BMP6-Engineered MSCs Induce Vertebral Bone Repair in a Pig Model: A Pilot Study

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Osteoporotic patients, incapacitated due to vertebral compression fractures (VCF), suffer grave financial and clinical burden. Current clinical treatments focus on symptoms' management but do not combat the issue at the source. In this pilot study, allogeneic, porcine mesenchymal stem cells, overexpressing the BMP6 gene (MSC-BMP6), were suspended in fibrin gel and implanted into a vertebral defect to investigate their effect on bone regeneration in a clinically relevant, large animal pig model. To check the effect of the BMP6-modified cells on bone regeneration, a fibrin gel only control was used for comparison. Bone healing was evaluated in vivo at 6 and 12 weeks and ex vivo at 6 months. In vivo CT showed bone regeneration within 6 weeks of implantation in the MSC-BMP6 group while only minor bone formation was seen in the defect site of the control group. After 6 months, ex vivo analysis demonstrated enhanced bone regeneration in the BMP6-MSC group, as compared to control. This preclinical study presents an innovative, potentially minimally invasive, technique that can be used to induce bone regeneration using allogeneic gene modified MSCs and therefore revolutionize current treatment of challenging conditions, such as osteoporosis-related VCFs.

1. Introduction

Osteoporosis is a debilitating systemic disorder with severe clinical and financial implications. It is estimated that osteoporosis affects 44 million Americans and more than 200 million people worldwide [1, 2]. In the United States alone, the direct annual cost of osteoporotic fractures was 19 billion dollars in 2005 and with the increase in life expectancy it is predicted to surge to an estimated $60 billion by 2030 [3–5]. Osteoporotic vertebral compression fractures (OVCFs) are the most common type of fractures of osteoporotic patients [6]. It is estimated that, in the United States alone, 1.5 million people suffer from OVCFs every year [7]. Current treatments aim at preventing OVCFs in osteoporotic patients by inhibiting bone resorption using various drugs. These drugs, however, are only efficient for a limited duration and are also associated with severe side effects [8–11]. Once OVCF occurs, nonsurgical treatments fail to treat the source or eliminate the pain and oftentimes lead to physical disability. The limitations of nonoperative management have fostered an increasing interest in new, minimally invasive, surgical approaches. Still, minimally invasive techniques, such as vertebroplasty and kyphoplasty, are associated with a number of serious clinical complications, such as infection, leakage into the spinal canal, and increased incidence of fracture to adjacent vertebrae [12–14]. Furthermore, in two recent
2. Materials and Methods

2.1. Cell Preparation

2.1.1. Isolation of BM-Derived MSCs. Allogeneic porcine MSCs were isolated from the bone marrow (BM) of donor Yucatan minipigs as previously described [22]. Briefly, costal BM of euthanized minipigs (35–40 kg with a mean age of 1.5 years) was harvested aseptically. The BM-containing ribs were scraped, flushed with PBS, and centrifuged at 900 g for 10 min. The pellet was resuspended in PBS, after which it was layered on lymphocyte separation medium (ICN Pharmaceuticals, Bryan, OH, USA). Following cellular isolation, the mononuclear cells were plated at a density of 0.4×10^6 cells per cm^2 and culture medium was changed every 3–4 days. Upon confluence, the cells were replated at a density of 7×10^6 cells per cm^2 for expansion. For this study, up to fifth passage cells were used for in vivo implantation.

2.1.2. Genetic Modification of MSCs. Genetic modification of porcine MSCs was performed with the aid of a Nucleofector device (Amaxa Biosystems, Cologne, Germany), as previously described [21, 23, 24]. Briefly, 2×10^6 MSCs were transfected with 10 μg of cDNA3-pCMV-rhBMP-6 (rhBMP-6) plasmid. Immediately after transfection, the cells were placed in complete growth medium (including 20% fetal calf serum) and maintained in culture for 24 hrs.

To estimate the amount of rhBMP-6 that was secreted after nucleofection, MSCs from three different donors were grown in confluent conditions in culture for 14 days. Media from the flasks were collected on Day 2 after nucleofection. To quantify the rhBMP-6 secreted during 24 hrs, the media were changed 24 hours before sampling. An enzyme-linked immunosorbent assay (ELISA) (R&D Systems, Minneapolis, MN, USA) was performed to measure the amount of rhBMP-6 protein secreted into the culture media by the nucleofected porcine MSCs and compared to the cells nucleofected with GFP reporter gene. After sampling the media, we lifted and counted cells so that we could normalize the secreted protein to 10^6 cells.

2.2. Vertebral Defect Model. The Institutional Animal Care and Use Committee of Cedars-Sinai Medical Center approved all animal procedures used in this study. Six minipigs were used to generate vertebral defects. In each pig, one vertebral defect was generated. Three pigs were treated with fibrin gel (FG) alone (Tisseel, Baxter, IL, USA), and three were treated with BMP6-MSCs (4×10^6) suspended in fibrin gel.

2.2.1. Pre-Op Care and Surgical Anesthesia. Following an 18-hour preoperative fast, each pig was sedated with intramuscular drugs (acepromazine 0.25 mg/kg, ketamine 20 mg/kg, and atropine 0.02–0.05 mg/kg), following which the animal was injected intravenously with propofol (2 mg/kg) to induce full anesthesia. After this had been achieved, the trachea was intubated and anesthesia was maintained using 1–3.5% isoflurane inhaled via the tracheal tube for the duration of the procedure.

Surgical Procedure. A 20-cm posterolateral skin incision was made over the lumbar region (L1–5), which was then exposed by a lateral transposae retroperitoneal approach. One critical-size cylindrical bone defect, 15-mm in depth and 4-mm in diameter, was created in one lumbar vertebra using a surgical drill bit (4.0 JAC 100 SC, Veterinary Orthopedic Implants, St. Augustine, FL). After surgery, the subcutaneous tissue was closed with an absorbable subcutaneous suture and the skin with an absorbable subcuticular suture.

The cDNA3-pCMV-rhBMP-6 nucleofected cells were lifted, counted, and divided to aliquots. Four million cells were suspended in 400 μL fibrin gel (FG; Tisseel kit, Baxter, IL, USA) and then implanted in each vertebral defect. As a control three vertebral defects were treated with 400 μL fibrin gel only (FG only). Next, the subcutaneous tissue layer was closed in a continuous pattern using an absorbable/coated suture and the skin was closed in a subcuticular pattern using a nonabsorbable interrupted over-and-over suture, which was removed 2 weeks after surgery. Finally, the skin area was cleansed with sterile gauzes and 0.5% chlorhexidine gluconate.
2.2.2. Post-Op Care. Buprenorphine (0.1 mg/kg) was administered IM for pain relief immediately after the surgery and every 12 hours during the first day (if needed). Body temperature, pulse, and respiration were closely monitored for potential complications, such as infection.

2.3. Evaluation of Bone Regeneration. Bone regeneration was monitored in vivo at 6 and 12 weeks for the duration of the study (i.e., 6 months) using a clinical computed tomography (CT) scanner [Biograph PET-CT, Siemens] to verify that the defect had not spontaneously repaired. After 6 months, the pigs were immobilized with intramuscular ketamine (20 mg/kg) and acepromazine (0.25 mg/kg) and euthanized using a veterinary euthanasia solution (1 cc/10 lbs) injected intravenously into an ear vein. Next, the vertebrae were examined ex vivo using high-resolution microcomputed tomography ($\mu$CT) and histological techniques.

2.3.1. Ex Vivo Microcomputed Tomography Analysis. Ex vivo, high-resolution assessment of bone regeneration was performed at the end of the study (i.e., 6 months). For this purpose, the animals were euthanized and their spines were subsequently removed. Bone regeneration at the defect site was evaluated using a preclinical cone-beam $\mu$CT imaging system (vivaCT 40; Scanco Medical AG, Brüttisellen, Switzerland). Microtomographic slices were acquired using an X-ray tube with a 55 kVp potential and reconstructed at a voxel size of 35 $\mu$m. The $\mu$CT analysis used for this study was described in detail by Kallai et al. [25]. Briefly, after locating the defect region, the defect margins were aligned to a standard position and a cylindrical volume of interest (VOI; 5 mm in diameter, 15 mm in length) was defined for a 3D evaluation. A constrained 3D Gaussian filter ($\sigma = 0.8$ and support $= 1$) was used to partly suppress the noise in the volumes. The bone tissue was segmented from marrow and soft tissue by using a global thresholding procedure [26]. In addition to visual assessments of the structural images, morphometric indices were determined on the basis of microtomographic datasets by using direct 3D morphometry [27]. Evaluation of regenerated bone tissue was made by a qualitative assessment of bone structure based on 2D cross sections and 3D images and a quantitative assessment of bone structure based on microtomographic datasets created using direct 3D morphometry. The following morphometric indices were determined for newly formed bone in the regeneration sites: (1) volume of mineralized bone tissue [BV, mm$^3$] and (2) connectivity density [Conn-Dens, 1/mm$^2$], which was derived from the Euler number [28], a topologic measure used to describe the porosity of the bone sample and to show the extent of branching in the bone structure.

2.3.2. Histological Analysis. Histological evaluation of bone formation was performed on operated vertebrae after 6 months of study, as previously described [24, 29]. Briefly, the vertebrae were fixed in 4% formalin, decalcified with 0.5 M EDTA in saline (pH 7.4), and embedded in paraffin. Tissue sections were cut at a thickness of 5 microns and subsequently stained using haematoxylin and eosin (H&E) and Masson's Trichrome (MTC) stain.

2.4. Statistical Analysis. The vertebral defect model was created in 6 minipigs consisting of 3 pigs/vertebral defects per group. Results are presented as means ± standard error of means. Statistical tests for significance were performed using unpaired, one tail, Student's t-test (GraphPad Prism, San Diego, CA), and the minimal criterion for significance was determined to be a probability level less than 0.05.

3. Results
Ventral defects, 4 mm in diameter and 15 mm in depth, were successfully generated in the vertebral bodies of minipigs (Figure 1). Porcine BM-MSCs were nucleofected with rhBMP-6 encoding plasmid and the secretion of BMP6 protein was verified using an immunoassay (ELISA, Figure 2).

Following defect establishment and construct implantation, clinical CT imaging at 6 and 12 weeks revealed considerable repair in the vertebral defect treated with MSCs overexpressing BMP6 compared to those treated with fibrin gel only (Figures 3(a), 3(b), 3(e), and 3(f)). High-resolution $\mu$CT analysis of the vertebrae harvested 6 months after surgery showed that the vertebrae architecture was almost completely regenerated in the BMP6-MSCS treated defects. Conversely, the control (fibrin gel only) treated defects less bone repair was visible (Figures 3(c), 3(d), 3(g), and 3(h)). Although significance was not reached, most likely due to the small sample size ($n = 3$), quantitative data derived from the $\mu$CT analysis showed higher connectivity density and bone volume indices in defects treated with BMP6-MSCs when compared to control (Figure 4). In addition, histological evaluation of the two treatment groups using H&E and MTC staining further demonstrated the favorable effect of the BMP6-modified MSCs on bone regeneration; that is, after 6 months of study, nearly complete defect closure was evident in the BMP6-MSCS group while less bone regeneration was seen in the control group (Figure 5).

4. Discussion
The most common type of injury in osteoporotic patients is vertebral compression fracture (VCF). Current surgical, and nonsurgical, interventions for VCF do not provide appropriate clinical responses to repair and induce new bone formation, frequently leaving patients in pain and/or disability. Bone formation and fracture repair are dependent on the appropriate number and function of resident MSCs. This has inspired investigators to utilize MSCs in a number of preclinical animal models to increase the number of functional MSCs and thereby enhance bone regeneration in vivo [30–40]. Indeed, the field of stem cell therapy has been expanding rapidly over the last two decades. MSCs have also been used in conjunction with gene delivery methods to express various bone inducing factors, such as BMPs, to further augment the bone formation process. Various studies have reported promising results involving genetically
modified stem cells as therapeutic vehicles to induce bone regeneration in vivo [18, 19, 21, 29, 41–50]. Our group has previously reported the accelerated repair of a vertebral defect in rats using a local injection of BMP6-modified stem cells [19]. The amounts of BMP6 secreted in the current study were comparable to our previous studies in porcine ASCs [21, 24] and BM-MSCs [21]. Based on these promising results, we hypothesized that BMP6-MSCs administration will accelerate bone formation in a vertebral defect of minipigs. To our knowledge, this is the first study that attempts to promote vertebral bone regeneration via implantation of allogeneic, gene modified stem cells in a large animal model.

To test our hypothesis, a 4 mm in diameter and 15 mm deep cylindrical defect was first created in the lumbar vertebral body of minipigs (Figure 1). Following defect establishment, BMP6-MSCs suspended in fibrin gel, or fibrin gel alone, were implanted into the defect for 6 months. At 6 and 12 weeks after surgery, bone regeneration was evaluated via in vivo clinical CT. Already at 6 weeks, it was shown that the defect treated with the BMP6-MSCs began to heal while the defect treated with fibrin gel alone exhibited less new bone growth (Figure 3). At the study completion (i.e., 6 months), the vertebrae were excised for ex vivo analyses using μCT and histology. Similar to the in vivo CT results, defects treated with the BMP6-modified MSCs were almost entirely repaired whereas those treated with control had less bone growth. Quantitative data obtained from CT scans showed a twofold increase in connectivity density as well as an increase (though not significant) in bone volume in defects treated with the BMP6-MSCs, as compared to control (Figure 3).

One limitation of the study is the small sample size used \( n = 3 \). We believe that differences between groups were not significant due to that reason; hence future studies should incorporate more animals to reduce experimental error. Optimization of the technique, such as varying cell loading density and/or type of biomaterial used, may also improve therapeutic outcome. For example, fibrin gel, which has low biomechanical properties, might be replaced by a different biomaterial that may offer an option to increase the vertebral height, while the engineered MSCs induce new bone formation.

An alternative approach might include a systemic treatment. Since MSCs intrinsically home and engraft at sites of injury and inflammation [51, 52], systemic (versus local) delivery of cells may also improve clinical outcome; this is because, unlike local transplantation, cells can migrate toward the wound site in response to physiological cues from the body (such as inflammatory mediators), at the appropriate time and quantity.

One more limitation of the study is the use of healthy animals. Clearly further studies should be conducted in osteoporotic animals. In addition, the feasibility of an image-guided injection of engineered MSCs to the injured vertebra should be demonstrated.

In summary, current treatments for VCFs in osteoporotic patients provide poor patient outcome, an outcome that is not better than sham treatment, which calls for a fresh clinical approach [30]. This pilot preclinical study aims at...
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**Figure 3:** BMP6-MSCs induce vertebral bone repair. Bone regeneration in lumbar vertebral defects was monitored using clinical CT imaging on 6 ((a) and (e)) and 12 weeks after surgery ((b) and (f)). Animals were sacrificed on week 24 (i.e., 6 months) after surgery and excised vertebrae were subjected to μCT imaging. Bone formation was quantified based on μCT scans (analyzed region is highlighted in red, (c) and (g)). Marked differences in bone regeneration can be seen in defects treated with BMP6-MSCs ((g) and (h)) versus fibrin gel only ((c) and (d)).

**Figure 4:** Quantification of bone regeneration in lumbar vertebral defects. Bone volume (BV) and connectivity density (Conn-Dense) parameters were higher in lumbar defects treated with BMP6-MSCs compared to defects treated with fibrin gel only. Statistical significance difference was not reached, probably due to the small sample size (bars indicate SE, \(n=3\)).

bridging this gap by presenting a potentially noninvasive technique that delivers BMP6-modified MSCs via local injection to augment bone formation in a large animal model. Our results elude that this technique may be an attractive alternative to substitute or supplement current therapeutic approaches, particularly because the gene delivery method used is nonviral and considered clinically safe [23, 53]. Still, more preclinical studies are needed in order to verify our results and translate this research method into clinical practice.
Figure 5: Histological sections of vertebral bone defects. Treated vertebrae were harvested and processed for histology. Sections were stained with H&E and Masson’s Trichrome and imaged using a light microscope; representative images showing advanced defect closure in the BMP6-MSC group versus the control (fibrin gel only) group.

Conflict of Interests

The authors declare that there is no conflict of interests regarding the publication of this paper.

Authors’ Contribution

Gadi Pelled, Dmitriy Sheyn, and Wafa Tawackoli contributed equally to this work.

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