially near the ileocecal junction) is known to be the site at which most spontaneous tumors arise in conventional-raised rodents (24).

Histological Findings

Of the 48 lesions identified by stereomicroscopy, a total of 43 lesions were confirmed histologically. Ten lesions were classified as foci of hyperplasia, 20 as adenomas and 13 as adenocarcinomas. The remaining 5 lesions (all very small) were missed during histological processing, possibly because of the difficulty in orientation of such small lesions. The histological distribution of the lesions across the different diet groups is shown in Table 2.

Foci of hyperplasia consisted of elongated crypts with maintenance of normal glandular structure and without compression of adjacent mucosa. The adenomas consisted of sessile (flat) or pedunculated masses of elongated crypts with piling of basal epithelium and compression of adjacent mucosa. Herniation of glands into the submucosa was sometimes present but the

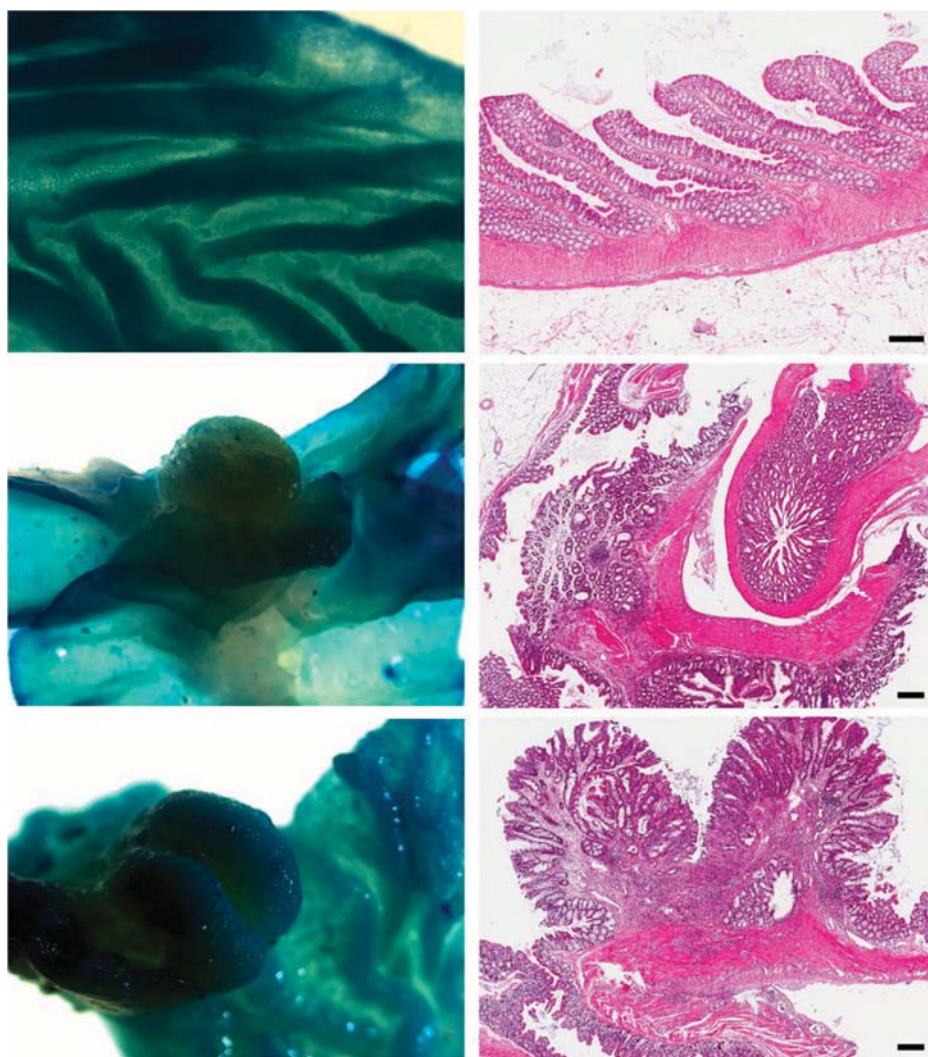


FIG. 3. Stereomicroscopic and histological appearance of colon. Upper panels: Normal colonic mucosa. Middle panels: mucosal adenoma. Lower panels: Adenocarcinoma of colon. Both tumors were from female mice on high-fat Western diet. (Histology: hematoxylin and eosin; bar = 100 μ m.) (Color figure available online).

basement membranes appeared to be intact. Adenocarcinomas consisted of dysplastic epithelial glands with invasion into the submucosa and loss of a well-demarcated basement membrane. Intravascular or intralymphatic foci were not seen and there was no gross evidence of metastasis. Fig. 3 shows stereoscopic pictures and corresponding histology of a normal colonic mucosa (upper panels), a representative adenoma (middle panels) and an adenocarcinoma (lower panels).

270 Tumors in the Small Intestine and Stomach

As part of the study, the stomach, duodenum, jejunum, and ileum were also examined stereoscopically for tumors. A total of 16 overt masses were identified. Of these, 14 were on the serosal surface and, when examined histologically, proved to be dense clusters of lymphoid cells [i.e., either lymphomas

or gut associated lymphoid tissue (GALT)]. Lymphoma of the gastrointestinal system is a common finding in aged C57BL/6 mice (25). Lymphoid lesions were equally divided among the diet groups. The 2 lesions on the mucosal surface were identified histologically as adenomas—one in the stomach of a male mouse on the high-fat diet and one in the duodenum of a female mouse on the mineral-supplemented high-fat diet (Fig. 4). With so few lesions, nothing can be said concerning dietary influence.

285 Systemic and Bone Calcium Levels in Mice on the High-Fat Diet with and Without the Multimineral Natural Product

Serum calcium levels were assessed in mice sacrificed at the 18-mo time-point. Values are shown in Fig. 5. Female mice

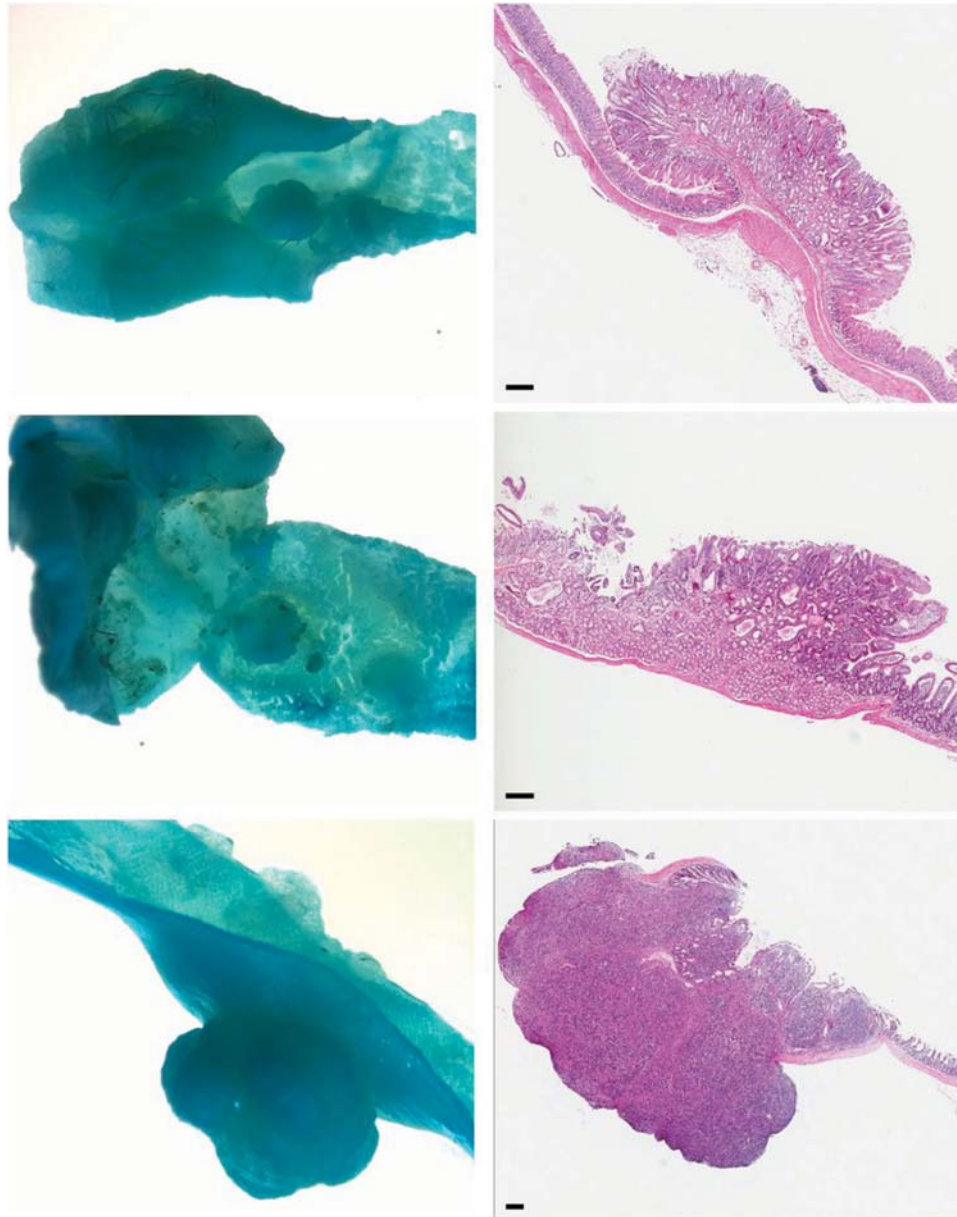


FIG. 4. Stereomicroscopic and histological appearance of three extra-colonic lesions. Upper panels: mucosal polyp (adenoma) in stomach. Middle panels: mucosal polyp (adenoma) in duodenum. Lower panels: Inflammatory cell nodule (lymphoma) on the serosal surface of ileum. (Histology: hematoxylin and eosin; bar = 200 μm .) (Color figure available online).

290 had a slightly higher serum calcium level than males but differ-
 ences were not significant. Diet appeared to have no effect on
 serum calcium levels for either female or male mice. Because
 bone serves as a reservoir for calcium in the body, bone calcium
 content was assessed. Differences were slight. Supplementa-
 295 tion increased bone calcium compared to the respective control
 groups by 2%–7% (Fig. 5). Taken together, these findings sug-
 gest no major changes in systemic calcium levels in mice given
 the multimineral supplement.

DISCUSSION

The data presented here demonstrate a virtually complete 300
 inhibition of colon polyp formation in mice on a HFWD sup-
 plemented with a natural product from the red marine algae,
Lithothamnion calcareum. The natural product consists of the
 inorganic minerals accumulated from seawater in the algae
 fronds. Calcium is the major constituent, but in addition, there 305
 is a high level of magnesium and detectable levels of 72 addi-
 tional trace elements (see Supplement Table 1 for the complete

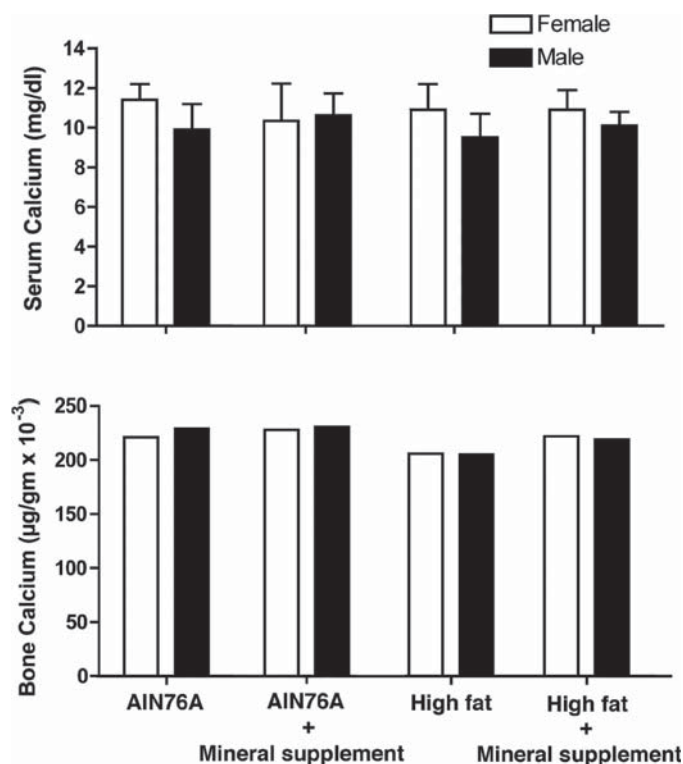


FIG. 5. Calcium levels in serum and bone. Serum calcium values are means and standard deviations. For bone calcium measurements, bones from all mice in a group were pooled to provide a single value.

analysis). What minerals in the algae product contribute to colon polyp prevention and the mechanism(s) underlying their chemopreventive activity are not fully understood. Calcium is, undoubtedly, critical, but it is unlikely that calcium alone is responsible for all of the activity in the natural product. The level of protection seen with the multiminer product was greater than that reported previously for calcium alone (16). Furthermore, protection was greater than seen in our control group of mice on the low-fat rodent diet, AIN 76 A, which had a comparable amount of calcium to that provided in the multiminer material. Both epidemiological studies (1–6) and interventional studies (7,8) in humans have demonstrated that calcium has the capacity to reduce polyp formation in the colon. However, its effectiveness is far from complete. A multiminer approach to polyp prevention may have efficacy not observed with calcium alone. There may be another benefit to a multiminer-based approach. A recent meta-analysis of the calcium supplementation literature came to the conclusion that calcium supplement use was associated with an increase in cardiovascular events (26). The inclusion of multiple minerals along with calcium in a supplement could, perhaps, reduce the calcium concentration required for efficacy. Only controlled clinical studies will be able to address this issue. The multiminer algae product used here is a GRAS-listed agent and is already present in a number of products sold for human consumption in Asia, Australia,

Europe, and North America. A small-scale clinical study for osteoarthritis symptom-relief has been conducted (27). A clinical study focused on colon polyp reduction could be carried out and could provide valuable additional information on the potential clinical benefit of this product with respect to the colon.

The multiminer natural product contains a number of cationic metals (in addition to calcium) that may contribute to reduced polyp formation in the colon—either alone or in conjunction with calcium. Among these, magnesium may be especially important. A recent study demonstrated that although magnesium supplementation by itself had little chemopreventive activity, the ratio of calcium:magnesium was as important as the level of calcium itself (28). Other minerals present in the natural product, including copper, chromium, manganese, molybdenum, selenium, and zinc have all been shown to reduce tumor formation or suppress other types of tissue injury in the gastrointestinal tract (29–32). Several of these are components of antioxidant enzymes (33) and thought to function by reducing oxidant-driven mutations that lead to dysregulated growth. Each of these elements might exert some degree of protection against polyp formation by itself, or they might function synergistically with one another or with calcium. It is unlikely, however, that any one of these components is the major sole contributor to protection for the simple reason that all of these essential trace elements are present in both the high-fat diet and the AIN 76A diet used here (see footnote in Supplement Table 2). It is hard to envision how the additional amounts contributed in the algae product could provide a benefit not seen with the same dietary minerals already present.

The rare earth (lanthanoid) minerals comprise another component of the multiminer natural product that may be beneficial. The lanthanoids are cationic metals with atomic numbers between 57 and 71. With the exception of promethium, which is a manmade element, all of the other lanthanoids are present in detectable levels in the natural product (Supplement Table 1). These closely related elements have an ionic radius similar to that of calcium, but with a higher overall charge density (34,35). Lanthanoid ions bind to calcium-binding sites on proteins, and in some cases the binding affinity is greater than that of calcium itself (36). The extracellular calcium-sensing receptor (CaSR) is a lanthanoid-responsive protein in epithelial cells (37–40). Lanthanoids also impact the functioning of calcium channels, including receptor-activated channels, voltage-gated channels, and those responsive to mechanical stress (41–43). Other important regulatory molecules influenced by lanthanoids include calcineurin and certain calcium-dependent ATPases (44). With multiple regulatory molecules affected by lanthanoid ions, it is not unlikely that cellular activity would be altered in their presence. A past study (45) demonstrated that gadolinium, one of the lanthanoid metals, was able to activate certain features of the differentiation program in human colon epithelial cells. In a recent study (46), we demonstrated epithelial cell growth inhibition with several individual lanthanoid metals (including gadolinium) under conditions in which other major cationic

mineral components of the multiminer natural product were ineffective. Finally, we have also recently found that micromolar amounts of gadolinium synergize with calcium to induce apoptosis in colon carcinoma cells under conditions in which either alone is ineffective (unpublished). Based on these observations, we hypothesize that the lanthanoid metals in the multiminer natural product contribute to reduced colon polyp formation by modulating calcium effects on epithelial cell growth and differentiation.

This is not to suggest that other minerals in the multiminer rich product are inert. This is also not to suggest that a direct effect on colonic epithelial cell growth and differentiation is the only possible mechanism for the multiminer supplement. Rather, there are likely to be multiple individual minerals contributing to growth control in a number of different ways. Among possibilities for ways in which alterations in dietary minerals could alter the carcinogenesis process is pH buffering to foster a change in the microbial content of the gut (47) and reduce inflammation (48). Bile salt chelation is another possibility (49). Finally, alterations in dietary minerals might directly influence the transit time through the gastrointestinal tract and lead to differential exposure to carcinogens in the diet. All of these possibilities will need to be examined separately.

It is of interest that the majority of colon polyps in both control and high-fat diets occurred in female mice with a preponderance occurring in aged animals. The increased incidence of colonic polyps in female mice as compared to males, and in aged female mice as compared to the younger cohort, may suggest the possible role of estrogen as an inhibitory factor in polyp development and the loss of estrogen during menopause as a contributor to colon cancer development. Although mice do not undergo a precipitous drop in estrogen levels at menopause, there is a gradual decline in estrogen production beginning at about 12 mo of age (50), such that by 16 mo, circulating estrogen levels are essentially zero. In support of this, estrogen receptors have been identified in gastrointestinal epithelial cells (51). In addition, the Women's Health Initiative trial of estrogen plus progestin in post-menopausal women identified a reduced risk of colon cancer in association with the hormone replacement therapy (52). Finally, a small-scale study in rodents found that estrone in conjunction with soy reduced tumor formation in azoxymethane-treated animals more effectively than soy alone (53). At this point, it is not clear how mineral supplement might overcome the loss of estrogen in the aged mice.

In conclusion, this study shows that a multiminer natural product obtained from marine algae is able to reduce colon polyp formation in C57BL/6mice both on a high-fat diet and a low-fat diet. Based on the results presented here, we suggest calcium alone cannot explain the protective effects of the multiminer supplement, and that a multiminer approach to colon polyp chemoprevention may prove to be more efficacious than an approach based on the use of calcium alone. Our past studies showing that the same multiminer natural product also reduces bone loss in female mice (54) and our recent studies showing

reduced liver tumor formation (55) suggest that the beneficial effects of the multiminer supplement may extend beyond the colon.

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