The Outcomes of Cemented Femoral Revisions for Periprosthetic Femoral Fractures in the Elderly: Comparison with Cementless Stems

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Introduction: The present study compares the outcome of the long cemented stem and the revision uncemented stem used in periprosthetic femoral fractures. We propose that the revision with a long stem cemented prosthesis does not compromise fracture healing.

Patients and Methods: A consecutive series of 37 patients, operated between 2010 and 2017, were enrolled in a retrospective analysis. A long cemented stem was implanted in 21 patients (study group; age at operation: 63 to 89 years). A distally tapered fluted uncemented stem was used in 16 patients (control group; age at operation: 35 to 77 years). The clinical outcome was evaluated with Merle d’Aubigné and Postel scoring system. Standard radiographs were taken before surgery, at 3, 6, and 12 months postoperatively, and last follow-up. Any and all complications during the follow-up period were recorded.

Results: Although a significant difference ($P = 0.006$) was observed in the post-operative Merle d’Aubigné score over the 12-month follow-up period, no significant difference ($P = 0.066$) was found in the post-operative pain score between the study and control groups. Periodic radiographic assessments showed the disappearance of radiolucent lines and the diaphyseal part of the fracture was healed in all 34 followed-up cases during the first annual follow-up. Early surgical complications were seen in both groups, the medical complications were observed only in the study group.

Conclusion: Based on our results, periprosthetic fractures of the femur after a total hip arthroplasty were associated with significant morbidity and increased mortality in elderly patients. Revision with a long-stem cemented prosthesis provided early pain-free weight-bearing without compromising the healing of femoral fractures in elderly patients with osteoporotic bone, altered mobility, poor balance, and reduced cognitive capacity.

Keywords: revision total hip arthroplasty, periprosthetic hip fractures, cemented stem

Introduction

The management of periprosthetic fractures of the femur after a total hip arthroplasty can be challenging. The rising incidence of primary total hip arthroplasty, along with an ageing population that is more susceptible to falling trauma, results in a substantial increment of revision operations. From this perspective, orthopedic surgeons will face a greater number of periprosthetic fractures in elderly patients with osteoporotic bone, altered mobility, poor balance, and reduced cognitive capacity.

Elderly patients with periprosthetic fractures have a higher mortality risk, as well as greater surgical and medical complications. Therefore, the goals in primary management for elderly patients are vastly different from those observed in younger
patients. In brief, surgery seeks an early, pain-free, weight-bearing status without compromising the healing of femoral fractures. Revision with a long-stem cemented prosthesis can be used in these cases as it often allows an earlier weight-bearing ability. However, there are few reports available on the outcome of long cemented stems in elderly patients with periprosthetic fractures.

This study evaluates a consecutive series of patients with periprosthetic fractures of the femur after total hip arthroplasty treated with a long cemented stem, focusing on the benefits and risks of surgery. Thus, we compared (1) the hip function, (2) the radiographic outcome of fracture healing, and (3) the complications after revisions performed for periprosthetic fractures with the long stem cemented prosthesis and the distally tapered fluted uncemented stem to determine the efficacy of this technique. Our hypothesis was that revision with a long stem cemented prosthesis does not compromise fracture healing. Further, we theorized that both morbidity and mortality would be significantly pronounced in elderly patients with a long cemented stem.

Patients and Methods

Patients

All patients with a periprosthetic fracture of the femur after a total hip arthroplasty, having femoral revision at our institution between January 2010 and December 2017, were enrolled in this retrospective analysis (Supplementary Table 1). In total, 37 revision hip arthroplasties were done in 16 men (43%) and 21 women (57%). A long cemented stem was implanted in 15 elderly patients with considerable comorbidity level (ASA score ≥3, Charlson comorbidity index ≥4) and 6 patients with expected non-compliance with postoperative non-weight bearing (Figure 1). These 21 patients, with a mean age at operation 79 years (63 to 89), comprised the study group. Uncemented femoral revision was performed in 16 patients without significant comorbidity and that were deemed compliant with protected weight bearing after surgery; these patients constitute the control group, mean age at operation 63 years (35 to 77) (Table 1). The Ethics Committee of the University Hospital in Hradec Kralove approved the study protocol (reference number 202103P06) and waived the requirement for informed consent due to the retrospective design. The accessed patient data complied with relevant protection, privacy guidelines and regulations. The study was performed in accordance with the ethical standards described in the Helsinki Declaration.

Surgical Procedure

All the procedures were performed in one specialized arthroplasty medical center by four experienced orthopedic surgeons using a direct lateral approach in supine position of the patient. First, the hip joint was opened and dislocated; the cup was assessed for fixation stability and the polyethylene liner for wear. The lateral approach was extended distally; the fracture site was then used for removal of the stem and cement. After additional reaming of the diaphysis, the trial stem was inserted in the medullary canal bypassing the distal fracture site by at least 2 diaphyseal diameters to achieve sufficient stability. The fracture fragments were reduced and held with clamps, followed by trial reduction. After evaluation of the extremity length and implant stability, at least 2 cerclage strips were attached to the femur and the long stem was set in place with the correct anteversion. In the study group, the medullary canal was filled with cement in a retrograde manner before stem insertion and the fracture site was double-checked for cement leakage, with the subsequent removal of any extruded material.

Postoperatively, all patients received prophylactic intravenous antibiotics for 24 hours. Venous thromboprophylaxis was done with low-molecular-weight heparin for 5 weeks. Postoperative physiotherapy consisted of physical exercise focused on muscle strengthening and mobilization of the hip. Mobilization was started on the first postoperative day in a high vertical walker. Patients with a cemented long stem were allowed to bear weight as tolerated; on the other hand, protected weight bearing was recommended for up to 12 weeks until evidence of radiological healing occurred after uncemented femoral revision.

Outcome Assessments

The clinical outcome was evaluated with Charnley’s modified Merle d’Aubigné and Postel scoring system. The pain score component in this method was evaluated separately because the result of the original scoring system includes objective clinical parameters other than pain. The functional status was determined through the independent ability to walk, making a clear distinction between the following two observations:

1. Maintained independent ambulatory ability: able to walk without support from another person (with aid if necessary).
2. Not maintained: assistance from another person or use of a wheelchair required.

The standard radiographs were evaluated separately by three orthopedists, and the data revised for inter-observer agreement. In case of any observed difference, the patient’s radiograph was studied by all three reviewers together. Fracture union was characterized by the presence of bone bridging in both the anteroposterior and lateral radiographs.

Complications were registered throughout the follow-up period. All patients were under follow-up before surgery, at 3, 6, and 12 months postoperatively, and at the time of the latest appointment. The mean follow-up period was 55 (0–99) months for the study group (n = 21), whereas that the mean follow-up was 62 (7–124) months for the control group (n = 16).

**Figure 1** The study flow-chart.

**Statistical Analysis**

The measurement data were processed with the help of MS Excel 2007 (Microsoft Corp, Redmond WA, USA), NCSS 2007 (Hintze, J. (2007). NCSS 2007. NCSS LLC, Kaysville, UT, USA), and PASS 13 Power Analysis and Sample Size Software (Hintze, J. (2014). NCSS, LLC. Kaysville, UT, USA). With respect to the structure of the tested data, we opted for a Wilcoxon Signed-Rank Test (Paired Difference) or a Mann–Whitney U-test. The parameters with low number of observations in the tested groups were evaluated with a Freeman-Halton modification (Freeman and Halton, 1951) of Fischer’s exact test. In all the tests, a value of $P < 0.05$ was considered as significant.
Results

Study Design and Patients

The scheduled clinical and radiographic follow-up was accomplished in 34 patients within the first year after surgery. Two patients in the study group died within the first month after surgery; one patient in the control group was lost to follow-up after 7 months.

Of the 19 remaining patients in the study group, eight had died at time of review, whereas that of the remaining 15 patients in the control group, one patient was deceased from causes unrelated to the surgery (pulmonary tumor).

Clinical Results

Whereas no significant difference was found in the post-operative pain score ($P = 0.066$), a significant difference in the post-operative Merle d’Aubigné score was observed between the study and control groups ($P = 0.006$) over the 12-month follow-up period (Table 2).

The functional status of the patients in the study group deteriorated over time. Further, all but one of the patients were able to walk independently before surgery; in contrast, at the time of the latest follow-up, 8 out of 11 living patients (73%) were still able to walk independently with the aid of forearm crutches. All followed-up patients in the control group maintained independent ambulatory capacity at the time of review.

Radiography

The radiographs obtained immediately after surgery in all 37 patients proved the correct position of the revision prosthesis and adequate stability of the femoral fracture. Periodic assessments showed that, during the 3–12 months period, the radiolucent lines disappeared and the fractures were united at the first annual follow-up in all but one of the 34 followed-up cases (Figures 2 and 3). One patient in the study group had a pain-free non-union of the greater trochanter, showing a healed diaphyseal section of the fracture during the final follow-up, 5 years after the surgery.

Complications

The observed complications are shown in Table 3. Whereas early surgical complications were seen in both groups, the medical complications were observed only in the study group.

Dislocations happened in two patients within the first months postoperatively in the study group. One patient underwent an efficient closed reduction of a correctly implanted prosthesis and one patient had a successful conversion to a constrained liner for recurrent instability. Two patients in the control group had a deep infection; however, they were successfully treated with surgical debridement and the mobile parts were exchanged, this was followed by 6 weeks of antibiotic therapy. One patient in the control group suffered from prolonged unrecovered peroneal nerve palsy, which was managed conservatively. Following surgery, 6 patients in the study group (28.5%) suffered medical complications, the most common of which was pneumonia ($n = 3$).

Two patients of the study group and one patient of the control group had late hip-related complications. In the study group, one late deep infection occurred 30 months after revision surgery and antimicrobial suppression was chosen because the existing comorbidities did not allow additional surgery. In one patient of the study group, a cemented stem had been removed because of recurrent dislocation developed after a fatal stroke 6 years postoperatively. One patient in the control group had late deep infection; however, she was successfully treated with two-stage exchange arthroplasty.

Two patients in the study group died – both of pneumonia, within one month postoperatively. The overall 1-year mortality in patients with periprosthetic fractures of the femur treated with the long cemented
stem was estimated in 9.5%. Eight patients in the study group and one patient in the control group were deceased at the time of review from causes unrelated to the surgery.

**Discussion**

Although most authors advise revision with a long stem prosthesis used as an intramedullary rod in the treatment of Vancouver B2 and B3 periprosthetic femoral fractures, there is a continuous debate on whether cemented or uncemented implants should be used.9–14 Our study demonstrates a clear improvement in hip/thigh pain relief following revision total hip arthroplasty in patients with periprosthetic fractures. We suppose that the significant difference in post-operative Merle d’Aubigné and Postel scores found between the study and control groups was

![Figure 2](https://example.com/figure2.png) **Figure 2** Preoperative anteroposterior radiograph of a Vancouver type B1 periprosthetic femoral fracture in a 77-year-old female (A) treated with a long cemented femoral stem (B). At 5 years of follow-up, the fracture was healed and the stem did not show any sign of aseptic loosening.

![Figure 3](https://example.com/figure3.png) **Figure 3** Preoperative anteroposterior radiograph of a Vancouver type B2 periprosthetic femoral fracture in a 61-year-old man (A) treated with an uncemented femoral revision (B). Fracture was healed and uncemented modular tapered fluted revision stem was well osteointegrated after 1 year of surgery.

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<thead>
<tr>
<th>Table 2 Clinical Results of Patients</th>
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</thead>
<tbody>
<tr>
<td>Follow-up (mean; range; months)</td>
</tr>
<tr>
<td>55 (0–99)</td>
</tr>
<tr>
<td>Preoperative pain score ± the postoperative increase at 12 months postoperatively (mean; points)</td>
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<td>Hip flexion at 12 months postoperatively (mean; range; degrees)</td>
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<tr>
<td>Hip abduction at 12 months postoperatively (mean; range; degrees)</td>
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<tr>
<td>Preoperative Merle d’Aubigné and Postel score ± the postoperative increase at 12 months postoperatively (mean; points)</td>
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<td>Functional status preoperatively</td>
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<tr>
<td>Independent ability to walk</td>
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<td>Support from another person or use of wheelchair</td>
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<tr>
<td>Functional status at the time of latest follow-up</td>
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<td>Independent ability to walk</td>
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**Notes:** *Two patients lost to further follow-up within the first month after surgery.** One patient lost to further follow-up 7 months after surgery.
affected by the deteriorated functional status of elderly patients with considerable high comorbidity level in the study group. The radiographic evaluation showed that revision with a long stem cemented prosthesis did not compromise fracture healing because the diaphyseal section of the fracture was found completely healed during the first annual follow-up meeting in the 19 cases evaluated in this period. Although early surgical complications were recorded in both groups, morbidity and increased medical complications were predominant in our study group.

However, this case-control study is limited by the small number of patients and short follow-up duration. Regardless, all patients with a periprosthetic fracture of the femur after a total hip arthroplasty having femoral revision at our tertiary referral hospital during an 8 years period were enrolled in a retrospective survey. Regrettably, the available literature is rather lacking in long-term follow-up and short follow-up of larger series comparing long stem cemented prosthesis and distally tapered fluted uncemented stem in periprosthetic femoral fractures. Nevertheless, the follow-up period in our analysis was sufficient to reveal the healing of fractures. Furthermore, the number of patients and follow-up period is comparable to other published surveys on long cemented stem in patients with periprosthetic fractures. We are aware of the significant differences in baseline characteristics between the groups. Fractures in the control group with the revision uncemented implant occurred at a younger age than the study group with a long cemented stem. The shorter follow-up of the cemented group reflects the frailty of the elderly patients, who were significantly older, in comparison with those in the uncemented group.

Several disadvantages have been previously reported concerning the use of long cemented stems. Cement may be extruded into the fracture site, which could be the reason of non-union or refracture. Whereas Corten et al published a healed diaphyseal section of a fracture in sixteen followed-up patients after more than 1 year, Springer et al reported three (9%) out of thirty-four unrevised cemented stems with radiographic non-union after 68 months of follow-up. The non-union rate could be diminished with thorough handling of the surrounding soft tissues to prevent bone devascularization; further, careful inspection and removal of excess cement and the use of autogenous bone grafts or cortical strut allografts could also alleviate this issue. Aseptic loosening is another concern regarding cemented stems. On the one hand, Corten et al demonstrated no revision of sixteen patients due to aseptic loosening; in contrast, Springer et al found twenty-five (60%) out of forty-two cemented stems with a well-fixed, unrevised implant and healed periprosthetic fracture. In the present study, we identified seventeen (81%) out of twenty-one well fixed, unrevised long cemented stems during the last follow-up. Finally, there is an indication for greater early mortality if cement is used. However, the high perioperative mortality rate seems to be rather patient- and procedure-related rather than implant-related.

In our study, the overall 1-year mortality in elderly patients with periprosthetic fractures treated with the long cemented stem was of 9.5%. In this regard, Young et al reported a six-month overall mortality of 0.9% after revision total hip arthroplasty compared to 7.3% after revision surgery for periprosthetic fracture. After revision hip arthroplasty, Yao et al found a greater six-month, one-year, and five-year mortality rate for periprosthetic fractures than for loosening/wear and/or infection. Recently, Gibbs et al found that hospital-acquired pneumonia and dislocation were associated with increased one-year mortality rates after revision hip arthroplasty for periprosthetic fractures. Compared to this study, our dislocation rate (10.5%) in the elderly patients with long cemented stems was similar.
We observed pneumonia in two patients deceased within the first month after surgery.

Revision surgery is a major procedure that involves an extended operating time, substantial blood loss, and considerable physiological stress.\textsuperscript{20,21} Revision hip arthroplasty for periprosthetic femoral fracture differs from other common indications, where the elective procedure is performed after preoperative optimization of the patient.\textsuperscript{22,23} Based on recent studies, surgery time was not identified as a risk factor for mortality in periprosthetic fractures.\textsuperscript{19,24} This suggests that the selection of appropriate surgical and anesthetic options may be crucial for the preoperative optimization of the patient. For elderly patients with poor balance, postoperative partial weight bearing after hip revision might be more difficult.\textsuperscript{25} Therefore, the use of long-stem cemented prosthesis could allow early weight bearing in elderly and morbid patients or in patients non-compliant with partial weight bearing after hip revision.

Conclusions

Based on the obtained results in our study, the periprosthetic fractures of the femur after a total hip arthroplasty are associated with significant morbidity and increased mortality in the elderly. Revision with a long-term cemented prosthesis provides early pain-free weight-bearing without compromising the healing of the femoral fracture in elderly patients with osteoporotic bones, altered mobility, poor balance, and reduced cognitive capacity. However, the management of periprosthetic fractures around the femoral stem requires extensive surgical expertise.

Ethical Statement

Informed consent regarding surgery was obtained from all patients before intervention. The Ethics Committee of the University Hospital in Hradec Kralove approved the study protocol (reference number 202103P06) and waived the requirement for informed consent due to the retrospective design. The patient data accessed complied with relevant patient data protection and privacy regulations. The study was performed in accordance with the ethical standards of the Helsinki Declaration.

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Disclosure

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References


