Effect of coffee drinking on platelets: inhibition of aggregation and phenols incorporation

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Coffee is among the most widely consumed beverages in the world. The relationship between coffee drinking and CVD has been extensively studied. Recently several studies indicate a moderate coffee consumption can be protective against atherosclerosis and CVD(9). Moreover, ex vivo platelet aggregation is related to CVD mortality(10).

Epidemiological studies indicate a J-shaped relationship linking coffee consumption and cardiovascular risk, suggesting that moderate coffee consumption can be beneficial. Platelet aggregation is of critical importance in thrombotic events, and platelets play a major role in the aetiology of several CVD. The aim of this study was to evaluate the effect of coffee drinking on platelet aggregation ex vivo, using caffeine as control. A crossover study was performed on ten healthy subjects. In two different sessions, subjects drank 200 ml coffee, containing 180 mg caffeine, or a capsule of caffeine (180 mg) with 200 ml water. Platelets were separated from plasma at baseline and 30 and 60 min after coffee drinking.

Platelet aggregation was induced with three different agonists: collagen, arachidonic acid and ADP. Coffee drinking inhibited collagen (P<0.05 from baseline at time 30 min) and arachidonic acid (P<0.05 from baseline at time 60 min) induced platelet aggregation. Caffeine intake did not affect platelet aggregation induced by the three agonists. Coffee consumption induced a significant increase of platelet phenolic acids (likely present as glucuronate and sulphate derivatives), caffeic acid, the principal phenolic acid in coffee, raising from 0·3 (SEM 0·1) to 2·4 (SEM 0·6) ng/ml (P<0.01). Caffeine was not detectable in platelets. Coffee drinking decreases platelet aggregation, and induces a significant increase in phenolic acid platelet concentration. The antiplatelet effect of coffee is independent from caffeine and could be a result of the interaction of coffee phenolic acids with the intracellular signalling network leading to platelet aggregation.

Cardiovascular diseases: Coffee: Human: Phenolic acids: Platelet aggregation

Abbreviations: PRP, platelet-rich plasma; TxA2, thromboxane A2; TxB2, thromboxane B2.

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Subjects and methods

Subjects and study design
Ten volunteers (five males and five females), aged 24–35 years, participated in a cross-over study. Subjects, free from known diseases and moderate coffee drinkers (two to four cups/d), were instructed to refrain from consuming coffee and phenol-rich foods and beverages for the 2 d prior to the experiments. Subjects were also asked to abstain from any platelet inhibitor and dietary supplements for ≥10 d prior to the experiments. Subjects reported to the laboratory on two separate occasions, 2–4 weeks apart, after an overnight fast (10–12 h). A venous blood sample was taken at time 0. Immediately after the first blood collection subjects were provided with a cup of freshly prepared American coffee (200 ml) or a capsule containing 180 mg caffeine with 200 ml water. This dose was equivalent to the caffeine contained in the cup of coffee. Further blood collections were made 30 and 60 min after coffee or caffeine consumption. Venous blood samples were collected into evacuated tubes containing sodium citrate. Platelet-rich plasma (PRP) was separated by low-speed centrifugation (180 g, 15 min) at room temperature.

All subjects gave informed consent before entering the study, and all procedures were approved by the Ethical Committee of the National Institute for Food and Nutrition Research.

Coffee brew preparation and analyses
Coffee brew was prepared by using a commercial automatic brewing machine, using 60 g roasted and ground coffee per litre of water.

Caffeine was measured in coffee by HPLC as previously described by Blanchard et al. (24), and the amount found after three replicate analyses (180 (SEM 9) mg/cup) was used for caffeine capsule formulation.

Measurement of plasma caffeine concentrations
Caffeine was detected in plasma and platelets by HPLC as previously described (24).

Measurement of platelet aggregation
Platelet aggregation was measured ex vivo on PRP using an aggregometer, with constant stirring at 1000 rpm and at 37°C. PRP was separated from blood after 20 min centrifugation at 3000 g. The pellet was resuspended in water, sonicated, acidified to pH 3 with 1 M-HCl and stored at −80°C. The presence of phenolic acid into platelets was measured in samples untreated (free phenolic acids) and in samples subjected to alkaline hydrolysis (total phenolic acids) (6).

No treatment. Platelet suspension (0.5 ml) at pH 3 was added with 50 ng o-coumaric acid as internal standard and 300 mg NaCl, then extracted three times with ethyl acetate (× 4 volumes) by vortexing for 5 min. After each extraction, samples were centrifuged (3000 g, 10 min, room temperature) and the supernatants collected. The organic phase was dried under nitrogen flow. The residue was dissolved in 0.5 ml water, vortexed for 5 min, then the pH was brought to pH 7–8 with 0.1 M-NaOH and sample passed through 1 ml Supelclean LC-SAX tube preconditioned with 1 ml absolute methanol and 2 ml water. The tube was then washed with 1 ml water. Phenolic acids elution was obtained with 1 ml buffer containing 1 M-acetic acid–methanol (90:10). The eluant was immediately brought to pH 3 with 6 μl 4 M-NaOH, filtered and an aliquot (100 μl) was injected into the HPLC system.

Alkaline hydrolysis treatment. Platelet suspension (0.5 ml) containing 50 ng o-coumaric acid as internal standard had the following added in order: 55 μl H2O, 20 μl 0.5 M-EDTA, 200 μl 5% ascorbic acid and 225 μl 8 M-NaOH, and was then incubated for 30 min at 30°C. At the end of incubation, the pH was brought to 3 with 8 M-HCl and 600 mg NaCl added. Samples were extracted three times with ethyl acetate (× 4 volumes) as reported earlier. The residue was dissolved in 0.5 ml water, vortexed for 5 min, then processed for solid-phase extraction as described earlier.

The overall procedures allow an almost complete recovery of the phenolic acids under study, as found by recovery experiments performed adding known amounts of pure compounds to platelet suspension samples. Recovery was 98.6 ± 0.4 (SD 9.9), 95.4 ± 0.4 (SD 13.6), 87.6 ± 0.4 (SD 4.4), 100.7 ± 0.7 (SD 5.7) (n = 5) for caffeic, p-coumaric, ferulic and isoferulic acids, respectively.

Samples were analysed by HPLC-electrochemical detector as previously described (25) with minor modifications concerning the elution gradient.

Measurement of platelet thromboxane B2 formation
Thromboxane B2 (TxB2) production was measured in platelets separated at time 0, 30 and 60 min after coffee drinking, on a sub-group of six subjects. PRP was incubated with collagen (3 μg/ml). The platelet activation was stopped after 10 min and thromboxane A2 (TxA2) production was determined using TxB2 (the stable breakdown product of TxA2) ELISA kits (R&D System).

Measurement of platelet phenolic acid concentrations
Phenolic acids in biological samples are routinely detected in our laboratory by HPLC-electrochemical detector (26). For the study of incorporation of phenolics in platelets, PRP was added with acid citrate dextrose and further centrifuged for 7 min at 780 g. The pellet was washed once with PBS containing acid citrate dextrose, centrifuged and the final pellet was resuspended in water, sonicated, acidified to pH 3 with 1 M-HCl and stored at −80°C. Caffeine was detected in plasma and platelets by HPLC as previously described (24) and the number of platelets in the PRP was standardized as percentage of maximal aggregation.

aggregation was induced by collagen (3 μg/ml) and measured as described earlier.
Protein was measured by the method of Lowry et al.\(^26\), using bovine serum albumin as standard. The concentration of phenolic acids is expressed as ng/mg protein.

Free forms of phenolic acids in platelets were detected only at very low amounts (traces), therefore all the results presented refer to the total (free + bound) phenolic acid content.

**Statistics**

All data are presented as mean values with their standard errors. Statistical analysis was carried out using repeated-measures ANOVA, followed by Tukey’s test for multiple comparisons. Analyses were performed with KaleidaGraph software version 3.6 (Synergy Software, Reading, PA, USA). \(P<0.05\) was considered statistically significant.

**Results**

**Effect of coffee and caffeine consumption on ex vivo platelet aggregation**

Coffee drinking significantly inhibited platelet aggregation induced by arachidonic acid and collagen (Fig. 1), while no statistically significant differences were observed when ADP was used as agonist (Fig. 1).

Coffee drinking also inhibited collagen-induced TxB2 formation (Fig. 2).

Caffeine intake did not affect platelet aggregation induced by the three different agonists (Fig. 1).

**Effect of coffee and caffeine supplementation on plasma caffeine concentrations**

As expected, a statistically significant increase in plasma caffeine concentrations was detectable after both coffee and caffeine consumption (Fig. 3). However, the time course of caffeine increase was different in the two sessions, being slower after caffeine administration than after coffee consumption. Caffeine plasma concentrations measured after 30 min of caffeine intake (15.2 (SEM 7.8) \(\mu\)M) were, in fact, statistically significantly different from those observed 30 min after coffee drinking (20.6 (SEM 8.1) \(\mu\)M).

**Effect of coffee consumption on platelet phenolic acid concentration**

Coffee drinking induced a significant increase in phenolic acid platelet concentration (Table 1). All phenolic acids under study increased significantly, but they showed different incorporation kinetics. In fact, the maximum incorporation peak was at 30 min for \(p\)-coumaric and ferulic acids, and at 60 min for caffeic and isofericulic acids. The maximum concentration reached after coffee drinking was: 2.4 ng/mg protein for caffeic acid, 1.9 ng/mg protein for ferulic acid, 1.8 ng/mg protein for \(p\)-coumaric acid and 0.9 ng/mg protein for isofericulic acid.

The most relevant increases were observed for caffeic acid (8-fold at peak time with respect to time 0) and isofericulic acid (4.5-fold at peak time with respect to time 0). \(p\)-Coumaric and ferulic acids increased more slightly, the increment of their concentration at peak time being about 60 and 90\%, respectively.

Phenolic acids were present in platelets almost exclusively as conjugated forms, free phenolic acids being present only as traces.

**Effect of phenolic acids on in vitro platelet aggregation**

To evaluate the capacity of phenolic acids in inhibiting platelet aggregation, a mix of phenolic acids was added to PRP at the concentrations measured in platelets after coffee drinking. After 30 min of incubation at 37°C aggregation was induced by collagen (3 \(\mu\)g/ml). As shown in Fig. 4, the mix of phenolic acids was able to significantly inhibit platelet aggregation and collagen-induced TxB2 formation.

**Discussion**

The first aim of the present study was to determine the effects of coffee on ex vivo platelet aggregation induced by three different agonists (collagen, arachidonic acid and ADP), using caffeine as control. While coffee drinking inhibited collagen and arachidonic acid-induced platelet aggregation,
caffeine intake did not affect platelet aggregation induced by any of the agonists (Fig. 1). Coffee and caffeine intake induced a similar increase in plasma caffeine concentrations, which reached the same peak concentration 60 min after both supplementations (Fig. 3). Even if the kinetic of absorption was different, we can exclude some effect of caffeine on platelet aggregation. In fact, 60 min after caffeine intake (when plasma caffeine concentration was equal to the concentration reached after coffee drinking), no effect of caffeine on platelet aggregation was observed. Similar results were obtained by Rein et al. (21), who did not find any effect on ADP-induced platelet aggregation after acute intake of caffeine (17 mg). On the contrary, Varani et al. (22) observed that chronic caffeine intake could affect ADP-induced platelet aggregation, but only when the intake lasted longer than 1 week and was higher than 400 mg/d. According to the present results, Varani et al. (22) did not observe any effect of caffeine on platelet aggregation after acute intake.

The capacity of coffee to inhibit platelet aggregation has already been demonstrated, but only in an animal model. Bydlowski et al. (19) demonstrated that an intravenous administration of coffee extracts inhibited ex vivo arachidonic acid and ADP-induced platelet aggregation in rabbits. In man, Polagruzo et al. (20) did not find significant modification of platelet function 2 and 6 h after coffee drinking. However, their results are scarcely comparable with the present data, as platelet function was estimated using a completely different methodology (the measure of the adrenalin/collagen or ADP/collagen induced clotting time) and time-points.

Apart from coffee, several papers report an effect of specific food items on platelet aggregation in vivo in man, both in acute and chronic studies. An antiplatelet effect has been demonstrated for fruit juice (20), red wine (18), onion (27), tea (17, 28) and tomato (29) (for a review, see Nardini et al. (30)). These findings support the role of the diet (as a whole and/or its single components) in the modulation of the platelet network and confirm the important role of diet in the prevention of CVD.

Looking for the molecular mechanism through which coffee can affect platelet aggregation, we measured the capacity of coffee drinking to affect collagen-induced platelet TxA2 formation. The present results demonstrated that coffee inhibited both collagen and arachidonic acid-induced platelet aggregation, but it had no effect on ADP-induced platelet aggregation. Differently from ADP, the platelet activation pathways induced by arachidonic acid and collagen are both mediated by cyclo-oxygenase, through the formation of TxA2, which further stimulates aggregation. As shown in Fig. 2, platelets collected 60 min after coffee drinking and activated with collagen released significantly less TxB2 than those collected before drinking. The present result is in agreement with literature data reporting that coffee inhibits in vitro TxB2 generation in rabbit platelets (19). Thus, coffee affects the signal transduction pathway, which leads to aggregation through the inhibition of one of the enzyme activities that lies upstream of the formation of TxB2 in the signalling cascade.

It is well known that phenolic compounds can inhibit in vitro and ex vivo platelet aggregation (11, 23). Their anti-aggregating

![Fig. 2. Thromboxane B2 (TXB2) production in platelets separated at time 0, 30 and 60 min after coffee consumption (n = 6). Platelets were activated using collagen (3 μg/ml). Values are means with their standard errors depicted by vertical bars. Mean values were significantly different from those of time 0 (repeated-measures ANOVA followed by Tukey’s test): *P < 0.05.](https://www.cambridge.org/core/terms)

![Fig. 3. Caffeine concentration at baseline, 30 and 60 min after coffee (••••) and caffeine (−−) consumption (n = 10). Values are means with their standard errors depicted by vertical bars. Mean values were significantly different from those of time 0 (repeated-measures ANOVA followed by Tukey’s test): **P < 0.01.](https://www.cambridge.org/core/terms)
activity has been attributed to several mechanisms, among them the capacity to inhibit TXA2 generation \(^{(31,32)}\). Coffee contains several phenolic compounds, which are absorbed and rapidly metabolized in man \(^{(6,33)}\). In a previous study we demonstrated that coffee drinking induces a significant increase in plasma caffeic acid, which reaches micromolar concentrations 1 h after coffee intake \(^{(6)}\).

The incorporation of polyphenols into circulating cells after the intake of a polyphenol-rich food has not been demonstrated yet. In the present study, we demonstrate for the first time that metabolites of phenolic acids are incorporated in human platelets after a single oral dose of coffee.

It is important to underline that the evaluation of the actual bioavailability of polyphenols and their subsequent interaction with cells and tissues is a prerequisite for the real understanding of their physiological effect. In fact, many of the effects claimed for these compounds are exerted inside the cells \(^{(34,35)}\), but, unfortunately, no direct evidence exists about the "physiological" presence of these compounds in cells in \textit{in vivo} studies. Information on polyphenol tissue distribution is still very scarce, and available data concern only animal models. Animal studies, mostly using radio-labelled compounds, have shown that phenolic compounds are able to penetrate tissues, particularly those in which they are metabolized (intestine and liver \(^{(36-38)}\); for a review, see Manach \textit{et al.} \(^{(39)}\). In tissues phenolic compounds occur prevalently in conjugated forms, indicating that the ingested phenolics are extensively and rapidly metabolized \(^{(40,41)}\).

To our knowledge, only three studies report data on the occurrence of polyphenols in human cells \textit{in vivo} \(^{(42-44)}\), and, among these, only one study evaluates their presence in human blood cells. In the present study, apigenin metabolites were found in human erythrocytes, but no increase in intracellular concentration was observed after apigenin-rich parsley consumption \(^{(44)}\).

With the exception of isoferulic acid, all the phenolic acids measured in platelets after coffee drinking were present in coffee brew, even if in bound forms as esters of quinic acid (data not shown). In coffee, the most abundant of these esters is chlorogenic acid, whose phenolic moiety is represented by caffeic acid \(^{(45)}\). Isoferulic acid is a metabolite of caffeic acid and is not present, as such, in coffee.

Its increase after coffee drinking indicates that caffeic acid is extensively metabolized, but we cannot speculate if its metabolism takes place before or after the incorporation into platelets. All the phenolic acids measured into platelets were present in conjugated forms, indicating a wide and rapid metabolic process. Yet, preliminary evidence obtained with an enzymatic hydrolytic procedure seems to indicate that most of the phenolic acids are present in platelets as glucuronates and sulphonates (data not shown).

It must be evidenced that phenolic acid concentration inside platelets is sufficiently high to explain coffee anti-aggregative effects. The sum of platelet phenolic acid concentration, in fact, reaches about 6 ng/mg protein at 30 and 60 min. A raw calculation (based on a mean platelet volume and protein content) lets us estimate that caffeic acid alone can reach an intracellular concentration of about 2 mM. To test if phenolic acids, at the concentration observed in platelets after coffee drinking, were able to affect platelet aggregation, we ran \textit{in vitro} experiments, using a mix of the different phenolic acids at the concentrations measured \textit{in vivo}. As shown in Fig. 4, the mix is capable of significantly inhibiting collagen-induced platelet aggregation and collagen-induced TXB2 formation. It is well known that phenolic acids are efficient antioxidants \(^{(45)}\) and caffeic acid is a good inhibitor of lipoxygenase \(^{(46)}\), cyclo-oxygenase \(^{(47)}\) and kinase \(^{(48)}\). All these activities could be critical for the anti-aggregative capacity of phenolic acids.

We are aware that phenolic acids incorporated into platelets are not in their free form, but in conjugated forms, likely glucuronates and sulphates. Since at the moment the appropriate standard compounds are not commercially available, we cannot determine if these forms are more or less active than their respective free forms. Scarc information is available concerning the potential activity of the metabolites of polyphenols. Conjugated forms of quercetin, one of the most abundant flavonoids in the human diet, retain antioxidant activity, although to a lesser extent in respect to quercetin \(^{(49)}\). The antioxidant activity of ferulic acid glucuronide is stronger than that exhibited by ferulic acid \(^{(50)}\). However, some polyphenol metabolites (quercetin tetrasulphate and genistein sulphates) have been reported to have lower antiplatelet activities than their parent molecules \(^{(52-53)}\). The possibility exists that the free forms penetrate into the cells, where a subsequent conjugation reaction takes place. This assumption is plausible since studies have shown the presence of UDP-glucuronyltransferase \(^{(52)}\) and phenolsulphotransferase \(^{(53)}\) in platelets and in \textit{in vitro} experiment demonstrates that metabolites of radio-labelled-hydroxytyrosol are rapidly formed after the incubation of hydroxytyrosol in whole human blood \(^{(54)}\).

Data presented here indicate that drinking 200 ml coffee (one cup) inhibits platelet aggregation in man. We also demonstrated that coffee phenolic acids penetrate into platelets, and that they are present in platelets at concentrations that are able to inhibit platelet aggregation \textit{in vitro}. Clearly, we cannot exclude that other non-phenolic coffee compounds, different from caffeine, could contribute to the antiplatelet effect of coffee observed \textit{ex vivo}.

For a long time the relationship between coffee and cardiovascular risk has been controversial. However, recent epidemiological studies strongly suggest the existence of a J-shaped relationship linking coffee consumption and CVD.
risk\(^{(1–3)}\). It is reasonable to hypothesize that such J-shaped correlation is the direct result of the combined positive and negative action of different molecules present in coffee. As it has been demonstrated that ex vivo platelet aggregation is related to CVD mortality\(^{(10)}\), the anti-aggregative action of coffee phenolic acids could represent the bright side of coffee.

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