Anti-Obesity Effects of a Mixture of Fermented Ginseng, *Bifidobacterium longum* BORI, and *Lactobacillus paracasei* CH88 in High-Fat Diet-Fed Mice

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Introduction

Ginseng and probiotics have anti-obesity effects in mice fed a high-fat diet (HFD). Absorption of ginsenoside and colonization of probiotics occur in the intestine. In this study, a mixture of fermented ginseng and two probiotics, *Bifidobacterium longum* BORI and *Lactobacillus paracasei* CH88, was administered to HFD-fed mice for 9 weeks. The mixture significantly suppressed weight gain (*p* < 0.05, *n* = 8) and lipid deposition in the liver and adipose tissues as well as increased the mice’s food intake. The adipocyte size of the adipose tissue was significantly decreased in the mixture-fed group, especially when 0.5% fermented ginseng and 5 × 10⁸/ml of the two probiotics were used (*p* < 0.05, *n* = 10). The expression of TNF-α in adipose tissue was efficiently downregulated in the mixture-fed group (*p* < 0.05, *n* = 4). The supplement also improved the mice’s fasting blood glucose levels (*p* < 0.05, *n* = 8) and total cholesterol feces excretion (*p* < 0.05, *n* = 8). The mixture of fermented ginseng and *B. longum* BORI and *L. paracasei* CH88 could have an anti-obesity effect and suppress lipid deposit in the liver and adipose tissues.

**Keywords:** Ginseng, lactobacteria, bifidobacteria, anti-obesity, lipid metabolism
composition of the intestinal microflora via attachment and colonization of the intestinal mucosa, resulting in healthful effects [10]. However, few studies have researched the mechanism of FG and probiotics when consumed together.

Therefore, the purpose of this study was to evaluate whether a FG and probiotics mixture had an anti-obesity effect on mice fed a HFD.

Materials and Methods

Fermented Ginseng and Probiotics

The FG and probiotics used in this study were in freeze-dried powder form and provided by Bifido Co., Ltd (Korea). The probiotics mix (PM) was prepared by mixing Bifidobacterium longum BORI and Lactobacillus paracasei CH88 at a ratio of 1:1.

Ginsenoside Analysis of Fermented Ginseng

To determine the ginsenoside content, the FG was ultrasonically extracted with 80% methanol for 1 h. The pre-treated samples were analyzed using an Ultimate 3000 HPLC system (Thermo Dionex, USA) and 210 nm UV detector. A VSD C-18 column (250 × 4.6 mm, 5 μm) was used for all separations. The mobile phase solvent consisted of distilled water (A) and acetonitrile (B). The gradient elution conditions were as follows: 0–50 min (5% B), 50–56 min (95% B), and 56–60 min (5% B). The flow rate and temperature were set to 1 ml/min and 20°C, respectively. Ginsenosides Rb1, Rb2, Rd, Rg1, and F1 were purchased from Biotech (China). Ginsenosides Rc, Re, Rg2, Rg3, Rh1, Rh2, and cK were obtained from Cogon Biotech (China).

Animals and Diets

In our study, male ICR mice (7 weeks old) were purchased from Central Lab. Animal (Korea). The animal breeding environment was adjusted to a cycle of 12 h light/12 h dark at a temperature of 23 ± 1°C and a humidity of 40–60%. After a 1-week adaptation period, the mice were randomly divided into the following five groups (n = 8): LFD (Low-fat diet; 10% fat of total calories), HFD (High-fat diet; 60% fat of total calories), or HFD supplemented with FG and PM (FG+PM) divided into LOW (0.25% FG + 2.5 × 10⁶ CFU/ml PM), MID (0.5% FG + 5 × 10⁶ CFU/ml PM), and HIGH (1.0% FG + 1 × 10⁷ CFU/ml PM).

The mice were fed the experimental diets for 9 weeks, following which they were anesthetized with Zoletil (Virbac Lab., France) and Rompun (Bayer, Germany) after fasting for 12 h. Cardiac blood, liver tissue, epididymal fat, and mesenteric fat were removed from the specimens and weighed. All procedures were approved by the Institutional Animal Care and Use Committee (IACUC; SNU-161108-7) (Seoul National University, Korea).

Histological Analysis

Morphological analysis of the liver and epididymal fat was performed to measure the hepatic fat accumulation and adipocyte size in the adipose tissue (AT). The liver and epididymal fat were fixed in 10% formalin (Wako Pure Chemical Industries, Japan) for 1 day and then stored in paraffin. Sections were then cut, stained with hematoxylin and eosin, and examined with an optical microscope.

Biochemical Analysis

The plasma triacylglycerol (TAG), total cholesterol (TC), and high-density lipoprotein cholesterol (LDL-C) were analyzed using a kit purchased from Asanpharm (Korea). The low-density lipoprotein cholesterol (HDL-C) was calculated by subtracting LDL-C from TC. The fasting blood glucose (FBG) was measured after 12 h of fasting in the 9th week. The plasma insulin was analyzed using a mouse insulin ELISA kit (AKRIN-011T; Shibayagi, Japan).

Hepatic Lipid Analysis

According to Li et al. [5], we used a modified Folch method [11] for the liver lipid analysis. The liver lipid was measured using a TAG test kit (Asanpharm, Korea).

RNA Extraction and Quantitative Real-Time Reverse Transcriptase Polymerase Chain Reaction (RT-PCR)

The total RNAs of the liver and AT were extracted using a TaKaRa MiniBEST Universal RNA Extraction kit (TaKaRa Bio, Japan) and RNeasy Lipid Tissue Mini Kit (Qiagen, Germany), respectively. The RNA was reverse transcribed to cDNA using a cDNA kit from TaKaRa Bio. mRNA expression analysis of the cDNA and SYBR premix Ex Taq (TaKaRa Bio, Japan) was performed with a StepOne real-time PCR system. The analysis conditions were denaturation at 95°C for 30 sec, followed by 40 cycles of 95°C for 5 sec and 60°C for 33 sec. The data were normalized for the housekeeping gene GAPDH and compared with the expression levels of the HFD group.

TC in Feces

The collected feces was ground into a powder and the subsequent process was the same as for the hepatic lipid analysis. The TC in feces was measured using a TC test kit (Asanpharm, Korea).

Statistical Analysis

The results were analyzed using the SPSS statistical package ANOVA test. The statistical differences were examined via Duncan’s multiple range test. The statistical significance was set at p < 0.05.

Results

Effect of FG+PM on Body Weight and Food Intake

The main ginsenosides of the experimental FG were protopanaxadiol-type Rd (19.1%), Rc (4.5%), and protopanaxatriol-type Re (3.3%). The others were Rb1
(1.4%), Rb2 (0.7%), Rg1 (0.9%), Rg2 (0.7%), and Rg3 (1.0%).

Table 1 shows the changes in mouse body weight and food consumption. The mice’s weight at the 9th week was significantly increased in the HFD group compared with the LFD group. Compared with the control group, the MID group showed a significant decrease in body weight and weight gain.

The HFD group had a significant increased food intake compared with the LFD group, and the experimental groups consumed more calories than did the control. The food efficiency of the MID group was significantly lower than that of the HFD group.

Effect of FG+PM on the Weights of the Liver, Epididymal Adipose Fat, and Mesenteric Adipose Fat

Table 1 shows the liver and the epididymal and mesenteric AT (EAT and MAT) weights. The liver weight was significantly decreased in the MID group compared with the control group. The EAT and MAT weights were significantly increased in the HFD group compared with the LFD group.

Effect of FG+PM on the Fasting Blood Glucose and Insulin Levels

The HFD group had a higher FBG level than that of the LFD group. The FBG levels of the experimental groups were lower than those of the HFD group, and those of the MID group were significantly lower (Fig. 1A). Although the MID group had the lowest insulin level, there was no significant difference (Fig. 1B).

Effect of FG+PM on the Plasma Lipid Profiles

As shown in Table 2, the mice fed the FG+PM had similar plasma TAG levels. Although the plasma TC and LDL-C

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Table 1. Effects of fermented ginseng plus probiotic mixtures on weight parameters of mice.

<table>
<thead>
<tr>
<th></th>
<th>LFD</th>
<th>HFD</th>
<th>LOW</th>
<th>MID</th>
<th>HIGH</th>
</tr>
</thead>
<tbody>
<tr>
<td>Initial BW (g)</td>
<td>38.6 ± 1.1</td>
<td>38.9 ± 1.4</td>
<td>37.4 ± 0.7</td>
<td>38.2 ± 1.9</td>
<td>37.6 ± 1.9</td>
</tr>
<tr>
<td>Final BW (g)</td>
<td>35.2 ± 3.5</td>
<td>67.2 ± 6.2</td>
<td>65.1 ± 8.9</td>
<td>58.0 ± 8.6</td>
<td>65.0 ± 7.1a</td>
</tr>
<tr>
<td>BW gain (g)</td>
<td>13.5 ± 4.4</td>
<td>28.4 ± 5.6</td>
<td>27.8 ± 8.5</td>
<td>19.8 ± 8.1</td>
<td>27.4 ± 7.2a</td>
</tr>
<tr>
<td>Food intake (kcal)</td>
<td>3,589.0 ± 29.3a</td>
<td>4,641.0 ± 1,150.5b</td>
<td>5,671.2 ± 179.3bc</td>
<td>6,021.8 ± 493.3bc</td>
<td>5,879.9 ± 1,260.2bc</td>
</tr>
<tr>
<td>Food efficiency</td>
<td>0.0038 ± 0.001a</td>
<td>0.0064 ± 0.002b</td>
<td>0.0049 ± 0.002ab</td>
<td>0.0034 ± 0.002bc</td>
<td>0.0048 ± 0.002bc</td>
</tr>
<tr>
<td>Liver</td>
<td>1.9 ± 0.24a</td>
<td>2.9 ± 0.77b</td>
<td>2.4 ± 0.66ab</td>
<td>2.2 ± 0.75b</td>
<td>2.5 ± 0.44ab</td>
</tr>
<tr>
<td>EAT</td>
<td>2.0 ± 0.65a</td>
<td>3.1 ± 0.60bc</td>
<td>3.3 ± 0.90b</td>
<td>2.5 ± 0.88bc</td>
<td>3.0 ± 0.52bc</td>
</tr>
<tr>
<td>MAT</td>
<td>0.75 ± 0.26a</td>
<td>1.59 ± 0.38ab</td>
<td>1.34 ± 0.68b</td>
<td>1.1 ± 0.43bc</td>
<td>1.6 ± 0.48bc</td>
</tr>
</tbody>
</table>

1Food efficiency = food intake (kcal)/BW gain (g).
2EAT, Epididymal adipose tissue.
3MAT, Mesenteric adipose tissue.
abcd Means in the same row not sharing a common letter are significantly different at p < 0.05 (n = 8).

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Fig. 1. Effects of fermented ginseng plus probiotic mixtures on fasting blood glucose (FBG) and insulin levels.

(A) Plasma FBG levels (n = 8). (B) Plasma insulin levels (n = 7). abcd Means not sharing a common letter are significantly different at p < 0.05.

levels of the LOW group were significantly lower than those of the HFD group, the others were significantly higher. The plasma HDL-C levels of the experiment groups were lower than those of the HFD group; however, there

### Table 2. Effects on FG+PM on the plasma lipid profile.

<table>
<thead>
<tr>
<th></th>
<th>LFD</th>
<th>HFD</th>
<th>LOW</th>
<th>MID</th>
<th>HIGH</th>
</tr>
</thead>
<tbody>
<tr>
<td>TAG (mg/dl)</td>
<td>27.3 ± 9.7</td>
<td>19.4 ± 9.8</td>
<td>24.5 ± 10.8</td>
<td>20.2 ± 4.3</td>
<td>22.2 ± 8.5</td>
</tr>
<tr>
<td>TC (mg/dl)</td>
<td>125.5 ± 9.4</td>
<td>109.2 ± 9.3</td>
<td>91.2 ± 11.4</td>
<td>138.3 ± 18.7</td>
<td>123.5 ± 6.3</td>
</tr>
<tr>
<td>LDL-C (mg/dl)</td>
<td>75.6 ± 7.3</td>
<td>62.6 ± 6.6</td>
<td>51.0 ± 7.1</td>
<td>98.4 ± 13.6</td>
<td>82.5 ± 9.0</td>
</tr>
<tr>
<td>HDL-C (mg/dl)</td>
<td>49.8 ± 4.1</td>
<td>46.5 ± 8.3</td>
<td>40.2 ± 6.3</td>
<td>39.9 ± 12.8</td>
<td>41.0 ± 9.3</td>
</tr>
</tbody>
</table>

*abcd* Means in the same row not sharing a common letter are significantly different at *p* < 0.05 (*n* = 8).

**Fig. 2.** Effects of fermented ginseng plus probiotic mixtures on the histology of liver and adipose tissue.

(A) H&E staining of liver. (B) H&E staining of adipose tissue. (C) Adipocyte size of adipose tissue. *abcd* Means not sharing a common letter are significantly different at *p* < 0.05.
Effect of FG+PM on the Histology of the Liver and AT

The lipid accumulations in the liver and AT were determined by H&E staining. The HFD group had higher lipids than the LFD group, and the lowest lipid accumulation of the experimental groups was in the MID group (Fig. 2A). The adipocyte sizes in the EAT were confirmed (Fig. 2B). The HFD group had a significantly larger adipocyte size than that of the LFD group, and the adipocyte size of the MID and HIGH groups were significantly smaller than that of the HFD group. (Fig. 2C)

Effect of FG+PM on the Hepatic Lipid Profiles

The TAG content was lower in the experimental group than in the HFD group, and the lowest in the MID group (Fig. 3).

Effect of FG+PM on the mRNA Expression in Liver Tissue

Figs. 4A and 4B show the mRNA expression of the lipids and glucose metabolism in the liver. The expression of peroxisome proliferator-activated receptor-γ (PPAR-γ) was significantly higher in the HFD group than in the LFD group. The expression of LDL receptor (LDLR) was significantly higher in the LOW group than in the control group. The HFD group had a significantly higher expression of glucokinase (GK) than that of the LFD group.

Effect of FG+PM on the mRNA Expression in AT

Fig. 5A shows the mRNA expression of AT inflammation. The expression of tumor necrosis factor (TNF)-α was significantly higher in the HFD group than in the LFD group and significantly lower in the MID and HIGH groups than in the control group.

Fig. 5B shows the mRNA expression of AT lipid metabolism. The expression of PPAR-γ was increased in the MID group compared with the control; however, there was no significant difference. The expression of lipoprotein lipase (LPL) in the MID group was higher than that in the control group but not significant.

Effect of FG+PM on the TC in Feces

The TC content in the plasma was significantly higher in the HFD group than in the LFD group. The TC content in

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Fig. 3. Effects of fermented ginseng plus probiotic mixtures on the TAG content in the liver (n = 4).

*abc* Mean not sharing a common letter are significantly different at p < 0.05.

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Fig. 4. Effects of fermented ginseng plus probiotic mixtures on mRNA expression in liver.

(A) Effects on mRNA expression related to lipid metabolism (n = 4).
(B) Effects on mRNA expression related to glucose metabolism (n = 4).

*abc* Mean not sharing a common letter are significantly different at p < 0.05.
Anti-Obesity Effects of a Mixture of Fermented Ginseng, Bifidobacterium, and Lactobacillus

The feces was higher in all the FG+PM groups than in the control group, and the LOW and HIGH groups had a significantly higher TC feces content (Fig. 6).

Discussion

It has been reported that a HFD induces obesity and impairs leptin and insulin signaling in the hypothalamus, which results in excessive weight gain [12]. Liu et al. [13] reported that the ginsenosides Rd, Rb1, Rb2, and Rc are pancreatic lipase inhibitors that prevent obesity by increasing fat excretion. Similarly, probiotics have reduced fat accumulation and weight gain by competing with other probiotic strains in the intestines for nutrients and by reducing the gut microbial diversity [14]. The results of our study are similar to those found in earlier experimental studies. The intake of FG+PM reduced weight gain, especially in the MID group, and significantly suppressed weight gain despite increased food intake.

Obesity is associated with dyslipidemia, which increases TAG and free fatty acids (FFA) and decreases HDL-C. In previous studies, ginseng [15] and probiotics [16] have been reported to improve the lipid profile by inhibiting hyperlipidemia. In the present study, there was no significant difference between TAG and HDL-C. In contrast, the TC was decreased in the HFD groups and increased in the FG+PM groups, with the exception of the LOW group. The biosynthesis of cholesterol is regulated by HMG-CoA reductase (HMGCR). Therefore, most cholesterol in a HFD inhibits HMGCR feed-back and prevents cholesterol biosynthesis. In the present study, FG+PM seemed to promote cholesterol biosynthesis through the positive feedback of HMGCR. In fact, previous studies have shown that ginsenoside Rb1 improved cholesterol biosynthesis [17]. Although it stimulated cholesterol biosynthesis, FG+PM also promoted the excretion of cholesterol into the feces (Fig. 6). Cholesterol can be converted into coprostanol by probiotics and then excreted directly into the feces, which decreases the amount of absorbed cholesterol [18]. However, both the control and the experimental groups in this study were within the normal cholesterol range; thus, it was difficult to state that dyslipidemia actually occurred.

Obesity causes hyperglycemia by counteracting FBG homeostasis. Obesity is associated with insulin resistance, and elevated FBG changes lipogenesis and lipolysis. According to the results of this study, although the FBG level was increased in the HFD group, the FG+PM level was lower and effectively inhibited in the MID group. This is consistent with the results that ginseng [15] and probiotics [19] decreased blood sugar. In fact, ginsenoside Rd has been shown to enhance glucose utilization through the PI-3 kinase-dependent pathway [20], and both ginsenosides Re and Rc have been reported to significantly
increase glucose uptake in cells through AMPK activation [21].

Because the liver is one of the major organs responsible for lipid metabolism, damage to hepatic lipid metabolism leads to an abnormal accumulation of lipids, which is called hepatic steatosis. HFD-induced obesity increases FFA uptake into the liver, which often leads to hepatic fat accumulation resulting in non-alcoholic fatty liver disease (NAFLD) [22]. Ginseng and probiotics are well known as NAFLD-ameliorating agents. The results of this study show that the liver weight and hepatic TAG content were effectively suppressed in the FG+PM groups, especially in the MID group. Moreover, the smallest lipid droplet size was observed in the MID group (Fig. 2A).

Excessive energy intake results in the fat storage processes of hypertrophy and hyperplasia, which are controlled by PPAR-γ [23]. The expression of PPAR-γ was decreased by 50% in the MID group compared with the HFD group (p = 0.15). The results of the present study are similar to those found in earlier experimental studies on ginseng [24] and probiotics [25].

LDLR in the liver is known to take up LDL-C in the blood, which is involved in blood lipid clearance. LDLR expression was upregulated in the FG+PM groups and consistent with the decreased plasma LDL-C level in the LOW group. The results of our study concur with those found in a previous study [26]. Thus, FG+PM intake may have a role in decreasing the LDL-C concentration in the blood.

NAFLD is strongly associated with hepatic insulin resistance, type 2 diabetes, and obesity. Although insulin is a major regulator of hepatic lipogenesis, glucose is also known to contribute to the coordinated regulation of carbohydrate and fat metabolism in the liver [27]. The expression of G6P, which is involved in gluconeogenesis, was lower in the experimental groups. Compared with the control group, the expression of G6P in the FG+PM groups was lower, especially in the HIGH group by 90% (p = 0.1). In contrast, the expression of GK, a glycolytic enzyme, was significantly elevated in the HFD group compared with the LFD group, which has the potential to promote the long-term storage of carbohydrates and TAG.

Taken together, this FG+PM combination effectively inhibited hepatic weight gain, TAG content, and liver fat accumulation, especially in the MID group.

Obesity is associated with hypertrophy in AT, which can cause adipocytes to move away from blood capillaries and result in adipocyte hypoxia [28]. Dead adipocytes form crown-like structures (CLS), and most macrophages that infiltrate the AT are found around the CLS [29]. AT macrophages promote insulin resistance and AT inflammation [30]. As found in previous studies [31, 32], FG+PM significantly suppressed the adipocyte size. In addition, FG+PM significantly inhibited the expression of TNF-α in the FG+PM groups (Fig. 5A), suggesting that FG+PM can inhibit hypertrophy by preventing the excessive formation of CLS in AT. In fact, Kim [33] reported that ginsenoside Re inhibited the expression of TNF-α in 3T3-L1.

PPAR-γ is involved in the regulation of genes involved in lipid accumulation and insulin sensitivity [34]. PPAR-γ is also used as a major target in the treatment of insulin resistance with thiazolidinediones (TZDs), PPAR-γ activators. TZDs increase lipid accumulation in AT and decrease hepatic and muscular fat, ultimately inhibiting the fatty liver [35]. We observed that the MID group had an increased expression of PPAR-γ compared with the control group (p = 0.14), which shows that FG+PM may have a role similar to TZDs. PPAR-γ activation is associated with the expression of LPL, and fatty acid synthetase (FAS). LPL carries fat and degrades lipoprotein, which was approximately 50% higher in the MID group than in the control group (p = 0.15). The upregulation of LPL lowered FFA in the circulation and liver by increasing the fat inflow into AT, which decreased the hepatic lipotoxicity and improved insulin sensitivity [36]. The results of our study correspond well with those found in earlier studies showing that ginsenoside Re, Rg1, and Rb2 increase LPL expression [37], and ginsenosides Rd and Rb1 promoted adipose differentiation and lipid accumulation in 3T3-L1 cells [38]. FAS expression was suppressed by the consumption of FG+PM, especially in the HIGH group compared with the control group (p = 0.05). Therefore, FG+PM can improve insulin sensitivity through PPAR-γ activation and inhibit fatty liver development. Thus, FG+PM exhibited an anti-obesity effect.

In summary, the combination of fermented ginseng, B. longum BORI, and L. paracasei CH88 efficiently suppressed weight gain, hepatic and adipogenic fat accumulation, and TNF-α secretion in AT. This combination also improved the FBG level and increased cholesterol excretion into the feces. It is suggested that the FG+PM mixture could have an anti-obesity effect in HFD-fed mice.

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Conflict of Interest

The authors have no financial conflicts of interest to declare.

References


