

Custom-made Articulating Spacer in Two-stage Revision Total Knee Arthroplasty. An Early Follow-up of 14 Cases of at Least 1 Year After Surgery

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Published online: 8 May 2007
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Abstract Total knee replacement (TKR) infection represents only a small percentage of all the potential complications in joint replacement, but one that can lead to disastrous consequences. Two-stage revision, which has been proven to be the most effective technique in eradicating infection, includes prosthesis removal, positioning of an antibiotic-loaded spacer, and systemic antimicrobial therapy for at least 6 weeks. It has been suggested that there is better performance in terms of range of motion, pain, extensor mechanism shortening, and spacer-related bone loss if articulating spacers are used instead of fixed spacers. In this paper, we describe our results in two-stage revision of infected total knee arthroplasty with a minimum follow-up of 12 months on 14 patients treated by antibiotic-loaded custom-made articulating spacer as described by Villanueva et al. (Acta Orthop 77(2):329–332, 2006). The mean flexion achieved after the second stage of the revision was 120°, ranging from 97° to 130°. The mean Hospital for Special Surgery score was 84. At 1 year after surgery, none of the knees showed any evidence of recurrence of the infection. Articulating spacers are a suitable alternative to fixed spacers with good range of motion after reimplantation and effectiveness against total knee replacement deep infections.

Introduction

Deep infection rates in total knee joint replacement vary in the published literature between 0.3% and 2.9% [2, 3]. Although newer surgical techniques and modifications in preoperative and postoperative care have lowered the overall rate of infections, infection remains a devastating complication for a patient. There are several options to treat an infected knee replacement, the choice of which depends on the time of onset, the microorganism, and radiological exams. These options range from prosthesis retention [13] for early infections up to revision arthroplasty, resection arthroplasty, and arthrodesis.

Literature data [4–7, 10, 16] support the effectiveness of two-stage revision using cement antibiotic combined with intravenous therapy spacers. Fixed spacers are not without their own potential complications including bone loss, pain, and muscle and extensor mechanism contracture.

In an attempt to obviate these problems, surgeons have considered using mobile spacers, which allow partial weight bearing and a limited range of motion up to 90° of flexion [11]. Mobile spacers can be fashioned in different manners. In one technique, the removed femoral component is steam-sterilized and loosely reimplanted with antibiotic-loaded cement [10]. On the tibial side, a new polyethylene insert is again loosely implanted with antibiotic-loaded cement. In another technique [12], pre-molded antibiotic spacers (Stage One [Biomet, Warsaw, IN] or Prostalac) are used. In a third technique, first described by Villaneuva et al. [1], Ha [14], and Goldstein [15], custom-molded spacers were used. This is the technique that we used in our series.

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Materials and methods

Between June 2005 and February 2006, we performed 14 (11 females, 3 males) two-stage revision arthroplasties



Fig. 1. Bone debrided ready for spacer implantation



Fig. 4. Posterior condyles



Fig. 2. Preparing femoral component in a doughy phase



Fig. 5. Tibial preparation



Fig. 3. Refinishing femur with curved osteophome



Fig. 6. Tibial refining



Fig. 7. Micromilling trochlear groove

(TKR) revisions using custom mobile spacers. Each patient was diagnosed as having a deep infection, supported by joint fluid aspiration, leukocytes bone scan, and radiological signs of loosening.

Patients' mean age at the time of revision was 68 years (range 60–76 years). The mean period since index knee replacement was 2.3 years, ranging from 12 months to 3 years. The mean flexion was 73.5° with a mean flexion contracture of 4°.

The isolated microbial agent was *Staphylococcus epidermidis* in 10 cases, whereas the others were *Staphylococcus aureus* and *Pseudomonas aeruginosa*. Every agent was found sensible to Tobramycin, except for the two cases of *S. aureus* isolation that were found sensible to Vancomycin.

Each patient was treated with a two-stage revision using an articulating antibiotic-loaded spacer as described by Villanueva et al. [1]. Simplex bone cement (Kalamazoo, Michigan, USA) was mixed with powdered antibiotic, the choice of which was dependent on the microbial isolation. We added 1 g/dose of Vancomycin to the cement in those cases in which microbial contamination was resistant to Tobramycin (already in the powered part of Simplex added cement). The cement is then molded on the debrided bone using curved retractors and a drill to reproduce the trochlear groove (Figs. 1, 2, 3, 4, 5, 6, 7, 8, 9, 10, and 11).



Fig. 8. Final spacer



Fig. 9. Intraoperative image

The postoperative rehabilitation protocol was the same for each patient. On the first postoperative day, patients were started on continuous passive motion. They were also allowed to begin ambulating bearing weight on the affected extremity, using a brace to lock the knee in full extension. No patient was allowed to flex the knee above 90° till the revision. Parenteral organism specific intravenous antibiotics therapy was started as soon as joint aspiration confirmed infection, and continued for at least 6 weeks. Before revision, a new aspiration was performed, confirming no microbial contamination and no signs of inflammation in joint fluid. The second stage consisted of the removal of the articulating spacer and then reimplantation of a new prosthesis. The minimum time for reimplantation was 6 weeks and the maximum was 11 weeks with a mean value of 9 weeks. In four patients, a rotating RHM hinge (Zimmer, Warsaw, IN) was used and in three patients, Genesis II prosthesis (Genesis II, Smith & Nephew, Memphis, TN) was implanted, whereas in seven further patients, a constrained condylar CCK (Zimmer, Warsaw, IN) was used.

At the time of the second stage of the revision, we observed neither further bone loss nor patellar tendon



Fig. 10. Knee flexion at wound closure

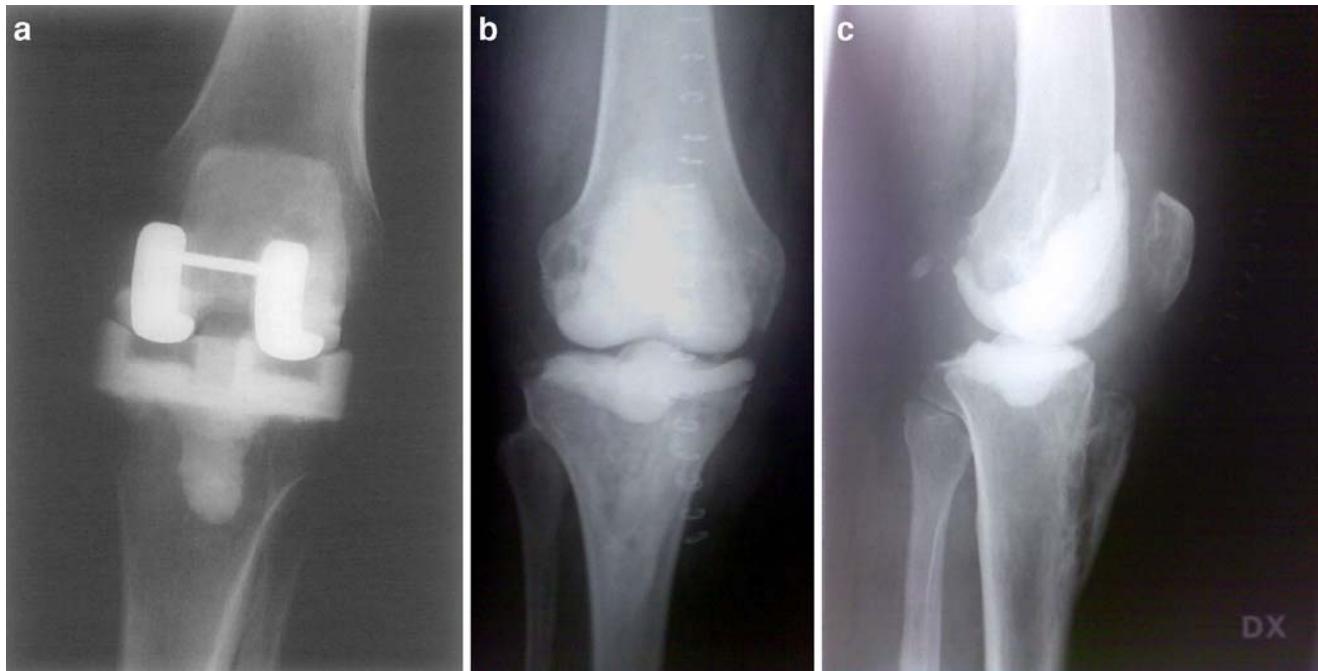


Fig. 11. a, b, c X-ray showing commercial PMMA spacer and the one described in this paper

shortening in any case. The mean flexion achieved after the second stage of the revision was 120°, ranging from 97° to 130°. The mean Hospital for Special Surgery score was 84. At 1 year after surgery, none of the knees showed any evidence of recurrence of the infection. Flexion after the revision was increased over the prerevision situation in all cases. In 1 case, there was a residual flexion contracture (10°) (Figs. 12 and 13).

Discussion

The infected total knee replacement is a difficult challenge for the orthopedic surgeon. The soft tissues and bone are

often necrotic; there is loss of bone; and exposure of the knee is difficult because of scar tissue. The golden standard in the treatment of an infected total knee arthroplasty is a two-stage protocol using an interval antibiotic spacer and intravenous administering of antibiotics for a period of 6 weeks [14].

Articulating spacers have been proposed as a way of preserving range of motion, bone stock, and length of soft tissues in two-stage revision total knee arthroplasty [2, 3, 8]. Premolded spacers have been described as being effective [2, 9, 10]. The problem with premolded spacers is that they are expensive and many times do not fit an individual patient. For that reason, we have considered using the method of molding an articulating spacer as

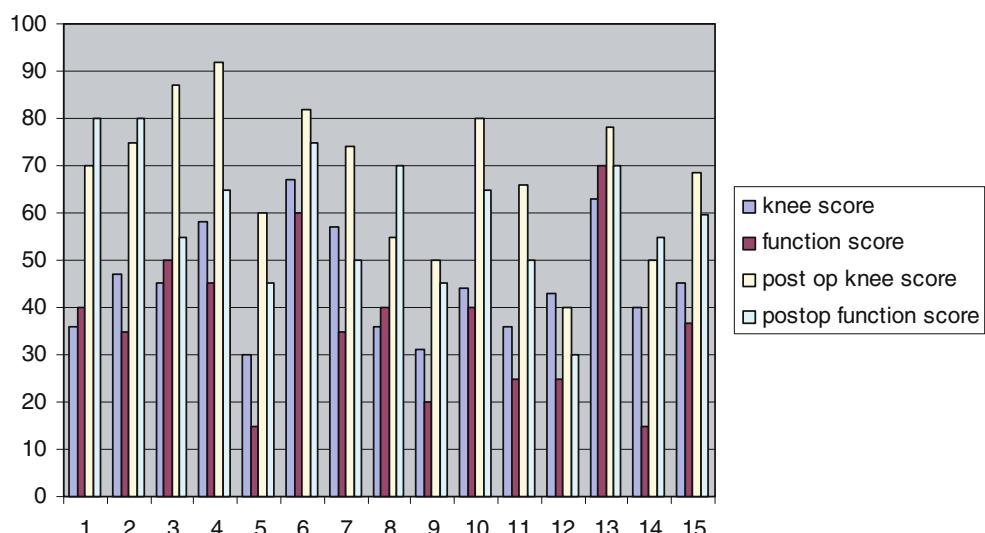


Fig. 12. Comparison of preoperative and postoperative knee and function scores

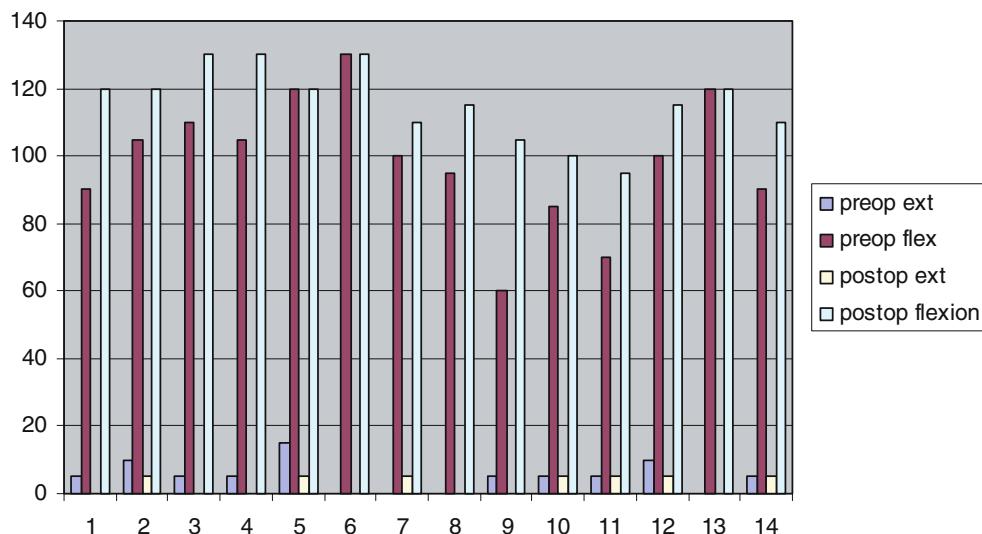


Fig. 13. Comparison of preoperative range of motion and 1 year postoperative range of motion

described by Villanueva et al. [1]. However, there is no agreement in the literature about the best fixed or mobile spacer to be used. [4–6, 8, 17–20]

This technique allows the surgeon to mold a spacer on single case anatomy. The advantage is perfect matching with residual bone (no further bone loss) and gap filling, so postoperative motion is preserved as collateral ligaments and patellar tendon length are maintained.

In this prospective study, although we have followed-up our patients for only 1 year after surgery, we have noted no recurrence of infection and good range of motion. We will continue to follow-up these patients over the next years; but for the present, we feel that this is a viable option for the patient who requires a two-stage reimplantation. However, a larger number of patients and a more prolonged follow-up are necessary to confirm the positive impressions obtained in this study.

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