Reindeer BMP extract in the healing of critical-size bone defects in the radius of the rabbit

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Background Native BMP extracts from reindeer effectively induce ectopic new bone formation in vivo, but their bone healing properties have not yet been evaluated. We investigated the effect of reindeer BMP extracts on the healing of long bone defects.

Methods The implants tested contained 5 mg or 10 mg of unsterilized BMP extract from reindeer and 10 mg of gamma-sterilized BMP extract administered with collagen carrier (Lyostypt, B. Braun, Germany). 70 µg of rhBMP-2 with collagen carrier (InductOs; Wyeth Europa) served as positive control, and collagen implants (Lyostypt) and untreated defects served as negative controls. New Zealand White rabbits with 1.5 cm of critical-size radius bone defects were used, with 8 weeks of follow-up.

Results Radiographic analysis showed bone formation (BF) to be higher in all groups containing BMPs than in the untreated controls. BF was also higher in the rhBMP-2 group, and marginally higher in the group treated with 10 mg of unsterilized reindeer BMP extract (p = 0.06) as compared to the collagen controls. Bone union (BU) was better in the unsterilized BMP extract groups and rhBMP-2 group than in the untreated controls. BU was also better in the implants with 10 mg of unsterilized reindeer BMP extract and rhBMP-2 than in the collagen-treated implants. The mean area of new bone at the site of the defect proved to be higher in all implants containing BMP than in the untreated defects. It was also higher in the groups with 10 mg of unsterilized BMP extract and rhBMP-2 than in the collagen-treated controls. Mechanical tests showed torsional stiffness of the bones to be higher in the group with 10 mg of unsterilized BMP extract than in the collagen group. The mean cross-sectional bone area measured by pQCT densitometry was higher in the rhBMP-2 group than in the collagen group. The mean bone density at the defect area was higher in the group with 10 mg of unsterilized BMP than in the rhBMP-2 group.

Interpretation We conclude that both reindeer BMP extract and rhBMP-2 induced improved healing of the rabbit radius bone defects at the doses used. Gamma sterilization of reindeer BMP extract reduced osteoinductivity slightly, but not significantly.

Bone morphogenetic proteins (BMPs) stimulate regeneration of bone and cartilage (Jortikka et al. 1993a,b, Cook 1999, Govender et al. 2002, Wozney 2002, Cheng et al. 2002, Pekkarinen et al. 2003). The bone-healing properties of different recombinant and extracted BMPs have been studied using long bone defect models in rabbits (Hollinger et al. 1998, Kokubo et al. 2003), dogs (Sciadini and Johnson 2000, Tuominen et al. 2000), rats (Takagi and Urist 1982, Chen et al. 2002, Matsuo et al. 2003) and sheep (Gao et al. 1996, 1997). Favorable results in preclinical and clinical trials have also led to the regulatory approval of different commercial recombinant BMP products for some clinical purposes by government agencies (Burkus et al. 2002a,b, Govender et al. 2002, Valentin-Opran et al. 2002, Matsuo et al. 2003). Considering that there are about 20 BMPs known, products consisting of single recombinant BMPs cannot be regarded as being ideal. Also, our knowledge
of their long-term effects and of how safe they are is still limited. BMPs occur as a mixture in living organisms, and native BMPs are also a mixture when extracted. Thus, studies aiming at safe and effective products from native BMPs are still needed. We have shown previously that reindeer BMP extracts effectively induce ectopic new bone formation in vivo (Pekkarinen et al. 2003, 2004, 2005a,b). However, the bone healing properties of reindeer BMP extract have not yet been evaluated.

We investigated the effects of reindeer BMP extracts on the healing of critical-size bone defects (CSDs) in the rabbit radius. The bone-inducing capacity of reindeer BMP extracts was compared to untreated controls and collagen-treated controls, while rhBMP-2 (InductOs) was studied as a reference because its bone-healing capacity is well documented (Hollinger et al. 1998, Sciadini and Johnson 2000, Yudell and Block 2000, Govender et al. 2002).

Methods

BMP materials

Native reindeer (Rangifer tarandus) BMP extract was prepared from diaphyseal bone of the reindeer. Cortical bones from each animal were chilled immediately after death. The epiphyseal ends, bone marrow and periosteum were mechanically removed, and after freezing in liquid nitrogen, the cleaned cortical bones were ground to a particle size of 1.0 mm3. The pulverized bone was demineralized in 0.6 M HCl and extracted in 4 M guanidine hydrochloride (GuHCl) at 4ºC. The GuHCl-extracted solution was filtered with a tangential flow system and concentrated. The concentrated solution was dialyzed against deionized water and the water-insoluble material was collected. After redissolving in 4 M GuHCl solution, the water-insoluble material was dialyzed against 0.25 M citrate buffer, pH 3.1. The citrate-buffer-insoluble material was washed with deionized water and lyophilized (Jortikka et al. 1993a).

We used the commercial product of recombinant human bone morphogenetic protein-2 (InductOs; Wyeth Europa, Maidenhead, UK). InductOs is authorized for the treatment of tibial fractures and consists of recombinant human bone morphogenetic protein-2 (rhBMP-2, diboterminalfa), applied to an absorbable collagen sponge (ACS) matrix (type I bovine collagen) (Govender et al. 2002).

Reconstitution of implants and sterilization

5 or 10 mg of reindeer BMP extract was dissolved in 100 µL sterile water and pipetted onto the collagen sponge (25 × 20 mm, Lyostypt compress from native collagen of bovine origin, mainly consisting of type IV collagen; B. Braun, Tutlingen, Germany). The doses were chosen according to our previous studies (Pekkarinen et al. 2003, 2004, 2005b, 2005c). Finally, implants were lyophilized in test tubes for 48 h.

The InductOs kit (Wyeth Europa, Maidenhead, UK) for implant contains 12 mg rhBMP-2, solvent (10 mL sterile water), and absorbable collagen sponge matrix (ACS). The solvent containing 12 mg of rhBMP-2 was made according to manufacturer’s instructions before operation. 46.7 µL of the solvent, thus containing 70 µg of rhBMP-2, was pipetted onto the collagen matrix (10 × 20 mm) to form the implant. The dose of rhBMP-2 was determined from the literature (Hollinger et al. 1998, Bessho et al. 1999, Li et al. 2002).

Control implants were constructed in an identical fashion, but they contained only collagen carrier (Lyostypt). All implants were made under sterile conditions.

Gamma sterilization of the reindeer BMP extract implants was performed by a specialized company (Isotron Ltd., Swindon, UK). The dose of irradiation was 4.10 MRad.

Groups

26 eight-month-old male New Zealand White rabbits were used. Each animal was treated bilaterally with two different implants randomized from the following groups: (1) collagen + 5 mg of unsterilized reindeer BMP extract (n 8), the “5 mg BMP” group; (2) collagen + 10 mg of unsterilized reindeer BMP extract (n 10), the “10 mg BMP” group; (3) collagen + 10 mg of gamma-sterilized reindeer BMP extract (n 10), the “10 mg sterilized BMP” group; (4) collagen + 70 µg of rhBMP-2 (InductOs) (n 8), the “rhBMP-2” group; (5) collagen (n 8), the “collagen” group; and (6) no implant (n 8), the “untreated” group.
**Surgical procedure**

Surgery was performed under general anesthesia (Domitor 0.3 mg/kg, Ketalar 20 mg/kg). 1 mL of prophylactic antibiotic (Procapen, 300,00 IU/mL; Orion, Espoo, Finland) was given to each rabbit preoperatively. Longitudinal skin incisions were made bilaterally over the radial bones at the middle one-third of the front leg. The periosteum was separated from the surrounding muscle. A 15-mm critical-size defect (CSD) located about 2–2.5 cm proximal to the radiocarpal joint was created using a circular saw attached to a dental handpiece. After that, the defect was checked and any remaining periosteum was removed. A thorough wash with saline was done and the implant was applied to the defect (Figure 1). No fixation was used because of the support from the ulna, and unrestricted weight bearing was normally allowed.

After 8 weeks, the rabbits were killed with an overdose of pentobarbital (Mebunat), 60 mg/kg intravenously. The radii were dissected out and the soft tissues were removed. The bones were wrapped in saline and frozen at −20°C until analysis.

**Radiographic evaluation of bone formation**

Radiographs of the radius (100 mA, 20 kV, 0.08 s/exp; Mamex de Maq, Soredex, Orion) were taken after 8 weeks of implantation (Figure 2). The percentage of new bone formation (BF) within the defect and the development of the union or non-union (BU) of the bone was estimated clinically by two independent investigators according to the scoring system presented by Sciadini et al. (1997). Any cases of disagreement were reviewed together. Interpretations between the different groups were done blind.

After clinical estimation, radiographic images were digitized using an optical scanner (HP ScanJet 4070). Bone formation was evaluated as the area (in mm²) of the calcified tissue visible on the region of the bone defect by using Scion Image Beta 4.02 (Scion Corp., MD, USA) software. The mean optical density (mmAl) of the defined area was measured with the same software. Calibration of the optical density was performed by using an aluminium wedge that was imaged with the bones.

**Computed tomography**

After thawing, the bones were scanned using a peripheral quantitative computed tomography (pQCT) device (Stratec XCT 960A, software version 5.21; Norland Stratec Medizintechnik GmbH, Birkenfeld, Germany). A voxel size of 0.148 × 0.148 × 1 mm³ was used. We scanned 3 slices of each sample: 1 slice in the middle of the implant and 1 at each end, 5 mm from the middle slice. The positions of the slices were defined by using the axial scout view of the pQCT system. Bone density (mg/cm³), and cross-sectional bone area (mm²) were recorded for each slice as given by the pQCT software, using an attenuation threshold of 0.4 c⁻¹ (169 mg/cm³) for bone. Average values for the 3 slices were used in the statistical analysis.
Bone formation in the defect area of rabbit radius evaluated by radiographic analysis, mechanical testing and pQCT densitometry. Data are presented as median (95% CI)

| Groups                | n   | Optical density (mmAl) | Bone formation (BF) | Bone union (BU) | Breaking load (Nm) | Stiffness (Nm/deg) | Energy to failure (Nm deg) | Cross-sectional area (mm²) | Bone density (mg/cm³) |
|-----------------------|-----|------------------------|---------------------|-----------------|--------------------|---------------------|---------------------------|--------------------------|---------------------|
| 5 mg of BMP           | 5   | 2.0 c                  | 4 a                 | 3 a             | 0.33               | 0.019 c             | 3.9                       | 30                       | 611                 |
| 95% CI                |     | (0.6–4.0)              | (2–4)               | (0–3)           | (0–0.49)           | (0–0.048)           | (0–24)                    | (11–47)                  | (422–722)           |
| 10 mg of BMP          | 7   | 1.7 c                  | 4 b c               | 3 a d           | 0.46               | 0.039 d             | 5.3                      | 22                       | 744                 |
| 95% CI                |     | (0.9–3.3)              | (3–4)               | (0–3)           | (0–0.88)           | (0–0.069)           | (0–9.5)                  | (13–31)                  | (650–828)           |
| 10 mg of st. BMP #    | 7   | 1.5 d                  | 4 b                 | 3 a             | 0.21               | 0.011               | 3.2                      | 27                       | 716                 |
| rhBMP-2               | 7   | 2.5 d                  | 4 b d               | 3 a d           | 0.46               | 0.031               | 3.2                      | 18–39                    | (583–806)           |
| 95% CI                |     | (0.8–4.3)              | (4–4)               | (0–3)           | (0–0.46)           | (0–0.031)           | (0–11)                   | (18–39)                  | (511–697)           |
| Untreated             | 5   | 1.6                    | 2                   | 0              | 0.23               | 0.020 e             | 2.0                      | 33 d                     | 611                 |
| 95% CI                |     | (0.7–2.5)              | (1–3)               | (0–2)           | (0–0.43)           | (0–0.037)           | (0–8.3)                  | (19–49)                  | (514–874)           |
| Collagen              | 5   | 1.7                    | 1                   | 0              | 0.36               | 0.057               | 4.8                      | 14–37                    | 722                 |
| 95% CI                |     | (0.0–3.4)              | (0–4)               | (0–3)           | (0–0.64)           | (0–0.020)           | (0–24)                   | (5.4–29)                 | (340–827)           |

a p < 0.05, b p < 0.01 vs. untreated.

Mechanical tests
The radii of the rabbits were thawed to room temperature for torsional testing. During the mechanical tests, the bones were kept moist. The ends were embedded into the molds with dental stone (GC Fujirock Improved Dental Stone; G-C Dental Industrial Corp., Tokyo, Japan). Torsional shaft was 3.5 cm. After hardening of the dental stone, the bones were placed in the torque machine (Lepola et al. 1993) and torsionally loaded at a constant angular speed of 6 degrees per sec until failure. Maximum breaking load (Nm), torsional stiffness (Nm/degree) and energy to failure (Nm degree) were recorded (Jämsä and Jalovaara 1996).

Statistics
Statistical analysis was performed using the SPSS statistical package and the Confidence Interval Analysis software (version 2.0.0; University of Southampton). The non-parametric Kruskall-Wallis test was used to evaluate the statistical differences between the groups, and the Mann-Whitney test for pairwise comparison between each BMP group and the untreated and collagen groups (negative controls), and between the reindeer BMP groups and the rhBMP group (positive control).

Values of p < 0.05 were considered statistically significant. Results are given as median and 95% confidence limits, using the Wilcoxon method for continuous variables and the binomial method for grading variables.

Results
Because of physeolysis, 7 rabbits had to be killed before the endpoint. Thus, since the animals were treated bilaterally, 3 samples from the 5 mg BMP group, the 10 mg BMP group, the 10 mg sterilized BMP group, and the untreated groups, and 1 sample from both rhBMP-2 and collagen groups were excluded from the study.

Bone formation and bone union evaluated clinically from radiographs
Bone formation (BF) at the defect site was higher in the groups with 5 mg BMP (p = 0.04), 10 mg BMP (p = 0.004), 10 mg sterilized BMP (p = 0.02), and rhBMP-2 (p = 0.001) than in the untreated group. BF was also higher in the rhBMP-2 group (p = 0.02) and marginally higher in the group with 10 mg BMP (p = 0.06) than in the collagen group (Table).
Bone union (BU) was higher in the groups with 5 mg BMP (p = 0.04), 10 mg BMP (p = 0.01), and rhBMP-2 (p = 0.01) than in the untreated group. BU was also higher in the groups with 10 mg BMP (p = 0.04) and rhBMP-2 (p = 0.04) than in the collagen group (Table).

There were no significant differences in bone formation and bone union between the groups containing BMPs (Table).

**New bone area and optical density evaluated by radiography**

Mean new bone area (mm²) at the defect site was higher in the groups with 5 mg BMP (p = 0.03), 10 mg BMP (p = 0.02), 10 mg of sterilized BMP (p = 0.04) and rhBMP-2 (p = 0.004) than in the untreated group. Also, the new bone area was higher in the groups with 10 mg BMP (p = 0.03) and rhBMP-2 (p = 0.04) than in the collagen group. There were no significant differences in area of new bone between the groups containing BMPs (Figure 3).

There were no significant differences in mean optical density (mm^2 Al) of new bone between the different groups (Table).

**Mechanical testing**

Mechanically unstable bones were not tested, and their values were considered to be zero in the statistical analysis. Stiffness of the bones (in Nm/deg) was significantly higher in the group with 10 mg BMP than in the collagen group (p = 0.02). There were no other statistically significant differences between the groups in the mechanical tests (Table).

**Cross-sectional bone area and bone density evaluated by pQCT**

Computed tomography (pQCT) showed cross-sectional bone area to be higher in the rhBMP-2 group than in the collagen group (p < 0.05). Bone density was higher in the 10 mg BMP group than in the rhBMP-2 group (p < 0.05).

**Discussion**

BMPs combined with different carrier materials have been shown to heal experimental bone defects in many studies (Moore et al. 1990, Gao et al. 1996, 1997, Viljanen et al. 1996, Hollinger et al. 1998, Sciadini and Johnson 2000, Tuominen et al. 2000, Yudell and Block 2000, Chen et al. 2002, Kokubo et al. 2003, Barboza et al. 2004). Until now, there have been no studies investigating the ability of reindeer BMP extract to heal long bone defects. We have shown previously that reindeer BMP extract is an effective inducer of new bone formation in a muscle pouch model in mice (Pekkarinen et al. 2003, 2004, 2005b, c). In the present study, we compared the bone healing properties of the reindeer BMP extract to results with untreated and collagen controls, while rhBMP-2 (InductOS) was also included in the comparisons because the bone healing capacity of rhBMP-2 has been well documented in experimental animal and clinical studies (Hollinger et al. 1998, Sciadini and Johnson 2000, Yudell and Block 2000, Govender et al. 2002, Li et al. 2002).

Many previous studies concerning the ability of BMPs to promote healing of long bone defects have been performed using synthetic polymer carriers or composite implants containing collagen and frames such as coral, tricalcium phosphate or hydroxyapatite (Gao et al. 1996, 1997, Tuominen...
et al. 2000, 2001, Hu et al. 2003, Kokubo et al. 2003). In the present study we did not use composite implants because the rabbit radius is a relatively small bone and its critical-size bone defect does not require extra fixation or a carrier that is mechanically stronger than collagen. BMPs have also been used successfully in the healing of fractures with collagen carrier alone (Bax et al. 1999, Blokhuis et al. 2001, Govender et al. 2002, Einhorn et al. 2003).

Hollinger et al. (1998) used rhBMP-2 with collagen carrier in the healing of rabbit radial critical-size defects. After 8 weeks, rhBMP-2 treatment regenerated osseous contours and new bone formation was significantly higher in the rhBMP-2 or autograft groups than in the collagen or untreated groups. Using a radial bone defect in dogs, Sciardini and Johnson (2000) reported that rhBMP-2 with a collagen sponge carrier had potential osteoinductive activity as measured by biomechanical parameters when evaluated 12 and 24 weeks postoperatively. Defects treated with rhBMP-2 implants were comparable to those treated with autograft, and significantly stronger than those treated with placebo. Li et al. (2002) enhanced bone consolidation of distraction osteogenesis in a rabbit model by using rhBMP-2 with an absorbable collagen sponge carrier. They concluded that rhBMP-2 enhances the consolidation stage of distraction osteogenesis in the rabbit tibia. Also, Yudell and Block (2000) showed that rhBMP-2 with absorbable collagen sponge carrier has the potential to stimulate bone gap healing in the zygomatic arch osteotomies.

Our results are in line with those of previous studies. Here, rhBMP-2 effectively stimulated the healing of the rabbit critical-size bone defect. Reindeer BMP extract also induced bone healing effectively and its effect was comparable to that of rhBMP-2 at the doses used. These findings confirm our previous results showing that reindeer BMP extract has good osteoconductivity in the muscle pouch model of mice (Pekkarinen et al. 2003, 2004, 2005a, b). Bessho et al. (1999) showed the osteoinductivity of rhBMP-2 to be less than one-tenth of the osteoinductivity of human BMP extract, but that conclusion does not match the findings of our study.

Previous studies on the bone healing capacity of purified BMP extracts have been done using different extracts and different carrier types than used here, but the results have still been more or less in line with ours (Moore et al. 1990, Gao et al. 1996, 1997, Viljanen et al. 1996, Tuominen et al. 2000, 2001). Moore et al. (1990) showed bovine BMP extract and Viljanen et al. (1996) moose BMP extract with collagen carriers to be effective in the healing of experimental bone defects. Gao et al. (1996) healed segmental tibial defects of sheep by using composite implants containing sheep BMP extract, type IV collagen, and tricalcium phosphate cylinder. They concluded that this composite implant possessed good osteoinductivity, osteoconductivity and mechanical strength. Tuominen et al. (2000) used a canine ulnar defect model to evaluate the bone healing capacity of composite implants containing bovine BMP extract, coral and collagen. They concluded that the BMP composite enhanced bone healing but the effect was not as good as that of corticocancellous autograft.

Our previous studies have shown that gamma sterilization of the reindeer BMP extract does not reduce its osteoinductivity significantly in vivo (Pekkarinen et al. 2004, 2005a, c). This study is by and large in line with our previous findings, and no significant difference between unsterilized or gamma-sterilized reindeer BMP extract could be seen. However, even though the gamma-sterilized reindeer BMP extract healed bone defects well, it appeared to be slightly inferior than unsterilized reindeer BMP extract or rhBMP-2.

In this animal model, it is advisable to use adult rabbits to prevent epiphyseal slipping. We used 8-month-old rabbits, but some animals underwent physeolysis during the evaluation time. These animals had to be killed before the endpoint, and they were therefore excluded from the study. Also, we observed only slight differences between the study groups in the mechanical properties of the bones. This was mainly due to the small sample number caused by physeolysis and the limited ability of mechanical testing to discriminate bones with non-union. In spite of this, the results were quite obvious.

The theoretical risk of transmissible spongiform encephalopathy has limited the use of materials of natural origin. Thus, there might be concern about the use of native reindeer proteins. However, prion disease has never been described in reindeer. Fur-
thermore, the manufacturing process and gamma sterilization of the BMP extract destroys prions and virus particles (Miekka et al. 2003).

We conclude that both reindeer BMP extract and rhBMP-2 (InductOs) effectively induced the healing of rabbit radius bone defects at the doses used. Gamma sterilization of the reindeer BMP extract did not reduce its osteoinductivity substantially, but the results were slightly inferior to those with unsterilized reindeer BMP extract or rhBMP-2.

Contributions of authors

PJ, TP and TJ: planned the study. PJ and TJ: conducted the study. OH: participated in the preparation of the BMP extract. TJ and OH: prepared the implants. TP: performed the surgery. TP and PJ: radiographic evaluation. TP, MM and TJ: performed the pQCT measurements, mechanical testing and statistical analyses. TP, PJ, TJ, MM and OH: interpreted the results and wrote the manuscript.

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