Phytoestrogens Regulate Vitamin D Metabolism in the Mouse Colon: Relevance for Colon Tumor Prevention and Therapy

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ABSTRACT Soybean products are highly represented in the traditional Asian diet. Major components of soy proteins are phytoestrogens, such as isoflavones. They may be responsible for the extremely low incidence of prostate and mammary tumors and possibly also of colon cancer in countries such as China and Japan. Serum 1,25-dihydroxyvitamin D₃ level is inversely related to incidence of some cancers. Levels are determined by skin exposure to ultraviolet light or, to a minor extent, nutritional uptake and by subsequent conversion of the precursor vitamin D to the active hormone by the cytochrome P450 hydroxylases CYP27A1, CYP27B1 (responsible for synthesis) and CYP24 (responsible for catabolism) in liver and kidney. However, vitamin D synthesis is also found in colonocytes and is enhanced during incipient malignancy. This may indicate an autocrine/paracrine role for this differentiation-inducing hormone in defense against progression. We were able to demonstrate that either a single large oral dose of genistein or feeding soy protein for 4 mo elevated CYP27B1 and decreased CYP24 expression in the mouse colon. Our data therefore suggest that an inverse correlation of soy product consumption with colon tumor incidence may be consequent to enhanced colonic synthesis of the antimitotic hormone 1,25-dihydroxyvitamin D₃. J. Nutr. 132: 3490S–3493S, 2002.

KEY WORDS: • soy protein feeding • genistein • 1,25-dihydroxyvitamin D₃ synthesis • 25-hydroxyvitamin D₃-1α-hydroxylase (CYP27B1) • 25-hydroxyvitamin D₃-24-hydroxylase (CYP24)

Large geographic gradients in mortality occur for a number of cancers in the United States: some rates (e.g., for breast, colon and prostate) are approximately twice as high in the North and Northeast as in the Southwest. Although colon cancer incidence may depend in part on dietary constituents, this cannot completely explain the gradient. Incidence has been observed to be inversely related to solar radiation (1). In the skin ultraviolet energy forms 90% of requisite vitamin D from the precursor 7-dehydrocholesterol. Only in fish-consuming populations is a major part of vitamin D ingested in fish oil. Vitamin D is 25-hydroxylated in the liver by the cytochrome P450 enzyme CYP27A1 and, according to conventional thought, 1α-hydroxylated by CYP27B1 only in the kidney to result in the active hormonal metabolite 1,25-dihydroxyvitamin D₃ (1,25-D₃).⁺

A serious lack of 25-hydroxyvitamin D₃ (25-D₃) in human serum, a measure of readily available vitamin D resulting from constitutive 25-hydroxylation of the precursor in the liver, arises commonly during the winter months because of reduced sunlight at higher latitudes as well as all year in the elderly population because of reduced outdoor activity: levels < 50 nmol/L are frequently measured, whereas optimal levels should be ~125 nmol/L. Adequate levels of 1,25-D₃ maintain human serum calcium homeostasis by increasing calcium uptake in the small intestine, improving calcium resorption in the kidney and regulating bone formation. A chronic lack of the active hormonal vitamin D metabolite leads to inadequate mineralization of bone: rickets in children and osteomalacia and osteoporosis in men and women. Although previous in vitro (2,3) and in vivo (4) studies demonstrated that 1,25-D₃

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4 Abbreviations used: CK, cytokeratin-8; CYP24, 25-hydroxyvitamin D₃-24-hydroxylase; CYP27B1, 25-hydroxyvitamin D₃-1α-hydroxylase; 1,25-D₃, 1,25-dihydroxyvitamin D₃; RT-PCR, reverse transcription-polymerase chain reaction; 25-D₃, 25-hydroxyvitamin D₃.
and some of its synthetic analogs (5) may act as antimitotic and prodifferentiating agents in, for example, colon cancer cells, the treatment of tumor patients with the active metabolite is still not feasible with most vitamin D analogs because of their hypercalcemic effect at the large pharmacologic doses necessary to achieve antimitotic action.

A completely different and unexpected physiological link between vitamin D and colon cancer treatment and prevention was recently discovered (7–9). This suggested the existence of an autocrine protective action of 1,25-D3 synthesized during incipient malignancy in colon cells that could slow down or even stop disease progression. In late-stage high-grade cancer tissue, the latter could include the consumption of vitamin D–containing foods or vitamin D supplements. However, regulation of the expression and activity of 1,25-D3 metabolic and catabolic enzymes such as CYP27B1 and CYP24 to maintain high steroid hormone levels would be even more effective.

A significantly lower prevalence of hormone-dependent cancers, such as prostatic and mammary cancers, occurs in Asian compared with Western industrialized countries (11), but a reduced incidence of colorectal cancer may occur as well. The reduced incidence has been linked to the consumption of the typical Asian diet, which contains high amounts of soy products and thus is rich in phytoestrogens, particularly in isolavonoids. One major isolavonoid is genistein, which has negative effects on tyrosine kinases (12) and topoisomerase I (13). Through their potential to act as selective estrogen receptor modulators, genistein and other phytoestrogens could conceivably affect vitamin D–related inhibition of tumor growth. Interactions between vitamin D and estrogen have recently been observed in human breast cancer cells (14) and in murine colon cancer models (15). Because genistein is an inhibitor of several members of the cytochrome P450 enzyme family (16–18), a likely site of interaction with the vitamin D system is at the regulation of expression and activity of the vitamin D–metabolizing cytochrome P450 enzymes, CYP27B1 and CYP24. We therefore initiated a study to address the questions of whether soy protein fed to mice as a normal food constituent for several months would alter CYP27B1 and CYP24 expression and whether a single gavage with 250 μg genistein would provide a similar result.

MATERIALS AND METHODS

Animals and diets. C57BL/6 mice were housed in the Centre for Laboratory Animal Care at the University of Vienna in a contained environment. Mice were weaned at age 2–3 wk and then fed ad libitum either the standard diet (basic AIN 76A) (19) or a soy diet (AIN 76A containing 200 g extracted soybean meal/kg instead of casein). Mice were anesthetized with ether and killed by cervical dislocation at age 16–17 wk. Another group was fed the standard diet and 250 μg genistein (Sigma-Aldrich, Vienna, Austria) in 70 μL H2O/5% ethanol or vehicle only was administered via gavage. Blood was drawn from the tail vein 6 h after and again 24 h after gavage when animals were killed and tissue samples were collected. Treatment groups consisted of at least 8 animals. The study protocols were reviewed and approved by the institutional committee of animal experimentation of the University of Vienna Medical School and by the Austrian Ministry of Science and Education.

Determination of tissue concentrations of genistein. Colon was extirpated from mice 24 h after oral administration of 250 μg genistein, freeze-dried and stored at −70°C. For the experiments, 50 mg of samples was dissolved in 300 μL H2O and sonicated for 10 min; 700 μL methanol was added for precipitation overnight at −20°C. After centrifugation and reprecipitation, methanol was evaporated and the water phase was extracted with hexane to remove lipids. The diluted water phase was incubated with the hydrolysis mixture (ascorbic acid, charcoal-stripped Helix Pomatia extract; Bisepra, France) for 2 h at 60°C. Isoflavonoids were extracted twice with diethyl ether. The ether phase was evaporated and dissolved in 300 μL of assay buffer (0.5% bovine serum albumin-Tris, pH 7.76). Samples were analyzed by time-resolved fluorescence immunoassay.

Determination of plasma concentrations of genistein. Blood was collected from mice 6 and 24 h after oral administration of 250 μg genistein. Plasma was prepared by conventional methods and freeze-dried. For analysis, samples were resolved in the same volume of distilled H2O as the original plasma volume. Enzymatic hydrolysis was performed as described (20). Briefly, 0.2 μL β-glucuronidase/mL (Boehringer Mannheim, Mannheim, Germany) and 0.1 μL sulfatase/mL (Sigma) were added, and samples were incubated overnight at 37°C. Isoflavonoids were extracted from plasma twice with diethyl ether. After evaporation of the ether phase, samples were suspended in 300 μL assay buffer (0.5% bovine serum albumin-Tris, pH 7.76). Levels of genistein were analyzed by time-resolved fluorescence immunoassay.

Time-resolved fluorescence immunoassay. Time-resolved fluorescence immunoassay was performed as indicated (20). Briefly, 15 μL of [3H]estradiol glucuronide (NEN Life Science Products, Wallac Oy, Turku, Finland) was added to tubes to measure recovery; 20 μL of hydrolyzed plasma was pipetted into prewashed goat anti-rabbit immunoglobulin G microtitre plate strips (Wallac Oy). To each well were added 100 μL antisera and 100 μL europium-labeled genistein. After incubation on a plate shaker at room temperature for 90 min, the strips were washed and 200 μL of dissociation-enhanced lanthanide fluorescence immunoassay enhancement solution (Wallac Oy) was added; after agitation for 5 min, an assay was performed in a VICTOR 1420 multilabel counter (Wallac Oy).

Western blot analysis. Western blot analysis was performed as described previously (4). Briefly, total protein, extracted from snap-frozen ascending and descending colon, was separated on a 12% SDS-polyacrylamide gel electrophoresis and subsequently blotted to a nitrocellulose membrane. The membranes were incubated overnight with sheep anti-CYP27B1 (The Binding Site, Heidelberg, Germany), sheep anti-CYP24 (kind gift from Dr. Moray Campbell, University of Birmingham, U.K.) and mouse anti-cytokeratin 8 (CK-8) (Cymbus Biotech, Chandlers Fort, Hant, U.K.) antibodies. Horseradish peroxidase–conjugated secondary antibodies (Amersham Life Sciences, Buckinghamshire, U.K.) were used. Subsequent detection was performed with the SuperSignal CL-HRP substrate system (Pierce, Rockford, IL). Bands were evaluated by densitometry with a video camera imaging system (Herolab, Wiesloch, Germany).

Semiquantitative reverse transcription–polymerase chain reaction. For analysis of CYP27B1 and CYP24 mRNA in relation to CK 8 mRNA expression by reverse transcription–polymerase chain reaction (RT-PCR), total RNA was extracted from snap-frozen mouse colon with TRIzol reagent (Invitrogen, Paisley, U.K.); 2 μg total RNA was used for synthesis of single-stranded cDNA (SUPERSCRIPT II kit, Invitrogen). Multiplex RT-PCR (i.e., simultaneous amplification of transcripts
Specific for either CYP27B1 or CYP24 and a transcript specific for the epithelial cell marker CK 8) was used for semiquantitative evaluation of mRNA expression levels.

PCR conditions were established for each individual primer pair and subsequently adapted for multiplex PCR with the CK 8 primers: 15 s at 94°C, 30 s at 66°C and 1 min at 72°C for 34 cycles using the GeneAmp PCR System 9600 (PE Applied Biosystems, Foster City, CA). Primers were used the following: CYP27B1 sense, 5’-CAA GCA GCC GGC TAT GCT GG-3’; CYP27B1 antisense, 5’-TGT CTG GGA CAC GGG AAT TGC-3’; CYP24 sense, 5’-AAG GAC ACA GAG GAA GAA GCC -3’; CYP24 antisense, 5’-GGA TGG CAC ACT TGG GGT AA-3’; CK 8 sense, 5’-GTG CCC AGT ACG AGG ACA TGG TTG-3’; and CK 8 antisense, 5’-TGT TGC GGT TCA TCT CGG AG-3’. PCR products were checked for correct size and fragment length through multiple digestions with restriction enzymes. Gels were scanned and analyzed with a video camera imaging system (Herolab), and band density was measured under ultraviolet light. The levels of CYP27B1 and CYP24 expression were correlated with that of the epithelial cell marker CK 8.

Statistical analyses. Data are presented as mean ± sd. Student’s t test was used for statistical group analysis. P values < 0.05 were considered statistically significant.

RESULTS

Effect of gavage with genistein

Plasma and colon tissue accumulation. When 250 μg genistein was administered via gavage and blood was drawn from the tail vein after 6 h, we found a >3000-fold accumulation of genistein in plasma compared with control animals. The second blood sample after 24 h showed a steep decline of genistein concentration to a 40-fold accumulation. Interestingly, genistein accumulation was also observed in colon tissue after 24 h, reaching fivefold higher concentrations than that measured in control animals subjected to gavage with 5% ethanol in water (Table 1).

CYP27B1 mRNA and protein expression. CYP27B1 is apparently equally expressed in the ascending and descending mouse colon (Fig. 1A). When the effect of a single high genistein dose was monitored, it became apparent that both mRNA and protein expression levels of CYP27B1 were significantly and equally increased in the proximal and distal part of the mouse colon (Fig. 1B). CYP24 expression, however, was reduced by genistein (data not shown).

Effect of soy protein supplementation in the AIN 76 diet

CYP24 mRNA and protein expression. There is significant variation of CYP24 mRNA expression in colon segments. Although CYP24 mRNA is more highly expressed in the ascending colon of control animals and is suppressed by soy feeding, control values are much lower in colon descendens and soy does not affect its expression (Fig. 2A). In contrast, CYP24 protein is about equally distributed in different parts of the mouse colon and is approximately equally repressed by soy feeding (Fig. 2B). CYP27B1 levels were enhanced by the soy diet (data not shown).

DISCUSSION

Many epidemiologic and migrant studies support the concept of cancer prevention by nutritional means. The diet prevailing in Western industrialized countries may be a main cause of the high incidence of “Western” diseases. Among these diseases are major types of cancer known to be hormone dependent and colorectal cancer. Isoflavones are naturally occurring plant chemicals with striking similarity to the chemical structure of mammalian estrogens and therefore are classified as phytoestrogens. Soybeans, in particular the protein fraction, are the most abundant source of isoflavones; much smaller amounts are found in other beans and plants. Thus the average daily dietary intake of isoflavones in Western populations is typically negligible. In Asian countries such as Japan or China, soy is traditionally a staple, and in these countries the incidence of hormone-dependent cancers such as breast and prostate cancer is exceedingly low. This has been attributed, at least in part, to phytoestrogen consumption.

After ingestion, soybean isoflavones are hydrolyzed by intestinal glucosidases, which release the aglycones daidzein,
genistein and glycitein. In Japanese who consume their traditional diet, plasma concentrations of genistein of up to 2960 nmol/L are measured (21). This far exceeds human normal plasma estradiol concentrations, which range between 147 and 294 nmol/L. Relative molar binding affinities of different estrogenic compounds revealed that phytoestrogens have significantly higher affinities for estrogen receptor β than for estrogen receptor α (22). Recently it was suggested that in women the colon major type of estrogen receptor may be estrogen receptor β (23).

Although genistein, the major isoflavonoid present in soybeans, was shown to inhibit protein tyrosine kinase activity, which links its action to growth factor signaling pathways, a definite mechanism for cancer prevention in vivo has not been determined. One mode of preventive action may be via the regulation of cytochrome P450 enzymes involved in the synthesis of 1,25-D3. This secosteroid has consistently been shown in many laboratories to have general antimitotic properties and to be effective in vitro (24). However, elevation of its serum levels would result in hypercalcemia. Thus the concept of extrarenal production of 1,25-D3 with localized tissue accumulation and action is very attractive.

We were able to demonstrate increased expression of CYP27B1 and decreased expression of CYP24 in proximal and distal mouse colon after a single high oral dose of genistein. CYP27B1 mediates 1α-hydroxylation of the precursor 25-D3, whereas CYP24 could, when highly expressed, conceivably degrade newly synthesized 1,25-D3 or preferentially 24-hydroxylate the 25-D3 precursor. Thus regulation of these hydroxylases could result in optimal autocrine/paracrine production of the secosteroid in colon cells. Interestingly, this hydroxylase regulation coincides with measurable accumulation of genistein in mouse colon tissue, which is not as substantial as that reported by us in the mouse prostate, where 50-fold tissue concentrations of genistein are reached via some unknown concentrative mechanism (25).

During human food consumption, even under optimal conditions, high concentrations of genistein such as we used for in vitro studies can be achieved by nutritional means, we fed mice the basic AIN-76A diet with 20% soy as the protein source. This resulted in consistent downregulation of CYP24 in the mouse colon, which again would lead to optimal autocrine/paracrine production of the antimitotic secosteroid 1,25-D3. Although a conclusive link among colonic 1,25-D3 synthesis, phytoestrogen consumption and reduced human colon cancer incidence still needs to be established, it is interesting in this respect that colon cancer morbidity and mortality rates are lower in women than in men (26) and that hormone replacement therapy has consistently been implicated in reduced risk of colon cancer occurrence (27).

LITERATURE CITED