

# Chlorogenic Acids from Green Coffee Extract are Highly Bioavailable in Humans<sup>1,2</sup>

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## Abstract

Chlorogenic acids (CGA) are cinnamic acid derivatives with biological effects mostly related to their antioxidant and antiinflammatory activities. Caffeoylquinic acids (CQA) and dicaffeoylquinic acids (diCQA) are the main CGA found in nature. Because green coffee is a major source of CGA, it has been used for production of nutraceuticals. However, data on the bioavailability of CGA from green coffee in humans are inexistent. The present study evaluated the pharmacokinetic profile and apparent bioavailability of CGA in plasma and urine of 10 healthy adults for 8 h after the consumption of a decaffeinated green coffee extract containing 170 mg of CGA. Three CQA, 3 diCQA, and caffeic, ferulic, isoferulic, and *p*-coumaric acids were identified in plasma by HPLC-Diode Array Detector-MS after treatment. Over 30% (33.1 ± 23.1%) of the ingested cinnamic acid moieties were recovered in plasma, including metabolites, with peak levels from 0.5 to 8 h after treatment. CGA and metabolites identified in urine after treatment were 4-CQA, 5-CQA, and sinapic, *p*-hydroxybenzoic, gallic, vanillic, dihydrocaffeic, caffeic, ferulic, isoferulic, and *p*-coumaric acids, totaling 5.5 ± 10.6% urinary recovery of the ingested cinnamic and quinic acid moieties. This study shows that the major CGA compounds present in green coffee are highly absorbed and metabolized in humans. J. Nutr. 138: 2309–2315, 2008.

## Introduction

Chlorogenic acids (CGA)<sup>5</sup> are phenolic compounds formed by the esterification of cinnamic acids, such as caffeic, ferulic, and *p*-coumaric acids, with (-)-quinic acid. A series of health benefits have been associated with the consumption of CGA in the last few years, such as reduction of the relative risk of cardiovascular disease, diabetes type 2, and Alzheimer's disease (1–3), and antibacterial and antiinflammatory activities (4,5). Their lactones also have been shown to exert positive effects in rats such as enhancement of insulin action (6).

Green (or raw) coffee is a major source of CGA in nature (5–12 g/100 g) (7). Recent studies demonstrated that the consumption of green coffee extracts produced antihypertensive effect in rats and humans (8,9), improvement in human vasoreactivity (10), inhibitory effect on fat accumulation and body weight in mice and humans (11,12), and modulation of glucose metabolism in humans (13). Such biological effects have been attributed to CGA present in green coffee. The major CGA in green coffee are 3-, 4-, and 5-caffeoylquinic acids (3-, 4-, and 5-CQA),

3,4-, 3,5-, and 4,5-dicaffeoylquinic acids (3,4-, 3,5-, and 4,5-diCQA), 3-, 4-, and 5-feruloylquinic acids (3-, 4-, and 5-FQA), and 3-, 4-, and 5-*p*-coumaroylquinic acids (3-, 4-, and 5-*p*-CoQA). Caffeoylferuloylquinic acids are minor CGA compounds also found in green coffee, especially in *C. canephora* species. Very small amounts of CGA lactones formed by heating during primary processing may be also observed (7,14). Even though green coffee has been used for production of nutraceuticals (12), data on the pharmacokinetic profile and bioavailability of CGA from this food matrix are still inexistent. Moreover, although we have recently reported the pharmacokinetic profile of the main CGA compounds in humans after roasted coffee consumption (15), the bioavailability of CGA compounds from a food matrix has not been reported in animals or humans so far. Therefore, the objective of this study was to evaluate the pharmacokinetic profile of CGA compounds and metabolites in human plasma and urine after the acute consumption of a decaffeinated green coffee extract and to estimate the apparent bioavailability of CGA in this food matrix.

## Subjects and Methods

**Subjects.** We recruited nonsmoking male ( $n = 5$ ) and female ( $n = 5$ ) participants (22–55 y old) by word of mouth among students and faculty at the Universidade Federal do Rio de Janeiro. They were healthy as judged by a medical questionnaire, with normal blood values for hemoglobin and hematocrit, and were not taking any medication or nutritional supplements. The study protocol was approved by the Ethical Committee of Clementino Fraga Filho University Hospital at Universidade

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<sup>5</sup> Abbreviations used: AUC, area under the curve; CFQA, caffeoylferuloylquinic acid; CGA, chlorogenic acid;  $C_{max}$ , maximum plasma concentration; CQA, caffeoylquinic acid; diCQA, dicaffeoylquinic acid; FQA, feruloylquinic acid; *p*-CoQA, *p*-coumaroylquinic acid;  $T_{max}$ , time corresponding to maximum plasma concentration.

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Federal do Rio de Janeiro and fully explained to the subjects who gave their written informed consent prior to participation.

**Green coffee extract.** A hydroalcoholic decaffeinated green coffee extract produced from *C. canephora* cv. Pierre beans was spray-dried and encapsulated (0.2 g in each capsule; SVETOL).

**Study design and sample collection.** Subjects were instructed to avoid consumption of phenolic-containing foods for the 48 h prior to the study. They were asked to eat only animal foods, refined cereal foods, and artificial beverages. On the day of the study, after 10–12 h overnight fasting, an i.v. catheter was inserted into the antecubital vein and a baseline heparinized blood sample was obtained. Two capsules of green coffee extract were offered to each subject and sequential blood draws were obtained 0.5, 1, 2, 3, 4, 5, 6, 7, and 8 h after the capsule consumption. Blood samples were collected into heparin-containing tubes. Baseline blood aliquots were used to determine hematocrit and hemoglobin levels by standard methods. Plasma samples were obtained by centrifugation of the blood samples immediately after being drawn. Urine samples were also collected at baseline interval (minus 2–0 h) and at intervals of 0–2 h, 2–4 h, 4–6 h, and 6–8 h after coffee consumption into appropriate plastic containers. Total urine volume was measured for each collection period. Plasma and urine aliquots for determination of CGA were acidified with HCl and kept frozen in liquid nitrogen until analyses. Urine aliquots for determination of creatinine were acidified with HCl and kept at  $-20^{\circ}\text{C}$  until analyses. Every hour, starting 1 h after green coffee extract consumption, subjects ate a CGA-free snack composed of white bread (25 g) with cream cheese (15 g) and 100 mL of a saline solution containing 0.21 g of NaCl, 2.28 g of glucose, 0.22 g of potassium citrate monohydrate, and 0.1 g of sodium citrate dihydrate, until the end of blood draws.

**Blood hemoglobin, hematocrit, and urinary creatinine.** Hematocrit was determined by conventional capillary centrifugation. Blood hemoglobin was measured by the cyanomethemoglobin method, using a commercial kit (BioClin). Urinary creatinine was measured by the Jaffe reaction, as previously described (16).

**Chromatographic analyses.** Analyses of CGA (including CGA lactones and caffeoyltryptophan) in the green coffee extract, plasma, and urine were performed by HPLC and LC-Diode Array Detector-MS gradient systems as described in detail by Farah et al. (14,17) and Monteiro et al. (15). The detection limit for 5-CQA (4-fold baseline noise) under the conditions used in this study was 0.01 mg/L. Urinary CGA and phenolic acids excretions were expressed relative to that of creatinine (15). Molar ratios of specific CGA compounds were calculated in green coffee extract as ratios of total amounts and, in plasma, as ratios of the corresponding area under the curve (AUC).

**Pharmacokinetics and apparent bioavailability calculations.** For each subject, plasma concentrations of cinnamic acids and of individual and total CQA, diCQA, and CGA compounds were plotted over 8 h after consumption of the green coffee extract. From this plotting, the following plasma pharmacokinetic parameters were calculated: AUC ( $\mu\text{mol}\cdot\text{h}\cdot\text{L}^{-1}$ ) of total and individual components, using the trapezoidal approach (GraphPad Prism software, version 4.0); maximum plasma concentration ( $C_{\text{max}}$ ) ( $\mu\text{mol}/\text{L}$ ) and time corresponding to  $C_{\text{max}}$  ( $T_{\text{max}}$ ) (h) of total and individual components. Molar ratios between AUC of specific components were also calculated.

For calculation of the apparent bioavailability of green coffee extract CGA based on plasma results, the body surface area was initially estimated as follows:

$$\text{Surface Area} = \frac{(\text{Weight}^{0.425} \times \text{Height}^{0.725}) \times 71.84}{10000},$$

where the subject's weight is given in kg and the height is given in cm. Surface area was used to obtain the corresponding red cell mass for the respective gender and to estimate the total blood volume as described by Frenkel et al. (18). Plasma volume was estimated as follows:

$$\text{Plasma Volume (mL)} = \text{Blood Volume (mL)} \times \left(1 - \frac{\text{Hematocrit}}{100}\right).$$

Finally, the plasma amount of CGA and cinnamic acids for the 8 h after green coffee consumption was calculated by multiplying the AUC of total CGA and cinnamic acids by the total plasma volume. The apparent bioavailability for total CGA was estimated as follows:

$$\text{Apparent Bioavailability \%} = \frac{\text{Plasma CGA } (\mu\text{mol}) + \text{Cinnamic acids } (\mu\text{mol})}{\text{CGA Consumed } (\mu\text{mol})} \times 100.$$

Urinary recovery calculations were made considering the total number of equivalent moieties of cinnamic and quinic acids consumed in the green coffee extract and the total number of phenolic acid moieties recovered in urine, as a percentage.

**Statistical analyses.** Results are means with corresponding SD. Associations between plasma AUC or  $C_{\text{max}}$  and urinary excretion of specific compounds were tested using Spearman correlations (GraphPad Prism). Differences were considered significant at  $P \leq 0.05$ .

## Results

**Green coffee extract.** Nine major and 14 minor CGA compounds were identified in the decaffeinated green coffee extract offered to the subjects. The 2 capsules (0.4 g) consumed on the test day contained 170 mg (451  $\mu\text{mol}$ ) of CGA (including lactones and caffeoyltryptophan) (Table 1). CQA represented 71.2% of CGA in the capsules, with 5-CQA, 4-CQA, and 3-CQA contributing to 26.6, 21.6, and 23% (wt:wt), respectively. DiCQA and FQA represented 9.6 and 13.2% of CGA compounds, respectively.

**Subject characterization.** Subjects were healthy and had normal blood biochemistry and BMI (Table 2).

**Plasma samples.** At baseline, 9 of 10 subjects had small amounts of 3-CQA, 4-CQA, 5-CQA, 3,4-diCQA, 3,5-diCQA, 4,5-diCQA,

**TABLE 1** Contents of the main CGA in 2 capsules of the decaffeinated green coffee extract<sup>1</sup>

Compound	Dose, <sup>2</sup> $\mu\text{mol}/0.4 \text{ g}$
3-CQA	103.3 $\pm$ 0.14
4-CQA	97.4 $\pm$ 1.20
5-CQA	119.8 $\pm$ 0.23
3,4-diCQA	16.8 $\pm$ 0.01
3,5-diCQA	10.2 $\pm$ 0.06
4,5-diCQA	16.2 $\pm$ 0.32
3-FQA	20.7 $\pm$ 0.69
4-FQA	16.4 $\pm$ 0.30
5-FQA	22.1 $\pm$ 0.02
3- <i>p</i> -CoQA	1.1 $\pm$ 0.28
5- <i>p</i> -CoQA	1.7 $\pm$ 0.01
Diferuloylquinic acid (1 isomer)	1.3 $\pm$ 0.01
Caffeoylferuloylquinic acids (6 isomers)	11.8 $\pm$ 1.33
Caffeoyltryptophan <sup>3</sup>	10.5 $\pm$ 0.15
4-Caffeoylquinide <sup>3</sup>	0.3 $\pm$ 0.01
3-Feruloylquinide <sup>3</sup>	0.3 $\pm$ 0.05
4-Feruloylquinide <sup>3</sup>	0.1 $\pm$ 0.03
3,4-Dicaffeoylquinide <sup>3</sup>	0.03 $\pm$ 0.004

<sup>1</sup> Values are means  $\pm$  SD,  $n = 3$ .

<sup>2</sup> Portion = 0.4 g = 2 capsules.

<sup>3</sup> Caffeic acid derivatives computed as CGA.

**TABLE 2** Characteristics of the subjects participating in the study<sup>1</sup>

Subjects	Age	Weight	Height	BMI	Hematocrit	Blood hemoglobin
	<i>y</i>	<i>kg</i>	<i>m</i>	<i>kg/m<sup>2</sup></i>		<i>g/L</i>
Female	34.2 ± 15.3	54.2 ± 6.3	1.58 ± 0.02	22.0 ± 2.2	0.40 ± 0.03	129.4 ± 8.7
Male	28.0 ± 4.4	73.8 ± 8.7	1.73 ± 0.1	25.0 ± 1.5	0.44 ± 0.04	145.0 ± 8.7

<sup>1</sup> Values are means ± SD, *n* = 5.

and caffeic acid in plasma. Ferulic acid was identified in 6 subjects, isoferulic acid in 5 subjects, and *p*-coumaric acid in 3 subjects. After green coffee extract consumption, 3-CQA, 4-CQA, 5-CQA, 3,4-diCQA, 3,5-diCQA, and 4,5-diCQA were identified in the plasma of all subjects. Together, these compounds comprised ~82% of the total CGA in the green coffee extract. No FQA was detected in plasma of any subjects before or after extract consumption. Caffeic, ferulic, isoferulic, and *p*-coumaric acids, which were not detected in the encapsulated extract, were present in the plasma of different subjects after green coffee extract consumption, contributing 6.6, 6.2, 6.1, and 1.4% of total phenolics in plasma, respectively.

A large inter-individual variation was observed in the pharmacokinetic profile of all CGA compounds and cinnamic acids in plasma (Fig. 1). However, the plasma kinetic profiles of the individual isomers within the subclasses of CGA followed a similar pattern for most subjects. Therefore, for clarity, only the plasma pharmacokinetic profiles of total CQA, total diCQA, total CGA, and total cinnamic acids at baseline and for 8 h after coffee consumption are shown.

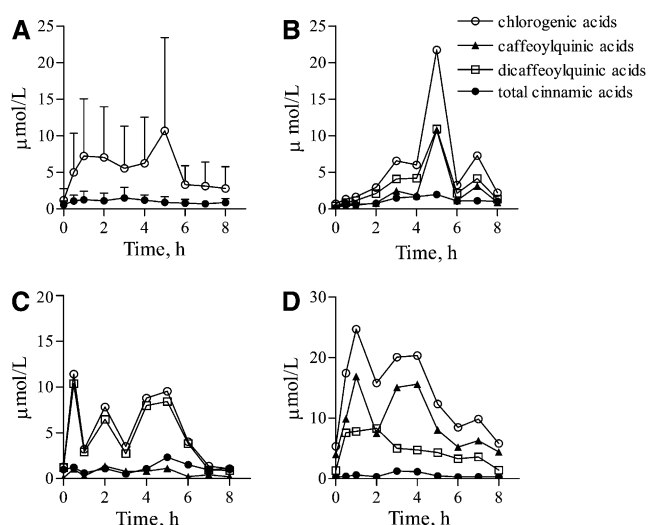
CGA  $C_{max}$  and  $T_{max}$  varied substantially among the subjects. The  $C_{max}$  of total CQA varied from 0.6 to 16.9  $\mu\text{mol/L}$  and that of total diCQA varied from 0.3 to 22.8  $\mu\text{mol/L}$ , whereas the  $C_{max}$  of total CGA varied from 1.2 to 39.7  $\mu\text{mol/L}$ .  $T_{max}$  for total CQA, total diCQA, and total CGA varied from 0.5 to 8 h (Table 3). The  $T_{max}$  of the isomers did not differ within the CQA and diCQA subclasses.

Of the individual compounds, 5-CQA was the major CGA identified in the plasma of all subjects at all time points after

green coffee extract consumption, as indicated by both its  $C_{max}$  and AUC. Based on AUC, 5-CQA, 4-CQA, and 3-CQA comprised 31.3, 7.5, and 5.2% of the total phenolic compounds in plasma.

Molar ratios among CGA compounds were calculated using their content in the green coffee extract and the AUC in plasma. For CQA, the ratio of 5-CQA:4-CQA:3-CQA in the green coffee extract was 1.2:1.0:1.1, whereas their corresponding ratio in plasma was 6.0:1.4:1.0. The molar ratio of 3,5-diCQA:4,5-diCQA:3,4-diCQA in the coffee extract was 1.0:1.6:1.7, whereas in plasma is was 1.7:1.4:1.0. Moreover, when the CGA classes were compared, the diCQA:CQA molar ratio in plasma was 6.2 times that in the green coffee extract (Tables 1 and 3).

**Urine samples.** Most subjects had phenolic compounds in their urine at baseline. Trace amounts of 5-CQA were observed in 5 subjects. Sinapic, gallic, *p*-hydroxybenzoic, and dihydrocaffeic acids were the major phenolic compounds at baseline, representing ~82% of the total amount of the identified phenolic compounds. The urinary excretion of phenolic compounds increased in 9 of 10 subjects after extract consumption (Table 4). As with plasma, there was a large inter-individual variation in the urinary excretion of all compounds after green coffee extract consumption. The only intact CGA compounds identified in urine after the extract consumption were 5-CQA and 4-CQA. Not only at baseline but also after extract consumption, sinapic, gallic, *p*-hydroxybenzoic, and dihydrocaffeic acids were the major phenolic compounds, representing ~85% of the total amount of phenolic compounds identified in urine. Protocatechuic, dihydroferulic, benzoic, and hippuric acids, which have been previously identified in urine after CGA consumption, were not



**FIGURE 1** Pharmacokinetic profiles of total CGA, CQA, diCQA, and total cinnamic acids in plasma of all subjects (A) and of subjects 4 (B), 5 (C), and 8 (D) 8 h after decaffeinated green coffee extract consumption. Values in A are means ± SD, *n* = 10.

**TABLE 3** Pharmacokinetic parameters of CGA and cinnamic acids identified in plasma after decaffeinated green coffee consumption<sup>1</sup>

Compound	$C_{max}$ $\mu\text{mol/L}$	$T_{max}$ <i>h</i>	AUC $\mu\text{mol}\cdot\text{h}\cdot\text{L}^{-1}$
3-CQA	0.9 ± 1.4	4.0 ± 2.6	3.0 ± 4.5
4-CQA	1.4 ± 1.1	3.6 ± 2.2	4.3 ± 5.4
5-CQA	5.9 ± 4.2	3.3 ± 2.4	17.9 ± 15.3
3,4-diCQA	1.5 ± 1.6	2.6 ± 1.8	5.0 ± 4.9
3,5-diCQA	2.7 ± 2.7	3.2 ± 2.5	8.7 ± 8.3
4,5-diCQA	2.5 ± 3.0	3.3 ± 2.5	6.8 ± 5.7
Total CQA	8.2 ± 6.3	3.3 ± 2.4	25.2 ± 24.4
Total diCQA	6.6 ± 6.9	3.2 ± 2.5	20.4 ± 17.5
Total CGA	14.8 ± 11.7	3.1 ± 2.6	45.6 ± 37.1
Caffeic acid	1.1 ± 0.9	3.6 ± 2.1	3.8 ± 3.2
Ferulic acid	0.8 ± 0.3	2.9 ± 1.8	3.6 ± 1.5
Isoferulic acid	0.9 ± 0.2	2.9 ± 1.8	3.5 ± 1.9
<i>p</i> -Coumaric acid	0.4 ± 0.03	2.5 ± 1.8	0.8 ± 0.2

<sup>1</sup> Values are means ± SD, *n* = 10.

**TABLE 4** Total urinary excretion of CGA and metabolites in each subject after decaffeinated green coffee consumption<sup>1</sup>

Subject	Gallic acid	<i>p</i> -Hydroxy-benzoic acid	Dihydrocaffeic acid	Vanillic acid	Siringic acid	Sinapic acid	5-CQA	4-CQA	Caffeic acid	Ferulic acid	Isoferulic acid	<i>p</i> -Coumaric acid	Total phenolics
	<i>μmol</i>												
1-Baseline	9.24	4.94	2.98	2.61	0.04	1.26	Nd	Nd	Nd	0.45	0.01	0.01	21.54
0–8 h	43.47	32.58	29.97	10.38	3.85	10.50	Nd	Nd	0.11	2.24	0.92	0.13	134.15
2-Baseline	3.55	1.08	0.25	1.86	0.47	0.97	Nd	Nd	Nd	0.13	0.02	0.01	8.34
0–8 h	18.67	24.14	15.81	16.37	11.07	17.52	0.42	Nd	0.86	3.33	1.02	0.14	109.35
3-Baseline	1.86	2.56	Nd	3.31	3.52	2.22	0.02	Nd	0.01	0.23	0.07	0.03	13.81
0–8 h	9.41	27.94	56.05	24.27	16.51	13.34	0.41	Nd	0.07	4.96	2.16	0.07	155.18
4-Baseline	0.70	0.28	0.16	0.41	1.62	1.43	Nd	Nd	0.03	0.55	0.02	Nd	5.22
0–8 h	7.98	4.69	13.60	9.47	16.63	12.08	0.60	Nd	0.21	4.60	1.98	0.03	71.86
5-Baseline	10.19	1.78	5.02	9.42	13.11	20.28	Nd	Nd	Nd	1.94	0.14	Nd	61.89
0–8 h	25.45	16.65	15.09	21.20	9.35	47.67	0.46	Nd	0.40	6.43	1.87	Nd	144.56
6-Baseline	10.57	12.03	4.25	2.73	0.34	19.55	0.04	0.04	0.09	Nd	Nd	Nd	49.63
0–8 h	115.87	189.74	61.33	12.94	1.52	130.47	2.33	0.90	1.41	1.76	4.68	2.84	527.23
7-Baseline	0.25	4.12	2.13	0.19	1.36	0.98	Nd	Nd	Nd	Nd	Nd	Nd	9.03
0–8 h	14.03	26.72	19.10	5.63	15.08	19.94	4.02	0.91	3.28	1.17	2.15	0.92	116.06
8-Baseline	6.79	12.12	27.54	Nd	3.64	0.87	0.35	Nd	0.40	0.09	0.73	0.03	52.57
0–8 h	68.21	50.45	67.45	0.58	25.62	15.06	1.22	0.83	3.60	2.39	11.80	0.10	249.35
9-Baseline	1.68	1.49	3.00	3.08	1.58	5.32	0.01	Nd	0.11	0.03	0.51	Nd	14.31
0–8 h	57.69	48.45	97.73	2.26	7.94	161.54	1.67	0.86	5.35	5.80	17.00	1.81	408.92
10-Baseline	25.25	19.91	12.71	0.86	Nd	39.11	0.21	Nd	1.72	0.21	4.83	0.05	104.86
0–8 h	112.22	146.38	73.92	3.20	Nd	178.79	1.75	1.22	9.43	3.83	18.78	9.76	559.29

<sup>1</sup> Nd, Not detected (< 0.02  $\mu\text{mol}$ ).

identified in the urine of any of the subjects before or after the extract consumption.

**Apparent bioavailability of CGA.** As with plasma and urinary values, the apparent bioavailability of CGA varied considerably among subjects. The apparent bioavailability based on consumed cinnamic moieties ranged from 7.8 to 72.1% in plasma, with a mean of  $33.1 \pm 23.1\%$ . Urinary recovery of phenolic compounds based on consumed cinnamic and quinic acid moieties varied from  $-15.6$  to  $25.9\%$  in urine when expressed as an increment from baseline values and from  $7.5$  to  $58.4\%$  when expressed as absolute values, with means of  $5.5 \pm 10.6\%$  and  $25.8 \pm 19.1\%$ , respectively. There was a negative correlation between apparent bioavailability and urinary recovery of CGA ( $r = -0.76$ ;  $P = 0.01$ ). Apparent bioavailability and urinary recovery were not correlated with gender, age, weight, height, or BMI.

## Discussion

This is the first study, to our knowledge, evaluating the pharmacokinetics of CGA compounds after the consumption of a green coffee extract. Also, the apparent bioavailability of CGA compounds from a food matrix and the urinary recovery of their metabolites were estimated in humans.

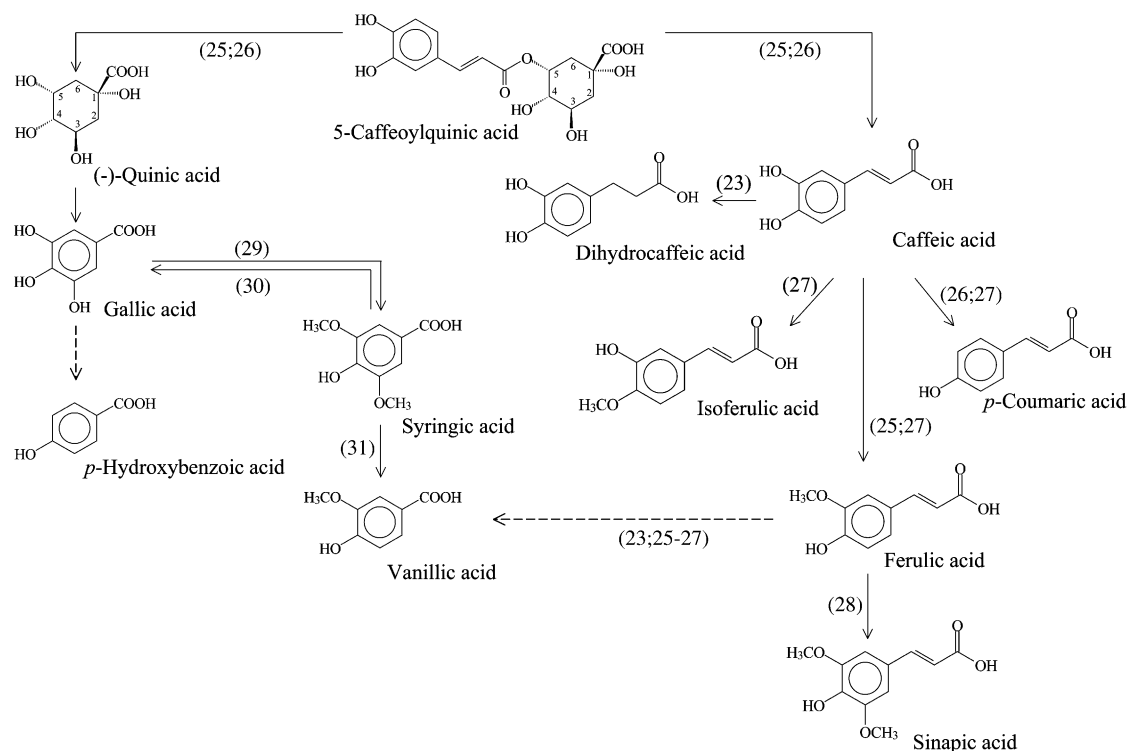
Although subjects consumed a low-phenolic diet for 2 d prior to the study and were fasting for 10–12 h when baseline samples were collected, all subjects had phenolic compounds in their baseline plasma and/or urine. This is in agreement with the fact that CGA and other phenolic compounds have been observed in saliva, gastrointestinal fluids, and urine of fasting subjects (19–21) and corroborates the hypothesis of storage and recycling of these

compounds through excretion and reabsorption suggested by Baer-Dubowska and Szaefer (22) and Farah et al. (20).

After green coffee extract consumption, 6 major CGA compounds were identified in the plasma of all subjects, as previously observed after roasted coffee consumption (15), accounting for almost 90% of phenolic compounds in plasma. Despite similar amounts of FQA and diCQA isomers in the extract, FQA was not detected in the plasma of any subject. This result is consistent with evidence of a poor absorption of FQA and/or of a rapid uptake by organs such as liver (23,24) and adipose tissue. In fact, a human hepatic cell study found that FQA uptake was favored compared with CQA and diCQA uptake (24). Another possible explanation would be demethylation of the ferulic acid moiety of the ester and conversion of FQA into CQA (Fig. 2).

Additionally, small amounts of caffeic, ferulic, isoferulic, and *p*-coumaric acids were identified in the plasma of different subjects after the green coffee extract consumption. Considering that no unesterified cinnamic acids were present in the extract, that only 4.4% of 5-CQA were hydrolyzed into caffeic acid in the analytical recovery tests, and that isoferulic acid could not be formed from CGA hydrolysis, our results suggest that most cinnamic acids in plasma originated from metabolization of CGA from the coffee extract, possibly in the intestinal lumen/mucosa and liver (28,33). Ferulic and isoferulic acids may be formed by methylation of caffeic acid (27). Conversely, caffeic acid may also derive from hydrolysis of FQA (34) (Fig. 2).

As previously observed after roasted coffee consumption (15), there was a large inter-individual variation in the pharmacokinetic profile of all CGA compounds and phenolic acids in plasma and urine after green coffee extract consumption, although the kinetic profiles of the individual CGA compounds within the subclasses were similar for each subject. This variability may be attributed to inter-individual differences in



**FIGURE 2** Proposed simplified scheme of 5-CQA metabolism, as representative of CQA and diCQA classes, based on results from the present study. Support references for proposed metabolic routes are shown. The authors adopted the IUPAC numbering system (32) for CGA.

digestive transit time, preferential site of absorption, and metabolism of cinnamates as reported in the literature for other phenolic compounds (15,20,35). According to Manach et al. (35), important inter-individual differences in the capacity to metabolize phenolic compounds originate from differences in the activity of the cytochrome P450 and carrier systems that may be influenced by genetic polymorphisms.

In contrast to the pharmacokinetic profile of flavonoids such as catechin (36) and isoflavones (37), various plasma concentration peaks were observed from 0.5 up to 8 h after the green coffee extract consumption (Fig. 1), suggesting a complex and dynamic process of absorption and metabolism. This irregular pharmacokinetic pattern makes the calculation of the CGA compounds half-life difficult. Longer duration studies would be necessary for this measurement.

Considering that liquid food may take up to 1 h to reach the small intestine (38,39) and that preliminary tests showed that the extract capsules take <15 min to solubilize in a simulated gastric condition (data not shown), the initial plasma concentration peaks confirm an early absorption of CGA in the stomach and jejunum followed by absorption along the small intestine (15,40–43). The later plasma peaks (>5 h) in some subjects may indicate absorption through the large intestine (44) and/or recycling through digestive fluids (20).

In the present study, the higher diCQA:CQA molar ratio in plasma compared with the green coffee extract supports previous evidence of differential mechanisms of absorption and/or metabolism for CQA and diCQA (15), with favored tissue uptake of CQA compared with diCQA and/or favored diCQA absorption that could be related to the higher lipophilicity of diCQA compared with CQA, possibly favoring diCQA diffusion through the intestinal mucosa cells. Acyl migration from CQA forming diCQA during analysis is not

possible, because adding green coffee extract to CGA-free plasma in a recovery test yielded similar CQA and diCQA isomer profiles in both the spiked plasma and the green coffee extract alone.

The higher mean molar ratio of 5-CQA:4-CQA and 3-CQA in plasma than in green coffee extract supports preferential absorption of 5-CQA compared with 4-CQA and 3-CQA isomers or of rapid metabolism and/or uptake of 3-CQA and 4-CQA by organs such as liver (24,15) and adipose tissue. The higher molar ratios of 3,4-diCQA:4,5-diCQA and 3,5-diCQA in plasma than in the green coffee extract also suggests mechanisms favoring plasma levels of 3,5-diCQA as described for 5-CQA.

Although the amount of CGA in decaffeinated green coffee in the present study was ~13% of that in decaffeinated roasted coffee in our previous study (15), the plasma total CGA AUC in the present study at 4 h after treatment, the duration of our previous study, was 1.4 times the AUC in that study. Although the 2 studies cannot be compared directly due to various differences, including subjects, the influence of the matrix in which CGA is consumed deserves investigation.

The 4 major urinary phenolic compounds excreted after green coffee consumption (sinapic, gallic, *p*-hydroxybenzoic, and dihydrocaffeic acids) are probably preferential metabolic products of the cinnamates identified in plasma. Vanillic and syringic acids are probably derived from a secondary metabolic pathway (Fig. 2). Protocatechuic, dihydroferulic, benzoic, and hippuric acids, which were identified in studies evaluating the urinary products of cinnamates from different food sources (25,27), were not identified in our study although they are consistent with the metabolic pathway presented (Fig. 2).

This is the first study to our knowledge investigating the bioavailability of CGA as a family of compounds and also the



first evaluating the bioavailability of CGA using plasma AUC values. The apparent bioavailability of CGA from the green coffee extract varied from 7.8 to 72.1% among the subjects, with a mean of  $33 \pm 23\%$ . Possibly, a higher bioavailability would have been obtained in a longer study, because in 3 of the subjects, plasma CGA concentrations were still high 8 h after green coffee extract consumption. The mean apparent bioavailability of 5-CQA, the only CGA evaluated in previous bioavailability studies, was  $33 \pm 27\%$ . This result is in contrast with the very low absorption of 5-CQA estimated in rat and human studies (as low as 0.1%) (42,45,46). On the other hand, our results are in agreement with the  $33 \pm 17\%$  absorption estimated by Olthof et al. (47), who analyzed ileostomy fluids from colostomized subjects for 24 h after the consumption of 2.8 mmol of 5-CQA. The results by Olthof et al. were considered an overestimation (48), because the amount of 5-CQA possibly lost during digestion was not taken into account and because previous studies (21,48,49) had been unable to identify intact CGA in plasma. However, the results obtained by Olthof et al. (47) are actually very close to those in the present study and their estimate was probably correct, because only a small portion of 5-CQA seems to be hydrolyzed in the digestive tract and only a minor fraction is absorbed in the form of caffeic acid (15, present study).

The low recovery of total phenolic compounds in urine when values were expressed as increments from baseline was downgraded by the high amount of phenolic compounds in the baseline urine period of 1 subject (subject 8; Table 4), producing a negative recovery of  $-15.6\%$ . Excluding this subject would bring the recovery, corrected to baseline, to  $7.8 \pm 8.1\%$ . This case, along with the irregular pharmacokinetic patterns observed in plasma, indicates that increments over baseline values may not be the most appropriate way to evaluate total urinary excretion of CGA and metabolites in response to acute ingestion. On the other hand, the use of absolute urinary values (not corrected by baseline) may overestimate excretion.

Low recoveries in urine were also found in previous human studies evaluating the metabolism of phenolic compounds, supporting evidence that urine may not be the preferential excretion route for intact CGA and metabolites (19,20,26,50). On the other hand, it should be noted that many urinary metabolites derived from CGA were probably not identified in this study, because the 12 urinary compounds identified represented  $\sim 50\%$  of the total chromatogram peak area. Olthof et al. (25) identified 23 phenolic acids by GC in human urine after consumption of 5-CQA.

The negative correlation between apparent bioavailability of CGA and corrected urinary recovery ( $r = -0.76$ ;  $P = 0.01$ ) may be explained by inter-individual differences in metabolic rates and/or preferential excretion routes.

In conclusion, the present study confirms that CQA and diCQA, which are major CGA compounds in coffee, are absorbed in the human body, being differentially absorbed and/or metabolized throughout the whole gastrointestinal tract. This study also supports evidence that urine is not a major excretion pathway of intact CGA compounds and their metabolites and identifies sinapic, gallic, *p*-hydroxybenzoic, and dihydrocaffeic acids as major urinary metabolites of CGA in humans. In addition, this study shows that the main CGA compounds present in the green coffee matrix are highly bioavailable in humans. A large inter-individual variation clearly exists in CGA absorption, metabolism, and kinetics in humans and requires further investigation into differences in genetic polymorphisms.

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