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Running Title: Phenolic Compounds Showed Adipogenesis Activity

Phenolic Compounds from Leaves of *Casimiroa edulis* Showed Adipogenesis Activity

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Abbreviations: LC/MS, liquid chromatography mass spectrometry; PPAR, peroxisome
proliferator-activated receptor; FA, fatty acid

*Casimiroa edulis* is known as *cochitzapotl*, and it belongs a species of tropical
fruiting tree in the family Rutaceae, native to eastern Mexico and Central America south
to Costa Rica. In this study, we isolated two furocoumarins and two
polymethoxyflavones from leaves of *Casimiroa edulis* and evaluated the functions of
glucose and lipid metabolism activity with 3T3-L1 adipocytes. We discovered that the
addition of furocoumarins increased glucose uptake and lipid accumulation in 3T3-L1
adipocyte. These results suggest that furocoumarin compounds can be used as
functional food-derived compounds, to regulate adipocyte functioning for the
management of metabolic syndrome, which is associated with dysfunctions of glucose
and lipid metabolism.

Key words: liquid chromatography/mass spectrometry (LC-MS); adipocyte;
peroxisome proliferator-activated receptor (PPAR); fatty acid (FA) syntheses; phenolic
compounds

Casimiroa edulis is a species of tropical fruiting tree. Zapotin in Casimiroa edulis
has been found to induce both cell differentiation and apoptosis in cultured human
promyelocytic leukemia HL-60 cells. This compound inhibits 12-O-tetradecanoylphorbol 13-acetate (TPA)-induced ornithine decarboxylase (ODC)
activity in human bladder carcinoma cells (T24 cells) and TPA-induced nuclear
factor-kappa B (NF-κB) activity in human hepatocellular liver carcinoma cells (HepG2
cells). The data suggest that zapotin deserves further investigation as a potential cancer
suppressive agent. Crude plant extracts of seeds and of leaves of Casimiroa edulis have
been found to affect blood pressure, cardiac activity, and aortic muscular tone, and
to possess anticonvulsant activity, but previous study has focused mainly on seed
components and leaf components that were not studied well.

Adipocytes play a major role in the maintenance of energy balance. The amount
of glucose uptake into the adipocytes is significant in the maintenance of serum glucose
levels, because adipocyte-specific knock-out of an insulin-dependent glucose
transporter (glut4) results in diabetic symptoms. Lipid synthesis (lipogenesis) is also a
vital function of adipocytes. In vivo studies have suggested that adipose tissue is responsible for more than 50% of fatty acid synthesis in mammals.\textsuperscript{10} Fatty acid (FA) synthesis is a cytosolic process utilizing acetyl CoA as precursor. It is generally accepted that fatty acid is synthesized from various substrates (glucose, pyruvate, lactate and acetate).\textsuperscript{11} It is indispensable for the management of adipocyte functions to evaluate complicated metabolisms, including lipogenesis and glucose uptake.

In this study, we tried to isolate phenolic compounds from \textit{Casimiroa edulis} leaves, and investigated other effect, adipogenesis, including glucose uptake, fatty acid synthesis activity, and lipid accumulation with 3T3-L1 adipocytes. PPAR\textgreek{g} synthesized ligand, pioglitazone, was used as positive control, since PPAR\textgreek{g} is abundantly expressed in 3T3-L1 adipocytes and plays a central role in metabolism and differentiation.\textsuperscript{12} Pioglitazone, a major insulin sensitizer, promotes preadipocyte metabolism including glucose uptake and FA synthesis through PPAR\textgreek{g} activation\textsuperscript{13} and is widely used in the treatment of type 2 diabetes.

\textbf{Materials and Methods}

Leaves of \textit{Casimiroa edulis} were collected at the Botanical Garden of Nippon Shinyaku Co., Kyoto, Japan. The voucher specimen was deposited at Gifu Pharmaceutical University. \textsuperscript{13}C\textsubscript{6} glucose and pioglitazone was purchased from Sigma (St. Louis, MO). [1,2,3,4,5,6,7,8,9,10,11,12,13,14,15,16-\textsuperscript{13}C] palmitic acid was from Watari (Tokyo). Unless otherwise indicated, all chemicals were purchased from Nacalai Tesque (Kyoto, Japan) or Wako Pure Chemicals (Osaka, Japan) and were of guaranteed reagent grade or tissue culture grade.
General experimental procedures. $^1$H- and $^{13}$C-NMR spectra were recorded on an EX400 ($^1$H at 400 MHz and $^{13}$C at 100 MHz, JEOL) spectrometer. Chemical shifts appeared as $\delta$ values with trimethylsilane (TMS) as internal reference. Peak multiplicities were quoted in Hz. EI-MS was recorded on a JMSDX-300 (JOEL) spectrometer. Kiesel gel 60 (60-230 mesh, Merck, Japan) and Fuji Silysia Chemical Chromatorex ODS (100—200 mesh) were used for column chromatography. Kiesel-gel 60 F256 (Merck, Tokyo) at 0.25 mm was used for analytical and 0.5 mm for preparative TLC.

Extraction and isolation of compounds. Dried leaves (550 g) extracted with MeOH (1 L x 5 times) at room temperature and was evaporated in vacuo. The MeOH extract (52 g) obtained was chromatographed on Silica gel (250 mm $\times$ 30 mm) with a mixture of chloroform–MeOH by increasing the polarity from 0% to 10 % methanol. The chloroform-MeOH (10:1) (3.5g) was further chromatographed on an ODS column (250 mm $\times$ 20 mm) eluted with H$_2$O, followed by increasing the concentration of MeOH to give four fractions. Fraction 2 (263 mg, 70% methanol) was repeatedly purified by preparative TLC ($n$-hexane-EtOAc 1:4) to give 1 (5 mg), 2 (7 mg), 3 (12 mg), and 4 (10 mg).

8-(3'-Hydroxymethyl-but-2-enyloxy)-psoralen acetate (compound 1). White amorphous solid. Negative ion FABMS [M-H]$^-$ m/z 327. $^1$H NMR(CDCl$_3$) $\delta$: 1.81 (3H, br s, H-5'), 2.04 (3H, s, acetyl-Me), 4.66 (2H, br s, H-4'), 5.09 (2H, br d, $J = 7.0$ Hz, H-1'), 5.86 (1H, br t, $J = 7.0$ Hz, H-3'), 6.38 (1H, d, $J = 10.0$ Hz, H-4), 6.82 (1H, d, $J = 2.5$ Hz, H-9), 7.38 (1H, s, H-5), 7.69 (1H, d, $J = 2.5$ Hz, H-10), 7.77 (1H, d, $J = 10.0$ Hz, H-3): $^{13}$C NMR (CDCl$_3$) $\delta$: 160.43 (C-2), 114.83 (C-4), 144.27 (C-4), 116.54 (C-4a), 113.81 (C-5), 125.95 (C-6), 148.32 (C-7), 131.39 (C-8), 143.67 (C-8a), 106.76 (C-9), 146.76 (C-10), 69.12 (C-1'), 125.13 (C-2'), 136.57 (C-3'), 62.79 (C-4'), 21.42 (C-5'), 20.83 (acetyl-Me), 170.85 (acetyl C=O).
8-(3'-Hydroxymethyl-but-2-enyloxy)-5-methoxypsoralen acetate (compound 2).

White amorphous solid. Negative ion FABMS [M-H]⁻ m/z 357. ¹H NMR(CDCl₃) δ:
1.80 (3H, s, H-5'), 2.03 (3H, s, acetyl Me), 4.18 (3H, s, C-5 OMe), 4.66 (2H, br s, H-4'), 4.91 (2H, br d, J = 7.0 Hz, H-1'), 5.86 (1H, br t, J = 7.0 Hz, H-2'), 6.28 (1H, d, J = 10.1 Hz, H-3), 6.99 (1H, d, J = 2.5 Hz, H-9), 7.64 (1H, d, J = 2.5 Hz, H-10), 8.12 (1H, d, J = 10.1 Hz, H-4); ¹³C NMR (CDCl₃) δ: 160.42 (C-2), 112.91 (C-3), 139.55 (C-4), 107.61 (C-4a), 144.26 (C-5), 114.61 (C-6), 150.57 (C-7), 136.51 (C-8), 125.63 (C-8a), 105.12 (C-9), 145.16 (C-10), 69.36 (acetyl Me), 170.84 (acetyl C=O), 60.79 (3H, s, C-5 OMe).

5,6,2',3',5',6'-Hexamethoxyflavone (compound 3). White amorphous solid: exhibited spectral data (¹H NMR and EIMS) comparable to published data.¹⁴

5,6,2'-Trimethoxyflavone (compound 4). White amorphous solid: ¹H NMR data was comparable to published data;⁶ ¹³C NMR (CDCl₃) δ: 159.05, 157.90 (C-2, C-2'), 111.68 (C-2), 178.41 (C-4), 119.07, 119.02 (C-4a, 8), 147.82 (C-5), 149.65 (C-6), 113.33 (C-7), 151.82 (C-8a), 120.65 (C-1'), 112.96 (C-3'), 132.13 (C-4'), 120.62 (C-5'), 129.07 (C-6')

LC/MS system. The Agilent Series 1100 LC system (Agilent Technologies, Waldbronn, Germany) consisting of a G1323B control module, a G1312A quaternary pump, a G1322 degasser, a G1329A, a G1330A autosampler, a G1316A column oven, and a G1314A detector, was used. The column used was an Xbridge™ (C18 3.5 μm, 2.1×150 mm, Waters, Milfold, MA). The LC pump gradient was as follows: 85% mobile phase A (water in 5 mM ammonium acetate) and 15% mobile B (methanol in 5 mM ammonium acetate) from 0 to 20 min, 99% mobile phase A and 1% mobile phase B from 20 to 20.1 min, and 85% mobile phase A and 15% mobile phase B from 20.1 to 35 min.
The injection volume, flow rate, and column temperature were 5 μL, 0.2 mL, and 40°C, respectively. MS conditions were selected ion monitoring (SIM); polarity negative; 10 psi nebulizer gas, 150 V fragment ion voltage; 3,000 V Capillary voltage, 350°C; dwell time 50 msec. Unless otherwise specified, all reagents used were of analytical grade, and the water was deionized. Solvent: LC/MS grade methanol. [1,2,3,4,5,6-\textsuperscript{13}C] glucose and [1,2,3,4,5,6,7,8,9,10,11,12,13,14,15,16-\textsuperscript{13}C] palmitic acid were detected at 185 and 271, m/z respectively.\textsuperscript{15)

Spectra of furocoumarins were determined in LC/MS scan mode, polarity negative (50- 400 m/z).

\textit{GC/MS system.} Gas-chromatographic analyses were performed using an Agilent 6890 quadrupole gas chromatograph (GC) equipped with an Agilent 5973B mass spectrometer (MS). The GC column was a DB-5MS+DG capillary type (30m × 0.25 mm i.d., 0.25 μm film thickness; J&W Scientific, Folsom, CA). A 1-μL extract was injected in splitless mode at an injection temperature of 250°C. The oven temperature was programmed to increase from an initial 60°C (held for 1 min) to 300°C (held for 2 min) at a rate of 20°C/min. The temperatures of the quadrupole and ion source were 150 and 230°C. The GC-MS system was operated in scan mode (50-500 m/z) with the electron multiplier tune value.

\textit{Cell culture.} 3T3-L1 preadipocytes, purchased from the American Type Culture Collection (Rockville, MD), were grown in DMEM-high glucose medium supplemented with 10% fetal bovine serum, 100 U/mL of penicillin, and 100 μg/mL streptomycin at 37°C in 5% CO\textsubscript{2} using a 12-well plate (5 × 10\textsuperscript{4} cell/well). Post-confluent preadipocytes were differentiated into adipocytes by adding 10 μg/mL insulin, 0.1 μg/mL dexamethasone, and 112 μg/mL isobutylmethylxanthine to the medium with 5 μM pioglitazone, and 5 μM each test compound in DMSO, as previously described.\textsuperscript{12) The differentiation medium was removed after 2 days and replaced with
DMEM-high glucose medium containing 5 μg/mL insulin with TZD or the test compound every 2 days. Ten days after the addition of the differentiation medium, the cells were plated in culture dishes for a given experiment. After differentiation, they were washed with warm PBS, and the medium was replaced with glucose-free DMEM containing 4.5 mg/mL [1,2,3,4,5,6-13C] glucose. They were incubated for a further 48 h. This was repeated twice.

Extract and saponification. The media were collected after incubation for specific durations and diluted with methanol. Diluents were filtered with a 0.22 μm filter membrane, and were analyzed by the LC/MS system. The cells were dissolved in 5 mL KOH 20% (ethanol:water 4:6; v/v), and hydrolyzed for 60 min at 80°C, followed by acidification with 6 mL 5 N HCl. The FAs were extracted twice with 4 ml diethylether, dried under N2 at 40°C, dissolved in 5 ml methanol, and analyzed by the LC/MS system.

Oil Red O staining. Cells were fixed with 10% formaldehyde/PBS and stained with Oil Red O solution (0.5% Oil Red O-isopropyl alcohol/H2O (3:2, v/v)).

Statistical analysis. Results are expressed as mean and standard error of mean (SEM). Statistical significance of differences among groups was evaluated by Student’s t-test. Differences were considered significant at \( p < 0.05 \).

Results and Discussion

NMR analysis. The ¹H NMR of compound 1 showed the presence of two sets of
coupled protons that $\delta 6.38$ and $7.77$ ($J = 10.0$ Hz) and $\delta 6.82$ and $7.69$ ($J = 2.5$ Hz).

The data indicated compound 1 was a furocoumarin. An aromatic singlet ($\delta 7.37$) was also observed that correlated with the C-4 signal of the coumarin skeleton in the HMBC spectrum. Other HMBC correlations indicated that the furan ring was attached to C-6, 7, like psoralen. $^1$H and $^{13}$C NMR showed the presence of two oxymethylene ($\delta_H 4.66$ and 5.09, and $\delta_C 62.79$ and 69.12), a tri-substituted double bond ($\delta_H 5.09$; $\delta_C 125.13$ and 136.57), a methyl attached to a double bond ($\delta_H 1.81$; $\delta_C 21.41$), and an acetyl group ($\delta_H 2.04$; $\delta_C 20.83$ and 170.85). In the HMBC spectrum, an oxymethylene proton was correlated with C-8 of a coumarin. These data are characteristic of a prenyloxy group, in which one of methyl groups had been oxidized to a hydroxyl methyl and acetylated. The hydroxymethyl group was of the cis form with respect to the oxymethylene group (H-1’) of the prenyl substituent on the basis of the NOESY spectrum (H-1’/H-4’ and H-2’/H-5’).

Therefore, the structure was determined to be as described in Fig. 1. This compound was first isolated from leaves of *Casimiroa pringlei*. $^{16,17}$

The $^1$H NMR spectrum of compound 2 was very similar to that of 1, but an aromatic proton disappeared and a methoxy group ($\delta 4.18$) was observed. An NOE was observed between the methoxy group and H-4 of the coumarin skeleton. By detailed analysis of NMR by 2D methods, the structure of 2 was determined to 5-methoxylated compound of 1. A deacetylated coumarin of 2 was isolated from *Casimiroa tetrameria*.

To the best of our knowledge, compound 2 was a new compound.

The structures of compounds 3 and 4 were determined and identified by comparison to previous data as 5,6,2’,3’,5’,6’-hexamethoxyflavone (3)$^{18}$ and 5,6,2’-trimethoxyflavone (4)$^{14,18}$ respectively.

Figure 2 shows spectra of furocoumarins and polymethoxyflavones. Parent ions of them as determined by NMR were obtained by LC/MS and GC/MS analysis (327, 357, 402, 312 m/z). This result confirms the NMR analysis data.

*Glucose uptake and glycerol output activity.* To elucidate the effects of
compounds 1-4 on adipocyte functioning, we examined the insulin-dependent glucose uptake and glycerol output of differentiated 3T3-L1 adipocytes. As shown in Fig. 3A, the addition of 5 μM compounds 1 and 2 enhanced the insulin-dependent uptake of glucose (1.2-1.3 fold increases as compared to control). The addition of the other compounds did not enhance insulin-dependent uptake of glucose. On the other hand, the addition of 5 μM compounds 1-4 did not enhance glycerol output or lipolysis (Fig. 3B). These compounds at the concentrations used in this study did not affect cell viability (data not shown).

De novo FA synthesis activity and lipid accumulation of adipocytes. To assess the effects of compounds 1-4 on lipid metabolism in 3T3-L1 adipocytes, we analyzed de novo FA syntheses activity with compounds 1-4. Figure 3C shows new synthesis of palmitic acid in 3T3-L1 adipocytes treated with compounds 1-4 for 48 h. The addition of pioglitazone at (5.0 μM) increased the amounts of synthesized FA, by approximately 2.2-fold in the 3T3-L1 adipocytes as compared with control. The addition of 5.0 μM compounds 1 and 2 enhanced FA syntheses by approximately 1.5-fold as compared with control. Finally, to investigate the dose-dependent effects of furocoumarins on the lipid accumulation of adipocytes, we performed Oil Red O staining. The addition of furocoumarins increased lipid accumulation dose-dependently (Fig. 4).

Obesity and associated disorders are the major noncommunicable public health problems of the 21st century. Studies indicate that high levels of body fat are associated with an increased risk of developing numerous adverse health conditions.\textsuperscript{19,20} Hence, effective therapeutic approaches to obesity and obesity-induced metabolic syndrome are of wide interest. In this study, we isolated two furocoumarins and two polymethoxyflavones from leaves of \textit{Casimiroa edulis} fruit and evaluated the functions of glucose and lipids metabolism activation. We found that the addition of furocoumarins (compounds 1 and 2) caused increased glucose uptake, FA synthesis, and
lipid accumulation in 3T3-L1 adipocytes. Furanocoumarins are known to have biological effects in humans. For example, bergamottin and dihydroxybergamottin are attributable to the grapefruit juice effect, in which these furanocoumarins affect the metabolism of certain drugs, but the concentration of furanocoumarins used in our assays had no influence on cell viability. They might lower blood glucose levels by orally ingested in a meal or an ordinary drink, and this calls for further evaluation in vivo.

References
**Figure legends**

**Fig.1.** Structures of Compounds Isolated from Leaves of *Casimiroa edulis*.

Furocoumarins (compounds 1 and 2), polymethoxyflavones (compounds 3 and 4)

**Fig.2.** Mass Spectra of the Isolated Compounds.

Mass patterns of compounds 1-4 are shown. Compounds 1-2 and Compound 3-4 were analyzed by GC/MS and LC/MS respectively.

**Fig.3.** $^{13}$C$_6$-Glucose Uptake (A), Glycerol Output (B), and *De Novo* $^{13}$C-FA Synthesis (C) in 3T3-L1 Adipocytes after the Addition of Compounds As Determined by LC/MS Analysis of the Medium at 48 h. Incubation.

$^{13}$C$_6$-Glucose uptake and glycerol output of 3T3-L1 adipocytes after addition of coumarins and polymethoxyflavones were determined by LC/MS. The concentrations of $^{13}$C$_6$-glucose and glycerol were determined by the LC/MS system at 185 m/z ($^{13}$C$_6$-glucose) and 91 m/z (glycerol) respectively. *De novo* synthesis of palmitic acid was monitored by analyzing 271 (m/z). Lane 1, control; lane 2, pioglitazone; lanes 3-4, compounds 1 and 2; lanes 5 and 6, compounds 3 and 4 respectively. Lane 1, control; lane 2, pioglitazone; lanes 3 and 4, compounds 1 and 2; lanes 5 and 6, compounds 3 and
4 respectively. Concentrations (in μM) at all data points is 5.0 **p < 0.05 compared to control.

4 **Fig. 4. De Novo \(^{13}\)C-FA Synthesis and Lipid Accumulation of 3T3-L1 Adipocytes.**

Adipocytes treated with pioglitazone or compounds 1,2 (furocoumarins) (1-100 μM) were fixed with 10% formaldehyde/PBS and stained with Oil Red O solution (0.5% Oil Red O-isopropyl alcohol/H\(_2\)O 3:2, v/v). Microscopy views of representative 3T3-L1 cells (original magnification, x 100).
Figure 2, Nagai
Figure 3, Nagai
Figure 4, Nagai