Chlorogenic and phenolic acids are only very weak inhibitors of human salivary α-amylase and rat intestinal maltase activities

Hilda Nyambe-Silavwe, Gary Williamson

School of Food Science and Nutrition, University of Leeds, Leeds LS2 9JT, UK

1. Introduction

Phenolic acids occur at high levels in many foods, including coffee, apples, potatoes, artichokes and prunes, and are predominantly found in the form of chlorogenic acids, where the phenolic acid moiety is attached to a quinic acid to form various isomers (Clifford, 1999). In foods, the most abundant is 5-cafeoylquinic acid (IUPAC numbering; 5-CQA, Fig. 1) and this isomer has also been the most studied. Numerous papers and reviews have been published on the potential health effects of phenolic acids (Van Dam & Hu, 2005; Higdon & Frei, 2006). In coffee drinkers, by far the most common source of chlorogenic acids, the isomer 5-CQA, which confers this antioxidant effect is not very clear (Van Dam & Hu, 2005), but among the mechanisms proposed is attenuation of carbohydrate digestion, as suggested for other polyphenols (Hanhineva & Hu, 2005), but among the mechanisms proposed is attenuation of carbohydrate digestion, as suggested for other polyphenols (Hanhineva & Hu, 2005). The most effective inhibition was with 3,4-dimethoxy-cinnamic acid (plateau at maximum 32% inhibition of human α-amylase at 0.6 mM), but this compound is found in coffee in the free form only at very low concentrations. Espresso coffee contains the highest levels of 5-CQA among all commonly consumed foods and beverages with a typical concentration of ~5 mM, and much lower levels of free phenolic acids. We therefore conclude that inhibition of carbohydrate-digesting enzymes by chlorogenic or phenolic acids from any food or beverage is unlikely to be sufficient to modify post-prandial glycaemia, and so is unlikely to be the mechanism by which chlorogenic acid-rich foods and beverages such as coffee can reduce the risk of developing type 2 diabetes.

Abbreviations: 5-CQA, 5-cafeoylquinic acid; PBS, phosphate-buffered saline; DNS, 3,5-dinitrosalicylic acid
* Corresponding author.
E-mail address: g.williamson@leeds.ac.uk (G. Williamson).

https://doi.org/10.1016/j.foodres.2018.07.038
Received 25 May 2018; Received in revised form 27 June 2018; Accepted 28 July 2018
Available online 30 July 2018
0963-9969/ © 2018 The Authors. Published by Elsevier Ltd. This is an open access article under the CC BY license (http://creativecommons.org/licenses/by/4.0/).
IC₅₀ values < 100 μM (Oboh et al., 2015). Two other studies showed that porcine α-amylase was inhibited by 5-CQA with IC₅₀ values also < 100 μM (Karim et al., 2017; Narita & Inouye, 2009) with caffeic and quinic acid being weaker inhibitors with IC₅₀ values of > 0.3 and > 25 mM respectively. However most of the studies used porcine, and > 25 mM respectively. However most of the studies used porcine, not human, α-amylase. Inhibition of different sources of enzyme varies widely (Nyambe-Silavwe et al., 2015; Pyner, Nyambe-Silavwe, & Williamson, 2017) and hence the current study aimed at using α-amylase from humans (salivary α-amylase) to reassert this activity of phenolic acids. We also determined the effects on α-glucosidase using a rat intestinal extract as the enzyme source, which has comparable inhibition properties to the human intestinal enzyme (Pyner et al., 2017).

2. Materials and methods

2.1. Reagents and standards

Caffeic acid, ferulic acid, 3,4-dimethoxycinnamic acid, 5-cafeoylquinic acid, 3,5-dinitrosalicylic acid, potassium sodium tartrate, amylose and human salivary α-amylase type IX-A were all purchased from Sigma-Aldrich. Co., Ltd., Dorset, UK. Oasis MAX cartridge 1 mL (30 mg) and 3 mL (60 mg) were purchased from Waters Ltd., Milford, MA, U.S.A. All the reagents were of the highest purity and standards were ≥ 98 %. The colour reagent was prepared by mixing 20 mL of 96 mM of 3,5-dinitrosalicylic acid with 8 mL of 5.3 M (12 g in 8 mL of 2 M sodium hydroxide) and 12 mL Millipore water. Human salivary amylase type IX-A stock concentration of 1.25 U/mL was prepared in PBS (0.01 M, pH 6.9) to give 0.5 U/mL in the assay according to the optimized assay (Nyambe-Silavwe et al., 2015).

2.2. α-Amylase inhibition assay

Amylose (1 mg/mL) was used as the substrate and the assay was conducted according to the optimized assay (Nyambe-Silavwe et al., 2015). A total assay volume of 500 μL was used consisting of 200 μL each of amylose and enzyme, 50 μL PBS and 50 μL of potential inhibitor at different concentrations. The potential inhibitor was replaced by an equal volume of PBS for the control. The reaction was carried out at 37 °C for 10 min upon addition of 200 μL of pre-incubated enzyme at 37 °C to a mixture of substrate, PBS and varying concentrations of inhibitor, also pre-incubated at 37 °C. To end the reaction, the samples were placed in the water bath at 100 °C for 10 min, cooled on ice and centrifuged for 5 min. Solid phase extraction (SPE) was carried out on the sample for removal of polyphenols that have been shown to interfere with the colour reagent solution containing 3,5-dinitrosalicylic acid (DNS). DNS reagent was added to the sample in a ratio of 2:1 and heated at 100 °C for 10 min. From each sample, 250 μL was placed in a 96 well plate and the absorbance was recorded at 540 nm. The rate of enzyme inhibition was calculated as a percentage of the control (without inhibitor) using the formula:

\[
\% \text{Inhibition} = \left(\frac{\text{Abs Control} - \text{Abs sample}}{\text{Abs control}}\right) \times 100.
\]

Where inhibition was obtained above 50%, IC₅₀ was calculated graphically by dose-dependent inhibition.

2.3. α-Glucosidase inhibition assay

The method used to assess rat α-glucosidase inhibition was adapted from Adisakkwattana, Charoenlertkul, & Yibchok-anun, 2009 as modified by Nyambe-Silavwe & Williamson, 2016. An assay volume of 500 μL was used and consisted of 50 μL of sodium phosphate buffer (10 mM, pH 7), 50 μL of potential inhibitor, 200 μL of acetone-derived protein intestinal extract from rat intestine (4 mg solid/mL for maltose) and 200 μL of substrate (3 mM maltose) (Nyambe-Silavwe & Williamson, 2016). Sodium phosphate buffer (50 μL) was put in place of the potential inhibitor for the control sample. The reaction was carried out at 37 °C for 20 min by adding the enzyme source to a mixture of sodium phosphate buffer, potential inhibitor and substrate. The reaction was stopped by heating in a water bath at 100 °C for 10 min, cooled to room temperature, polyphenols removed by solid phase extraction, hexokinase reagent added and absorbance read at 340 nm in a plate reader. Inhibition in the samples was calculated as a percentage of the control.

2.4. Statistical analysis

Statistical analysis was performed by one-way analysis of variance using the Number Cruncher Statistical System version 6.0 software (NCSS, LLC). Significant differences were assessed with Tukey-Kramer multiple comparison test (p ≤ .05). The data are expressed as the mean ± standard deviation (n = 3).

3. Results

3.1. Inhibitory effect on human salivary α-amylase and rat maltase activity

5-CQA only weakly inhibited human salivary α-amylase and rat intestinal α-glucosidase activities with maximum of 20.5 and 13.9 % respectively at the highest (5 mM) tested concentration (Fig. 2A). As a positive control, acarbose (a well-known carbohydrate inhibitor) exhibited an IC₅₀ value of 3.5 ± 0.3 μM for human salivary α-amylase and 0.40 ± 0.01 μM for rat intestinal maltase activities respectively (Fig. 2B and C), as expected (Nyambe-Silavwe et al., 2015).

Against human salivary α-amylase, ferulic acid, caffeic acid and 3,4-dimethoxycinnamic acid showed some dose-dependent inhibition, but the extent of inhibition was very low compared to acarbose (Fig. 2B). The most effective was 3,4-dimethoxycinnamic acid which gave maximum inhibition of 32% at 0.6 mM, but with no further change at increasing concentrations (p ≥ .05). Caffeic acid and ferulic acid both showed inhibition of < 20% at the highest concentration tested of 1 mM. None of the free phenolic acids inhibited rat intestinal maltase activity (Fig. 2C) even at the highest concentration tested.

4. Discussion

The aim of the present study was to determine whether the anti-diabetic properties attributed to drinking coffee (and possibly to other foods or beverages containing chlorogenic acids) are due to an acarbose-like action, a drug used to attenuate hyperglycaemia through inhibition of carbohydrate-digesting enzymes. In a detailed survey of...
espresso coffees, the strongest form of coffee normally consumed and commercially available in a major city of the UK, a typical serving of espresso was in the range of 24–422 mg per serving in an average serving size of 43 mL (Crozier, Stalmach, Lean, & Crozier, 2012). Since 5-CQA was on average 51% of the total content, then this corresponds to a 5-CQA concentration of 4.8 mM. This is diluted in the mouth with saliva and in the intestine with various digestive juices (Williamson, 2013). At this concentration, we would predict < 20% inhibition of human salivary α-amylase, which is not sufficient to exert an effect in vivo on carbohydrate digestion and post-prandial glucose concentrations, since we previously found that oleuropein, with IC₅₀ values superior to 5-CQA (0.56 mM for rat intestinal α-glucosidase and 1.4 mM for human salivary α-amylase), did not attenuate post-prandial blood glucose after consumption of bread as a carbohydrate-rich food (Kerimi et al., 2018). According to phenol-explorer (Neveu et al., 2010), filter coffee contains considerably less 5-CQA (Fig. 1), at ~2 mM. In addition, for most other commonly consumed foods, the concentration of chlorogenic acid is lower or much lower than in coffee (Fig. 1). Raw potato, for example, does not contain enough chlorogenic acid to exert any significant inhibition of α-amylase, and after cooking, the amount decreases; for oven baking or French fries, all chlorogenic acid is lost, for boiled potatoes, only 35% is left, or 55% after microwaving (Dao & Friedman, 1992). Free phenolic acids are present at very low levels in coffee (Encarnação, Farrell, Ryder, Kraut, & Williamson, 2015) and in most other foods (Neveu et al., 2010).

We found here that 5-CQA and free phenolic acids are very weak inhibitors of human salivary α-amylase, and even at high concentrations, 25% inhibition was generally not reached. We have specifically used a naturally-occurring substrate (amylase from starch) rather than a synthesized dye-linked substrate, which would have markedly different affinity for the enzyme, have also used a human source of α-amylase, and have ensured that the phenolic acids do not interfere in the DNS product determination. Several studies using the porcine pancreatic enzyme (Funke & Melzig, 2005; Karim et al., 2017; Narita & Inouye, 2009; Narita & Inouye, 2011; Oboh et al., 2015) have reported that chlorogenic acids inhibited α-amylase, and obtained an IC₅₀ value of 0.08 mM for 5-CQA (Narita & Inouye, 2011) and of 0.026 mM for caffeic acid (Oboh et al., 2015). It is now well established (Nyambe-Silavwe et al., 2015; Pyner et al., 2017) that the use of different enzyme sources for the inhibition assays as well as different substrates (Nyambe-Silavwe et al., 2015) can yield very different results.

For inhibition of rat intestinal maltase, 5-CQA was a weak inhibitor, but none of the free phenolics showed inhibition. Other research (Iwai, Kim, Onodera, & Matsue, 2006Kamitani, Iwai, Fukunaga, Kimura, & Nakagiri, 2009) also reported inhibition of α-glucosidase which is in contrast to our results demonstrating minimal inhibition. Human maltase is less susceptible to inhibition than rat maltase (Pyner et al., 2017), and hence we would have expected even lower inhibition in volunteers in vivo. We therefore conclude that the anti-diabetic effects of coffee consumption are not due to inhibition of carbohydrate-hydrolysing enzymes. However, several studies have shown that consumption of coffee reduces postprandial blood glucose levels. There was a significant reduction in total area under the glucose curve in a rat model after consumption of a standardised meal containing carbohydtrate with chlorogenic acid (Tunncliffe, Eller, Reimer, Hittel, & Shearer, 2011). In healthy males, it was shown that consumption of coffee polyphenol extract significantly reduced peak postprandial blood glucose as well as improving postprandial blood GLP-1 response which is associated with anti-diabetic effects (Jokura, Watanabe, Umeda, Hase, & Shimotoyodome, 2015). It was also shown, both in humans (Sarriá, Martínez-López, Mateos, & Bravo-Clemente, 2016) and in a rat model (Budryn et al., 2017), that consumption of green/roasted coffee blend led to lowering of blood glucose. Hence due to overwhelming evidence that coffee and its phenolic acids are associated with anti-diabetic properties via modulation of glucose metabolism, other mechanisms may be involved which include inhibition of intestinal glucose transport. In this respect, when consumed with a glucose bolus, coffee exhibited some effect on post-prandial glycaemia, although not directly on plasma glucose concentration (Johnston, Clifford, & Morgan, 2003), and this could be at least partly due to effects on glucose transporters or on hormonal response to food. These aspects should be addressed in future studies.

---

**Fig. 2.** Inhibition of enzyme activities by phenolics.
A: Inhibition of α-amylase (■) and of rat intestinal maltase (●) activities by 5-cafeoylquinic acid (5-CQA). B: Inhibition of α-amylase activity using amylase as substrate by acarbose (■), 3,4-dimethoxycinnamic acid (▲), caffeic acid (●) and ferulic acid (▼). C: Inhibition of rat intestinal maltase activity using maltose as substrate by acarbose (■), 3,4-dimethoxycinnamic acid (○), caffeic acid (△), and ferulic acid (□).
Acknowledgements

Some of this work was funded by the BBSRC, UK (Biotechnology and Biological Sciences Research Council, DRINC grant number BB/M027406/1).

References


