■ RESEARCH

High tibial osteotomy using polycaprolactone-tricalcium phosphate polymer wedge in a micro pig model

H.-C. Lim,
J.-H. Bae,
H.-R. Song,
S. H. Teoh,
H.-K. Kim,
D.-H. Kum

From the Korea
University College of
Medicine, Seoul,
Republic of Korea

Medial open-wedge high tibial osteotomy has been gaining popularity in recent years, but adequate supporting material is required in the osteotomy gap for early weight-bearing and rapid union. The purpose of this study was to investigate whether the implantation of a polycaprolactone-tricalcium phosphate composite scaffold wedge would enhance healing of the osteotomy in a micro pig model. We carried out open-wedge high tibial osteotomies in 12 micro pigs aged from 12 to 16 months. A scaffold wedge was inserted into six of the osteotomies while the other six were left open. Bone healing was evaluated after three and six months using plain radiographs, CT scans, measurement of the bone mineral density and histological examination.

Complete bone union was obtained at six months in both groups. There was no collapse at the osteotomy site, loss of correction or failure of fixation in either group. Staining with haematoxylin and eosin demonstrated that there was infiltration of new bone tissue into the macropores and along the periphery of the implanted scaffold in the scaffold group. The CT scans and measurement of the bone mineral density showed that at six months specimens in the scaffold group had a higher bone mineral density than in the control group, although the implantation of the polycaprolactone-tricalcium phosphate composite scaffold wedge did not enhance healing of the osteotomy.

■ H.-C. Lim, MD, PhD,
Professor
■ H.-R. Song, MD, PhD,
Professor
■ H.-K. Kim, PhD, Student
Department of Orthopaedic
Surgery
Korea University Guro
Hospital, 80 Guro-Dong, Guro-
Gu, Seoul 152-703, Korea.

■ J.-H. Bae, MD, PhD, Assistant
Professor
■ D.-H. Kum, MD, Resident
Department of Orthopaedic
Surgery
Korea University Ansan
Hospital, Gojan 1-Dong,
Danwon Gu, Ansan-Si,
Gyeonggi-Do 425-707, Korea.

■ S. H. Teoh, PhD, Professor
Department of Mechanical
Engineering
National University of
Singapore, 9 Engineering Drive
1, Singapore 117576.

Correspondence should be sent
to Professor J.-H. Bae; e-mail:
osman@korea.ac.kr

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Recently, in order to try and overcome the disadvantages of autograft and allograft,¹ attempts have been made to find adequate supporting material for augmentation after open-wedge high tibial osteotomy. Hydroxyapatite,^{2,3} β -tricalcium phosphate⁴⁻⁷ or a combination of both⁸ are currently the most commonly used synthetic augments. Good clinical results have been reported with biocompatibility, absence of toxicity and non-inflammatory reactions, but relatively slow biodegradation and brittleness limit their application to load-bearing sites.² A more recent approach is to use a composite of bioceramic and a resorbable polymer.^{9,10}

Porous polymer scaffolds are promising materials for tissue engineering because they offer a physical, three-dimensional (3D) support and serve as a template for cell proliferation and ultimately the ingrowth of new tissue. Recently, polycaprolactone-based composite scaffolds fabricated by fused deposition modelling have been investigated,¹¹⁻¹⁴ and shown to have a highly reproducible internal structure based on layer-by-layer deposition of thermoplastic material. Such scaffolds were designed to retain their mechanical properties for five to six months and then gradually degrade over a period of two

years.¹⁵ These scaffolds have a fully interconnected network of pores, which supports ubiquitous proliferation of cells^{16,17} while retaining their mechanical properties after implantation in experimental animal models.¹⁸⁻²⁰

Rapid bone healing and early weight-bearing are clinically important, especially in patients with simultaneous bilateral open-wedge osteotomies. Many research teams have suggested that the use of supporting material could protect against collapse of the opening osteotomy and accelerate bony healing. However, to our knowledge, there are no reports which compare non-augmented with augmented medial open-wedge high tibial osteotomy. The purpose of this study was to compare a non-augmented group with a polycaprolactone-based composite scaffold implantation group after open-wedge high tibial osteotomy in a micro pig animal model, in order to assess whether the implantation of a polycaprolactone-based composite scaffold would produce more rapid bone healing.

Materials and Methods

Scaffolds were fabricated with polycaprolactone-tricalcium phosphate (ratio of composition 80:20) microfilaments (260 μ m to 370 μ m) by

Fig. 1

Photograph showing the scaffold designed to fit the osteotomy site with a height of 9 mm, length of 20 mm and width of 20 mm.

fused deposition modelling as previously described.^{15,17,19,21} Fused deposition modelling, a rapid prototyping technology, was used to produce scaffolds with a honeycomb-like structure, and a fully interconnected network of channels, with controllable porosity and size of channel. The adhesion of each layer is achieved in a hydrostatic pressure chamber (from 0.01 to 10 atmospheres) by subjecting the block scaffold to a temperature just below the melting point of the polymer (55°C to 60°C) for 30 to 45 minutes to allow proper sintering of each layer. Each scaffold manifests a lay-down pattern of 0°/60°/120° with a porosity of 70% and average pore size of 0.515 mm as determined by micro-CT analysis. The scaffolds have a compressive modulus of 23.1 MPa (SD 6.16) and a compressive strength of 6.38 MPa (SD 0.82). A lay down pattern of 0°/60°/120° was used to make a honeycomb array of interconnected equilateral triangles with a regular porous shape. In order to fit the osteotomy site, the scaffold was designed with a height of 9 mm, a length of 20 mm and a width of 20 mm (Fig. 1).

The research protocol was approved by our institution. Medial open-wedge osteotomies were performed on the proximal tibia in 12 micro pigs (12 to 16 months old) weighing 30 kg to 50 kg. The control group (n = 6) had no augmented materials at the site of the osteotomy but the scaffold group (n = 6) had a polycaprolactone-tricalcium phosphate scaffold wedge implanted into the opening gap (Fig. 2). At three months and six months after operation, three pigs in each group were killed and the osteotomies evaluated by CT scan, bone densitometry and histological evaluation.

Operative procedure. All procedures were performed under general anaesthesia and aseptic conditions. An oblique skin incision 5 cm to 7 cm long, was made parallel to the upper border of the sartorius fascia. A guide pin was positioned 3 cm below the joint line at the level of the tibial tuberosity and advanced obliquely from the anteromedial aspect of the proximal tibia to 1 cm below the joint line at the posterolateral cortex. The osteotomy was created using a

Fig. 2

Photograph showing the osteotomy surface after opening; the scaffold wedge has been implanted into the opening gap.

saw and osteotome, preserving the lateral cortex. The osteotomy surface was opened up and widened with a spreader. In the control group, a short 9 mm spacer plate was then placed at the anterior aspect of the superficial medial collateral ligament. The plate was fixed with a proximal and a distal screw. In the scaffold group, a custom-made polycaprolactone-tricalcium phosphate scaffold wedge was implanted into the gap. It was fashioned to be 9 mm high and 20 mm wide. Then a non-spacer plate was applied with a proximal and a distal screw. The soft tissues and skin were closed.

On the first post-operative day and at each month after operation, anteroposterior and lateral plain radiographs were taken under intravenous anaesthesia to assess the progression of bone union, as well as any loss of correction, failure of fixation or collapse of the osteotomy site. Non-union was defined as no progression of bone bridging across the opening site at three or six months after operation with sclerotic margins at the periphery of the wedge. Loss of correction at the osteotomy site was determined by the change of the anatomical medial proximal tibia angle on the serial plain radiographs.

The CT density was measured to assess the formation of new bone at three and six months after operation. All CT images were acquired using a 64-channel multidetector row CT scanner (Brilliance 64; Philips Medical System, Cleveland, Ohio) with detector collimation of 16 mm × 0.75 mm, and tube energy and current of 120 kV and 250 mAs, respectively, 0.7 mm beam pitch, and in the osteoscaning mode. The thickness of the display section was 2/1 mm (section thickness/section gap) and the image thickness for secondary raw data used in multiplanar reformatting was 1/0.5 mm.

Table I. Histological grade of healing of bone defect²²

Grade	Histological findings
0	No new bone
1	New bone only at the inner edge of opening gap
2	New bone at one-third of the opening gap
3	New bone bridging between one-third and two-thirds of the opening gap
4	New bone bridging the entire opening gap but not united
5	Clearly united and remodelling

Table II. The ratio of mean CT density (C1/C2)* in both groups

	Three months	Six months	p-value
Control group	1.41	1.33	0.021
Scaffold group	1.39	1.11	0.013
p-value	> 0.05	0.037	

*C1, new bone formation area in the opening gap; C2, proximal metaphysis above the osteotomy site

Eight or nine coronal CT images were obtained for each specimen. An independent musculoskeletal radiologist measured the CT density value (Hounsfield Units) of the area of new bone formation (C1) in the opening gap and in the proximal metaphysis (C2) above the osteotomy site using the Picture Archiving and Communications System (STARPACS; Infinitt Healthcare Co., Seoul, Korea). The average CT density value of C1 and C2 was calculated from all the coronal images in each specimen. The ratio of CT density of C1 to C2 was calculated and compared between the control and scaffold groups.

At three and six months after operation, the distribution of bone mineral density (BMD) was measured using dual energy x-ray absorptiometry (QDR 1000; Hologic, Bedford, Massachusetts). After scanning, the BMD of the two regions of interest was recorded; namely the area of new bone formation in the opening gap (R1) and the proximal metaphysis above the osteotomy site (R2). The ratio of the BMD value of R1 to R2 was calculated and compared between the control and scaffold groups.

Each specimen was fixed in 10% neutral-buffered formalin for two weeks and then processed for histology at three and six months after operation. Dehydration was accomplished using a graded series of ethyl alcohol (Harleco, Gibbstown, New Jersey) and cleared with xylene substitute (Thermo Electron Corporation, Waltham, Massachusetts). Infiltration was performed using a graded series of methylmethacrylate solutions (ACROS Organics, Morris Plains, New Jersey). Specimens were embedded using a catalysed mixture of Oste-Bed resin solution containing 2.5 g of benzoyl peroxide (Sigma-Aldrich, St Louis, Missouri) per 100 ml under vacuum for at least 48 hours. After polymerisation, specimens were placed in a freezer for 24 hours. They were then trimmed of excess plastic (Isomet 1000; Buehler, Lake Bluff, Illinois), ground to expose the relevant portions (Ecomet 3; Buehler) and sectioned longitudinally

(3 μ m to 4 μ m thick) using a microtome (Microm HM355S; Richard Allan Scientific, Kalamazoo, Michigan). Sections from each specimen were then stained with haematoxylin and eosin to evaluate new bone formation under a light microscope. The healing of regenerated bone was graded using a six point scale (Table I).²²

Statistical analysis. The differences in the pixel values, CT density and BMD values between the two groups was assessed using the Mann-Whitney U test. Statistical analyses were performed using the SPSS (SPSS for Windows Release 15.0; SPSS Inc., Chicago, Illinois). A p-value < 0.05 was considered statistically significant.

Results

Bone union was achieved at six months after operation in both groups. The trabecular continuity began at the inner portion of the osteotomy site and progression of trabecular continuity to the outer portion was observed in all specimens on serial plain radiographs. After operation the mean anatomical medial proximal tibial angle in the control group was 92° (SD 2.4), and 94° (SD 3.1) in the scaffold group. There was no significant change of angle at the final follow-up ($p > 0.05$) and valgus limb alignment was maintained well without loss of correction, failure of fixation or collapse of the osteotomy site in either group.

In both groups, partial bone formation was identified at the inner portion of the osteotomy site. The mean ratio of the CT density showed significant differences between the specimens at three and six-months in both the control and the scaffold groups ($p = 0.021$ and $p = 0.013$, respectively, Table II). At six months the specimens in the scaffold group showed higher levels of BMD ($p = 0.037$, Fig. 3).

The mean ratio of the BMD of new bone formation at three- and six-month specimens showed significant differences between the control and scaffold groups ($p = 0.033$ and $p = 0.045$, respectively, Table III). At six months the specimens showed a higher level of BMD in the scaffold group ($p = 0.027$, Table III).

The histological grading was consistent with the CT density and the BMD (Table IV). New bone formation increased with time and the mean grade in the scaffold group was higher in the control group. In the control group, the defect in the gap was filled with mature bone and loosely organised fibrous tissue. The six-month specimens showed more extensive formation of new bone. In the

Fig. 3a

Fig. 3b

Coronal CT scan images showing six-month specimens from a) the control and b) the scaffold group. Bone formation is more evident in the scaffold group.

Table III. The ratio of mean bone mineral density (R1/R2)* in both groups

	Three months	Six months	p-value
Control group	0.69	0.81	0.033
Scaffold group	0.62	0.91	0.045
p-value	> 0.05	0.027	

*R1, new bone formation area in the opening gap; R2, proximal metaphysis above the osteotomy site

Table IV. The mean histological grading data

	Three months	Six months	p-value
Control group	2.5	3.2	0.031
Scaffold group	2.9	4.1	0.035
p-value	> 0.05	0.021	

scaffold group, there was new bone formation within the interconnected pores of the scaffold and bone ingrowth from the host bone along the periphery of the scaffold (Fig. 4). There was no inflammatory reaction in either group.

Discussion

We wished to determine whether implantation of a polycaprolactone-based composite scaffold would produce faster bony consolidation compared with a non-augmented control group. Our CT scan and histological findings showed a higher level of BMD in the polycaprolactone-tricalcium phosphate scaffold group six months after open-wedge high tibial osteotomy. However, augmentation with a polycaprolactone-tricalcium phosphate scaffold wedge

Fig. 4a

Fig. 4b

Photomicrographs (haematoxylin and eosin, $\times 1.25$) of a six-month specimen from the scaffold group showing a) new bone formation within the interconnected pores between the equilateral rods of scaffold (dark brown) and b) bone ingrowth into the scaffold from the adjacent host bone.

did not show faster bony consolidation when compared with the non-augmented group.

In the context of proximal open-wedge tibial osteotomy, the limited availability of adequate autologous bone, and the morbidity associated with its harvest, has encouraged a search for appropriate substitute materials. Recent studies have reported that the use of hydroxyapatite alone or in combination with tricalcium phosphate is satisfactory.^{2,4,6} In these studies the histological and radiological findings showed a good resorption of the supporting material with complete incorporation and remodelling into new bone with increasing bony consolidation. However, there were no unaugmented control groups for comparison. One clinical study²³ demonstrated that hydroxyapatite/tricalcium phosphate

scaffolds promoted ingrowth of bone but showed no faster bony consolidation or any other additional benefits compared with the non-augmented osteotomy site with an opening angle $< 10^\circ$. Likewise we also found that the implantation of polycaprolactone-tricalcium phosphate scaffold had no additional beneficial effects on bony consolidation when compared with a non-augmented group using a 9 mm bone defect in an animal model.

Most surgeons do not perform a bone graft in proximal tibial osteotomies with a gap < 10 mm, because they believe that it is unnecessary. However, the rapid healing and early weight-bearing are clinically important, especially in patients with simultaneous bilateral osteotomy, even though the osteotomy gap may be small.

The scaffold used in the present study had a honeycomb array of interconnected equilateral triangles with a lay down pattern of $0^\circ/60^\circ/120^\circ$ and a porosity of 70%. The constructs exhibited bone formation, remodelling and maturation along with rapid vascularisation *in vivo*.^{17,18,24,25} This scaffold architecture seemed to have favoured bone maturation and remodelling.^{17,24} However, differences could not be found between two groups, but an increased level of BMD was observed at six months in the scaffold group. This fully interconnected architecture was filled in all scaffolds by new bone formation and bone ingrowth from host bone.

We observed that the polycaprolactone-tricalcium phosphate scaffold remained in all specimens six months after operation. It has been found that the degradation of polycaprolactone is remarkably slow, requiring three years for complete removal from the host body.²⁵ The addition of tricalcium phosphate particles increases the surface area available for hydrolysis, which may accelerate degradation of the scaffold. It is important to match the degradation rate to the rate of deposition of mineralised matrix in order to maintain the structural integrity of the defect while still promoting the remodelling of bone tissue, which takes at least 10 to 15 months *in vivo*. Rapid degradation can limit its application in engineering the induction of new bone tissue, especially at load-bearing sites.

This study has several limitations. The different types of plates (non-spacer plate *vs* spacer plate) between the two groups may have affected the results. The primary stability of fixation is important for complete bony consolidation. Clinically, a less rigid construct can result in nonunion and loss of correction. It is possible that the spacer plate provided more stability than a non-spacer plate. However, because of lack of biomechanical data, it is not known which construct was more stable in our animal model. The polycaprolactone-tricalcium phosphate scaffold gives additional mechanical strength to the osteotomy with a non-spacer plate construct.

The implantation of the polycaprolactone-tricalcium phosphate composite scaffold wedge did not enhance healing of the osteotomy or provide any other additional

benefits compared with the non-augmented osteotomy group at six months after open-wedge high tibial osteotomy in a micro pig model. The clinical relevance of this implant will require further study.

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