Research Article

Antidiabetic Effect of *Morinda citrifolia* (Noni) Fermented by Cheonggukjang in KK-Ay Diabetic Mice

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Antidiabetic effects of *Morinda citrifolia* (aka Noni) fermented by Cheonggukjang (fast-fermented soybean paste) were evaluated using a T2DM (type 2 diabetes mellitus) murine model. Six-week-old KK-Ay/TaJcl mice were randomly divided into four groups: (1) the diabetic control (DC) group, provided with a normal mouse diet; (2) the positive control (PC) group, provided with a functional health food diet; (3) the *M. citrifolia* (MC) group, provided with an MC-based diet; (4) the fermented *M. citrifolia* (FMC) group, provided with an FMC-based diet. Over a testing period of 90 days, food and water intake decreased significantly in the FMC and PC groups compared with the DC group. Blood glucose levels in the FMC group were 211.60–252.20 mg/dL after 90 days, while those in the control group were over 400 mg/dL after 20 days. In addition, FMC supplementation reduced glycosylated hemoglobin (HbA1c) levels, enhanced insulin sensitivity, and significantly decreased serum triglycerides and low-density lipoprotein (LDL) cholesterol. Furthermore, a fermented *M. citrifolia* 70% ethanolic extract (FMCE) activated peroxisome proliferator-activated receptor-γ (PPAR-γ) and stimulated glucose uptake via stimulation of AMP-activated protein kinase (AMPK) in cultured C2C12 cells. These results suggest that FMC can be employed as a functional health food for T2DM management.

1. Introduction

Diabetes mellitus (DM) is a chronic disorder characterized by hyperglycemia, with disturbances in carbohydrate, lipid, and lipoprotein metabolism stemming from defects in insulin secretion and insulin action, or both. About 90–95% of all cases of diabetes mellitus can be attributed to type 2 diabetes mellitus (T2DM, or adult-onset diabetes). T2DM is caused by the nonresponsiveness of cells to insulin rather than by defective insulin production. If not treated, T2DM can result in numerous severe complications, including coronary artery disease, peripheral vascular disease, and even death due to hyperglycemia [1]. T2DM currently affects more than 246 million people worldwide, and it is estimated that the number of patients with diabetes mellitus will increase to 360 million by 2030 [2]. Indeed, about 5–10% of the total health care budget has been used for treatment and management of T2DM in many countries [3].

Management of T2DM usually begins with a change in diet and exercise. Nonetheless, most patients ultimately require pharmacotherapy, such as injected insulin and/or oral antidiabetic drugs. Many oral antidiabetic drugs are available for the treatment and control of the symptoms of T2DM, such as sulfonylurea agents, biguanides (e.g., metformin), thiazolidinedione (TZD) drugs (also known as glitazones), α-glucosidase inhibitors, and glucagon-like peptide-1 (GLP-1) inhibitors. However, these drugs can cause serious adverse effects, among them hypoglycemia, hepatic toxicity, weight increase, physcosia (enlargement of the abdomen), and lactic acidosis [4]. Thus, there is an increasing demand for natural products (and in particular, nutraceuticals) that have fewer adverse effects and higher antidiabetic efficacy. Many studies have been conducted on natural products that efficiently and safely reduce blood glucose levels in diabetic patients, and several diabetes mellitus hundred plants have reported beneficial effects. For example, *Lagerstroemia speciosa* [5], *Gymnestis sylvestre* [6], *Mori folium* [7], and *Psidiium guajava* [8] are currently sold commercially as complementary and alternative medicinal materials for the treatment of diabetes.
Morinda citrifolia, also called Noni, is a tropical plant that grows widely throughout the Pacific. *M. citrifolia* has traditionally been used for the prevention, and treatment of arthritis, circulatory weakness, diabetes, cancer, and skin inflammation [9–12], and is widely known as a health food and a dietary supplement. A growing number of pharmacological studies on *M. citrifolia* have been published in recent years, but many of the reports are only accessible as congress abstracts. In addition, few peer-reviewed research manuscripts are available that discuss the antidiabetic activity of *M. citrifolia*, even though *M. citrifolia* has customarily used to treat diabetes. In fact, only a single study has presented evidence, limited at best, for the antidiabetic effect of *M. citrifolia* fruit juice in an animal model of T2DM [13]. The present study therefore focused on the antidiabetic properties of *M. citrifolia* and fermented *M. citrifolia* (FMC) in KK-Ay obese/diabetic mice, which are afflicted with a genetically induced form of diabetes that manifests itself as hyperinsulinemia, a result of insulin resistance [14]. *M. citrifolia* was fermented using Cheonggukjang, a traditional fast-fermented Korean soybean paste that was itself fermented using various Bacillus spp. After demonstrating the efficacy of FMC, we then aimed to elucidate the antidiabetic mechanisms of FMC.

### 2. Materials and Methods

#### 2.1. Preparation of FMC

Fifteen kilogram of dried *M. citrifolia* (MC) fruitpowder (Bobsaewoo Seoul, Korea) was heated in a hot kneaster (300–500 L, Daehan Food Machine, Inc., Gimpo, South Korea) at 82.5 ± 2.5°C for 4 h. *Cheonggukjang* (1%, employed to initiate fermentation) containing soybeans, Bacillus sp. (KCTC 11351BP), Bacillus subtilis (KCTC 11352BP), Bacillus sonolensis (KCTC 11354BP), and Bacillus circulans (KCTC 11355BP), were added to water or 70% ethanol (10X the sample amount w/v) with stirring for 24 h at 25°C. The extracts were centrifuged for 10 min at 3000 rpm, and the supernatants were filtered through Whatman no. 5 filter paper. The filtrates were evaporated by a rotary evaporator (RE200; Yamato Co., Tokyo, Japan) under reduced pressure at a temperature lower than 40°C. The resulting concentrates were freeze-dried and stored at −20°C prior to experimentation.

#### 2.2. Extraction of FMC

The dried and milled MC and FMC were added to water or 70% ethanol (10X the sample amount w/v) with stirring for 24 h at 25°C. The extracts were centrifuged for 10 min at 3000 rpm, and the supernatants were filtered through Whatman no. 5 filter paper. The filtrates were evaporated by a rotary evaporator (RE200; Yamato Co., Tokyo, Japan) under reduced pressure at a temperature lower than 40°C. The resulting concentrates were freeze-dried and stored at −20°C prior to experimentation.

#### 2.3. Animals, Diets, and Treatments

Twenty-eight KK-Ay/Tajcl mice (5 weeks old, male) with genetic obese type II diabetes were purchased from Clea Japan Inc. (Tokyo, Japan). After acclimatizing for 1 week, the mice were divided into four groups of seven animals each, as follows: diabetic control (DC group), banaba-fed animals (positive control (PC) group), MC-fed animals (MC group), and FMC-fed animals (FMC group). Banaba (*Lagerstroemia speciosa*) leaves are commercially sold as a functional health food intended for the regulation of blood glucose levels. Diabetic control animals were freely fed on a normal mouse diet (AIN-93G food). PC, MC, and FMC groups were freely fed on AIN-93G blended with freeze-dried banaba, MC, or FMC powder (0.4%). All mice were maintained in the same room under conventional conditions with a regular 12 h light/dark cycle and temperature and relative humidity maintained at 23 ± 2°C and 50 ± 5%, respectively. At the end of the experimental period (90 days), the animals were fasted overnight, anaesthetized with an intraperitoneal injection of zoletil 50 (30 mg/kg), and sacrificed for the procurement of tissue samples. The study was approved by the Ethics Committee of the Experimental Animal Research Laboratory, Korea Food Research Institute (approval code, KFRI-M-10020).

#### 2.4. Analysis of Blood Glucose Levels

Whole blood was sampled from the tail vein of mice without fasting at a fixed time
every 10 days over a period of 90 days, and the blood sample was analyzed using the glucose oxidase method, which utilizes a blood glucose sensor strip (ACCU-CHEK Sensor, Roche, Basel, Switzerland).

2.5. Analysis of Glycosylated Hemoglobin (HbA1c) Levels. At the end of the 90-day experimental period, mice were subjected to fasting overnight. Next, a fasting whole blood sample (4 μL) was obtained via retro-orbital sinus puncture. The blood sample was injected into an A1c test cartridge and then analyzed by using a Hemoglobin A1c Testing Analyzer (Asan Pharmaceutical, Seoul, Korea).

2.6. Serum Analysis. After 90 days of dietary treatment, KK-Ay/Talcl mice were sacrificed, and fasting blood samples were collected and centrifuged at 3000 rpm for 10 min to obtain serum samples for serum chemical analysis. Triglyceride, total cholesterol, high-density lipoprotein (HDL)-cholesterol, and low-density lipoprotein (LDL) cholesterol were measured by an ADVIA Automated Hematology Analyzer (ADVIA 1650, Bayer, USA). Fasting serum glucose levels were assessed using a glucometer, which employed the glucose oxidase/peroxidase reaction. This reaction allows the colorimetric detection of H2O2 for the estimation of free glucose in the serum. Fasting serum insulin levels were measured with a murine insulin TMB (tetramethyl benzidine) enzyme-linked immunosorbent assay (ELISA) kit (Shibayagi, Gunma, Japan). A homeostasis model assessment of insulin resistance (HOMA-IR), an index of insulin resistance, was calculated using fasting insulin and glucose concentrations according to the following equation:

\[
\text{HOMA-IR} = \frac{\text{fasting insulin (μIU/mL) \times fasting glucose (nmol/L)}}{22.5}
\]

2.7. Muscle Differentiation and Glucose Uptake Assay. C2C12 cells were grown in Dulbecco's modified eagle medium (DMEM) supplemented with 10% fetal bovine serum (FBS), penicillin (120 unit/mL), and streptomycin (75 μg/mL) in a 5% CO2 atmosphere. Differentiation medium (DMEM medium with 1% horse serum) was added to confluent C2C12 cells for 4 days. The cells were then incubated overnight in low-glucose, serum-free medium followed by 24-h treatment in the presence of 2-[N-(7-nitrobenz-2-oxa-1,3-diazol-4-yl) amino] -2-deoxy-d-glucose (2-NBDG). The fluorescence was read at excitation and emission wavelengths of 485 and 535 nm.

2.8. Western Blot Analysis. C2C12-derived myotubes were harvested with lysis buffer (50 mM Tris HCl, 1% Triton X-100, 0.5% sodium deoxycholate, 150 mM NaCl, 1 mM EDTA, 1 mM PMSF, 1 mM sodium orthovanadate, 1 mM NaF and 0.2% protease inhibitor cocktail; pH 7.2). Proteins (40 μg) were subjected to sodium dodecyl sulfate polyacrylamide gel electrophoresis (SDS-PAGE) followed by transfer onto a nitrocellulose membrane. Protein phosphorylation or expression was detected by Western blot analysis using primary antibodies specific for phospho-AMP protein kinase (AMPK) and β-actin (Cell Signaling Technology, Beverly, MA).

2.9. Peroxisome Proliferator-Activated Receptor (PPAR)-γ Transcriptional Activity Assay. Plasmids for PPAR-γ or retinoid X receptor (RXR)-α, as well as luciferase reporter vectors containing PPAR-response elements (PPREs), were the generous gift of Dr. Jae Bum Kim (Seoul National University, South Korea). For the PPAR-γ assay, human embryonic kidney (HEK) 293 cells were cotransfected with PPAR-γ or RXR-α plasmids and PPAR-PPRE luciferase reporter vectors by using a transfection reagent (Qiagen, Valencia, CA) for 18 h. Cells were then treated with fermented Morinda citrifolia ethanol extract (FMCE) for 24 h, and PPAR-γ activity was measured via a Luciferase Assay System (Promega, Madison, WI).

2.10. Statistical Analysis. Numerical data are expressed as means ± the standard deviation using SAS 9.1 software. The significance of differences between experimental conditions was analyzed using one-way analysis of variance (ANOVA) followed by Duncan's multiple range test. Values of \( P < 0.05 \) were considered significant.

3. Results

3.1. Blood Glucose and HbA1c Levels. Figure 2 shows non-fasting blood glucose levels measured every ten days over the experimental period of 90 days. Blood glucose levels were remarkably increased in the DC group compared with the PC (banaba fed) group after 20 days, and were maintained at more than 400 mg/dL throughout the remaining 70 days of the experiment. Blood glucose levels in the MC group also increased to the same extent as that observed for the DC group during the first 40 days of dietary evaluation. After 40 days, the levels gradually decreased to about 300 mg/dL at 70 days and were then maintained at this level for the next 20 days. In contrast, blood glucose levels did not increase in the FMC group relative to the PC group, and the levels were maintained at 211.60–252.20 mg/dL during the experimental period. The blood glucose levels in the FMC group were actually lower than those in the PC group.

Figure 3 shows HbA1c levels at 90 days. The mean Hb1Ac level in the FMC group was 7.62%, which was significantly lower than the Hb1Ac levels in the other three groups. The corresponding values were 9.45% in the DC group, 8.96% in the PC group, and 8.96 in the MC group.

3.2. Serum Glucose and Insulin Levels. Serum glucose (Figure 4) and insulin (Figure 5) levels were significantly decreased in the PC, MC, and FMC groups at 90 days relative to the DC group. The FMC group in particular showed decreased serum glucose and insulin levels by about 36% and 45%, respectively (Figures 4 and 5). The HOMA-IR index for
3.3. Serum Lipid Profiles. Triglyceride levels were significantly decreased in the MC (169.00 mg/dL) and FMC (148.40 mg/dL) groups compared with the DC (236.60 mg/dL) and PC (236.60 mg/dL) groups (Table 1). Triglyceride levels in the MC and FMC groups decreased by about 29% and 37%, respectively, relative to the DC group. No significant differences were observed in total cholesterol between the groups; however, HDL-cholesterol levels in the MC and FMC groups were significantly higher than those observed in the DC group, whereas LDL-cholesterol levels in the FMC group were significantly lower than those observed in the other three groups (Table 1).

3.4. Glucose Uptake. To establish the potential antidiabetic mechanisms of FMC, we evaluated the effect of an ethanol...
Table 1: Effect of FMC on total serum cholesterol, HDL-cholesterol, LDL-cholesterol and triglyceride levels (mg/dL) in KK-Ay mice after 90 days.

<table>
<thead>
<tr>
<th></th>
<th>Total cholesterol</th>
<th>HDL-cholesterol</th>
<th>LDL-Cholesterol</th>
<th>Triglycerides</th>
</tr>
</thead>
<tbody>
<tr>
<td>Diabetic control (DC)</td>
<td>141.25 ± 16.69</td>
<td>116.00 ± 6.12b</td>
<td>9.60 ± 2.06a</td>
<td>235.60 ± 46.73a</td>
</tr>
<tr>
<td>Positive control (PC)</td>
<td>144.80 ± 12.54</td>
<td>119.33 ± 9.48ab</td>
<td>9.75 ± 3.90a</td>
<td>221.50 ± 31.05ab</td>
</tr>
<tr>
<td>MC</td>
<td>137.80 ± 13.53</td>
<td>133.40 ± 11.20a</td>
<td>9.40 ± 2.15b</td>
<td>169.00 ± 17.02b</td>
</tr>
<tr>
<td>FMC</td>
<td>137.00 ± 12.47</td>
<td>130.40 ± 10.13a</td>
<td>7.80 ± 1.17b</td>
<td>148.40 ± 17.07b</td>
</tr>
</tbody>
</table>

Each value represents the mean ± SD (n = 7).

Means with different superscripts in the same column are significantly different from each other (P < 0.05, Duncan’s multiple range test).

3.5. PPAR-γ Agonist Activity. As shown in Figure 8, FMC stimulated the transcriptional activity of PPAR-γ in a dose-dependent manner as assessed by using a luciferase assay, but FMCW, MCW and MCE did not. The PPAR-γ agonist activity of FMC was comparable or even better than that of rosiglitazone, a known commercial PPAR-γ agonist.

3.6. Phosphorylation of AMPK. To examine whether FMCE can increase the phosphorylation of AMPK, western blotting analysis was performed. As shown in Figure 9, FMCE markedly stimulated the phosphorylation of AMPK in C2C12-derived myotubes in a dose-dependent manner. In contrast, FMCW did not induce AMPK phosphorylation (data not shown).

4. Discussion

The results of this study clearly indicate that the antidiabetic effects of FMC were stronger than those of the commercially available, functional health food, banaba. For example, blood glucose levels are an important indicator of diabetic status, and an FMC-based diet was more effective in reducing blood (Figure 2) and serum (Figure 4) glucose levels in KK-Ay mice than a normal diet. As reported in previous findings [13], MC also gradually reduced blood glucose levels after 40 days of experimentation; however, FMC was far more efficient in this regard compared with MC. It can be presumed that the antidiabetic activity of M. citrifolia was markedly improved through solid substrate fermentation, a bioconversion technique employed to increase the function of natural products. This result would be in agreement with recent studies [15, 16] demonstrating that the pharmacological effects of medicinal plants are increased through solid substrate fermentation.
Glycosylated hemoglobin is considered to be the gold-standard indicator for the accurate and reliable measurement of fasting glucose. Fasting glucose levels are strongly associated with the extent of ambient glycemia during the 3-month period prior to fasting, as well as with the degree of protein glycation. Parallelizing the results for blood glucose levels, the amount of glycosylated hemoglobin in the FMC group was lower than that in the other test groups (Figure 3). Hence, FMC can assist in the maintenance of low blood glucose levels and the control of blood glucose over the long term.

This hypoglycemic action of FMC was associated with a decrease in both insulin content (Figure 5) and insulin resistance (Figure 6). In the control group, insulin content in addition to blood glucose levels was high relative to the other groups, particularly the FMC group. This is indicative of hyperinsulinemia and hyperglycemia caused by insulin resistance in DC mice. Notably, the HOMA-IR index (an index of insulin resistance) was significantly decreased in the FMC group compared with the DC group. The HOMA-IR index provides a simple method to measure insulin sensitivity, and its use is widespread in both clinical and animal studies [17].

Insulin resistance, or the failure of normal insulin-induced actions in major tissues including the liver, fat, and skeletal muscle, plays a key role in the pathophysiology of T2DM. A primary feature of insulin resistance is the reduction of glucose uptake into muscle cells. We therefore used C2C12 skeletal muscle cells to evaluate whether FMC increases glucose uptake. Our results indicate that FMCE stimulated uptake of 2-NBDG into terminally differentiated C2C12-derived myotubes in a dose-dependent manner (Figure 7). Thus, stimulation of glucose uptake may be one mechanism by which FMC improved insulin resistance in KK-Ay mice. MCW and MCE also increased glucose uptake into muscle cells; however, the effects of these extracts were lower than the effect of FMC. These results indicate that the glucose disposal rate mediated by MC in muscle cells was increased by solid substrate fermentation.

Recently, many researchers have reported that various plant-derived flavonoids, anthraquinones, and terpenes stimulate glucose uptake in cells [18–20], reducing insulin resistance. In addition, insulin resistance has been reported to be associated with inflammation and oxidative stress [21]. Consequently, compounds such as flavonoids and lignans, which possess powerful antiinflammatory and antioxidant properties, may be useful in the control of T2DM. It has been reported that M. citrifolia contains numerous anthraquinones, flavonol glycosides and terpenoids, which possess antioxidant, antiinflammatory and antidyslipidemic effects [22–24]. Therefore, these compounds were likely increased in FMC by fermentation of MC, leading to at least some of the antidiabetic actions of FMC. However, further detailed study will be required to identify antidiabetic compounds in FMC.

AMPK functions as a sensor of cellular energy that can be activated by glucose deprivation and high AMP-to-ATP ratios [25]. There are indications that the activation of this enzyme is beneficial for the treatment and prevention of T2DM and metabolic syndrome [26, 27], because its phosphorylation leads to increased glucose uptake and lipid oxidation in muscle cells [28, 29]. To investigate whether the effect of FMC on glucose uptake was mediated through AMPK activation, C2C12-derived myotubes were treated with FCME at various concentrations. As shown in Figure 9, FMCE dose-dependently increased the phosphorylation of AMPK. The actions of FMCE were similar to those
of AICAR (5-amino-4-imidazolcarboxamide riboside), an AMPK-specific activator. These data are in agreement with recent reports by Lee et al. [30] and Benhaddou-Andaloussi et al. [31], which demonstrated an increase in cellular glucose uptake as a result of AMPK activation.

PPAR-γ is a transcription factor that regulates gene expression in the liver, adipose tissue, vascular endothelium, and muscle [32]. PPAR-γ can improve insulin sensitivity and glucose tolerance via the regulation of lipid storage, glucose homeostasis, and adipokine production [33, 34]. We found that FMCE functioned as a PPAR-γ agonist to increase PPAR-γ-dependent luciferase activity in vitro. This suggests that FMC acts on PPAR-γ as well as AMPK to augment glucose uptake. FMCE (400 μg/mL) showed enhanced PPAR-γ agonist relative to the TZD drug rosiglitazone (20 μM). TZDs are widely used as PPAR-γ agonists in the treatment of diabetes, and TZD-mediated activation of PPAR-γ results in marked improvement in blood glucose levels and insulin sensitivity [35]. PPAR-γ agonists also significantly decrease total cholesterol and triglyceride in type 2 diabetic patients and model animals [36, 37]. The current results indicate that FMC can now be considered a new type of PPAR-γ agonists.

Hypertriglyceridemia and hypercholesterolemia are central components of diabetic dyslipidemia, occurring in 20–60% of all diabetic patients. Dyslipidemia characterized by high plasma levels of total cholesterol and triglycerides, in addition to low plasma levels of HDL cholesterol, is among the many risk factors for atherosclerotic cardiovascular disease. The latter is in turn a major cause of morbidity in T2DM patients [38]. In this study, dietary supplementation with FMC caused a significant decrease in serum triglycerides and LDL-cholesterol and a concomitant increase in HDL-cholesterol. This improvement of the lipid profile may be attributed to the PPAR-γ agonist activity of FMC, at least in part.

In summary, the decrease in blood glucose level mediated by FMC was associated with a significant reduction in insulin resistance, as revealed by the plasma insulin content and blood glucose levels in vivo in a KK-Ay mouse model of T2DM. This enhanced insulin sensitivity was related to elevated glucose disposal rates in C2C12 myotubes via the activation of PPAR-γ and AMPK. In addition, FMC ameliorated dyslipidemia in KK-Ay mice, most likely by stimulating PPAR-γ. These pleiotropic antidiabetic actions provide strong support for the use of FMC for the treatment of diabetes. However, high-quality clinical studies are essential to determine the optimal conditions for complementary or alternative treatment in diabetic patients.

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References


