Combined treatment with caffeic and ferulic acid from 
Baccharis uncinella C. DC. (Asteraceae) protects against metabolic syndrome in mice

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Abstract

Fractionation of the EtOH extract from aerial parts of Baccharis uncinella C. DC. (Asteraceae) led to isolation of caffeic and ferulic acids, which were identified from spectroscopic and spectrometric evidence. These compounds exhibit antioxidant and anti-inflammatory properties and have been shown to be effective in the prevention/treatment of metabolic syndrome. This study investigated whether the combined treatment of caffeic and ferulic acids exhibits a more significant beneficial effect in a mouse model with metabolic syndrome. The combination treatment with caffeic and ferulic acids was tested for 60 days in C57 mice kept on a high-fat (40%) diet. The data obtained indicated that treatment with caffeic and ferulic acids prevented gain in body weight induced by the high-fat diet and improved hyperglycemia, hypercholesterolemia and hypertriglyceridemia. The expression of a number of metabolically relevant genes was affected in the liver of these animals, showing that caffeic and ferulic acid treatment results in increased cholesterol uptake and reduced hepatic triglyceride synthesis in the liver, which is a likely explanation for the prevention of hepatic steatosis. In conclusion, the combined treatment of caffeic and ferulic acids displayed major positive effects towards prevention of multiple aspects of the metabolic syndrome and liver steatosis in an obese mouse model.

Key words: Obesity; Caffeic acid; Ferulic acid; Metabolic syndrome

Introduction

Metabolic syndrome is a prothrombotic and proinflammatory state characterized by visceral obesity, insulin resistance, dyslipidemia and hypertension (1). As a result of increased synthesis and uptake of cholesterol and triglycerides from plasma as well as reduction in the export of these lipids to the circulation (2,3), patients with metabolic syndrome frequently exhibit steatosis and non-alcoholic fatty liver disease (2). Given that metabolic syndrome affects millions of individuals worldwide, there is increased interest in the development of molecules that can mitigate the metabolic consequences of obesity and liver disease. In this regard, caffeic (CA) and ferulic (FA) acids are natural organic compounds present in large amounts in the aerial parts of Baccharis uncinella C. DC. (Asteraceae) (4). These compounds function as key intermediate molecules in the biosynthesis of lignin, one of the principal components of plant biomass and its residues (5), and exhibit promising beneficial effects on metabolism when used in experimental models of metabolic syndrome. For example, treatment with CA has been reported to improve hyperglycemia and hepatic steatosis in animals kept on a high calorie diet, but obesity, hypertriglyceridemia and hypercholesterolemia were not always prevented (6–9). In addition, multiple studies indicate that treatment with CA improves dyslipidemia and minimizes fasting hyperglycemia, but does not always prevent diet-induced obesity (10–13).
Both CA and FA molecules only exhibit partial beneficial effects on metabolism; therefore, we investigated whether combined treatment with CA and FA could present a therapeutic advantage in the treatment of metabolic syndrome. Our results show that this combined approach with mice kept on a high-fat diet (HFD) successfully prevents obesity, dyslipidemia and liver steatosis.

**Material and Methods**

**Plant material**

Aerial parts of *B. uncinella* C. DC., were collected from Campos de Jordão, São Paulo State, Brazil in 2005. Botanical identification was made by Prof. Dr. Oriana A. Fávero (UPM). A voucher specimen (number SP382050) has been deposited at the Herbario da Prefeitura Municipal de São Paulo (PMSP), São Paulo, Brazil.

**Instruments**

Sephadex LH-20 (Amersham Biosciences, England) was used for column chromatographic separation while silica gel 60 PF254 (Merck, USA) was used for analytical TLC (0.25 mm). The $^1$H and $^{13}$C NMR spectra were recorded on a Bruker Ultrashield Avance II spectrometer operating at 300 and 75 MHz, respectively. Spectra were performed with CD$_3$OD or DMSO-$d_6$ (Tedia, Brazil) using the residual solvent peak as the internal standard. The chemical shifts ($\delta$) are given in parts per million and coupling constants (J) in Hz. LRESIMS was measured with a Micromass Platform mass spectrometer, operating in negative mode. Semi-preparative high-performance liquid chromatography was performed using a Dionex Ultimate 3000 chromatograph equipped with a Luna C18 column (250 × 10 mm, 5 $\mu$m id; Phenomenex, USA) and a UVD-DAD detector.

**Extraction and isolation**

Dried and powdered aerial parts of *B. uncinella* (400 g), popularly known as *vassoura*, were extracted with EtOH at room temperature. After solvent elimination under reduced pressure, 14 g of crude extract was obtained. Part of this material (5 g) was re-suspended in EtOAc and extracted using NaOH 4% until pH 10 was attained. The alkaline phase was made acidic (pH 1) with HCl 2% and extracted using NaOH 4% until pH 10 was attained. Part of this material (5 g) was re-suspended in EtOAc at room temperature. After solvent elimination under reduced pressure, silica gel 60 PF254 (Merck, USA) was used for analytical TLC was used for column chromatographic separation while silica gel 60 PF254 (Merck, USA) was used for analytical TLC. The animal livers were dissected and total RNA extracted using Trizol® (Life Technologies Inc., USA), according to the manufacturer’s instructions, and quantified by spectrophotometry (Nanodrop 2000c). For the reverse transcriptase reaction, 1.0 $\mu$g of total RNA was

| Components | Control | HFD |
|------------|---------|-----|
| Protein (%) | 22      | 20  |
| Carbohydrates (%) | 49     | 25  |
| Fat (%) | 4       | 40  |
| Others (%) | 25      | 15  |
| Calories (kcal·g$^{-1}$) | 1.7    | 5.4 |

Others: vitamins, minerals and fiber.
used in the SuperScript™ First-Strand Synthesis System for reverse transcription-polymerase chain reaction (Life Technologies Inc.) with a Mastercycler thermocycler (Eppendorf, Germany). Based on the reaction efficiency, approximately 120 ng of cDNA was used for amplification. Quantitative real time PCR was performed using QuantiTect™ SYBR® Green PCR (Qiagen, Valencia CA with program ECO by Illumina, USA). The cycle conditions were: 15 min at 94°C, 15 s at 94°C, 30 s at 60°C, and 30 s at 72°C for 50 cycles followed by the melting curve protocol to verify the specificity of amplicon generation. Gene expression was determined by the ΔΔCt method as described by Christoffolete (14). The housekeeping gene GAPDH and beta-actin were used as internal reference. Primer sequences are available upon request. The target genes were related to cholesterol (LDL-R, SREBP 2, LXR, ACAT-1, HMG-CoA), triglycerides (SREBP-1c, ChREBP, FAS, DGAT 2, MTTP, ATGL) and glucose (G6pase, PPAR-α) metabolism.

Western blotting
Brown adipose tissue was processed for mitochondrial isolation. Mitochondrial proteins were then size-fractionated by 12% SDS-PAGE and probed with UCP1 (Santa Cruz, Biotechnology, USA) (15).

Histology
After dissection, tissues (liver and white adipose tissue) were immersed in buffered formaldehyde solution (10%) and fixed for 24 h. Paraffin-embedded tissues were sectioned and processed as described for staining with hematoxylin-eosin or Masson’s trichrome. The area of adipocytes was estimated by analysis of images photographed at 100 × amplification with optical microscopy (Axioskop 2 plus, Zeiss, Germany). The images were analyzed by the program AxionVision Rel. 4.6, which estimated the area of 40 adipocytes per animal.

Statistical analysis
The statistical analyses were done by one-way analysis of variance followed by the Student-Newman-Keuls post-test when P<0.05. For all tests, P<0.05 was considered to be statistically significant. Data are reported as means ± SE.

Results
Characterization of caffeic and ferulic acids
Isolated compounds from aerial parts of B. uncinella were characterized as derivatives of cinnamic acid by analysis of their 1H NMR spectra. Typical signals of the trans-alkene system of C6-C3 derivatives at δ 7.43/7.50 (d, J=16.0 Hz, H-3) and 6.25/6.30 (d, J=16.0 Hz, H-2), as well as multiplets of range δ 6.70–7.10, assigned to aromatic hydrogens H-5, H-8, and H-9, were observed. Additionally, the spectrum of FA was observed to have an intense peak at δ 3.67, which was assigned to one methoxyl group. The 13C NMR spectra of CA and FA also displayed signals ranging from δ 115 to δ 150, corresponding to aromatic ring carbon atoms (C-1 to C-6), aliphatic sp2 carbon atoms at δ 116 (C-8) and δ 145 (C-7), and one carboxyl group at δ 168 (C-9). With FA, an additional peak corresponding to the methoxyl group linked to C-3 was observed at δ 56.1. LRESIMS showed a deprotonated ion (M-H)+ peak at m/z 179 and 193, corresponding to the molecular formula C9H8O4 and C10H10O4 of CA and FA, respectively. Following analysis of our recorded data and of a description in the literature (16), identification of CA and FA was achieved (Figure 1).

HFD and combined treatment with caffeic and ferulic acids
Keeping mice on HFD for 20 days increased caloric intake (Figure 2A) and accelerated body weight gain (Figure 2B and C). The combined treatment with CA and FA was started on the 21st day of HFD after the establishment of obesity. This regimen did not affect caloric intake (Figure 2A) but it did prevent body weight gain associated with HFD. Notably, the combined treatment reduced body weight gain to levels below those observed in the Control animals (Figure 2C). In addition, the fasting hyperglycemia induced by HFD (223±17.9 vs Control 150±12.1 mg/dL, with P<0.05) was improved with the combined treatment with CA and FA (180±19.9 mg/dL) and normalized the glucose intolerance caused by the HFD (Figure 3).

Investigation of the epididymal adipocyte area of the animals is shown in Figure 4. Feeding on the HFD increased the adipocyte area by approximately 2.5-fold whereas combined treatment with both CA and FA prevented such increase (Figure 4D). Brown adipose tissue was also studied by assessing the UCP1 protein levels, but the elevation
caused by the HFD was similar in all groups despite treatment with both CA and FA (Figure 5). HFD induced liver steatosis in control animals but this was prevented in the animals receiving combined treatment with CA and FA (Figure 6A–C). In addition, animals on HFD exhibited an elevation in plasma levels of cholesterol and triglycerides, but this increase was also prevented by the combined treatment with CA and FA (Table 2). These changes in liver and serum lipids were mediated by modifications in the key hepatic genes involved in lipid metabolism. Levels of FAS mRNA are decreased in animals treated with CA and FA. The gene expression for DGAT-2, the enzyme that catalyzes the synthesis of triglycerides, was also reduced. In addition, we also observed that the MTTP gene expression, the protein that transfers triglycerides to VLDL molecules, was significantly reduced by treatment with CA and FA. At the same time, the adipose triacylglycerol lipase (ATGL) mRNA levels increased with CA and FA treatment. This combined treatment also increased PPAR-α mRNA levels, but did not affect levels of mRNA for ChREBP or SREBP-1c mRNA (Figure 6D).

The genes involved in cholesterol metabolism were also evaluated. We found that the expression of SREBP-2, LXR and LDL-R genes all increased with the combined treatment of CA and FA. Unexpectedly, ACAT and HMG-CoA reductase mRNA levels were similar among the different groups in contrast to the data showing that cholesterol synthesis was inhibited by CA (Figure 6E).

Discussion

The present study shows that the combination therapy with CA and FA in mice with HFD-induced metabolic syndrome prevents obesity and reverts hyperglycemia,
dyslipidemia and hepatic steatosis, all conditions typically observed in mice treated with HFD only. Previous studies have shown that when used separately, CA prevented HFD-induced glucose intolerance but failed entirely in correcting obesity and dyslipidemia (6,9). Also, we found that FA alone corrected the HFD-induced dyslipidemia but failed to prevent obesity and glucose intolerance (10,11), which is in contrast to a previous report (9). In addition, studies have shown that treatment with either molecule alone prevented HFD-induced liver steatosis (6,17).

The ability of both CA and FA to prevent HFD-induced hypertriglyceridemia is remarkable and likely to involve i) a direct liver effect to coordinate reduction in the FAS and DGAT-2 expressions during HFD as well as to coordinate the induction of ATGL and PPARα expression and/or ii) an indirect effect mediated via prevention of obesity. Both CA and FA have been reported to decrease FAS enzymatic activity when administered separately (8–10). However, it is conceivable that the beneficial effect of treatment with CA on liver steatosis is predominantly indirect, via prevention of obesity, even though treatment with CA has previously been shown to induce PPARα in the liver (9). Therefore, the mechanism by which CA and FA prevent HFD-induced obesity remains to be clarified. Thermogenesis in brown adipose tissue does not seem to be involved given that UCP-1 levels were not affected by treatment with both molecules. The improvement in glucose tolerance is likely to be another byproduct of obesity prevention given the known negative correlation between obesity and insulin sensitivity.

The induction of the LDL-R, SREBP-2 and LXRα mRNA levels in liver by combined treatment with CA and FA is striking. These changes suggest that both the uptake and export of cholesterol in the liver were increased by the action of CA and FA, resulting in increased flow of hepatic cholesterol without causing hepatic steatosis and hypercholesterolemia. This is in agreement with a previously reported in vitro study showing that FA facilitates the capture of cholesterol.
and degradation of LDL cholesterol by isolated hepatocytes (18). The increased expression of LXR stimulated by the treatment of CA and FA suggests increased bile acid synthesis. It is difficult to reconcile these results with previous reports that suggest treatment with CA decreases the expression of SREBP-2 protein as assessed by western blotting, which is not supported by the elevation in LDL-R mRNA observed in the present study.

It is notable that in recent studies the combined treatment with CA and FA failed to increase mRNA levels of HMGCoA and ACAT-1, especially as both molecules, when administered separately, stimulate the expression of these genes (8,9).

In conclusion, this study provides compelling experimental evidence that combination therapy with CA and FA, isolated from aerial parts of B. uncinella C. DC. (Asteraceae), is highly effective in preventing the multiple aspects of metabolic syndrome in a HFD mouse model. The effects of these molecules are likely to take place in the liver as evidenced by changes in the expression of key genes involved in lipid metabolism. In addition, it is likely that there are direct effects of both molecules in the adipose tissue because of their efficacy in reducing diet-induced obesity.

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