Antioxidant Beverages: Green Tea Intake and Coronary Artery Disease

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ABSTRACT: Coronary artery disease (CAD) is recognized as an inflammatory disease. In the present study, we investigated the effect of green tea consumption on plasma inflammatory markers and the association between green tea consumption and CAD. In 22 healthy volunteers, green tea consumption (7 cups/day) significantly decreased serum malondialdehyde-modified low-density lipoprotein (MDA-LDL) concentrations, whereas green tea consumption tended to decrease plasma C-reactive protein and interleukin (IL)-6 concentrations. In 725 patients undergoing coronary angiography, the percentage of patients drinking <1 cup/day of green tea was higher in patients with myocardial infarction (MI) than in CAD patients without MI and patients without CAD (29% vs. 15% and 18%, P < 0.01). Green tea consumption was found to be inversely associated with MI in Japanese patients. The protective effect of green tea against atherosclerosis is more likely to be because of the inhibitory effect of LDL oxidation than because of anti-inflammatory effect.

KEYWORDS: antioxidant, green tea, coronary artery disease, myocardial infarction, inflammation

SUPPLEMENT: Inflammation, Atherosclerosis and Coronary Artery Disease

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Introduction

Inflammation has been recognized to play an important role in both the initiation and progression of atherosclerotic diseases, such as coronary artery disease (CAD). Several prospective studies demonstrated that plasma high-sensitivity C-reactive protein (hsCRP) levels, which are some of the markers of systemic inflammation, are powerful predictors of future myocardial infarction (MI) and cardiac death among apparently healthy individuals. The high hsCRP levels were also reported to be associated with an increased risk of further coronary events in patients with CAD. We previously reported plasma hsCRP levels to be associated with the presence and extent of coronary stenosis in patients with stable CAD. Moreover, we showed that plasma hsCRP levels correlated with the severity of both coronary and aortic atherosclerosis using magnetic resonance imaging (MRI).

Low-density lipoprotein (LDL) oxidation plays a key role in the development of atherosclerosis. Several enzyme-linked immunosorbent assay (ELISA) methods had been developed to measure oxidatively modified LDL levels in blood. Elevated levels of malondialdehyde-modified LDL (MDA-LDL), an oxidized LDL, have been reported in patients with CAD, especially in those with MI.

The intake of dietary antioxidants such as polyphenols is a potential therapy to prevent LDL oxidation and atherosclerosis progression. Polyphenols are mainly found in fruits, vegetables, and beverages. In Japan, green tea, which is very rich in catechins, is the most popular beverage and the major source of polyphenol intake (>80% of polyphenol intake). We previously reported that the daily consumption of green tea decreased serum MDA-LDL concentrations in 22 healthy Japanese male volunteers. Using animal models, green tea
was also shown to have anti-inflammatory effect in addition to the inhibitory effect against LDL oxidation.\textsuperscript{13–15} However, anti-inflammatory effect of green tea has not yet been elucidated \textit{in vivo}. Moreover, we previously reported that green tea consumption was inversely associated with MI in 393 Japanese patients undergoing coronary angiography.\textsuperscript{16} The present study extends our previous two reports\textsuperscript{12,16} by demonstrating the changes in plasma inflammatory markers by green tea consumption in 22 healthy volunteers and by increasing the number of study patients (from 393 to 725 patients) who underwent coronary angiography to elucidate the association between antioxidant beverages and CAD.

**Methods**

**Study design and subjects.** The study design was reported previously.\textsuperscript{12} Briefly, the study consisted of a one-week run-in period, a two-week water intake period, and then a subsequent two-week green tea intake period with 22 healthy Japanese male volunteers. Over the course of the study, we asked all study subjects to maintain their regular dietary habits but not to consume any kind of tea, wine, citrus liquor, or vitamin supplements. After the run-in period, all subjects consumed seven cups/day of water for two weeks (water period) and then consumed seven cups/day of green tea for the next two weeks (green tea period). During the water or green tea periods, they drank four cups (breakfast and lunch, two cups each) and three cups (dinner) of water or green tea after each meal (a total of seven cups/day comprising 700 mL). The green tea used in our study was a commercially available freeze-dried tea, sarasara ryokucha (Itoen Co.), which is on sale for drinking in Japan. One stick (0.9 g) of the freeze-dried tea can be dissolved in 100 mL of hot water and then be drunk as one cup of green tea. As a result, seven cups of green tea contain 542.5 mg of catechins. At the end of each period, subjects fasted overnight and blood samples were taken at 9:00 am on the next morning. Plasma hsCRP concentrations were measured using a BN II nephelometer (Dade Behring), and plasma interleukin (IL)-6 concentrations were measured by ELISA with a human IL-6 immunoassay kit (BioSource International). Plasma LDL-cholesterol concentrations were measured by the direct enzymatic method with a commercially available kit (Cholestest LDL, Daiichi Pure Chemicals), and serum MDA-LDL concentrations were measured using a BN II nephelometer (Dade Behring), any differences between two groups were evaluated by the unpaired $t$-test for continuous variables and by the chi-square test for categorical variables. Any differences among three groups were evaluated by repeated-measures ANOVA. A Bonferroni adjustment (0.05/3) was undertaken because the three periods were compared. A $P$ value of $<0.05$ was considered statistically significant.

Regarding the data of 22 healthy volunteers, to test whether or not the distributions of variables are deviating from a normal distribution, the $F$-test was applied to measure variables. Since plasma hsCRP, IL-6, and catechin concentrations were considered as non-parametric variables, these results are expressed as the median value, and any differences in these variables among the three periods were evaluated by the Friedman’s test. For parametric variables, results are expressed as the mean value $\pm$ SD, and any differences among the three periods were evaluated by repeated-measures ANOVA. A Bonferroni adjustment (0.05/3) was undertaken because the three periods were compared. A $P$ value of $<0.05$ was considered statistically significant.

**Results**

**Green tea intake and inflammatory markers.** Of the 22 study subjects, 20 had been in the habit of drinking green tea before the study. As reported previously,\textsuperscript{12} plasma total catechin concentrations significantly decreased at the end of the water period and then increased at the end of the green tea period (Table 1). Although there was no change in LDL-cholesterol concentrations, serum MDA-LDL concentrations (84 $\pm$ 45 vs. 76 $\pm$ 40 IU/L, $P < 0.05$) and the MDA-LDL/
Table 1. Plasma catechins, hsCRP, and IL-6 concentrations at the end of each period in 22 healthy subjects.

| RUN-IN | WATER | GREEN TEA |
|--------|-------|-----------|
| Total catechins (nmol/L) | 17.6 | 12.6* | 42.0** |
| EGCG (nmol/L) | 8.0 | 4.8* | 21.2** |
| GCG (nmol/L) | 7.0 | 7.0 | 8.6 |
| ECg (nmol/L) | 2.5 | 1.6 | 7.6|
| LDL-cholesterol (mg/dL) | 113 ± 32 | 110 ± 33 | 113 ± 28 |
| MDA-LDL (IU/L) | 81 ± 38 | 84 ± 45 | 76 ± 40a |
| MDA-LDL/LDL-cholesterol | 0.69 ± 0.18 | 0.74 ± 0.21 | 0.65 ± 0.20 |
| hsCRP (mg/L) | 0.21 | 0.28 | 0.19 |
| IL-6 (ng/L) | 0.16 | 0.18 | 0.15 |

Notes: Data are presented as the median value (n = 22) except for LDL-cholesterol and MDA-LDL concentrations, and MDA-LDL/LDL-cholesterol ratio that are presented as the mean value ± SD. *P < 0.05 vs. the run-in period; **P < 0.001, †P < 0.02, and ‡P < 0.05 vs. the water period. Total catechins concentration was defined as the sum of EGCG, GCG, and ECg values.

LDL-cholesterol ratio (0.74 ± 0.21 vs. 0.65 ± 0.20, P < 0.02) significantly decreased at the end of the green tea period. Regarding inflammatory markers, plasma hsCRP and IL-6 concentrations tended to decrease at the end of the green tea period (0.28 vs. 0.19 mg/L and 0.18 vs. 0.15 ng/L, respectively) (Table 1). However, these changes did not reach statistical significance in either inflammatory marker.

Green tea intake and CAD. Of the 725 study patients, 517 (71%) were found to have CAD on coronary angiograms, of whom 225 (43%) had MI. Clinical characteristics of study patients are shown in Table 2. Compared with patients without CAD, CAD patients with and without MI were predominantly male and had lower HDL-cholesterol levels and higher rates of diabetes mellitus and smoking. With regard to the intake of antioxidant foods, there were no significant differences between patients without CAD and CAD patients with or without MI (Table 3). However, the percentages of patients with <3 times/week intake of soybeans and fruits were significantly higher in CAD patients with MI than in those without MI (38% and 41% vs. 26% and 32%, P < 0.05). With regard to the intake of antioxidant beverages, there were no significant differences between patients without CAD and CAD patients without MI. Notably, the percentage of patients drinking <1 cup/day of green tea (green tea nondrinkers) was significantly higher in CAD patients with MI than in those without MI and in patients without CAD (29% vs. 15% and 18%, P < 0.01). The multivariate analysis revealed that green tea intake of <1 cup/day and soybeans intake of <3 times/week were significant factors associated with MI, independent of traditional risk factors and the intakes of other antioxidant foods and beverages (Table 4). The odds ratio for MI was 1.76 (95% confidence interval (CI) = 1.20–2.58) for green tea intake <1 cup/day and 1.43 (95% CI = 1.02–2.02) for soybeans intake <3 times/week.

Discussion
As shown in our previous report, plasma catechin concentrations significantly increased with green tea consumption (seven cups/day). Although there was no change in plasma LDL-cholesterol concentrations, green tea consumption significantly decreased serum MDA-LDL concentrations and the ratio of MDA-LDL/LDL-cholesterol, thus suggesting that green tea has an inhibitory effect against LDL oxidation in vivo. In addition to the inhibitory effect against LDL oxidation, several studies reported that green tea, which is very rich in catechins, has anti-inflammatory effect in animal models. Li et al. showed that EGCG inhibited angiotensin II- and IL-6-stimulated CRP expression in macrophages by interfering with reactive oxygen species generation. In the present study, we investigated the changes in plasma hsCRP and IL-6 concentrations after two weeks of green tea consumption. The daily consumption of green tea (seven cups/day) tended to decrease these inflammatory markers, but the differences did not reach statistical significance. Therefore, the protective effect of green tea against atherosclerosis is more likely

Table 2. Clinical characteristics in patients with and without CAD.

|                | CAD(-) (N = 208) | CAD(-) VS. MI(-) | MI(-) (N = 292) | MI(-) VS. MI(+) | MI(+) (N = 225) | MI(+) VS. CAD(-) |
|----------------|------------------|-----------------|----------------|----------------|----------------|-----------------|
| Age (years)    | 61 ± 10          | <0.001          | 65 ± 9         | <0.02          | 63 ± 10         | NS              |
| Male           | 132 (63%)        | <0.001          | 230 (79%)      | <0.05          | 195 (87%)       | <0.001          |
| Hypertension   | 114 (55%)        | <0.005          | 203 (70%)      | <0.05          | 133 (59%)       | NS              |
| Systolic BP (mmHg) | 130 ± 17     | <0.001          | 136 ± 19       | <0.002         | 130 ± 21        | NS              |
| Hyperlipidemia | 71 (34%)         | <0.001          | 161 (55%)      | <0.01          | 94 (42%)        | NS              |
| TC (mg/dL)     | 204 ± 35         | NS              | 202 ± 34       | <0.01          | 194 ± 36        | <0.0005         |
| HDL-C (mg/dL)  | 57 ± 16          | <0.001          | 50 ± 15        | <0.05          | 47 ± 13         | <0.001          |
| Diabetes mellitus | 33 (15%)    | <0.001          | 101 (35%)      | NS             | 68 (29%)        | <0.0005         |
| Cigarettes smokers | 75 (36%)    | <0.05           | 135 (46%)      | <0.05          | 125 (56%)       | <0.001          |

Notes: Hypertension was defined as >140/90 mmHg or on drugs. Hyperlipidemia was defined as TC >240 mg/dL or on drugs.
Abbreviations: NS, not significant; BP, blood pressure; TC, total cholesterol; HDL-C, high density lipoprotein-cholesterol.
to be because of the inhibitory effect of LDL oxidation than because of anti-inflammatory effect.

We previously reported that green tea consumption was inversely associated with MI in 393 Japanese patients undergoing coronary angiography. The present study extended our previous report by increasing the number of study patients (from 393 to 725 patients) to elucidate the associations between antioxidant beverages and foods, and CAD. In line with our previous report, we confirmed that green tea consumption was inversely associated with MI in 725 patients. A recent meta-analysis including our data from 393 patients indicated that routine green tea consumption was significantly associated with a decreased risk (17%) of CAD.

In hypercholesterolemic rabbits, green tea ingestion was reported to reduce aortic atherosclerosis, whereas the ingestion of black tea or vitamin E did not, suggesting the stronger anti-atherogenic effect of green tea than those of black tea or vitamin E. In our present study, the consumption of soybeans products was also associated with MI, but this association showed only borderline significance. Therefore, our results suggest that green tea consumption is more likely to play a major role in the prevention of CAD, especially MI, in Japanese patients than other antioxidant beverages and foods.

Our study has some limitations. First, one of our study limitations is the small number of study subjects (22 healthy subjects). Moreover, our study had no control group without

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**Table 3. Intakes of antioxidant beverages and foods in patients with and without CAD.**

| Antioxidant beverages | CAD(−) (N = 208) | CAD(−) VS. MI(−) P VALUE | MI(−) (N = 292) | MI(−) VS. MI(+) P VALUE | MI(+) (N = 225) | MI(+) VS. CAD(−) P VALUE |
|-----------------------|------------------|--------------------------|----------------|--------------------------|----------------|-------------------------|
| Green tea (cups/d)    |                  |                          |                |                          |                |                         |
| <1                    | 37 (18%)         | NS                       | 45 (15%)       | <0.001                   | 66 (29%)       | <0.01                   |
| 1–3                   | 100 (48%)        | NS                       | 140 (48%)      | <0.05                    | 87 (39%)       | NS                      |
| >3                    | 71 (34%)         | NS                       | 107 (37%)      | NS                       | 72 (32%)       | NS                      |
| Coffee (cups/d)       |                  |                          |                |                          |                |                         |
| <1                    | 115 (56%)        | NS                       | 157 (54%)      | NS                       | 207 (48%)      | NS                      |
| 1–3                   | 75 (36%)         | NS                       | 112 (38%)      | NS                       | 90 (40%)       | NS                      |
| >3                    | 17 (8%)          | NS                       | 23 (8%)        | NS                       | 28 (12%)       | NS                      |
| Black tea (cups/d)    |                  |                          |                |                          |                |                         |
| <1                    | 183 (88%)        | NS                       | 263 (90%)      | NS                       | 207 (92%)      | NS                      |
| 1–3                   | 23 (11%)         | NS                       | 28 (10%)       | NS                       | 18 (8%)        | NS                      |
| >3                    | 1 (1%)           | NS                       | 1 (0%)         | NS                       | 0 (0%)         | NS                      |
| Cocoa (cups/d)        |                  |                          |                |                          |                |                         |
| <1                    | 201 (98%)        | NS                       | 279 (96%)      | NS                       | 220 (98%)      | NS                      |
| 1–3                   | 5 (2%)           | NS                       | 13 (4%)        | NS                       | 4 (2%)         | NS                      |
| Wine (glasses/d)      |                  |                          |                |                          |                |                         |
| <1                    | 202 (97%)        | NS                       | 286 (98%)      | NS                       | 214 (95%)      | NS                      |
| 1–3                   | 6 (3%)           | NS                       | 6 (2%)         | NS                       | 11 (5%)        | NS                      |
| Soybeans (times/wk)   |                  |                          |                |                          |                |                         |
| <3                    | 62 (30%)         | NS                       | 77 (26%)       | <0.01                    | 85 (38%)       | NS                      |
| 3–4                   | 69 (33%)         | NS                       | 115 (39%)      | NS                       | 73 (32%)       | NS                      |
| >4                    | 77 (37%)         | NS                       | 100 (34%)      | NS                       | 67 (30%)       | NS                      |
| Vegetables (times/wk) |                  |                          |                |                          |                |                         |
| <3                    | 37 (18%)         | NS                       | 41 (14%)       | NS                       | 44 (20%)       | NS                      |
| 3–4                   | 37 (18%)         | NS                       | 68 (23%)       | NS                       | 53 (24%)       | NS                      |
| >4                    | 133 (64%)        | NS                       | 183 (63%)      | NS                       | 128 (57%)      | NS                      |
| Fruits (times/wk)     |                  |                          |                |                          |                |                         |
| <3                    | 70 (34%)         | NS                       | 92 (32%)       | <0.05                    | 93 (41%)       | NS                      |
| 3–4                   | 38 (18%)         | NS                       | 61 (21%)       | NS                       | 38 (17%)       | NS                      |
| >4                    | 100 (48%)        | NS                       | 138 (47%)      | NS                       | 94 (42%)       | NS                      |

Abbreviation: NS, not significant.
green tea period, and our study was a sequential but not crossover design, thereby making it impossible to separate period effects from treatment effects. To clarify the results of our study, a further study should be done using a crossover design in a larger number of study subjects. Regarding the association between green tea consumption and CAD, our study is cross-sectional. Such a study cannot establish causality. It only shows some association and is hypothesis generating. Therefore, a prospective study is required to elucidate the preventative effect of green tea on CAD and MI.

Conclusion

Green tea consumption was found to be inversely associated with MI in Japanese patients. Green tea consumption significantly decreased serum oxidized LDL concentrations, but green tea consumption tended to decrease inflammatory markers; thus, suggesting the protective effect of green tea against atherosclerosis is more likely to be because of the inhibitory effect of LDL oxidation than because of anti-inflammatory effect.

Author Contributions

Conceived and designed the experiments: RO, YM. Analyzed the data: YM. Wrote the first draft of the manuscript: RO. Contributed to the writing of the manuscript: YM, KK. Made critical revisions and approved the final version: RO, YM, KK. Agree with the manuscript results and conclusions: RO, YM, KK. All authors reviewed and approved the final manuscript.

REFERENCES

1. Ross R. Atherosclerosis: an inflammatory disease. N Engl J Med. 1999;340:115–26.
2. Ridker PM, Cushman M, Stampfer MJ, Tracy RP, Hennekens CH. Inflammation, aspirin, and the risk of cardiovascular disease in apparently healthy men. N Engl J Med. 1997;336:703–9.
3. Koenig W, Sund M, Frohlich M, et al. C-reactive protein, a sensitive marker of inflammation, predicts future risk of coronary heart disease in initially healthy middle-aged men: results from the MONICA (monitoring trends and determinants in cardiovascular disease) Augsburg Cohort Study, 1984 to 1992. Circulation. 1999;99:237–42.
4. Haverkate F, Thompson SG, Pyke SD, Gallimore JR, Pepys MB. Production of C-reactive protein and risk of coronary events in stable and unstable angina. European concerted action on thrombosis and disabilities angina pectoris study group. Lancet. 1997;349:462–6.
5. Taniguchi H, Momiyama Y, Ohmori R, et al. Associations of plasma C-reactive protein levels with the presence and extent of coronary stenosis in patients with stable coronary artery disease. Atherosclerosis. 2005;178:173–7.
6. Momiyama Y, Ohmori R, Tsukiyama K, et al. Associations between plasma C-reactive protein levels and the severities of coronary and aortic atherosclerosis. J Atheroscler Thromb. 2010;17:460–7.
7. Steinberg D, Parthasarathy S, Carew TE, Khoo JC, Witztum JL. Beyond cholesterol. Modifications of low-density lipoprotein that increase its atherogenicity. N Engl J Med. 1989;320:915–24.
8. Holvoet P, Vanhaecke J, Janssens S, Van de Werf F, Collen D. Oxidized LDL and malondialdehyde-modified LDL in patients with acute coronary syndromes and stable coronary artery disease. Circulation. 1998;98:1487–94.
9. Tanaka K, Bjo R, Inoue M, et al. Increased circulating malondialdehyde-modified LDL concentrations in patients with coronary artery disease and their association with peak sizes of LDL particles. Arterioscler Thromb Vasc Biol. 2002;22:662–6.
10. Fukushima Y, Ohie T, Yonekawa Y, et al. Coffee and green tea as a large source of antioxidant polyphenols in the Japanese population. J Agric Food Chem. 2009;57:1253–9.
11. Hertog MG, Kromhout D, Aravanis C, et al. Flavonoid intake and long-term risk of coronary heart disease and cancer in the seven countries study. Arch Intern Med. 1995;155:351–8.
12. Hirano-Ohmori K, Takahashi R, Momiyama Y, et al. Green tea consumption and serum malondialdehyde-modified LDL concentrations in healthy subjects. J Am Coll Nutr. 2005;24:342–6.
13. Cunha CA, Lira FS, Rosa Neto JC, et al. Green tea extract supplementation induces the lipolytic pathway, attenuates obesity, and reduces low-grade inflammation in mice fed a high-fat diet. Mediators Inflamm. 2013;2013:635470.
14. Uchiyama Y, Imaizumi T, Mohizuki K, Guda T. Dietary supplementation with a low dose (-)-epigallocatechin-3-gallate reduces pro-inflammatory responses in peripheral leukocytes of non-obese type 2 diabetic GK rats. J Nutr Sci Vitaminol. 2013;59:541–7.
15. Heber D, Zhang Y, Yang J, Ma JE, Henning SM, Li Z. Green tea, black tea, and oolong tea polyphenols reduce visceral fat and inflammation in mice fed high-fat, high-sucrose obese diets. J Nutr. 2014;144:1385–93.
16. Hirano R, Momiyama Y, Takahashi R, et al. Comparison of green tea intake in Japanese patients with and without angiographic coronary artery disease. Am J Cardiol. 2002;90:1150–3.
17. Kotani K, Maekawa M, Kanno T, Kondo A, Toda N, Manabe M. Distribution of immunoreactive malondialdehyde-modified low-density lipoprotein in human serum. Biochim Biophys Acta. 1994;1215:121–5.
18. Li M, Liu JT, Pang XM, Han CJ, Mao JJ. Epigallocatechin-3-gallate inhibits angiotensin II and interleukin-6-induced C-reactive protein production in macrophages. Pharmacol Rep. 2012;64:912–8.
19. Wang ZM, Zhou B, Wang YS, et al. Black and green tea consumption and the risk of coronary artery disease: a meta-analysis. Am J Clin Nutr. 2011;93:506–15.
20. Tjibburg LBM, Wiseman SA, Meijer GW, Westrate JA. Effects of green tea, black tea and dietary lipophilic antioxidants on LDL oxidizability and atherosclerosis in hypercholesterolemic rabbits. Atherosclerosis. 1997;135:37–47.

Table 4. Factors associated with MI (multiple logistic regression analysis in 725 patients).

| VARIABLES                        | ODDS RATIO (95% CI) | P VALUE |
|----------------------------------|---------------------|---------|
| Gender (male)                    | 1.82(1.18–2.81)     | <0.01   |
| HDL-cholesterol (mg/dL)          | 0.98 (0.97–0.99)    | <0.002  |
| Soybean intake (<3 times/week)   | 1.43 (1.02–2.02)    | <0.05   |
| Green tea intake (<1 cup/day)    | 1.76 (1.20–2.58)    | <0.005  |

Notes: The dependent variable was MI. The analysis included age, gender, hypertension, hyperlipidemia, HDL-cholesterol, diabetes mellitus, smoking, and the intakes of green tea, soybeans, vegetables, and fruits.