Ingestion of Green Tea Rapidly Decreases Prostaglandin E₂ Levels in Rectal Mucosa in Humans

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Abstract

The objective of this Phase I/II study was to assess the potential for green tea to be used as a colorectal cancer chemopreventive agent. This study measured the dose-related biological effects of administration of a single dose of green tea on the rectal mucosa of normal volunteers. Volunteers were admitted to the Robert Wood Johnson Medical School Clinical Research Center for 24 h. Baseline blood and rectal biopsy samples were obtained before the volunteers drank 0.6, 1.2, or 1.8 g of green tea solids dissolved in warm water. Blood samples were taken 2, 4, 8, and 24 h after the tea administration. Rectal biopsies were obtained at 4, 8, and 24 h. Prostaglandin E₂ (PGE₂) levels were analyzed by ELISA. Tea polyphenol levels in the blood, urine, and rectal tissue were measured by high-performance liquid chromatography using a Coulochem electrode array detection system. Statistical comparisons were made using ANOVA. Decreased levels of PGE₂ in rectal mucosa were observed at 4 and 8 h after consumption of green tea. There was no correlation between inhibition of PGE₂ and tissue or plasma levels of tea polyphenols. Ten of 14 subjects demonstrated a response rate, we believe these data support the study of green tea as a colorectal chemopreventive agent in more long-term Phase II trials.

Introduction

In the United States, colorectal cancer is the third most common cancer in men and the second most common cancer in women; it is also the second leading cause of cancer deaths overall (1). Although the mortality rate has been falling over the past 5 decades, the incidence of colorectal cancer has increased slightly since the early 1970s (2). The frequency of colorectal cancer in the American population and the personal and economic impact of treatment and associated morbidity make it a potentially rewarding target for preventive strategies. Although early detection and treatment have been demonstrated to reduce colorectal cancer mortality, primary prevention potentially offers far greater benefits by reducing morbidity and the costs of colorectal cancer treatment (3).

Tea (Camellia sinensis) has been suggested as a possible cancer-preventive agent (4). Most tea is consumed in the form of black tea (78%, primarily in Western countries) or green tea (20%, primarily in Asian countries; Ref. 4). Green tea contains a variety of polyphenols, including flavanols, flavandiols, flavanoids, and phenolic acids. Flavanols (catechins) are the major tea polyphenols. The major components responsible for the putative anticarcinogenic activities of tea are not clearly known. In green tea, however, the catechins EGCG,3 EGC, (-)-epigallocatechin-3-gallate, and EC are believed to be the active components because inhibition of carcinogenesis has been demonstrated with EGCG and mixtures of these polyphenols (4–9).

Data from animal models of carcinogenesis demonstrate a protective effect of green tea and catechins against carcinogenesis in skin, lung, esophagus, stomach, duodenum, liver, and pancreas (4–9). Several animal studies have demonstrated a protective effect of green tea on colon carcinogenesis (20–23). Rats that were chemically induced to form colon tumors were protected by low doses of green tea polyphenols (0.01% or 0.1% in the drinking water; Ref. 20). In another study using very low doses of green tea (0.05%, 0.01%, or 0.002%), rats ingesting the tea had a significantly lower incidence of colon carcinomas (21). Yet, a more recent study using a similar model did not find significant inhibition of colon carcinogenesis by black tea (24).

Epidemiological studies of the relationship between tea consumption and human cancer are plagued by multiple confounders and are not conclusive (25). The epidemiological evidence seems suggestive of a protective effect of tea consumption against the development of colorectal, uterine, and gastric cancers (4, 26, 27). Factors that confound the interpretation of these types of studies include influences of the temperature of the tea on the increased risk for esophageal cancer (26), type of tea consumed (green versus black), variations in methods of tea preparation, and lifestyle factors that may be associated with tea drinking.

A variety of mechanisms have been suggested for the anticarcinogenic effects of tea polyphenols. Tea polyphenols

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1 The abbreviations used are: EGCG, (-)-epigallocatechin-3-gallate; EGC, (-)-epigallocatechin; ECG, (-)-epicatechin-3-gallate; EC, (-)-epicatechin; PGE₂, prostaglandin E₂; HPLC, high performance liquid chromatography.

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are antioxidants with the ability to scavenge superoxide anions, singlet oxygen, and lipid peroxo-radicals, and to bind metal ions (4, 28–30). These antioxidant functions of tea polyphenols are generally considered to be important to the anticarcinogenic activities of tea. Inhibition of tumor promotion-related enzymes, including ornithine decarboxylase (31, 32), protein kinase C (33), lipoxygenase (34, 35), and cyclooxygenase (34, 35) by tea preparations may also be important. These activities may be particularly important when considering the potential use of tea preparations for prevention of colorectal cancer.

The effect of green tea on human colon carcinogenesis is unknown. To explore the potential of green tea as a colorectal cancer preventive, we initiated a Phase I trial of the effect of a single ingestion of green tea on a set of colorectal carcinogenesis biomarkers—basal PGE₂ levels, cyclooxygenase activity, crypt cell proliferation, and apoptosis—on the rectal mucosa of normal volunteers.

Materials and Methods

Subjects. A preliminary study to assess tissue penetration of tea polyphenols was carried out in 10 volunteer subjects scheduled to undergo abdominal surgery with resection of a portion of the intestine. Each of these preliminary subjects ingested 1.2 g of standardized green tea ~12 h before surgery. At the time of surgery, 1–10 g of full-thickness intestinal tissue (small bowel, colon, or rectum) was harvested, flash frozen in liquid nitrogen, and stored at ~70°C for subsequent determination of tissue polyphenol levels.

For the formal Phase I trial, 15 normal volunteers (ages 22–57) were admitted to the Robert Wood Johnson Medical School Clinical Research Center. They were instructed to start a clear liquid diet devoid of fruit and vegetable juices and tea 36 h before the study and to cease all food and beverage intake, except water, 12 h before the study. None of the volunteers were pregnant or lactating; had renal, hepatic, cardiac, or pulmonary dysfunction, or a coagulopathy; had a history of inflammatory bowel disease or colorectal cancer; were taking contraceptive medications; were smokers; had asthma; or had any other disorder, condition that occurred was one episode of persistent bleeding from a biopsy site requiring transport of the subject to an operating room for better visualization and suture ligation. Twenty-two of the volunteer subjects participated in the Phase I study. A total of 60 anoscopy biopsy specimens were obtained with biopsy forceps introduced through an anoscope. A single dose (0.6, 1.2, or 1.8 g) of standardized green tea powder (courtesy of Thomas J. Lipton Co., Englewood Cliffs, NJ) containing ~10% by weight of polyphenols (36) was mixed in warm water and drunk by the volunteers at 9:00 a.m. Blood was collected at each time point; half were snap frozen in liquid nitrogen, and stored at ~70°C. Urine was collected with each void, pooled into appropriate time periods, and was stored frozen at ~70°C.

The levels of tea polyphenols in plasma, urine, and rectal biopsies were determined by a HPLC procedure (37). In brief, the plasma, tissue, or urine was incubated with β-glucuronidase and sulfatase to convert conjugated polyphenol species to the free forms. The resulting free tea polyphenols were extracted into ethyl acetate, dried under nitrogen, and redissolved in buffer. The individual catechins were analyzed by HPLC using a Coulochem electrode array detection system. To determine basal PGE₂ levels, frozen colonic biopsy samples were homogenized at ice temperature in 0.3 ml of 0.1 M Tris-HCl buffer containing 5.6 μM indomethacin (pH 7.4) with a microtube pestle and vortexed thoroughly for 2 min. The homogenate was immediately analyzed for basal PGE₂ levels by ELISA according to the manufacturer’s instructions (Cayman Chemical).

After fixation, rectal biopsies were embedded in paraffin and serially sectioned at 5 mm and stained with H&E for histological analysis. The proliferation pattern was analyzed by immunostaining with anti-Ki67 (MIB-1) antibody according to a previously published procedure (38). The pattern of apoptotic cells was also analyzed. Apoptotic cells were visualized in histopathological sections that maintain crypt architecture by use of terminal deoxynucleotidyl transferase-mediated nick end labeling of apoptotic DNA strand breaks (39)

All biomarker parameters were assayed and interpreted by an observer blinded to the treatment dose and to the timing of the biopsy (baseline versus some time point after ingestion of tea).

Results

Preliminary Study of Tea Polyphenol Levels in Intestinal Mucosa. Ten subjects (ages 28–80, five females) received 1.2 g of standardized green tea ~12 h before laparotomy. Grossly normal intestinal tissue was harvested from the operative specimens and analyzed for tea polyphenol content. The mean ± SE of the polyphenol tissue concentration (ng/g wet tissue) was 382 ± 146 for EGCG, 89 ± 28 for EGC, and 39 ± 23 for EC. These data indicate that significant levels of tea polyphenols were detectable after ingestion of a single, moderate “dose” of tea, equivalent to two to three cups of brewed tea.

Phase I Study. Fifteen subjects (ages 22–57, two females) participated in the Phase I study. A total of 60 anoscopy biopsy procedures (4 per subject) were carried out. The only complication that occurred was one episode of persistent bleeding from a biopsy site requiring transport of the subject to an operating room for better visualization and suture ligation.

Tea Polyphenol Pharmacokinetics. Plasma concentration versus time curves for EGCG and EC are shown in Fig. 1, A and B, respectively. Due to an interfering peak, HPLC results for EGC were obscured. Maximal plasma concentration (Cₘₐₓ) was clearly dose dependent. Maximum plasma polyphenol concentrations occurred at 2 h after tea ingestion. The mean plasma...
half-life for each of the polyphenols was ~4 h for both EGCG and EC.

The highest EGC and EC levels were observed in urine samples collected between 0 and 4 h (Table 1). EGCG was not excreted in the urine, consistent with the previous results (37). Tissue concentration/time data for the tea polyphenols in rectal mucosa were variable and not clearly dose dependent (Table 2).

There was no evident correlation between the dose of tea ingested and the time-dependent concentration of polyphenols in the rectal mucosa.

Levels of PGE₂. Ingestion of green tea decreased rectal mucosal concentrations of PGE₂ at 4 and 8 h after tea administration (Fig. 2 and Table 3). At 24 h, the PGE₂ concentrations returned to the baseline levels. Although the high dose of tea (1.8 g) was more effective in lowering the PGE₂ levels than the low (0.6 g) and moderate (1.2 g) doses, a good dose-dependent effect was not observed because the moderate dose did not produce lower PGE₂ levels than the low dose. Seventy-one percent (10 of 14) exhibited a "response" to tea, as evidenced by a 50% reduction in basal PGE₂ levels at 4 h after ingestion.

Crypt Cell Proliferation and Apoptosis. Because of wide variability in the histological appearance of immunohistochemical stains of rectal biopsies, presumably reflecting the wide biological variability in individuals (40), there was no clearly evident effect of green tea ingestion on crypt cell proliferation or apoptosis (data not shown).

Discussion

One strategy to reduce cancer-related deaths is through chemoprevention. Chemoprevention can be defined as controlling the occurrence of cancer by slowing, blocking, or reversing the development of the disease by the administration of naturally occurring or synthetic compounds. Chemopreventive compounds should ideally have: (a) little or no toxic effect; (b) high

Table 1  Mean EGC and EC values* per green tea dose in urine over time

<table>
<thead>
<tr>
<th>Dose</th>
<th>Time interval (h)</th>
<th>EGC</th>
<th>EC</th>
</tr>
</thead>
<tbody>
<tr>
<td>0.6</td>
<td>0–4</td>
<td>1310±561</td>
<td>1310±465</td>
</tr>
<tr>
<td></td>
<td>4–8</td>
<td>552±392</td>
<td>492±276</td>
</tr>
<tr>
<td></td>
<td>8–24</td>
<td>189±154</td>
<td>304±130</td>
</tr>
<tr>
<td>1.2</td>
<td>0–4</td>
<td>2680±864</td>
<td>3100±996</td>
</tr>
<tr>
<td></td>
<td>4–8</td>
<td>300±84</td>
<td>480±147</td>
</tr>
<tr>
<td></td>
<td>8–24</td>
<td>600±383</td>
<td>660±320</td>
</tr>
<tr>
<td>1.8</td>
<td>0–4</td>
<td>4440±1580</td>
<td>7760±2560</td>
</tr>
<tr>
<td></td>
<td>4–8</td>
<td>600±205</td>
<td>1140±330</td>
</tr>
<tr>
<td></td>
<td>8–24</td>
<td>440±93</td>
<td>560±144</td>
</tr>
</tbody>
</table>

* Values are mean ng polyphenol/ml of urine ± SE of five subjects. EGCG is not excreted in the urine.

Table 2  Mean EGC, EC, and EGCG values* per green tea dose in rectal biopsies over time

<table>
<thead>
<tr>
<th>Dose</th>
<th>Time point</th>
<th>EGC</th>
<th>EC</th>
<th>EGCG</th>
</tr>
</thead>
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<tr>
<td>0.6</td>
<td>0</td>
<td>1.2±0.7</td>
<td>0.9±0.5</td>
<td>5.0±2.7</td>
</tr>
<tr>
<td></td>
<td>4</td>
<td>103±31</td>
<td>4.4±3.5</td>
<td>37.0±30.9</td>
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<tr>
<td></td>
<td>8</td>
<td>160±6.7</td>
<td>5.3±4.3</td>
<td>25.4±20.3</td>
</tr>
<tr>
<td></td>
<td>24</td>
<td>0.8±0.5</td>
<td>0.6±0.3</td>
<td>3.05±1.7</td>
</tr>
<tr>
<td>1.2</td>
<td>0</td>
<td>3.4±1.9</td>
<td>2.0±1.1</td>
<td>10.2±5.4</td>
</tr>
<tr>
<td></td>
<td>4</td>
<td>11.1±6.4</td>
<td>7.0±3.8</td>
<td>41.3±25.1</td>
</tr>
<tr>
<td></td>
<td>8</td>
<td>4.2±0.2</td>
<td>2.9±0.5</td>
<td>14.8±2.8</td>
</tr>
<tr>
<td></td>
<td>24</td>
<td>2.3±1.2</td>
<td>1.8±0.9</td>
<td>8.2±5.2</td>
</tr>
<tr>
<td>1.8</td>
<td>0</td>
<td>0.0±0</td>
<td>0.0±0</td>
<td>0.6±0.6</td>
</tr>
<tr>
<td></td>
<td>4</td>
<td>7.8±6.9</td>
<td>6.8±4.9</td>
<td>31.2±23.9</td>
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<tr>
<td></td>
<td>8</td>
<td>4.2±2.6</td>
<td>2.2±1.2</td>
<td>28.5±18.0</td>
</tr>
<tr>
<td></td>
<td>24</td>
<td>1.8±1.5</td>
<td>1.6±0.8</td>
<td>7.1±4.4</td>
</tr>
</tbody>
</table>

* Values are mean ng polyphenol/mg protein in rectal biopsy homogenate ± SE of three, three, and five subjects for doses 0.6 g, 1.2 g, and 1.8 g, respectively.

Due to problems with some samples, two subjects were excluded from the analysis from groups 0.6 and 1.2 doses. Thus, n = 3 for these groups.
Green Tea for Colorectal Cancer Prevention

efficacy against multiple types of cancer; (c) oral administration; (d) known mechanisms of action; (e) low cost; and (f) human acceptance (41–44). On the basis of these criteria, it is logical that compounds found naturally in foods and drinks have been investigated. Tea is thought to meet these criteria because it is widely consumed, has no known toxicity, and has demonstrated inhibitory activity against carcinogenesis in animal models (4, 22–25).

Our preliminary study demonstrated that measurable amounts of polyphenols are present in the gastrointestinal tissues. Our present and previous results (37) on plasma and urine concentrations of tea catechins indicated that these compounds are absorbed. The effect of p.o. administered tea was rapid; at 4 and 8 h after tea administration, PGE2 levels were significantly lower at all doses of tea. Although the 1.8-g dose seemed to be more effective in lowering PGE2 formation than the 0.6-g dose, a good dose-response relationship was not observed. This was probably due to the variability of the individuals and sampling procedure. There were no correlations between the inhibition of PGE2 formation and the tea polyphenol levels in the plasma or rectal tissue. This lack of correlation may be explained by individual variations and the difficulties caused by repeated rectal biopsies interfering with each other. The biopsys-induced inflammation may mask some of the effects of the tea.

Of interest, however, is that although increased inflammatory changes were seen in the immunohistochemical slides over time, lowering of PGE2 levels was observed. Whereas an increase in inflammation may be explained by serial anoscopies, inhibition of PGE2 may be the result of tea administration. This is believed to be due to the inhibition of cyclooxygenase-2 activity in the rectum. Preliminary experiments, however, indicate that the level of cyclooxygenase in the rectal mucosa, as determined by Western blot analysis, was not significantly affected by the tea treatment (data not shown).

The immunohistochemical slides regarding proliferation and apoptotic rates were difficult to interpret. Individual variations in these rates were very high and make it difficult to determine the effect of tea on these biomarkers. Recent analysis of the variability in rectal mucosal proliferation measurements calls into question the ability of proliferating cell nuclear antigen to show treatment effects (40). Potential sources of variation include the effect of bowel preparation, age, gender, diet, time of day, fluctuation over time, and biopsy location. Similar difficulties are likely to occur with the measurement of apoptotic cells, as with proliferation indices (45). Because subjects were able to act as their own control over time, certain variables were likely to play a role in this study’s difficulty in assessing proliferation and apoptosis. Proliferation indices are known to have circadian rhythm (46) that could affect our ability to demonstrate tea treatment changes over a 24-h period.

Green tea has biological activity in the colorectal mucosa. Because PGE2 inhibition in other settings is related to a lower risk of certain cancers and the response rate among the subjects was 10 of 14 (71%), these data suggest a potential role for green tea as a chemopreventive agent against colorectal cancer. The data support proceeding with a longer-term Phase II study.

**References**


**Table 3** Percent inhibition of PGE2 formation over time after green tea consumption in colorectal tissue

<table>
<thead>
<tr>
<th>Dose</th>
<th>4 h*</th>
<th>8 h*</th>
<th>24 h*</th>
<th>Response rate</th>
</tr>
</thead>
<tbody>
<tr>
<td>0.6</td>
<td>58.3 ± 12†</td>
<td>63.1 ± 12‡</td>
<td>17.2 ± 1§</td>
<td>4/5</td>
</tr>
<tr>
<td>1.2</td>
<td>49.6 ± 14†</td>
<td>41.0 ± 11‡</td>
<td>17.0 ± 4§</td>
<td>3/5</td>
</tr>
<tr>
<td>1.8</td>
<td>78.2 ± 10‡</td>
<td>60.5 ± 16‡</td>
<td>0 ± 25§</td>
<td>3/4</td>
</tr>
</tbody>
</table>

* Values are the mean percent inhibition compared with the baseline (0 h) time point ± SE for four or five subjects.
† Response rate is described as at least 50% inhibition at 4 h (responders/n).
‡ Inhibition is significantly different from the baseline time point.
§ Inhibition is significantly different from the other time points, but not the baseline point.

PGE2 levels were not determined in one subject because of lack of biopsy material.


