Research Article

Microbubble-Mediated Ultrasound Outweighs Low-Intensity Pulsed Ultrasound on Osteogenesis and Neovascularization in a Rabbit Model of Steroid-Associated Osteonecrosis

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Microbubbles magnify the acoustic pressure of low-intensity pulsed ultrasound (LIPUS) and may enhance its bioeffect for diagnostic and therapeutic purposes. This study compared the effect of this novel microbubble-mediated ultrasound (MUS) with that of the traditional LIPUS on osteogenesis and neovascularization in a rabbit model of steroid-associated osteonecrosis. We hypothesized that MUS might outweigh LIPUS on promoting osteogenesis and neovascularization in steroid-associated osteonecrosis. The bilateral femoral head necrosis was induced by lipopolysaccharide and methylprednisolone in the rabbits. The indices of bone mineral density (BMD), trabecular number, maximal loading strength, and mineral apposition rate were analyzed, demonstrating that the animal model of steroid-associated osteonecrosis was successfully established. Both the MUS group (GM) and the LIPUS group (GL) were sonated 20 min daily for six weeks. GM received an extra intracapsular injection of microbubbles before sonation every other day. Fluorescence bone labeling, Micro-CT Analysis, biomechanical test, quantitative real-time PCR, Western blot analysis, and histological evaluation were performed for comparing GM with GL. The results demonstrated a 39% higher mineral apposition rate in GM compared with GL. The BMD and the maximal loading strength of femoral head of GM increased by 4.3% and 27.8% compared to those of GL, respectively. The mRNA and protein expression of BMP-2 and VEGF were also significantly higher in GM. The number of blood vessels of GM was 65% greater than that of GL. MUS is more potent than LIPUS in enhancing osteogenesis, neovascularization, and biomechanical strength of femoral head in the animal model of steroid-associated osteonecrosis. Without increasing the intensity of sonation or the risk of tissue damage, MUS is better for inhibiting the process of steroid-associated osteonecrosis.

1. Introduction

Steroid usage is the most common nontraumatic cause of osteonecrosis [1]. Up to 40% of steroid users will develop some degree of osteonecrosis, known as the steroid-associated osteonecrosis [2, 3]. The precise mechanism in which corticosteroids induce osteonecrosis is currently unclear. Hypotheses include fat cell hypertrophy, fat embolization, intravascular coagulation, and osteocyte apoptosis [4–6]. To our knowledge there is no established preventative method for steroid-associated osteonecrosis.

Low intensity pulsed ultrasound (LIPUS) is understood as a noninvasive, feasible, and economic modality for the delayed union and the nonunion of bone fractures. It is approved by the Food and Drug Administration in 1994. Since then, the research exploring the prospects of its clinical application has not stopped. The therapeutic application of LIPUS is worldwide nowadays. Recently, our team for the first time reported the effect of LIPUS on steroid-associated osteonecrosis in a rabbit model of steroid-associated osteonecrosis [7]. The authors revealed that LIPUS could promote the bone repair and the vascularization in vivo [7, 8]. It provides us a new insight to preventing the progression of steroid-associated osteonecrosis.

Microbubbles have been increasingly used to improve the therapeutic effect of LIPUS [9]. Microbubbles, with a
gas-filled structure, are stabilized by a protein, lipid, or polymer shell. Microbubbles are initially developed as contrast agents for ultrasound imaging [10, 11]. They have the ability to produce mechanical actuations and shockwaves when they interact with the focal zone of LIPUS. More and more therapeutic applications of microbubbles are emerging at present [9, 12–14]. This novel ultrasound, known as microbubble-mediated ultrasound (MUS), has the ability to produce greater microstream and microjet than LIPUS [15, 16]. To our knowledge, whether MUS has more effect on the repair progress in steroid-associated osteonecrosis is unknown.

We hypothesized that MUS might outweigh the repair process of femoral head in steroid-associated osteonecrosis more efficiently than LIPUS. This study aimed to explore the enhancement effect of MUS on bone microstructure, mechanical strength, and vascularization compared with LIPUS.

2. Methods and Materials

2.1. Experimental Design. The experimental protocol was approved by the Animal Care and Use Committee of Zhejiang University. Twenty-four adult male New Zealand rabbits (mean weight: 2.69 kg, range 2.45 to 3.00 kg) were used. The rabbit model of steroid-associated osteonecrosis was prepared using a protocol similar to the previous studies [17, 18]. To begin with, all the rabbits received intravenous injections of 10 μg/kg body weight of lipopolysaccharide (Escherichia coli 011:B4; Sigma, St Louis, MO, USA). After 24 hours three injections of methylprednisolone (Pharmacia and Upjohn, Puurs, Belgium) (20 mg/kg body weight) were given intramuscularly every 24 hours. It has been reported that the osteonecrosis lesions began after two weeks and completed after six weeks [17]. The rabbits were randomly divided into three groups: the control group (G_C), the LIPUS group (G_L), and the MUS group (G_M) (n = 8). Both G_L and G_M received LIPUS. Additionally G_M received a 0.1 ml intracapsular injection (guided by B-ultrasound) of sulfur hexafluoride-filled microbubble solution (volume concentration: 7-8 μl/ml, mean diameter: 2.5 mm, 90%, 6.0 mm, 99%, and 11.0 mm; Sonovue, Bracco, Geneva, Switzerland) before insonation [9] (Figure 1). Three rabbits were randomly selected for MRI (T1-weighted image, TR 400 ms, TE 20 ms; T2-weighted image, TR 2000 ms, and TE 106 ms) two weeks after methylprednisolone injection to test whether osteonecrosis was induced.

2.2. Insonation. LIPUS (US 13; Cosmogamma Co, Bologna, Italy) was used at a pulse frequency of 1.0 MHz, an output intensity of 200 mW/cm², a 20% duty cycle, and a 100-Hz repetition rate [7]. Microbubble solution was prepared according to the manufacturer’s instructions. The skin around the hip was cleaned and ultrasound gel (KL-250; Kepler Co, Hangzhou, China) was used to couple the LIPUS transmission. The rabbits were anesthetized with 3% pentobarbital sodium before insonation. LIPUS therapy was administered to the bilateral femoral head 24 hours after the lipopolysaccharide injection and before the first methylprednisolone injection. The sham insonation was administered to the rabbits of G_C (the device was turned off). In this study, rabbits were insonated with 20 minutes per day for six weeks. Then all the rabbits were euthanized with an overdose intravenous...
injection of pentobarbital sodium. Bilateral femurs of the hind leg were collected for analyses.

2.3. Fluorescence Bone Labeling. All the rabbits received intramuscular injections of a fluorescence tracer, alizarin complexone (30 mg/kg, Merck, Darmstadt, Germany), and calcein (10 mg/kg, Sigma, St. Louis, MO, USA) at 14 and 4 days, respectively, before the euthanasia to label newly mineralized bone. The mineral apposition rate (μm/day) (MAR = mean vertical distance between two fluorescence bands/injection interval), which generally indicated the mineralization rate of the newly formed bone, was analyzed [19].

2.4. Micro-CT Analysis. The proximal parts of the bilateral femur were prepared for micro-CT (eXplore Locus SP; GE Co, Fairfield, CT, USA) scanning, with a spatial resolution of 45μm according to the protocol for animal studies. To evaluate the degree of plate-like and rod-like trabecular bone in the femoral head, a predetermined bone cylinder was used as the region of interest (3.0 mm diameter and 1.5 mm height) (Figure 4(d)) to quantify the mean volumetric bone mineral density (BMD) (mg/cc) and the trabecular architectural parameters, including the bone tissue volume density (bone volume/total volume, %), the trabecular number (/mm), the trabecular thickness (μm), the trabecular spacing (μm), and the connectivity density (/mm³) [20].

2.5. Biomechanical Evaluation. The loading strength of the femoral head was measured using an indentation test with a 2.8-mm-diameter indenter. The proximal femur was fixed and placed on a metal platform of a static materials tester (Z2.5; Zwick/Roell, Ulm, Germany). Alignment of the specimen was adjusted in accordance with the stainless steel rod in the upper holding device. The indenter was displacement-controlled at a constant rate of 5 mm/min and was stopped when the load markedly decreased. The maximum loading strength (N) was automatically recorded (TestXpert II; Zwick / Roell) [7] (Figure 5).

2.6. RNA Extraction and Quantitative Real-Time. mRNA levels of samples were determined by quantitative real-time PCR. In brief, the RNA of the samples was extracted with Trizol reagents (Invitrogen, USA). The primers (Table 1) were designed by Oligo 6.0 primer design software (Molecular Biology Insights, Colorado Springs, CO, USA) and synthesized by KeDuo (Hangzhou, China). Quantitative real-time PCR reactions were carried out by using a SYBR® Premix Ex Taq™ Kit (TaKaRa Biotech Co, Shiga, Japan). The cycle threshold (Ct) was measured for each mRNA in each sample. The relative quantifications of the gene expressions (BMP-2 and VEGF) were calculated by the formula $\Delta \Delta \text{Ct} = \Delta \text{Ct}_1 - \Delta \text{Ct}_2$, where $\Delta \text{Ct}_1$ is the value of the housekeeping gene 18S and $\Delta \text{Ct}_2$ indicates the Ct value of the target gene [21, 22].

2.7. Western Blot Analysis. Protein was isolated with radiomunoprecipitation assay buffer lysis buffer (Sangon Biotech Co, Ltd). Sodium dodecyl sulfate polyacrylamide gel (Keduo, Hangzhou, China) was formulated and loaded and electrophoresis occurred prior to transferring to a membrane (Sangon Biotech Co, Ltd), which was closed with skimmed milk powder. The samples were reacted with the primary β-actin antibody, secondary antibody (VEGF and BMP-2, Goat anti-rabbit, Cruz Biotec, Inc, Santa Cruz, CA, USA), and chemiluminescence was performed. The images were analyzed with Bandscan 5.0 (Glyko Inc, Novato, CA, USA) to calculate the intensity of the bands.

2.8. Histologic Evaluation and Immunohistochemistry. Specimens for histological evaluation were fixed in 4% paraformaldehyde solution. After decalcification in 10% EDTA, the samples were embedded in paraffin. Five-micron sections were cut and stained with hematoxylin and eosin for evaluation of necrosis and calculation of fat cell size. The α-smooth muscle actin staining was used to assess the number and the mean diameter of blood vessels [23].

The slices were examined using a light microscope (Olympus BX51, Tokyo, Japan). Five fields in the subchondral area of the femoral head on each section were chosen [7]. The first field was located at the approximate center of the femoral head and the other four were located on both sides of the first field. The following parameters were assessed: the average diameter of the fat cells [7, 17] and the number and the diameter of blood vessels. Different fields per slice were converted to digital pictures and quantitative analysis was performed using Image-Pro Plus 6.0 (Media Cybernetics Inc, Rockville, MD, USA).

2.9. Statistical Analysis. The data were expressed as the mean ± standard deviation. SPSS 20.0 software (SPSS, Chicago, IL, USA) was used to analyze the data by one-way analysis of variance with Scheffe’s multiple comparison method. p < 0.05 was considered statistically significant.

<table>
<thead>
<tr>
<th>Gene</th>
<th>GeneBank accession</th>
<th>Primer sequences (5’to 3’)</th>
<th>Size (base pairs)</th>
<th>Annealing (°C)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Rabbit BMP-2</td>
<td>NM 001082650</td>
<td>GGGGTGGAACGACTGGATGTGTCGACAGATGGCATGTTAGT</td>
<td>117</td>
<td>64</td>
</tr>
<tr>
<td>Rabbit VEGF</td>
<td>AY196796</td>
<td>GGGGGCTCGCTAGAATGAAAGCCTGCGCTGGTTGAGTTTGAT</td>
<td>97</td>
<td>65</td>
</tr>
<tr>
<td>Rabbit 18S</td>
<td>EU236969</td>
<td>GACCAGACAGGCAAGGACGCCATCGCAGCATGATTATG</td>
<td>119</td>
<td>64</td>
</tr>
</tbody>
</table>

3. Results

3.1. MRI Two Weeks after Methylprednisolone Injection. The image showed the appearance of the mild edema in the proximal femur (Figure 2).

3.2. Mineral Apposition Rate Evaluation. The mineral apposition rate of the femoral head was detected by the distance between the parallel fluorescent bands. The two fluorescent labels (green bands, calcein; red bands, alizarin complexone) were clear (Figure 3). According to the statistical analysis, there was a significant difference in the mineral apposition rate between \( G_M \) (2.660 ± 0.2390 \( \mu \)m/d) and \( G_L \) (1.913 ± 0.1457 \( \mu \)m/d) \( p = 0.008 \); Table 2).

3.3. Osteogenesis in Femoral Head. Micro-CT images indicated that the trabecular bone was more compact, thicker, and denser in the subchondral bone of \( G_M \) (Figure 4) compared with \( G_L \) and \( G_C \). The BMD of \( G_M \) (932.6 ± 6.861 mg/cm\(^2\)) was significantly higher than that of \( G_L \) (894.5 ± 12.84 mg/cm\(^2\)) \( p < 0.001 \); Table 2). The trabecular number of \( G_M \) (2.352 ± 0.06980 \( 1/\text{mm}^2 \)) was significantly higher than that of \( G_L \) (2.213 ± 0.07245 \( 1/\text{mm}^2 \)) \( p = 0.039 \); Table 2).

3.4. Mechanical Strength of Femoral Head. The maximal loading strength of femoral head of \( G_M \) (467.2 ± 65.96 N) was significantly greater than that of \( G_L \) (365.6 ± 57.40 N, \( p < 0.001 \)) (Figure 5).

3.5. BMP-2 and VEGF Expression. The mRNA expression of BMP-2, the protein expression of BMP-2, the VEGF mRNA expression, and the VEGF protein expression of \( G_M \) were significantly higher than those of \( G_L \) (1077 ± 8.527 vs 39.55 ± 3.141, \( p < 0.001 \); 5.753 ± 0.1206 vs 3.860 ± 0.2706, \( p < 0.001 \); 16.09 ± 2.911 vs 8.733 ± 0.5577, \( p = 0.006 \); 5.640 ± 0.2095 vs 3.467 ± 0.1656, \( p < 0.001 \); respectively) (Figure 6).

3.6. Histomorphometric Evaluation. The limbs of \( G_C \) were observed to have sparse trabeculae (Figure 7(a)) with massive empty lacunae, few osteoblasts, large marrow fat cells, and a disordered architecture of marrow tissue (Figure 7(c)). In contrast, no such changes were observed in \( G_M \) (Figures 7(g) and 7(i)). The average fat cell diameter of \( G_M \) (28.3 ± 8.4 \( \mu \)m) was lower than that in \( G_L \) (34.9 ± 7.5 \( \mu \)m) \( p = 0.075 \); Table 2).

To our knowledge, this is the first study to investigate the bioacoustic effect of MUS in a rabbit model of steroid-associated osteonecrosis. Our data support the hypothesis that MUS strengthens the repair process of femoral head more efficiently than LIPUS in steroid-associated osteonecrosis. The results provide radiological, biomechanical, molecular biological, and histological evidence to support our

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Table 2: Micro-CT, biomechanical and histomorphometric evaluations, and gene expression in the three groups.

<table>
<thead>
<tr>
<th></th>
<th>( G_C )</th>
<th>( G_L )</th>
<th>( G_M )</th>
</tr>
</thead>
<tbody>
<tr>
<td>BMD (mg/cm(^2))</td>
<td>883.4 ± 10.28</td>
<td>894.5 ± 12.84</td>
<td>932.6 ± 6.861*#</td>
</tr>
<tr>
<td>BV/TV (%)</td>
<td>40.30 ± 3.480</td>
<td>41.37 ± 4.612</td>
<td>41.95 ± 7.869</td>
</tr>
<tr>
<td>Tb.N (1/mm)</td>
<td>2.079 ± 0.08272</td>
<td>2.213 ± 0.07245*</td>
<td>2.352 ± 0.06980*#</td>
</tr>
<tr>
<td>Tb.Th (mm)</td>
<td>0.1705 ± 0.01974</td>
<td>0.1843 ± 0.02414</td>
<td>0.2064 ± 0.01200*</td>
</tr>
<tr>
<td>Tb.Sp (mm)</td>
<td>0.2672 ± 0.02893</td>
<td>0.2613 ± 0.02759</td>
<td>0.2241 ± 0.009524*#</td>
</tr>
<tr>
<td>Conn.D (1/ mm(^3))</td>
<td>10.79 ± 1.634</td>
<td>11.29 ± 1.053</td>
<td>14.39 ± 2.148*#</td>
</tr>
</tbody>
</table>

Values are mean ± SD.
* \( p < 0.05 \) vs. \( G_C \).
# \( p < 0.05 \) vs. \( G_L \).

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Figure 2: The edema sign (the yellow arrow) was observed in the proximal femur.
Figure 3: The bone of femoral head was labeled during bone regeneration and remodeling by calcine (green) and alizarin complexone (red). Representative labeling images from each group. The capacity of bone regeneration was the strongest in $G_M$ (f).
Figure 4: Representative micro-CT images from each group. The trabecular bone in G_M was thicker and denser than that in G_L and that in G_C. (d) Micro-CT-based trabecular architecture and sketches of the region of interest.

Figure 5: (a) Illustrating how the specimen was fixed in dental cement and the force was parallel to the long axis of the proximal femur. (b) Representative load-deformation curve obtained by the biomechanical test. (c) Statistical analyses of the biomechanical test data. ∗ p<0.05 vs. the blank group; # p<0.05 vs. G_L.
speculation that the microbubble use is a promising approach for improving the biological capabilities of LIPUS in bone repair [7].

We found that microbubble enhanced the bioacoustic effect of LIPUS on bone repair. Our study showed that the loading strength and BMD of the proximal femoral in GM was statistically higher than that in GL. This meant that MUS increased the biomechanical strength. Later, histopathology pictures also showed the smaller fat cells, the less bone and marrow necrosis, and the less empty osteocyte lacunae in the GM compared with GL. We failed to find the difference of the loading strength and BMD between GL and GC. This could be explained by the insonation duration of six weeks, which was a half of the insonation duration in our prior study [7].

The bioeffect of LIPUS, i.e., cavitation, are based on its mechanism of sonoporation [24]. Cavitation-related tissue effects arise from a complex array of interactions between microbubbles and tissue, which include the concomitant

Figure 6: (a) These band scans showed the Western blot analysis of BMP-2 and VEGF protein expression in GC, GL, and GM. (b) Statistical analyses of the mRNA expressions of BMP-2 and VEGF, protein expression of BMP-2, and VEGF data. *p < 0.05 vs. GC; #p < 0.05 vs. GL.
process of mechanical effects resulting from microstreaming and microbubble oscillation, growth, and collapse behaviors in the field. Sonoporation may directly increase the protein synthesis, the collagen synthesis, the membrane permeability, the integrin expression, and the cytosolic Ca\(^{2+}\) concentration [25]. Previous studies have found that the sonoporation could enhance the osteogenic differentiation and accelerated the osteoblast activity and the bone formation [26]. Our qPCR and Western blot results showed that BMP-2 expression was substantially higher in GM. It has been established that BMP-2 plays a central role in initiating the bone formation, stimulating the osteogenic differentiation, and increasing the osteoblast activity. The result of the mineral apposition also showed the dramatical new bone formation at the proximal of femoral head in GM. The data suggested that microbubbles amplified the osteogenetic effect of LIPUS.

Sufficient angiogenesis is a fundamental prerequisite of bone repair. The present study indicated that MUS had great potential in stimulating angiogenesis. Apart from the microbubble-enhanced cavitation, this bioacoustic effect may account for the promoted production of angiogenic growth factors by MUS. Yoshida et al. [27] found the ultrasonic destruction of microbubbles delivered to the ischemic limbs recruited inflammatory cells producing VEGF. In the present study, we found that the angiogenesis was more vigorous in GM compared with GL. Near-normal blood vessels were observed in GM. The minimal diameter of the blood vessels was also higher in GM than that in GL, which could better lead to an increase in blood perfusion.

Microbubbles form endogenously in tissues with the interaction between acoustic pulses and cavitation nuclei [28]. Microbubbles are produced in limited quantities endogenously in tissue. This indicated a need for more ultrasonic energy. However the excessive energy would increase the energy absorption by tissues, resulting in a damage to these tissues. Previous literatures indicated that the
cavitation could be amplified by the external microbubbles [14]. In our model, we injected 0.1 ml of microbubble solution intracapsular before sononation and exposed the proximal leg to 200 mW/cm² LIPUS for 20 min per day. Depending on this acoustic pressure, microbubbles presented a stable cavitation, where microbubbles oscillated over many cycles without collapsing. The stable cavitation stimulated by microbubbles was probably responsible for the increased efficiency of LIPUS.

Limitations definitely remained. First, the duration of MUS was set at six weeks. Further studies are required to reveal the saturation point, that is, the length of time after which the bioeffect of MUS is no longer significant. Second, the osteonecrosis lesion hardly led to joint collapse in rabbits. This could be attributed to the dissimilarities in weight bearing between quadrupedal animals and bipedal humans, particularly at the hip [17, 29]. Third, the cellular and the molecular mechanisms of osteogenesis and neovascularization by MUS need further investigation.

5. Conclusions

The present study indicated that MUS outweighed LIPUS on promoting the osteogenesis and the neovascularization in a
steroid-associated osteonecrosis model. Without increasing the intensity of insonation and the risk of tissue damage, MUS strengthened the femoral head which might prevent its further collapse. MUS holds more promise in inhibiting the pathological process of steroid-associated osteonecrosis than LIPUS.

**Data Availability**

The data used to support the findings of this study are included within the article.

**Conflicts of Interest**

All authors indicate they have no conflicts of interest.

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**References**


