Growth sensitivity in the epiphyseal growth plate, liver and muscle of SD rats is significantly enhanced by treatment with a fermented soybean product (cheonggukjang) through stimulation of growth hormone secretion

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Abstract. Cheonggukjang (CKJ), a fermented soybean product, has been reported to have beneficial effects on various chronic diseases, including cardiovascular disease, cancer and immune diseases. To investigate whether CKJ induces growth sensitivity in mammals, alterations of key parameters related to their growth were analyzed. Sprague-Dawley (SD) rats were treated with a high concentration of CKJ (H-CKJ) or a low concentration of CKJ (L-CKJ) for 10 days, and compared with vehicle-treated rats. The CKJ contained a high concentration of total flavonoids, phenolic compounds, daidzein and genistein, compared with the non-fermented soybean product. Body weight was higher in the H-CKJ-treated group compared with that in the vehicle- and L-CKJ-treated groups. Furthermore, the level of growth hormone (GH) was highest in the serum of the L-CKJ-treated group compared with the vehicle- and H-CKJ-treated groups. Moreover, the expression levels of the GH receptor increased in the liver tissue, but not in the muscle tissue, of the L-CKJ- and H-CKJ-treated groups. In the downstream signaling pathway of the GH receptor, the phosphorylation levels of Akt and Erk were upregulated in the liver tissue, but not in the muscle tissue, of the L-CKJ- and H-CKJ-treated groups. These results suggest that CKJ extract may enhance the sensitivity of the epiphyseal growth plate in SD rats, through the upregulation of GH secretion.

Introduction

Soybean is widely used to produce various fermented products, including cheonggukjang (CKJ), doenjang and gochujang in Korea. Of these products, CKJ is predominantly produced by fermentation with Bacillus subtilis for a short time. It contains numerous enzymes, microorganisms and bioactive compounds that are absent from unfermented soybean (1,2). In particular, flavonoid glycosides are converted into aglycones by hydrolysis, and a number of proteins are degraded into small peptides and amino acids during fermentation (3,4). CKJ exhibits diverse biological and pharmacological activities, including anti-obesity, antidiabetic and anti-inflammatory effects on human chronic diseases. In a study involving human subjects, CKJ supplementation was shown to significantly decrease visceral fat mass and apolipoprotein B/A1 levels (5). In another study, C57BL/6J mice exhibiting diet-induced obesity showed improvements in body weight, epididymal fat accumulation, and serum total cholesterol and low-density lipoprotein cholesterol levels, following CKJ treatment for 9 weeks (6). Furthermore, supplementation of CKJ has been shown to significantly reduce blood glucose and glycosylated hemoglobin levels, as well as improve insulin tolerance in C57BL/Ksj-db/db mice (a type II diabetic animal model) (7,8). In addition, when the anti-inflammatory activity of CKJ was investigated in the rat model of type I hypersensitivity and arachidonic acid-induced ear edema, CKJ treatment was
demonstrated to decrease passive cutaneous anaphylaxis (9). Moreover, ethanol extracts of CKJ were shown to significantly increase the viability of cultured mouse spleen and thymus cells, by suppressing apoptotic death (10). Although CKJ shows a variety of therapeutic activities in human disease, its effect on the sensitivity of body growth has not yet been investigated.

Therefore, in the present study, we investigated whether oral administration of CKJ significantly improves body growth via growth hormone (GH) secretion in young Sprague Dawley (SD) rats. The data presented provide strong evidence that CKJ is a potential candidate for the stimulation or enhancement of animal growth from infancy to adulthood.

**Materials and methods**

**Preparation of CKJ sample.** CKJ was manufactured using the Pungwon soybean strain that was kindly supplied by the National Institute of Crop Science in Miryang, Korea. The *B. subtilis* was supplied by the Applied and Environmental Microbiology Laboratory at Pusan National University (Miryang, Korea). To prepare the CKJ, the soybeans (100 g) were first washed, then soaked for 12 h with three volumes of tap water at room temperature. The soybeans were then treated with hot steam at 121°C for 30 min, before being allowed to cool to 37°C. Once cool, the steamed soybeans were inoculated with 5% (w/w) *B. subtilis* (1x10⁷ cells/ml), followed by fermentation at 37°C for 48 h. Powder of fermented soybeans (CKJ) was then prepared by freeze-drying, homogenization and sifting. The final CKJ sample extract was stored at -75°C prior to use.

**Principle component analysis of CKJ.** Freeze-dried CKJ powder (1 g) was subjected to extraction with 10 ml of distilled water at 70°C for 2 h. The sample was centrifuged (10 min, 102 x g) and the supernatant was collected for analysis. The concentration of total phenolics in CKJ was determined according to the Folin-Ciocalteu method (11). The sample (20 µl) was mixed with 100 µl of 0.2 N Folin-Ciocalteu reagent (Merck KGaA, Darmstadt, Germany) for 5 min, and then combined with 300 µl of 20% sodium carbonate. Following incubation at room temperature for 2 h, the absorbance of the reaction mixture was measured at 765 nm. Gallic acid (Sigma-Aldrich, St. Louis, MO, USA) was used as a standard to produce the calibration curve. Total phenolic content was expressed as gallic acid equivalents (mg) per gram of CKJ powder. The level of total flavonoids in CKJ was determined according to the method of Meda et al (12). The sample (200 µl) was added to test tubes containing 60 µl of 5% potassium nitrite, 600 µl of distilled water and 60 µl of 10% aluminum chloride. Following incubation at 25°C for 5 min, the absorbance of the reaction mixture was measured at 510 nm. The total flavonoid content was determined using a standard curve, with quercetin as a standard, and was expressed as quercetin equivalents (mg) per gram of CKJ powder. Furthermore, the total protein content was determined by the Bradford method (13), using bovine serum albumin as a standard. Total sugar content was determined according to the method of Dubois et al (14) with the modifications indicated below. The samples (0.5 ml) and the 5% (w/v) phenol solution (0.5 ml) were added to screw cap tubes, which were capped and vortex-stirred. Then 2.5 ml of concentrated sulfuric acid was added and mixed by vortexing. After incubation for 20 min at room temperature, the absorbance was measured at 490 nm using a spectrophotometer (BioSpec-mini; Shimadzu Corp., Kyoto, Japan).

**High-performance liquid chromatography (HPLC) analysis of daidzein and genistein concentration.** The concentration of daidzein and genistein in CKJ was analyzed, using a previously described method (15). Standard samples of 98% daidzein and 99% genistein were purchased from Sigma-Aldrich. The aqueous extracts of CKJ were mixed with 50% methanol and then this mixture was filtered with a 0.2-µm membrane filter (Wayers Co., Milford, MA, USA), before HPLC injection. The daidzein and genistein concentration was analyzed by the iLC 3000 HPLC system (Interface Engineering, Seoul, Korea) equipped with a Corona® CAD® Detector (ESA Biosciences, Inc., Chelmsford, MA, USA). The chromatographic separation was performed on a CAPCELL PAK MG C18 (4.6x250 mm; particle size,5 µm; Shiseido Co.,Ltd.,Tokyo,Japan). The mobile phase consisted of solvent A (deionized water) and solvent B (acetonitrile), using the gradient elution program: 0-25 min, 30-90% of solvent B and 25-40 min, 90% of solvent B. A flow rate of 1.0 ml/min was used for the sample analysis. The nebulizer gas was compressed nitrogen. The gas flow rate and gas pressure were maintained at 1.53 l/min and 35±2 psi, respectively. The output signal of the detector was recorded using Clarity™ chromatography software (DataApex, Prague, Czech Republic).

**Care and use of animals.** Four-week-old female SD rats (SamTacho, Osan, Korea) were bred at the Pusan National University-Laboratory Animal Resources Center (Korea Food and Drug Administration accredited; registration no., 231). The SD rats received an *ad libitum* diet of standard irradiated chow (Purina Mills, Seoungnam, Korea). The rats were maintained in specific pathogen-free conditions under a strict 12-h light/dark cycle (lights on at 6:00 a.m. and off at 6:00 p.m.), at a temperature of 23±2°C, with 50±10% relative humidity, according to the Guide for Laboratory Animals (Institute for Laboratory Animal Research, Washington, DC, USA). Animal experiment protocols were carefully reviewed and approved in accordance with the ethical and scientific care procedures of the Pusan National University-Institutional Animal Care and Use Committee (approval no. PNU-2012-0006).

**Experimental design and detection of body/organ weights.** A total of 15 four-week-old female SD rats were divided randomly into three groups, with five rats per group. The first [low concentration of CKJ (L-CKJ)-treated] and second [high concentration of CKJ (H-CKJ)-treated] groups of rats received 50 and 100 mg/kg body weight/day of CKJ extract diluted in water, respectively, via oral administration. The third group (vehicle-treated) received a daily administration of a comparable volume of water. The body weights of the rats were measured using an electronic balance (Mettler Toledo, Greifensee, Switzerland) at the indicated time points during the experimental period. At 10 days after the CKJ treatment, all animals were sacrificed using CO₂ gas and the weights of five organs collected from SD rats were also measured using the same method as that used to detect the body weight.
Furthermore, bone and tissue samples were acquired and stored in Eppendorf tubes at -70˚C until assay.

**Western blotting.** Protein prepared from liver and muscle tissues of vehicle-, L-CKJ- and H-CKJ-treated rats were separated by 4-20% sodium dodecyl sulfate-polyacrylamide gel electrophoresis for 3 h, following which, the resolved proteins were transferred to nitrocellulose membranes for 2 h at 40 V. Each membrane was incubated separately with the following primary antibodies overnight at 4˚C: Anti-GH receptor (ab78426; Abcam, Cambridge, UK), anti-Erk (sc-94; Santa Cruz Biotechnology, Inc., Santa Cruz, CA, USA), anti-pErk (sc-7383; Santa Cruz Biotechnology, Inc.), anti-Akt (#9272; Cell Signaling Technology, Inc., Danvers, MA, USA), anti-pAkt (#4058; Cell Signaling Technology, Inc.), and anti-actin (A5316; Sigma-Aldrich). The membranes were then washed with washing buffer (137 mM NaCl, 2.7 mM KCl, 10 mM Na₂HPO₄, 2 mM KH₂PO₄ and 0.05% Tween 20) and incubated with horseradish peroxidase (HRP)-conjugated goat anti-rabbit IgG (Zymed Laboratories, Inc., South San Francisco, CA, USA) at a 1:1,000 dilution, at room temperature for 2 h. Membrane blots were developed using a Chemiluminescence Reagent Plus kit (Pfizer, New York, NY, USA).

**Histological analysis and observation by optical microscopy.** Femur bones collected from the SD rats were fixed with 10% formalin, for at least two days at room temperature. The fixed femur bones were then demineralized for four days in 15% formic acid-prepared deionized water, embedded in paraffin wax and sectioned into 4-µm slices. Bone sections were subsequently stained with hematoxylin and eosin (Sigma-Aldrich) and observed by optical microscopy. The thickness of the epiphyseal growth plate was measured at three points using the Leica Application Suite (Leica Microsystems (Schweiz) AG, Heerbrugg, Switzerland).

**Quantification of GH by enzyme-linked immunosorbent assay (ELISA).** The concentration of GH in sera collected from vehicle- and CKJ-treated rats was determined by following the ultra-sensitive assay procedure, using reagents in the Growth Hormone Rat ELISA kit (KRC5311; Invitrogen Life Technologies, Carlsbad, CA, USA). The sera and standards were incubated in antibody-coated plates for 2 h at room temperature on a plate shaker at 51-61 x g. The wells were then washed six times using a PV100 automatic plate washer (Hoefer Inc., Holliston, MA, USA). HRP conjugates were subsequently added to all wells, followed by incubation for 30 min at room temperature on the shaker. The reaction was terminated by the addition of 50 µl of stop solution. Color alterations in the wells were read using a VMax® microplate reader (Molecular Devices, Sunnyvale, CA, USA) at 450 nm.

**Results**

**Concentration of key components in CKJ.** To identify changes in the functional components of CKJ, the concentration of several important components, including proteins, total flavonoids and total phenolic compounds, were measured using a number of traditional methods. The levels of these three aforementioned components significantly increased following fermentation of the soybean (Table 1). However, no significant alteration in the total sugar concentration was detected. In addition, the levels of two key components that were correlated with the putative beneficial effects for animal growth, were measured by HPLC analysis. The concentration of daidzein was detected as 57.19 µg/g, while the concentration of genistein was detected as 42.31 µg/g (Fig. 1). Therefore, the aforementioned results indicate that the CKJ used in this study contained high concentrations of protein, total flavonoids, total phenolic compounds, daidzein and genistein.

<table>
<thead>
<tr>
<th>Components</th>
<th>Non-fermented soybean</th>
<th>Fermented soybean (cheonggukjang)</th>
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<tbody>
<tr>
<td>Protein (mg/g)</td>
<td>489.060</td>
<td>565.030±12.141*</td>
</tr>
<tr>
<td>Total sugar (mg/g)</td>
<td>5.141</td>
<td>5.252±0.721</td>
</tr>
<tr>
<td>Total flavonoid (mg/g)</td>
<td>0.595</td>
<td>0.761±0.094*</td>
</tr>
<tr>
<td>Total phenolic compound (mg/g)</td>
<td>8.927</td>
<td>9.861±0.885*</td>
</tr>
</tbody>
</table>

CKJ, cheonggukjang. Data are expressed as the means ± SD. *P<0.05, compared with the non-fermented soybean.
Effect of CKJ on total body and organ weights. In order to determine the short-term effects of CKJ on animal growth, alterations of total body and organ weight in young SD rats were detected following CKJ administration for 10 days. The body weights of L-CKJ-treated rats were significantly higher compared with those of vehicle-treated SD rats from days seven to 11. However, in the H-CKJ-treated group, an increase in body weight was detected only on day three, and its level returned to that of the vehicle-treated group from day five (Fig. 2A).

Furthermore, of the five important organs, the weights of the brain, liver and kidney significantly increased in the L-CKJ-treated group compared with those in the vehicle-treated group. However, heart and spleen weights were maintained at constant levels in the L-CKJ- and H-CKJ-treated groups (Fig. 2B). These results suggest that the L-CKJ induced significant increases in total body and organ weights in young SD rats.

Effect of CKJ on the weight and length of the femur, and thickness of the epiphyseal growth plate. To investigate the effect of short-term CKJ treatment on skeletal growth, femur weight and length were measured in young SD rats treated with two different concentrations of CKJ. The femur is the most proximal bone of the leg in tetrapod vertebrates capable of walking or jumping (16). Therefore, we selected the femur as a skeletal growth indicator and investigated its characteristics in relation to the growth of young rats. Notably, significant alterations in the femur weight and length were not detected in SD rats treated with L-CKJ or H-CKJ for 10 days (Fig. 3A). However, a marked difference was observed in the epiphyseal growth plate of the distal femur. In the H-CKJ-treated group, the thickness of the epiphyseal growth plate significantly increased compared with that of the vehicle-treated group. The number of cells in the proliferation and hypertrophy zone of the growth plate significantly increased in this group. Furthermore, the thickness of the epiphyseal growth plate in the L-CKJ-treated group was marginally enhanced compared with that of the H-CKJ-treated group (Fig. 3B). The aforementioned results suggest that short-term CKJ treatment stimulated the growth of epiphyseal plate in young SD rats, although there was no direct effect on femur weight and length.

Effect of CKJ on GH secretion from the pituitary gland. To investigate the effects of short-term CKJ treatment on GH secretion ability, alteration of the GH concentration was examined in young SD male rats following CKJ treatment for 10 days. The GH concentration in the blood serum significantly increased in the L-CKJ-treated group compared with that of the vehicle-treated group. However, the H-CKJ-treated group showed a 37% lower GH level compared with that of the L-CKJ-treated rats, although its level remained markedly higher compared with that of the vehicle-treated group (Fig. 4A). These results show that the L-CKJ treatment induced GH secretion from the pituitary gland.
Effect of CKJ on GH sensitivity in the liver and muscle. To test whether the CKJ treatment has an effect on GH sensitivity in two target organs, the expression levels of the GH receptor and downstream proteins were analyzed in the liver and muscle of SD rats. The liver tissue showed a dose-dependent response to the CKJ treatment in terms of GH receptor expression, although the H-CKJ-treated group showed significantly higher GH receptor expression compared to the L-CKJ-treated group. However, the muscle tissue did not show any response to the CKJ treatment (Fig. 4B). Of the downstream signaling proteins in the GH receptor pathway, Akt and Erk were selected as key target proteins for the investigation of GH sensitivity. In the liver, the phosphorylation level of Akt significantly decreased in both the L-CKJ- and H-CKJ-treated groups compared with that of the vehicle-treated group. However, in the muscle, its level was lower only in the L-CKJ-treated group and there was no change in Akt phosphorylation in the H-CKJ-treated group.

Furthermore, the expression levels of pAkt and Akt simultaneously decreased in a CKJ concentration-dependent manner. The phosphorylation pattern of Erk differed from that of Akt. In the liver tissue, the phosphorylation level of Erk marginally increased in the L-CKJ-treated group, whereas it slightly decreased in the H-CKJ-treated group. In the muscle tissue, the phosphorylation level of Erk was dependent on the CKJ concentration (Fig. 5). These results indicate that the CKJ treatment differentially regulated the response to GH in liver and muscle.

Discussion

Fermented soybean includes different types of isoflavones, such as daidzein, genistein, enistein and glycetin. Their functions are closely associated with lowering the risk of breast and prostate cancer, the risk of cardiovascular disease and improving bone health (1). Generally, the concentration of
Daidzein and genistein, which was analyzed in this study, was found to be significantly increased in fermented soybean (data not shown). In particular, the concentration of the two isoflavones in the 60% ethanol extract of the fermented soybean was ~10-fold greater than that from the non-fermented soybean (15). However, in a study using 80% ethanol extracts of fermented soybean, the concentration of daidzein was markedly increased compared with that of the non-fermented soybean, yet genistein was undetected in the two forms of the soybean (10). In the present study, using aqueous extracts of the CKJ, the results were in agreement with the aforementioned study, although the rate of increase varied. Therefore, we suggest that the high concentration of daidzein and genistein may be a key contributory factor in the bone growth of CKJ-treated SD rats.

The synthesis and secretion of GH in somatotroph cells of the anterior pituitary gland may be induced by several natural products (17,18). For instance, administration of yeast extract for four weeks was shown to increase the body weight and GH secretion of young SD male rats (19). The bioassay-guided fraction of MeOH extract from the fenugreek (Trigonella foenum-graecum L.) seed stimulates the release of GH from rat pituitary cells. In particular, fenugreek saponin I and dioscin have been demonstrated to induce marked increases in GH levels, compared with other compounds (20). In the case of Glycyrrhiza radix, its MeOH extract and n-hexane fraction induce GH secretion by up to 1.89- and 4.59-fold, respectively, compared with the basal level (21). Recently, GH stimulation has received attention as a treatment for overcoming short stature (20,21). In an effort to identify medicinal foods that have a beneficial effect on human growth, the effect of CKJ on GH secretion in SD rats was investigated. As shown in Fig. 4, the results suggest that the CKJ was able to stimulate GH release from the pituitary gland.

Moreover, GH receptor exists in two forms: The full-length membrane-bound human receptor and the GH-binding protein. After GH binds to its receptor, dimerization of the GH receptor is followed by phosphorylation of JAK2 and the GH receptor in cells. The signal induced downstream of the GH-GH receptor is transferred into the nucleus, where transcription factors, including signal transducers and activators of transcription, regulate the expression of the GH and GH receptor (22). Moreover, the GH-GH receptor complex is known to activate the PI3K pathway via JAK2 phosphorylation of IRSs (23). In this downstream pathway, Akt/PKB are important in numerous cell processes, including proliferation, survival and metabolism (24). Following direct injection of GH into mice, it was observed that phosphorylation of Akt

Figure 4. Alterations of GH concentration and GH receptor expression. (A) After final CKJ administration, blood sera were collected from each group of Sprague-Dawley rats. GH concentration was detected using an ELISA kit with 0.5 ng/ml of sensitivity. (B) GH receptor level was analyzed in two target organs, the liver and muscle, as described in Materials and methods. Data are presented as the means ± SD of three replicates. *P<0.05, compared with the vehicle-treated group. GH, growth hormone; L-CKJ, low concentration of cheonggukjang; H-CKJ, high concentration of cheonggukjang.

Figure 5. Effects of cheonggukjang (CKJ) administration on growth hormone (GH) receptor downstream pathway proteins in (A) the liver and (B) the muscle using western blotting. Total cell lysates were prepared from tissues of vehicle-, L-CKJ-, and H-CKJ-treated groups, as described in Materials and methods. Fifty micrograms of protein per sample was immunoblotted using specific antibodies for each protein. Data are reported as the means ± SD of three replicates. *P<0.05, compared with the vehicle-treated group. L-CKJ, low concentration of cheonggukjang; H-CKJ, high concentration of cheonggukjang.
decreases slightly in the liver, whereas it increases in muscle tissue (23). The results of the present study using liver tissue are in agreement with previous results (23). However, the results of the current study using muscle differ in that Akt phosphorylation decreased or remained unchanged in the CKJ-treated groups.

In addition, GH has been demonstrated to control cell proliferation, differentiation and migration through the regulation of mitogen-activated protein kinases, which are activated by several mitogens and growth factors (25). A previous study demonstrated a lower level of Erk phosphorylation in the liver of rats following GH treatment (23). However, in the present study, the phosphorylation level of Erk significantly increased in the two organs. Therefore, these results differ from previous results, in which the phosphorylation level of Erk decreased in the liver and remained unchanged in the muscle of GH-treated mice (23). We therefore suggest that this difference may be attributable to the various compositions of CKJ.

Bone growth may be induced by several natural compounds and extracts. The Buguzhi extract applied to the collagen matrix has the effect of stimulating new bone formation in the parietal bone of New Zealand white rabbits (26). Fetal bone growth may be stimulated by maternal administration of Cissus quadrangularis petroleum ether extract during pregnancy (27). Additionally, SD male rats fed yeast extract exhibit increased tibial bone and femur bone growth (19). This study investigated the effects of fermented soybean extract on bone growth in SD rats. The results differed greatly from those of previous studies. The CKJ treatment did not induce an increase in femur weight or length, although thickness of the growth plate markedly increased. Such a significant difference in femur bone growth may be attributed to the dose and period of the CKJ administration, or the extract composition.

In conclusion, these results suggest that treatment with CKJ, containing enhanced flavonoid and phenolic compounds, for a short time period improves GH sensitivity in the epiphyseal growth plate, liver and muscle of SD rats through the upregulation of GH secretion.

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