Good stability but high periprosthetic bone mineral loss and late-occurring periprosthetic fractures with use of un cemented tapered femoral stems in patients with a femoral neck fracture

A 5-year follow-up of 31 patients using RSA and DXA

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Background and purpose — We previously evaluated a new un cemented femoral stem designed for elderly patients with a femoral neck fracture and found stable implant fixation and good clinical results up to 2 years postoperatively, despite substantial periprosthetic bone mineral loss. We now present the medium-term follow-up results from this study.

Patients and methods — In this observational prospective cohort study, we included 50 patients (mean age 81 (70–92) years) with a femoral neck fracture. All patients underwent surgery with a cemented cup and an un cemented stem specifically designed for fracture treatment. Outcome variables were migration of the stem measured with radiostereometry (RSA) and periprosthetic change in bone mineral density (BMD), measured with dual-energy X-ray absorptiometry (DXA). Hip function and health-related quality of life were assessed using the Harris hip score (HHS) and the EuroQol-5D (EQ-5D). DXA and RSA data were collected at regular intervals up to 4 years, and data concerning reoperations and hip-related complications were collected during a mean follow-up time of 5 (0.2–7.5) years.

Results — At 5 years, 19 patients had either passed away or were unavailable for further participation and 31 could be followed up. Of the original 50 patients, 6 patients had suffered a periprosthetic fracture, all of them sustained after the 2-year follow-up. In 19 patients, we obtained complete RSA and DXA data and no component had migrated after the 2-year follow-up. We also found a continuous total periprosthetic bone loss amounting to a median of –19% (~39 to 2). No changes in HHS or EQ-5D were observed during the follow-up period.

Interpretation — In this medium-term follow-up, the stem remained firmly fixed in bone despite considerable periprosthetic bone mineral loss. However, this bone loss might explain the high number of late-occurring periprosthetic fractures. Based on these results, we would not recommend un cemented femoral stems for the treatment of femoral neck fractures in the elderly.

In Sweden, cemented stems have been used primarily for patients with a femoral neck fracture (FNF), but with the introduction of modern hydroxyapatite- (HA-) coated implants, un cemented fixation has increased in popularity (Garellick et al. 2011). Excellent long-term results have been reported for patients with primary osteoarthritis (OA) of the hip (Boden et al. 2006a). The concept of inserting an un cemented femoral component is attractive to many surgeons, as the cementing procedure can induce cardiac arrhythmia and/or cardiorespiratory collapse (Parvizi et al. 1999). However, a recent report from the Swedish Hip Arthroplasty Register indicated that un cemented implants used for patients with an FNF are associated with a 20-fold higher risk of reoperation due to periprosthetic fracture than cemented matte stems (Leonardsson et al. 2012).

We have already published an evaluation of a new un cemented femoral stem designed for elderly patients with an FNF; this showed good clinical results and stable implant fixation up to 2 years after surgery despite substantial periprosthetic bone loss (Sköldenberg et al. 2011). We now present the medium-term follow-up from this study.
Patients and methods

This observational, prospective single-cohort study was carried out between October 2005 and August 2013 (inclusion period October 2005 to March 2008) at the Department of Orthopedic Surgery, Danderyd Hospital, Stockholm, Sweden. The study was approved by the Ethics Committee of Karolinska Institutet (Dnr 04-086/4) and the Radiation Safety Committee at Danderyd Hospital (Dnr 005-5).

Patients

We included 50 patients ≥ 70 years of age with a displaced femoral neck fracture (Garden III or IV) who had an intact cognitive function (at least 8 correct answers on a 10-item (SPMSQ) mental test) (Pfeiffer 1975), who could walk independently with or without the help of walking aids, and who were willing to participate in the study. All patients gave their written consent to participate. Exclusion criteria and details of the study protocol have been presented previously (Sköldenberg et al. 2011).

Stem design

The implant, the Biomet Fracture Stem (BFX; Biomet UK Ltd., Bridgend, UK), is a tapered, collared stem intended for uncemented fixation. It is made of titanium alloy (Ti-6Al-4V) with a grit-blasted surface roughness of 7.5–10 µm. It has a straight 3° proximal-to-distal taper in 2 planes and a taper from the lateral shoulder to the medial calcar area. It has plasma-sprayed hydroxyapatite (HA) on the entire surface (thickness 65–95 µm, crystallinity 50–70%, purity > 95%) to enable fast ingrowth into osteoporotic bone. The geometry of the stem, except for the collar, is identical to that of the Bi-Metric stem (Meding et al. 2004, Boden et al. 2006).

Surgery

The patients underwent THA surgery using the new stem articulating on a 32-mm cobalt-chrome head against a cemented cup (Müller; Biomet UK). A posterior approach, with repair of the posterior capsule and external rotator muscles, was used in all patients. The femur was reamed until cortical bone contact was obtained. Then the proximal femur was prepared with broaches of increasing size until rotational stability was achieved. Before the final implant was inserted, 5–9 tantalum marker beads (1.0 mm in diameter) were inserted in the cancellous bone of the proximal femur. The patients were mobilized using a standard physiotherapy program. They were encouraged to mobilize with full weight bearing using crutches for support.

Outcome measures

The outcome variables were migration of the stem measured with RSA, changes in bone mineral density (BMD) in 7 Gruen zones around the stem, the occurrence of periprosthetic fractures and other hip-related complications, and clinical outcome. 

Radiostereometric analysis. The RSA method followed the published guidelines for RSA (Valstar et al. 2005). We took digital calibrated radiographs (Bucky Diagnostic; Philips, Eindhoven, the Netherlands) using a fixed and a mobile X-ray (Roentgen) source (120 kV, 4–6 mAs), and a uniplanar calibration cage (Uniplanar digital 43; RSA Biomedical, Umeå, Sweden). All data were analyzed using UmRSA computer software (RSA Biomedical). Based on double examinations at 12 months, we calculated the precision of RSA. For translation along the x- (transverse), y- (vertical), and z- (anteroposterior (AP)) axes, this was 0.27, 0.19, and 0.52 mm, respectively. For rotation about the x- (flexion/extension), y- (ante-/retroversion), and z- (varus/valgus) axes, the values were 0.52°, 0.76°, and 0.27°, respectively, and for the maximal total point motion (MTPM) it was 0.74 mm. Further details of the RSA method in this study have been presented in the report on 2-year follow-up (Sköldenberg et al. 2011).

Bone densitometric analysis. BMD of the periprosthetic femur in the frontal plane was measured in the 7 Gruen zones in 1 plane using DXA (DPX-L; Lunar Co., Madison, WI). The change in frontal periprosthetic BMD ratio in all the individual zones, as well as in the entire periprosthetic region (zones 1–7), was calculated by dividing the BMD value at each follow-up visit by the postoperative BMD and converting it to a percentage change. We had previously conducted double measurements and validated the method. Postoperatively, the BMDs of vertebrae L1–L4 (lumbar spine) of all patients were measured to assess the patient’s general bone mass. The BMDs of the L1–L4 vertebrae were also measured at the 4-year follow-up.

Clinical outcome. Hip function was evaluated with the Harris hip score (HHS) at all follow-ups. This score has been validated for patients with FNFs (Frihagen et al. 2008). Health-related quality of life was assessed with the EQ-5D (EuroQoL) questionnaire. After inclusion, but before surgery, we asked the patients to estimate their HHS and EQ-5D for the week prior to the occurrence of the fracture.

Hip-related complications and reoperations. We used the Swedish unique personal ID-number to identify all hip-related complications during the study period. We searched digital medical charts at Danderyd Hospital, the Swedish Hip Arthroplasty Register, and the Swedish Patient Registry. All hip-related complications of the study were managed and registered at our department, and no other reoperations or complications were found to have occurred at other hospitals in Sweden.

Sample size

In the previously published paper (Sköldenberg et al. 2011), we used a sample size that indicated that we would need to recruit 50 patients to allow for loss to follow-up and to allow for analysis of subgroups of patients with high and low BMD. Before the 4-year follow-up, a new sample size analysis was performed to indicate how many patients would be needed.
for investigation of the migration pattern of the stems in the patients who were still living. With 18 patients, the study would have a power (2-sided, p = 0.01) of more than 99%, and of 93% to detect a continuous migration in MTPM and y-translation. These estimates were based on a previous RSA study with the HA-coated version of the Bi-Metric stem (Søballe et al. 1993) where mean MTPM was 1.9 (SD 1.3) mm and mean y-translation was 0.2 (0.2) mm.

Statistics
We used a simple linear regression analysis to investigate factors predictive of stem migration (expressed as MTPM) or bone loss (percentage change in BMD in zones 1–7). Due to the low number of patients available for migration and BMD analysis at 4 years (n = 19), we did not consider it appropriate to use a multivariate analysis. Factors investigated separately in the analysis were covariates which, from the results of previous studies, are known to influence migration or bone remodeling around implants. These factors are (1) postoperative periprosthetic BMD (zones 1–7) (Sköldenberg et al. 2011), (2) postoperative BMD of the lumbar spine (Alm et al. 2009), (3) age (Brodner et al. 2004), (4) gender (Brodner et al. 2004), and (5) stem size (Sköldenberg et al. 2006). We used the Shapiro-Wilk normality test to confirm the normality of residuals. Comparisons between migration, BMD, and clinical scores at 2 and 4 years and also comparisons of bone mineral loss in patients with and without periprosthetic fractures were analyzed with non-parametric tests. All p-values ≤ 0.05 were considered significant. We used SPSS software version 20 for Mac.

Results
Patients
At the final follow-up in August 2013, 12 of the 50 patients had died, 7 were unable to participate further in the study (due to ill-health or dementia (n = 6) or from the fact that they had moved out of the Stockholm area (n = 1)) and 31 were available for clinical follow-up (Figure 1 and Table 1). For 19 of these 31 patients, we were able to obtain complete RSA and DXA data at 4 years. The mean follow-up time for the whole cohort was 5.3 (0.2–7.5) years.

Outcome variables
Radiostereometry. We found no migration of the stems between 2 and 4 years. At 4 years, the median migration for translations compared to the postoperative examinations was small: –0.07, –0.06, and 0.06 mm on the x-, y-, and z-axes. Likewise, the median rotation was small at –0.6, –0.5, and –0.08 degrees on the x-, y-, and z-axes. The total migration (MTPM) was 1.5 mm at 4 years (Table 2). We found no correlation between migration and the initial periprosthetic or lumbar spine BMD, age, gender, or stem size. None of the patients who had suffered a dislocation showed any migration in any direction.

Bone densitometry. We found a decrease in periprosthetic BMD in all Gruen zones (1–7): median –19%, which was...
statistically significant compared to the 2-year value of \(-8.8\%\) (\(p = 0.01\), Wilcoxon signed-rank test). In individual zones, the decrease in periprosthetic BMD was continuous in all zones except zones 3 and 7 (Figure 2). The bone loss was greatest in zones 1 and 7, with a decrease of 36\% and 26\% (Table 2). The loss of BMD in zones 1–7 correlated with the initial BMD surrounding the implant \((R = 0.5, p = 0.03)\): the lower the initial BMD, the greater the bone loss (Figure 3). We found no correlation between bone loss and the BMD of the lumbar spine, age, gender, or stem size.

**Clinical outcome.** Mean HHS was 83 (95\% CI: 78–88), with no statistically significant change compared to the 2-year value. Similarly, the health-related quality of life (EQ-5D) remained unchanged compared to the 2-year value (mean 0.7, 95\% CI: 0.5–0.8). We found no correlation between clinical outcome data and migration or bone loss (data not shown).

**Hip-related complications and reoperations.** There were 2 periprosthetic joint infections, 1 early and 1 hematogenous. Both patients underwent debridement, antibiotics and implant retention (DAIR) and recovered successfully. 7 patients sustained a dislocation of the hip at a median of 24 (2–92) days after surgery. All were treated with closed reduction under sedation. 3 patients experienced a subsequent dislocation.

### Table 2. Migration and percentage change in BMD at 2 and 4 years for 19 patients with complete data at 4 years. The p-values were derived from the Wilcoxon signed-rank test

| Migration                | 2 years Median | 2 years Min | 2 years Max | 4 years Median | 4 years Min | 4 years Max | p-value |
|--------------------------|----------------|-------------|-------------|----------------|-------------|-------------|---------|
| **Translation, mm**      |                |             |             |                |             |             |         |
| Transverse (x)           | −0.08          | −0.2        | 0.5         | −0.07          | −0.3        | 0.3         | 0.3     |
| Vertical (y)             | −0.07          | −0.3        | 0.3         | −0.06          | −1.0        | 0.6         | 0.05    |
| Anterioposterior (z)     | 0.01           | −1.5        | 0.5         | 0.06           | −1.5        | 0.6         | 0.1     |
| Rotation (°)             |                |             |             |                |             |             |         |
| Flexion/extension (x)    | −0.5           | −3.3        | 0.5         | −0.6           | −3.1        | 0.5         | 1.0     |
| Ante-/retroversion (y)   | −0.5           | −4.7        | 0.2         | −0.5           | −4.3        | 1.0         | 0.1     |
| Varus/valgus (z)         | −0.03          | −1.9        | 0.5         | −0.08          | −1.8        | 0.5         | 0.2     |
| **Total migration**      | 1.6            | 0.8         | 5.5         | 1.5            | 0.6         | 5.2         | 0.4     |

**Percentage change in BMD versus postop**

| All zones (1 to 7)       | −8.8           | −39.1       | 1.3         | −18.8          | −36.9       | 2.4         | 0.01    |
| Zone 1                   | −33.5          | −59.6       | −13.7       | −36.2          | −59.6       | −13.7       | 0.01    |
| Zone 2                   | −12.0          | −61.1       | 10.6        | −21.6          | −47.0       | 17.7        | 0.03    |
| Zone 3                   | −4.4           | −60.5       | 13.6        | −9.6           | −39.0       | 14.2        | 0.4     |
| Zone 4                   | −2.3           | −37.8       | 17.5        | −6.2           | −64.1       | 4.3         | 0.07    |
| Zone 5                   | 0.0            | −31.6       | 21.7        | −8.5           | −29.8       | 13.4        | 0.05    |
| Zone 6                   | −16.7          | −43.9       | 3.3         | −24.1          | −52.1       | 3.0         | 0.02    |
| Zone 7                   | −26.4          | −43.8       | 11.0        | −25.9          | −56.9       | 8.0         | 0.2     |
| Lumbar spine             | 0.03           | −5.1        | 7.0         | 0.05           | −4.06       | 6.01        | 0.7     |

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Figure 2. Graphs showing periprosthetic bone remodeling in zones 1–7 with median percentage change in bone mineral density (BMD) around the implant.* \(p < 0.05\) compared to the 2-year follow-up (Wilcoxon signed-rank test).
location, one of which required a revision of the acetabular component. 6 patients sustained a periprosthetic fracture after low-energy trauma. These fractures occurred 2.1, 2.1, 2.2, 2.9, 5.5, and 5.7 years after surgery. 3 were classified as Vancouver B1 (undisplaced with stable stem) and were treated nonoperatively with protected weight bearing (Figure 4). 1 was a B2 fracture (loose stem), which was treated with a long revision stem, and 2 were C type fractures (distal to the stem); these were treated with open reduction and internal fixation. All fractures healed.

The patients who sustained a periprosthetic fracture after the 2-year follow-up but before the 4-year follow-up could not be evaluated with RSA or DXA at 4 years. However, we had BMD data at 2 years on 5 of the 6 patients. The median change in BMD in zones 1–7 at 2 years was –26% (–38% to –13%) compared to the initial postoperative value. For the remaining 44 patients, we had 2-year data on 38 patients with a median change in BMD of –14% (–49% to 1%).

Discussion

In this observational, prospective cohort study on a selected group of cognitively intact elderly patients, we found a 1-tenth incidence rate of periprosthetic fractures up to 5 years after surgery. The implant functioned well and was radiographically stable despite substantial progressive periprosthetic bone mineral loss. We believe that this bone mineral loss—in combination with fracture patients’ propensity for falls—would explain our findings.

Uncemented stems and femoral neck fractures

Our results are in keeping with the short-term (1-year) results of a randomized clinical trial (RCT) comparing an HA-coated stem (Corail) and a cemented stem (Spectron) in patients
with FNF, where the authors noted an incidence of postoperative periprosthetic fractures of 4% (Figved et al. 2009) in the uncemented group as compared to 1% in the cemented group. The same research group have recently published their 5-year data and found an even higher incidence of late-occurring postoperative periprosthetic fractures in the uncemented group (9.4%) compared to the cemented group (1%) (Langslet et al. 2014). In general, there appears to be an increased risk of intraoperative fractures and postoperative fractures associated with the use of uncemented stems in FNF patients (Parker and Gurusamy 2006). In a recent report from the Swedish Hip Arthroplasty Register (Garellick et al. 2011), a comparison of survival curves between cemented and uncemented stems used for patients with osteoarthritis of the hip showed a biphasic progression. During the first 5 years after surgery, the risk of stem revision, regardless of cause, was reduced by about 50% when using cemented fixation. During the second period (8–16 years), the survival of the uncemented stems was better than that of cemented stems due to the increasing revision rate resulting from aseptic loosening of the latter stem type.

We recently published a report of elderly patients with a displaced FNF with similar demographics (Chammout et al. 2012) as in the current study, and we found mortality rates of 75% after 11 years and of 87% after 17 years. This has also been reported by others, and it could therefore be argued that, in this patient group, the superior short-term results found with the use of a cemented stem are of greater relevance than long-term stem survival. The risk of periprosthetic fracture should be considered when choosing the type of stem to be used in elderly patients. A cemented stem is associated with substantially lower risk of fracture (Leonardsson et al. 2012), and this should outweigh the potential risks to the patient associated with the use of cement (Parvizi et al. 1999).

Migration and bone remodeling

We used RSA to evaluate the fixation of the stem, and although RSA has been used to study the healing of FNFs after internal fixation (Ragnarsson and Kärnholm 1991), only 3 other studies using RSA have been published on the subject of hip arthroplasty in this patient group. The first involved the 2-year results of the current study and the other 2 studies dealt with the subject of cartilage wear and implant stability following hemiarthroplasty (Figved et al. 2012, Schewelov et al. 2012). Similarly, bone remodeling around femoral stems after THA in patients with degenerative joint disease has been extensively studied using DXA (Kilgus et al. 1993, Ang et al. 1997, Boden et al. 2006b). However, to date there have been few reported clinical consequences (Hsieh et al. 2005). This periprosthetic bone loss may be one possible explanation for the increased risk of periprosthetic fractures found in patients who have undergone surgery for FNF with uncemented implants (Leonardsson et al. 2012), particularly as we found the BMD loss to be more pronounced in those who later sustained a periprosthetic fracture (Figure 4). This bone loss seems to be associated with larger stem size and a low initial BMD (Engh and Bobyn 1988, Rahmy et al. 2004, Sköldenberg et al. 2006) (Figure 3), though we were unable to verify the effect of stem size, possibly due to the low number of patients available for DXA at follow-up. The BMD gradually decreases even more over time, and generally more in the proximal areas than in those that are distal. 2 recent publications have further supported the hypothesis that there is indeed a lower reoperation rate for cemented stems than for uncemented stems (Gjertsen et al. 2012, Viberg et al. 2013).

Strengths and weaknesses of the study

To our knowledge, this has been the first study to measure periprosthetic BMD around a femoral stem in hip fracture patients. We could also correlate this to the incidence of periprosthetic fractures. With our extensive database search of all possible complications, we were able to identify periprosthetic fractures that would not normally have been reported to the Swedish Hip Arthroplasty Register. The greatest weakness was that we lacked a control group, and since there have been no previous publications on FNF patients and periprosthetic change in BMD, we can only compare our results with those for OA patients. In addition, our sample size calculation was performed for RSA data and not to estimate the incidence of complications. The study was also performed on a selected group of cognitively intact FNF patients, whose results may have differed from the general hip fracture population.

Conclusion

In conclusion, we found an important clinical effect of stress-shielding: a high incidence of periprosthetic fractures. Thus, even though the stem can easily achieve safe fixation even in osteoporotic bone and give good clinical results, we would not recommend use of this uncemented stem in the treatment of femoral neck fracture in elderly patients.

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