Original

The Effect of Additive Formula Diet on Bone Structure of the Femur in Ovariectomized Rats

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Abstract: Osteoporosis is a major problem in the elderly population worldwide. Low calcium intake and vitamin D blood level are risk factors for osteoporosis, and improving their intake is effective in patients with micronutrients deficiency. However, the effect of these interventions is ambiguous. Additive formula diet (AFD) contains fructo-oligosaccharide (FOS), isoflavone (ISO) and 1.0% citric acid Ca as a supplement for patients with osteoporosis. We aimed to investigate the effect of AFD on bone structure of the femur in ovariectomized rats. Sixteen 20-week old ovariectomized rats were randomly distributed into 2 groups; one group was fed normal diet (N, n = 8) and the second group was fed AFD (A, n = 8). Both groups were fed for 24 weeks, and body weight was measured at 8 and 24 weeks. After measuring the weight at 24 weeks, rats were euthanized using carbon dioxide. Lateral femur bone was extracted, and bone mineral density (BMD) and bone mineral content (BMC) were measured via micro computed tomography. Non-decalcified ground sections of the femur were examined via polarized-light microscopy. At 24 weeks, BMD and BMC were significantly higher for the A group than in the N group. The A group showed significantly better structural values with respect to Tb.Th, Tb.N, Tb.sp, Tb.spac and SMI than the N group. The A group showed significantly denser trabecular observations than the N group. Examination of the non-decalcified ground section from the A group showed strong polarized light properties of orange compared with sections from the N group. AFD may improve bone turnover in osteoporosis with the expectant decrease in the incidence of falls and bone fractures, which may enhance the quality of life of the elderly.

Key words: Additive formula diet, Osteoporosis, Menopause, ovariectomy, femur

Introduction

The world’s aging population is showing a marked increase from 5.1% in 1950 to 7.7% in 2010. This rapid increase is still in progress, and expected to lead to a “super-aging society”. The increase in Japan’s aging population can be observed when comparing it to other industrialized countries; up until the 1980s this population was not prominent, then it started to increase in the 1990s, and reached its highest level in 2005. Consequently, an increase in diseases specific to geriatrics is expected. Bone and joint diseases are common among the Asian population, with their prevalence tripling over the past 20 years, and with 80% of patients being menopausal women. In Japan, although there are many patients with osteoporosis, it was speculated that 80% of them did not receive the required medical care. Osteoporosis increases the risk of bone fractures in the elderly, and consequently, the need for nursing care. Moreover, the anticipated fear of falling limits movement with decreased exercise and bone density. This creates a vicious circle of reduced motor function and further decline of muscle strength. Sarcopenia, defined as the decline in muscle strength, results in decreased daily activity. A previous report showed that sarcopenia and bone volume are strongly correlated.

Low calcium intake and vitamin D level in blood were considered as risk factors for osteoporosis, and improving their intake is considered effective, especially for the elderly with micronutrient deficiency due to inadequate drug intake secondary to chronic disease. However, the effect of these interventions is ambiguous. Aiming at efficient intake and absorption of calcium,
Nakada et al. fed ovariectomized osteoporosis-model rats a supplement composed of fructo-oligosaccharide (FOS), isoflavone (ISO) and phosphate Ca (3.0% content), and found a significant effect on bone formation and strength, and quality of the femoral shaft. However, a Ca concentration of 3.0% is high, and may cause nutritional imbalance due to overdose.

Therefore, additive formula diet (AFD) was prepared as 1.0% citric acid Ca instead of 3%, and was fed to osteoporotic rats with low serum calcium. The results showed that AFD was safe and that it improved metabolism. However, its effect on menopausal osteoporosis is still unknown. Therefore, the aim of this study was to investigate the effect of AFD on bone structure of the femur in ovariectomized rats.

Materials and Methods

Animals

Sixteen female Wistar rats (Sankyo Labo Service Co., Tokyo, Japan), aged 20 weeks, were used. They were housed in individual metal cages at a room temperature of 23 ± 1°C and humidity of 50 ± 1%, with ad libitum access to food and water. The experimental protocol was approved by an animals’ experiment Ethics Committee (approval no. AP13-MD013). All experiments were conducted according to the Guidelines for the Treatment of Animals, Nihon University, Chiba Japan.

Preparation of rats

For ovariectomy, the rats were anaesthetized with isoflurane. The abdominal area was sterilized with 75% ethanol and was surgically opened. Both ovaries were elevated and completely excised, thus completing ovariectomy. Then, the uterus and adipose tissue were placed back into the abdomen, and the incision was sutured. Both groups of rats were on the prescribed the diet at the age of 20 weeks.

Measurement of body weight

Body weight of all rats was measured at the following times: at intervention initiation, intervention eight weeks later (28 weeks of age time) and intervention 24 weeks later (44 weeks of age time).

Micro-computed tomography (CT)

Micro-CT settings

Micro-CT of the femur was captured by the R_mCT 2® (Rigaku Co., Tokyo, Japan). The R_mCT 2® imaging conditions were as follows: tube current 160 iA, tube voltage 90 kV, FOV 10 mm.
shooting time 26 seconds, and voxel size 20 μm × 20 μm × 20 μm. Micro-CT images were taken for the femur, and phantoms for CT-value proofreading were taken after 24 weeks. Imaging was on the distal femoral metaphysis (6.0 mm × 5.0 mm × 3.0 mm) (Fig. 1).

**Measurement of bone mineral density (BMD), bone mineral content (BMC) and trabecular structure**

Digital images were converted to 16-bit gray-scale TIFF format using the Atlas TIFF Convertor® (Rigaku Co., Tokyo, Japan), then analysed using TRI/3D-Bon BMD (TRI/3D-Bon; Ratoc System Engineering Co., Tokyo, Japan). For BMD, BMC and trabecular structure measurement, a hydroxyapatite calibration curve was prepared from the phantom images (hydroxyapatite content: 200 mg/cm³, 300 mg/cm³, 400 mg/cm³, 500 mg/cm³, 600 mg/cm³, 700 mg/cm³, 800 mg/cm³, and 1550 mg/cm³); then, BMD, BMC and trabecular structure of the femur were calculated using the TRI/3D-Bon trabecular structure analysis routine (auto-detection mode) using the obtained CT values.

Trabecular structure measurement: The three-dimensional structure of the trabecular bone obtained by micro-CT imaging was calculated with structure model index (SMI). In this three-dimensional structural analysis, trabecular thickness (Tb.Th), trabecular number (Tb.N), trabecular separation (Tb.Sp), trabecular spacing (Tb.Spac), and Trabecular Bone Pattern factor (TBPf) were determined.

**3D-map analysis**

Bone situation (inferred from BMD values) was determined from a 3D-map showing BMD distributions, obtained by micro-CT, with the following colour coding: high BMD: red to orange; intermediate BMD: yellow to green; low BMD: light blue to purple as shown in Fig. 2. The analysis of the 3D image was conducted using TRI/3D (TRI/3D-BON; Ratoc System Engineering, Tokyo Japan) image analysis software from bone density values. 3D-map conditions were: 1 pixel, 30 μm; range, 300-1500 mg/cm³.

**Microscopic examination of the non-decalcified ground section**

After immersing the femur in alcohol, fat was removed using fixation, dehydration, and xylene with methyl methacrylate resin. Resin embedding (Osteo Resin™ Embedding kit, Wako Pure Chemical Industries Co., Japan) was performed to reduce the thighbone to a single unit.

The embedded bones were cut in sagittal sections along their major axis using an automatic exact cutting machine (ISOMET, Buehler, USA) to produce non-decalcified ground sections of 100
BMD values were colour coded in the range of 300 mg/cm$^3$ – 1500 mg/cm$^3$. Blue to light blue represent low BMD, yellow-green to yellow intermediate BMD, and orange to red high BMD.

The cut sections were examined using a polarized-light microscope (LEICA M60, Leica Microsystems GmbH, Germany) in Cross Nicol.

**Statistical analysis**

Variables were presented as mean ± standard deviation. Student’s t-test was used to compare body weight, BMD, BMC, and trabecular structure between both groups, with the null hypothesis of no difference between N and A groups. P values <
0.05 were considered statistically significant.

Results

Measurement of body weight
Table 2 shows body weight measurements for the 2 groups at each observation time. From 8 to 24 weeks after the intervention, the increase in body weight was more in A than in N groups.

Micro-CT
BMD measurement
BMD was significantly higher in A than in N group at 24 weeks.

BMC measurement
BMC was significantly higher in A than in N group at 24 weeks.

Measurement of trabecular structure
Trabecular structure values for the 2 groups at 24 weeks were as follows: N: Tb.Th 69.4 ± 4.0 μm, Tb.N 0.1 ± 0.04 1/mm, Tb.Sp 427.1 ± 48.3 μm, Tb.Spac 496.6 ± 50.5 μm, TBPF 25.7 ± 4.9 1/mm, and SMI 3.1 ± 0.05; A: Tb.Th 76.5 ± 2.0 μm, Tb.N 0.3 ± 0.02 1/mm, Tb.Sp 220.8 ± 23.3 μm, Tb.Spac 285.8 ± 29.3 μm, TBPF 21.5 ± 0.8 1/mm, and SMI 3.0 ± 0.05 (Table 4) (Tb.Th, Tb.N, Tb.Sp, Tb.Spac, TBPF, SMI p<0.001).

3D-map analysis
Figure 3 shows the 3D-map for the 2 groups after 24 weeks of the intervention. Red to orange, yellow to green, and light blue to purple in the 3D-map indicate high, intermediate, and low BMD, respectively. Most of the cortical bone section from N rats was shown in orange to yellow, denoting high to intermediate BMD, with occasional red areas reflecting a high BMD. The inside part of the cortical bone section from A rats appeared orange to yellow, with more red areas than in sections from N group. In group N, there was little observable cancellous bone (Fig. 2).

Table 4. Trabecular structure measurement quantifying the state of the trabecular bone in the cancellous bone.

<table>
<thead>
<tr>
<th></th>
<th>Tb.Th (μm)</th>
<th>Tb.N (1/mm)</th>
<th>Tb.Sp (μm)</th>
<th>Tb.Spac (μm)</th>
<th>TBPF (1/mm)</th>
<th>SMI</th>
</tr>
</thead>
<tbody>
<tr>
<td>OVX+Normal</td>
<td>69.4±4.0</td>
<td>0.1±0.04</td>
<td>427.1±48.3</td>
<td>496.6±50.5</td>
<td>25.7±4.9</td>
<td>3.1±0.05</td>
</tr>
<tr>
<td>OVX+AFD</td>
<td>76.5±2.0</td>
<td>0.3±0.02</td>
<td>220.8±23.3</td>
<td>285.8±29.3</td>
<td>21.5±0.8</td>
<td>3.0±0.05</td>
</tr>
</tbody>
</table>

Microscopic examination of the non-decalcified ground sections
Polarized light micrograms are shown in Fig.3. Sections from A group showed stronger orange polarized light in the bone as compared to sections from N group.

Discussion
The purpose of this study was to examine the benefits of administering AFD to ovariectomized rats, and to assess the improvements in bone structure due to FOS, ISO, and calcium citrate contained in AFD.

The weight of rats in both N and A groups exhibited an upward trend at 1 week of age, and there was no difference in weight gain between both groups. Therefore, there was no difference in meal intake stemming from differences in the feed. BMD and BMC measurements of trabecular bone revealed significantly higher values for A than for N group. Most of the cortical bone part of the N group was shown with orange. Cancellous bone shown with blue ~ light blue was observed near cortical bone in the medullary cavity. In A group, high BMD (shown in red) was observed in the major part of cortical bone, and intermediate to low BMD (shown in blue to green) was observed in parts of the marrow cavity. Numerical comparison of BMD revealed an approximate difference of 30 mg/cm³, but visual inspection of the 3D-map revealed a difference in the trabeculae number of trabecular bone by a measurable range. Furthermore, the following parameters were used for quantification of trabecular microarchitecture: trabecular thickness (Tb.Th (μm)), trabecular number (Tb.N (1/mm)), trabecular separation (Tb.Sp (μm)), trabecular spacing (Tb.Sp (μm)), Structure Model Index (SMI; used to assess whether trabecular structure has plate-like or rod-like geometry), trabecular bone pattern factor (TBPF; used to assess convex, concave, and honeycomb-like trabecular surfaces). In comparing A to N groups to quantify and inspect the state of the trabeculae, it shows that the trabeculae are larger in size, the space between the trabeculae is smaller, and there is a larger number of trabeculae in A group. These findings suggest that the components present in AFD increased the number and promoted the growth of trabeculae. However, the connection between trabeculae did not change, as it was apparent from the values of SMI and TBPF. This difference between both groups may indicate that the 3 components of AFD improved bone turnover and were beneficial for bone formation. In A group, the thighbone metaphyseal cancellous bone showed dense trabeculae.
as compared to N group, based on results of the bone quantity structure measurement as observed by polarized light microscopy. Detecting orange polarized light suggested the presence of collagen fibres. Ovariectomy in an animal model mimics many types of bone loss caused by oestrogen deficiency, known to cause trabecular bone loss, which is the same cause of postmenopausal osteoporosis. ISO present in AFD has a structural resemblance to oestrogen, possess an oestrogen-like action and exhibits an affinity to oestrogen receptors such as ER. A component that has affinity for these oestrogen receptors, ISO daidzein is converted to equol. It is converted into black-bellied plover and equol by ISO, and these ingredients have affinity to an oestrogen receptor, which inhibits bone loss. Furthermore, the fructo-oligosaccharides in AFD are non-digestible oligosaccharides known to enhance mineral absorption. Previously published reports suggest that calcium citrate, another component of AFD, is more easily absorbed than calcium carbonate.

Although calcium citrate is insoluble in water, chelation of calcium helps the calcium be absorbed by the body. Previous experiments suggest that Ca citrate inhibits BMD decline by suppressing elevated parathyroid hormone level in blood and thus blocking bone calcium outflow. The aforementioned findings suggest that the components of AFD exhibit inhibitory action on bone resorption and promote bone formation stemming from improved bone turnover.

Conoscopic imaging showed that A group exhibited stronger amber coloured polarization properties in cortical bone as compared to N group. Although a big difference in birefringence between both groups was not observed, A group exhibited denser trabecular bone than N group. These differences between both groups may be the result of active bone remodelling facilitated by components of AFD. Thus, the bones of the ovariectomized rats may have improved due to the action of AFD.

These findings suggest that AFD is effective for treating osteoporosis through its ability to increase the rate of bone turnover following calcium absorption in the body. This experiment took in AFD for long-term (24 weeks), and, as a result, improvement of the bone substance was found. Therefore, we thought that it was necessary to examine an optimal intake period on taking in AFD.

Also, we examined the metabolic pathway of AFD and thought that confirmation of the harm was important.

AFD may improve bone turnover in osteoporosis and expected to prevent the incidence of falls and bone fractures, and thus could be valuable for improving and maintaining the QOL of the elderly.

Conflict of Interest
The authors have declared that no COI exists.

References


