Abstract. Insulin resistance is a major factor in type II diabetes development, occurring when insulin levels are normal, but do not have normal interactions with adipose, muscle or liver tissue. The present study aimed to explore the hypoglycemic effect of Antrodia cinnamomea (AC) mycelium powder by evaluating its impact on insulin resistance and plasma free fatty acid (FFA) levels in steroid -induced insulin -resistant (SIIR) rats. Male Wistar rats were administered dexamethasone for 5 days to induce insulin resistance. The SIIR rats were subsequently randomly assigned into three experimental groups (EGs) and a control group (CG), where saline was orally administered. The EGs were orally administered different doses of AC (100, 200 or 500 mg/kg) and an optimal dose for further study was determined. Changes in plasma insulin and glucose levels were calculated to investigate the hypoglycemic effect of AC. To evaluate insulin resistance, the homeostasis model assessment-estimated insulin resistance of the SIIR rats was determined. Changes in plasma FFA levels were detected and levels of insulin signal proteins (IRS -1, GLUT-4 and PI3K) were analyzed by western blot to elucidate AC's mechanism of action. The SIIR rats exhibited significantly decreased plasma glucose levels in the first 30 min, with plasma FFA levels displaying a marked downward trend (P<0.05) when they were administered the optimal dose of AC (200 mg/kg). The decrease in plasma glucose and FFA levels was significantly larger in the EG compared to the CG, and insulin signal protein levels were also significantly increased (P<0.05). The hypoglycemic effect observed may be due to decreased plasma FFA levels and increased expression of intracellular insulin signal proteins. Furthermore, insulin sensitivity was enhanced, indicating that AC acts as an insulin sensitizer in insulin resistant animal models.

Introduction

Diabetes mellitus (DM) is one of the top 10 most prevalent diseases worldwide, ranking fourth in Taiwan. Individuals with DM are at an increased risk of developing eye disease, foot lesions, nerve degeneration, cerebrovascular disease, cardiovascular disease, hypertension and renal disease (1). The pathology of type II DM results from insulin resistance and/or a lack of insulin secretion, which ultimately increases plasma glucose levels. This hyperglycemic state stimulates insulin secretion and induces hyperinsulinemia (2). The clinical symptoms of insulin resistance include dyslipidemia, hypertension, glucose intolerance, hyperuricemia or gout, central obesity, impaired blood coagulation or hyper-coagulation, hyperandrogenism like symptoms, fatty liver and coronary vascular disease (3). These chronic complications not only threaten the life of those diagnosed with type II DM, but also puts a heavy burden on medical resources and the social economy (4).
Todate, the best methods for improving insulin resistance are exercise (5), smoking cessation (6) and medications including metformin, a biguanide (7) and thiazolidinediones (TZDs), a class of oral hypoglycemic agents (8). In complementary and alternative medicine, it has also been demonstrated that electroacupuncture on the Zhongwan (CV12) acupoint or the bilateral Zusanli (ST36) acupoints may lower plasma glucose and enhance insulin sensitivity (9-11). Among the aforementioned methods, the use of medication is the most effective and convenient for the management of type II DM, but can result in adverse side effects. Therefore, it is important to identify other methods with fewer side effects that can improve insulin resistance.

**Antrodia cinnamomea** (AC) is a Taiwanese fungus species that grows in the hollow trunk of the *Cinnamomum kanehirai* tree. Previous investigations have indicated that AC may suppress tumor formation (12), enhance the immune system (13), inhibit viral activity and protect the liver (14). It is often used in Taiwanese folk medicine. Some evidence has also indicated that AC can act as an anti-oxidant, improve hypertension and decrease plasma lipids (15-17). Ergostatrien-3β-ol (EK100) from the AC was evaluated for its hypoglycemic effects and was demonstrated to improve diabetes and dyslipidemia in mice fed a high-fat diet. EK100 treatment also resulted in decreased visceral adipocyte size and reduced the ballooning degeneration of hepatocytes. Levels of glucose transporter 4 (GLUT-4) protein and Akt phosphorylation in skeletal muscle are also significantly increased in EK100 treated mice (18). In addition, antroquinonol extracted from the mycelium of AC effectively inhibited dipeptidyl peptidase-4 activity and AMP-activated protein kinase (AMPK) activation (19). Dehydroeburicoric acid from AC prevented the development of diabetic and dyslipidemic states in streptozotocin-induced diabetic mice, through the regulation of GLUT-4, peroxisome proliferator activated receptor α, fatty acid synthase and AMPK phosphorylation (20). An extract made from the fruiting body of AC lowered liver triglyceride and total cholesterol levels (17) and enhanced the production of superoxide dismutase, catalase and glutathione peroxidase (21).

In addition to the aforementioned effects of AC, its therapeutic effects on insulin resistance have also been investigated (22-24), but the mechanism by which AC enhances insulin sensitivity has not been completely elucidated, particularly for steroid-induced insulin-resistance (SIR) (25,26). Steroids are widely prescribed and are known to cause insulin resistance. Patients with DM treated with steroids commonly require an increased dose of insulin. Thus, the aim of the present study was to use the SIR rat as a model to explore the hypoglycemic and insulin resistance improving effects of orally administered AC and to investigate the mechanisms underlying its hypoglycemic and insulin resistance improving properties.

**Materials and methods**

**Preparation of AC mycelium.** AC mycelium was cultured on solid-state cereal medium (provided by the Chair Professor Wai-Jane Ho, Da-Yeh University, Changhua, Taiwan) for 85-90 days at 21-23°C in the dark. The solid medium was mainly barley-supplemented with yeast extract and glucose. At the end of the culturing period, cultures were harvested, dried and ground into powder for subsequent experiments. The final yield from the solid-state culture was ~20% of the original dry weight of cereal medium (Fig. 1A). The powdered culture was then mixed with normal saline to produce oral solutions at the concentrations of 100, 200 and 500 mg/kg body weight (BW). The solutions were subsequently stored in a 4°C refrigerator ready for use.

**High-performance liquid chromatography (HPLC) assay.** Components of the AC powder methanol extract were assessed by HPLC assay with a configured module using a Waters Alliance 2695 with auto-sample injection (Waters X Terra column, 4.6x250 mm, 5 µm; mobile phase A, water; mobile phase B, acetonitrile; injection volume, 10 µl; detector, PDA; detector wavelength, qualitative 250 nm, quantitative 280 nm; temperature, 25°C; flow rate, 1.0 ml/min; Waters Corporation, Milford, MA, USA). The peaks of the retention time from the analysis of the AC methanol extract were compared with the peaks of the internal standard dissolved in methanol solution (provided by Professor Yueh-Hsiung Kuo, China Medical University, Taichung, Taiwan).

**Animal models.** A total of 24 Male Wistar rats aged 8-10 weeks, 250-300 g BW, were obtained from the BioLasCO (Taipei, Taiwan). They were housed in rooms at 25±1°C with relative humidity of 65±5%. The rats were acclimatized in an alternating 12 h light/12 h dark cycle with free access to water and standard rat chow. After 1 week of adaptation, the rats were randomly assigned into experimental and control groups. The Institutional Animal Care and Use Committee of Da-Yeh University approved the methods of this study according to the national guidelines for the Care and Use of Animals.

A previously published research protocol was followed to establish the SIR rat model (26). At 8 weeks, rats were intraperitoneally (i.p.) injected with dexamethasone at a dose of 1 mg/kg daily for 5 days. It was concluded that insulin resistance was successfully induced when their measured fasting plasma glucose levels were higher than 150 mg/dl. Tests were performed at 8:00 a.m. and all SIR rats were fasted before each test and anesthetized with pentobarbital (40 mg/kg i.p.) (11).

**Experimental protocols.** The SIR rats were randomly divided into four groups and administered the following treatments once: Group A, control group (n=6), 1 ml/kg oral saline; group B (n=6), 100 mg/kg AC; group C (n=6), 200 mg/kg; and group D (n=6), 500 mg/kg AC. The three experimental groups (EGs) were force-fed the AC solution. Every AC sample was then mixed with normal saline to produce oral solutions at the concentrations of 100, 200 and 500 mg/kg body weight of cereal medium (Fig. 1A). The powdered culture was then mixed with normal saline to produce oral solutions at the concentrations of 100, 200 and 500 mg/kg body weight (BW). The solutions were subsequently stored in a 4°C refrigerator ready for use.

**Assay of plasma free fatty acid (FFA) levels.** Plasma FFA levels were measured using a non-esterified fatty acid kit...
Fatty acid (FFA) levels were measured via an ELISA. As described in our previous study (25), plasma FFA was transformed into a purple adduct for subsequent detection by automatic spectrophotometer (COBAS MIRA Plus system; Roche Diagnostics, Basel, Switzerland).

**ELISA of plasma insulin levels and resistance test.** A Mercodia ultrasensitive rat insulin ELISA kit (cat. no. 10-1251-01; Mercodia AB, Uppsala, Sweden) was used to detect plasma insulin levels. The homeostatic model assessment of insulin resistance (HOMA-IR) was calculated using the following formula: [fasting plasma insulin levels (µU/ml) x fasting plasma glucose (mmol/l)]/22.5 (27,28).

**Western blot analysis.** The samples were minced coarsely and homogenized by an ultrasonic processor (VCX 750; Sonics and Materials Inc., Newtown, CT, USA) in a radioimmunoprecipitation assay lysis buffer with the protease inhibitor, phenylmethylsulfonyl fluoride (Santa Cruz Biotechnology, Inc., Dallas, TX, USA). The muscle extracts were centrifuged at 16,440 x g at 4°C for 1 h, and the supernatants were measured using a spectrophotometer. A total of 90 µg/ml protein was separated by 8% SDS-PAGE, and the proteins were transferred to a polyvinylidene fluoride (PVDF) membrane for western blotting. The PVDF membrane was then submerged in 5% non-fat milk to block the nonspecific binding sites in the membrane at 25°C for 1 h. The membrane was incubated overnight with anti-insulin receptor substrate-1 (IRS-1; 1:200, sc-559), anti-GLUT-4 (1:200, sc-7938) and anti-phosphoinositide 3-kinase (PI3K; 1:200, sc-376112) antibodies (all from Santa Cruz Biotechnology, Inc.) at 4°C in the refrigerator. Finally, the membranes were incubated with goat anti-rabbit immunoglobulin G-horseradish peroxidase antibodies (GTX213110-01, 1:2,000; GeneTex, Inc., Irvine, CA, USA) for at 25°C for 1 h, and specific bands were detected by an enhanced chemiluminescence kit (Clarity™...
and Clarity Max™ Western ECL Blotting Substrates; Bio-Rad Laboratories, Inc., Hercules, CA, USA. The bands were quantified using optical densitometry (Gel-Pro analysis version 4.0; Media Cybernetics, Rockville, MD, USA). The actin bands were used as an internal loading control, and the results are presented as a ratio of signal-to-actin (11,26).

**Statistical analysis.** The experimental results are presented as the mean ± standard error in each group (n=6). The statistical analysis of the results was performed using the Student’s t-test or one-way analysis of variance, with a least significant difference post hoc test. P<0.05 was considered to indicate a statistically significant difference.

**Results**

**HPLC assay.** HPLC analysis detected eburicoic acid + dehydroeburicoic acid (TR1+TR2), sulphurenic acid + dehydrosulphurenic acid (TR3+TR4) and EK100 (standard retention times: 36-38, 20-23, and 52.8 min, respectively). The AC sample contained TR1+TR2, TR3+TR4, and EK100 (Fig. 1B).

**Hypoglycemic effect of AC in SIIR.** The SIIR rats in the EGs were administered different doses of AC (100, 200 or 500 mg/kg) and the CG rats were administered normal saline. At 30 min after treatment, the plasma glucose levels in the rats administered 200 and 500 mg/kg of AC was significantly lower than those in the CG (P<0.05). After 60 min, the plasma glucose levels in all EGs was significantly lower than those in the CG (P<0.01; Fig. 2A). At 30 min after treatment administration, the hypoglycemic activity in the 200 and 500 mg/kg groups was 21.69 and 15.22% respectively, which was significantly greater than that of the CG (3.33%, P<0.05). After 60 min, the hypoglycemic activity of the 200 and 500 mg/kg groups was 26.59 and 26.86%, respectively, both of which were significantly greater than that of the CG (9.74%, P<0.05; Fig. 2B). No specific adverse events, including no mortality, normal activity, water and food intake were observed following each experiment at any dose.

**Hypoglycemic effect of the optimal dose of AC.** The above results indicated that 200 mg/kg AC was optimal to achieve a hypoglycemic effect. Therefore, this particular dose was used in the subsequent experiments. Thirty minutes after administration of 200 mg/kg of AC, the plasma glucose level decreased from 155.33±23.12 to 125.81±29.81 mg/dl (P<0.05 vs. baseline), and to 113.69±14.15 mg/dl after 60 min (P<0.05 vs. baseline).

**Plasma insulin and HOMA-IR levels in SIIR rats following administration of 200 mg/kg AC.** Following oral administration of 200 mg/kg AC, plasma insulin levels were detected in serum samples taken from the femoral vein using an ELISA kit. Plasma insulin levels increased from 0.35±0.41 to 0.51±0.30 pmol/l at 30 min after AC administration, with a significant elevating trend compared to those of the CG, in which plasma insulin levels changed from 0.11±0.07 to 0.07±0.02 pmol/l after 30 min. There was a significant difference between the levels of insulin in the EG and the CG at 30 min (Fig. 3A).

There were no significant differences in HOMA-IR levels between the EG and the CG at baseline and 30 min after oral administration of 200 mg/kg of AC. However, HOMA-IR levels in the EG were significantly decreased compared...
with the CG at 60 min after AC administration (1.53±0.22 vs. 0.83±0.50 respectively, P<0.05; Fig. 3B).

Plasma FFA levels in SIIR rats following 200 mg/kg AC. At 60 min, plasma FFA levels in the EG had decreased from 1.33±0.24 to 0.58±0.35 mmol/l, and were significantly decreased (15.69±3.97%) compared with in the CG, in which plasma FFA levels were increased from 1.25±0.19 to 1.30±0.09 mmol/l (5.5±0.80%). Additionally, the area under the curve of plasma FFA in the EG (1.01±0.15) was significantly smaller than that of the CG (1.16±0.25, P<0.05; Fig. 4).

Expression of insulin signal proteins following AC administration. The levels of insulin signal proteins, IRS-1, GLUT-4, and PI3K, were significantly elevated compared to the CG following AC administration. The fold changes (EG/CG) were 1.71, 2.10 and 1.87, respectively (Fig. 5).

Discussion

Metabolic syndrome has gained growing attention in recent years, particularly in developed countries. Overeating and an aging society contributes to the increasing incidence of metabolic syndrome, and insulin resistance may be a major factor that contributes to the development of type II DM (29). Therefore, preventing metabolic syndrome or correcting insulin resistance may help to slow the development of type II DM. There are many oral hypoglycemic agents and insulin sensitizers like TZDs available, but the side effects of these drugs limit their use (30). An agent with fewer side effects that can manage of plasma glucose levels is desirable for patients, particularly one that can also improve insulin resistance.

AC is a valuable medicinal fungus found in the forests of Taiwan. It grows naturally in the hollow trunk of the old Cinnamomum kanehirai tree or through artificial culture on host wood sections. Due to its versatile biomedical activities, AC is highly valued (12-16) and Taiwanese forests are threatened with the illegal logging of Cinnamomum kanehirai trees. This study developed a solid-state culture method for AC product to combat this serious illegal logging problem. Additionally, a standard operating procedure was developed for the AC solid-state culture method, to verify the quality of each batch (provided by Chair Professor Wai-Jane Ho from Da-Yeh University; Fig. 1A). The possible toxicity or potential side effects of the AC mycelium power may be relative to the dose, but in the present study, 100% of the rats survived following administration of the highest dose (500 mg/kg). Physiological signs including respiration, hair color and food and water intake, were all normal after 1 week of AC administration. As metabolic syndrome is a chronic condition requiring long-term treatment, the potential toxic effects of AC are very important and should be investigated. However, acute and subacute toxicity tests were not the aim of the present study.

The AC powder methanol extract was analyzed using HPLC to verify the presence of active hypoglycemic components, EK100, TR1+TR2 and TR3+TR4 (Fig. 1B). These components were previously reported to be bioactive and present in AC (18,20,31). There is concern that the solid-state culture method may result in variation of the components between batches and different strengths of bioactivity, but this HPLC analysis method allows for the quality control of each batch. Prior to the study, experimental doses were determined in preliminary tests by administering various doses to a SIIR rat once a week to establish a small, medium, and high dose (100, 200, and 500 mg/kg) appropriate for rats. Thus, the doses used are different than what would be appropriate in a clinical situation.

The hypoglycemic effect of AC was subsequently evaluated and supported by the findings of the present study. Prior to the study, experimental doses were determined in preliminary tests by administering various doses to a SIIR rat once a week to establish a small, medium, and high dose (100, 200, and 500 mg/kg) appropriate for rats. Thus, the doses used are different than what would be appropriate in a clinical situation.

The hypoglycemic effect of AC was subsequently evaluated and supported by the findings of the present study. The optimal oral dose to achieve a hypoglycemic effect was determined to be 200 mg/kg AC, and this was used to explore the mechanism of action of AC.

Previous research has confirmed the hypolipidemic effect of AC (17,20,31,32), supporting the results of the present study that indicated a decrease in plasma FFA levels. To establish the SIIR animal model, steroids were administered, causing an increase in plasma FFA levels and the development of insulin
resistance (11,25). AC improved insulin resistance by lowering plasma FFA levels in the SIIR state. Steroids are frequently used to treat inflammatory diseases and the impairment of insulin sensitivity is a problematic side effect, particularly in patients with type II DM. The results of this study demonstrate that AC may improve insulin resistance caused by the steroid administration.

The increase in plasma insulin in the EG was not significant at the 60 min time-point compared with the CG, which might have been due to the duration of plasma insulin secretion within this studied animal model due to oral administration of the AC powder. However, HOMA-IR at the 60 min time-point indicated an improvement in insulin resistance following AC treatment; enhanced insulin secretion induced by AC was recently reported in MIN6 cells (33). Therefore, this AC may elevate plasma insulin and improve insulin resistance.

With respect to signal transduction proteins, AC caused an increase in IRS-1, PI3K and GLUT-4. Previous studies have indicated that AC stimulates AMPK to enhance GLUT-4 translocation and activates the peroxisome proliferator activated receptor α (PPARα) to decrease plasma FFA levels, which may complement the insulin signaling pathway to result in a hypoglycemic effect and improvement in insulin resistance (18,20,32,34). In our previous studies, we also used the SIIR animal model to test the hypoglycemic effect of the Xylaria nigripes (Xn) fungus and the Gardenia jasminoides (GJ) plant (26,35). However, Xn was found to exert a serotonin-associated hypoglycemic effect, and PPAR activation was key to the hypoglycemic effect of GJ. This differs from the mechanism of AC identified in this study, in which a decrease in plasma FFA levels was observed.

Administration of 200 mg/kg AC to SIIR rats resulted in a decrease in plasma glucose levels, which was closely associated with a decrease in plasma FFA levels. Furthermore, an increase in the expression of insulin signaling proteins (GLUT-4, IRS-1 and PI3K) was observed with improved insulin resistance. These results indicate that AC acts as an insulin sensitizer in insulin resistant animals. Due to the use of an animal model in this study, results cannot be applied to a clinical situation. Thus, a randomized controlled trial of AC mycelium powder should be performed to determine the clinical dosage and enhance its effect on insulin resistance.

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