Low bone density and high morbidity in patients between 55 and 70 years with displaced femoral neck fractures: a case-control study of 50 patients vs 150 normal controls

Stefan Bartels 1,6*, Jan-Erik Gjertsen 2,3, Frede Frihagen 4, Cecilia Rogmark 5 and Stein Erik Utvåg 1,6

Abstract

Background: A displaced femoral neck fracture (FNF) in patients 55-70 years is a serious injury with a high risk of treatment failure and the optimal surgical treatment remains unclear. We aimed to describe characteristics of fracture patients compared to a sample from the normal population.

Methods: Fifty patients aged 55-70 years with a displaced FNF were gender- and age- matched with a control group of 150 persons without a hip fracture using computergenerated randomization and the Norwegian National Population Register. To reduce the risk of spurious selection bias, the sample size of the control group was trebled compared to the fracture group. Dual-energy x-ray absorptiometry (DXA) was performed. Demographics and hip function (Harris Hip Score, Oxford Hip Score, and Hip Dysfunction and Osteoarthritis Outcome Score) were collected.

Results: There were more than 75% women in both groups. The mean age was 64.5 years in the fracture group and 65.1 in the control group. Results for DXA measured for lumbar spine, total hip and the femoral neck showed that patients with displaced FNF were significantly more osteoporotic. Fracture patients had significantly lower body mass index, higher Charlson comorbidity index (CCI), and higher ASA (American Society of Anesthesiologists) score than the control group. No clinically relevant differences in hip function were found. There were 48% smokers in the fracture group compared to 10% in the control group. The odds ratio for obtaining a displaced FNF was high if the patients suffered from osteoporosis, smoked or had several comorbidities.

Conclusions: This study showed that patients aged 55-70 years with a displaced femoral neck fracture had lower bone density and higher comorbidity compared with a gender- and age-matched population without femoral neck fractures. This suggests that this patient group is epidemiologically similar to older patients with femoral neck fractures.

Keywords: Femoral neck fracture, Bone mineral density, Comorbidity, Osteoporosis
Background
Regardless of age, a displaced femoral neck fracture (FNF) is a severe injury and will almost always require hospitalization and surgery [1]. Patients with these fractures have a high risk of subsequent surgical complications, reduced function, hip pain and reduced health-related quality of life. The health economic aspect is a great challenge, even though the overall incidence of hip fractures has decreased in recent decades [1–7]. The literature on elderly patients older than 70 years with displaced FNFs is extensive and arthroplasty is clearly recommended as the treatment of choice [8–14]. The middle-aged patient group aged 55-70 years is less well described and the treatment for displaced FNFs is still controversial [11–13]. These patients are probably still working and demand a high level of activity. A Norwegian study has reported the overall hip fracture incidence in this particular age group to be 92 per 10 000 (53 women and 39 men) in the period 2009-2013 [15]. We found a specific incidence of about 6.1 fractures for both genders per 10 000 for displaced FNFs in patients 55-70 years in 2017 in Norway [16, 17]. Most of these FNFs are caused by a low-energy trauma, and the patients often have other diseases and factors, such as medication (steroids, anti-epileptic medication), alcoholism, other substance abuse, or osteoporosis, all of which may increase the risk of complications, including revision surgery [18–20]. Studies including bone density at the time of fracture are rare and often described a more geriatric population [21]. For patients under 60 years of age, internal fixation (IF) is usually recommended, as many surgeons endeavor to prevent replacement of the hip joint [12, 13, 22]. Studies investigating outcome after FNF in patients younger than 70 years have found a high risk of reoperation after IF due to mechanical failure, non-union or avascular necrosis [22–24]. Most of the investigated patients in this age group had symptomatic comorbidities and the 1-year mortality has been reported to be as high as 15% [24]. This may indicate that many hip fracture patients under 70 years of age are more osteoporotic and frail than individuals at the same age in the general population. Thus, their fractures may beneficially be treated mainly by arthroplasty, as in patients older than 70 years.

The aim of the present study was to describe differences in bone density and morbidity at the time of injury between a group of 50 patients aged 55-70 with a native intracapsular displaced FNF compared with a gender- and age- matched cohort of 150 participants from the general population without a fracture.

Methods
This pragmatic gender- and age- matched case-control included 50 consecutive patients aged 50-70 years presenting with a low-energy displaced FNF in the native hip joint and belonging to the catchment area of Akershus University Hospital, Norway from December 2013 to November 2017. The control group consisted of 150 participants from the population in the same catchment area. The Department of Data and Analytics at Akershus University Hospital was responsible for the recruitment of this group using computergenerated randomization lists and both the National Population Register and the unique national identification number assigned for each inhabitant in Norway. Patients with cognitive impairment were not included in either group. Controls were matched to the fracture group by loose matching [25]. For matching we divided the patients into three age groups for both genders (55-59, 60-64, 65-69 years) using the number of cases in the fracture group as the base. The tripled size of the control group was chosen to achieve statistical power [26]. Three hundred and forty-one potential participants were invited by mail, and one hundred and ninety-one persons declined or did not attend the agreed appointment. Information from the FNF patients was collected before discharge. DXA was performed postoperatively before discharge or within at least 6 weeks after injury. No anti-osteoporotic medication was given to the fracture patients group before DXA was performed. Interviews, questionnaires, and DXA for the control group were performed between November 2016 and June 2017 during a single outpatient appointment. All participants in the control group and in the fracture group signed an informed consent form.

The following variables were registered: age, gender, height and weight (measured in conjunction with DXA using a standard scale and a stadiometer), current smoking status, Charlson comorbidity index (CCI), American Society of Anesthesiologists (ASA) score and presence of diabetes [27, 28]. Furthermore, the Harris Hip Score (HHS), Oxford Hip Score and the Hip Dysfunction and Osteoarthritis Outcome Score (HOOS) were recorded [29–31]. For patients with displaced FNF, we asked for specific hip function scores the last week before the current fracture. Body Mass Index (BMI) was calculated as weight/height$^2$ in kg/m$^2$. Each participant in the control group and patients in the fracture group underwent a DXA measurement using lumbar spine (L1-L4), total hip and femoral neck. Five patients in the fracture group could not undergo DXA measurement in the hip, due to implants after recent fracture care and from previous surgery unrelated to the recent injury in the contralateral side.

The DXA scan was used to determine bone mineral density (BMD) measured in the lumbar spine (L1-L4), total hip and the femoral neck. All BMD measurements were performed by two independent DXA technicians on the same DXA scanner (Lunar iDXA™ GE Healthcare...
The definition of osteoporosis was that used by the World Health Organization, where a T-score ≤ -2.5 SD is osteoporosis, a T-score > -2.5 - < -1 SD is osteopenia and T-score ≥ -1 SD is normal bone [32].

### Statistical analysis
Continuous variables are presented as mean values and categorical variables were summarized by the number of subjects and percentage in each category. We used the independent samples t-test for continuous variables with normal distribution, and the Mann-Whitney U-test for categorical variables.

### Table 1 Baseline characteristics according to the two different groups

| Fracture group | Control group | p-values |
|----------------|---------------|----------|
| Mean age, years (95% CI) | 64.5 (63.5-65.6) | 65.1 (64.4-65.7) | 0.677 a |
| Gender, w/m (%) | 38 (76%) / 12 (24%) | 116 (77.3%) / 34 (22.7%) | 0.846 b |
| Mean BMI (kg/m²) (95% CI) | 24.2 (23.1-25.4) | 26.7 (26.0-27.4) | 0.001 c |
| Diabetes mellitus, n (%) | 5 (10%) | 5 (3.3%) | 0.061 b |
| Smoking, n (%) | 24 (48%) | 15 (10%) | <0.001 b |
| Mean Charlson Comorbidity Index Score total (95% CI) n (%) | 2.64 (2.39-2.89) | 2.14 (2.04-2.24) | 0.001 b |
| 1 | 3 (6%) | 19 (12.7%) |
| 2 | 20 (40%) | 94 (62.7%) |
| 3 | 21 (42%) | 34 (22.7%) |
| 4 | 4 (8%) | 3 (2%) |
| 5 | 2 (4%) | 0 |
| ASA score, n (%) | 6 (12%) | 73 (48.7%) | <0.001 b |
| 1 | 31 (62%) | 74 (49.3%) |
| 3 | 13 (26%) | 3 (2%) |
| 4 | 0 | 0 |

*Independent samples t-test

#Fig. 1 Differences in comorbidity presented by ASA classification in % for both groups
non-normally distributed outcomes. The Pearson Chi-square test was used for categorical variables. All tests were two-sided and results were considered statistically significant at a 5% level. To identify risk factors for a femoral neck fracture we used a logistic regression model with independent variables. The Odds Ratios (ORs) and the 95% Confidence Intervals (CIs) were reported. The analyses were performed using IBM SPSS, version 25.0 (IBM Corp., Armonk, NY, USA).

**Results**

**Comorbidity**
We found more comorbidities (higher CCI score and more patients with ASA class 2-3) in the fracture group (Table 1) (Fig. 1). Patients in the fracture group had lower BMI and there were more smokers in this group.

**Hip function**
No significant differences in hip function using the Oxford Hip Score and HOOS could be found, but the control group reported better HHS (97.1) compared to the fracture group (93.6) (Table 2) (Fig. 2).

**Bone mineral density**
BMD measurements were performed for all participants in the control group. Five participants in the fracture group had implants in both hips, making measurements in this area impossible. There were more patients with osteopenia and osteoporosis in the fracture group compared to the control group both when comparing results from lumbar spine, total hip, and the femoral neck (Table 3) (Fig. 3).

**Risk analysis**
A multivariate logistic regression analysis for unadjusted variables showed statistically significant differences in the OR for smoking, presence of osteoporosis, and CCI score (Table 4). The adjusted OR for obtaining a femoral neck fracture was 6.7 for smokers compared to non-smokers, and 7.5 for participants with osteoporosis compared to participants with normal BMD.

**Discussion**
The present study showed that patients aged 55-70 years with a displaced FNF were more osteoporotic and had more comorbidities compared to a gender- and age-

---

**Table 2** Differences in hip function between cases (before fracture) and controls

|                      | Fracture group n = 50 | Control group n = 150 | p-values |
|----------------------|-----------------------|-----------------------|----------|
| Mean Harris Hip Score (95% CI) | 93.6 (91.0-96.3)      | 97.1 (95.8-98.4)      | < 0.001<sup>a</sup> |
| Mean Oxford Hip Score (95% CI) | 46.2 (44.8-47.2)      | 46.0 (45.4-47.0)      | 0.173     |
| Mean HOOS (95%CI) Pain | 92.9 (89.1-96.7)      | 95.3 (93.5-97.1)      | 0.226     |
| Symptoms             | 94.3 (90.6-97.7)      | 95.2 (93.4-97.0)      | 0.226     |
| ADL                  | 92.4 (87.3-97.5)      | 95.7 (93.9-97.5)      | 0.244     |
| Sport                | 89.9 (83.6-96.2)      | 92.1 (89.3-94.8)      | 0.739     |
| QoL                  | 94.0 (89.7-98.4)      | 93.6 (90.9-96.4)      | 0.792     |

<sup>a</sup>Mann-Whitney U-test
matched control group. In addition, we found that fracture patients were more frequently smokers and had lower BMI. Thus, our data suggests that the risk of sustaining a displaced femoral neck fracture probably increases with the presence of osteoporosis, comorbidities and smoking. Regarding baseline hip function, the control group reported better scores in only one of the three hip scores used, suggesting that the difference was not clinically relevant.

There were statistically significantly more patients with low bone density in the fracture group. These findings agree well with other studies suggesting that young patients (<50 years of age) with hip fracture due to low-energy injury suffer from early osteoporosis [18, 20]. We found more comorbidities in the fracture group, measured by both the CCI score and the ASA score. These findings are supported by another study reporting comorbidities as an important and determinant factor in non-elderly patients with hip fractures [20]. We found a significantly higher proportion of smokers in the fracture group, where almost 50% were smokers at the time of injury compared to 10% in the control group. This has also been confirmed by a meta-analysis identifying smoking as a risk factor for any kind of fracture and for hip fractures in particular [33]. Other authors have postulated smoking and diabetes mellitus as the strongest independent impact factors for increased hip fracture risk [34, 35]. Diabetes has previously been described as an important risk factor for hip fractures in both women and men [35–37]. A statistically significant difference in the presence of diabetes between the fracture group and the control group was observed.

### Table 3 Differences in DXA-measures between 50 patients with FNF and the control group

| Dual-energy X-ray absorptiometry | Fracture group | Control group | p-values |
|----------------------------------|----------------|---------------|----------|
| Lumbar spine T-score             |                |               | 0.003 a  |
| T-score >-1                       | 19 (38%)       | 83 (55%)      |          |
| T-score -1 - -2.5                 | 17 (34%)       | 53 (35%)      |          |
| T-score ≤ -2.5                   | 14 (28%)       | 14 (9%)       |          |
| Hip total T-score b              |                |               | <0.001 a |
| T-score >-1                       | 8 (18%)        | 79 (53%)      |          |
| T-score -1 - -2.5                 | 22 (49%)       | 63 (42%)      |          |
| T-score ≤ -2.5                   | 15 (33%)       | 8 (5%)        |          |
| Hip femoral neck T-score b       |                |               | <0.001 a |
| T-score >-1                       | 4 (9%)         | 47 (31%)      |          |
| T-score -1 - -2.5                 | 21 (47%)       | 85 (57%)      |          |
| T-score ≤ -2.5                   | 20 (44%)       | 18 (12%)      |          |

*Pearson Chi-square test
b5 patients with implants after fracture care in both hips, making measurements in the hips impossible

Fig. 3 T-score values for fracture and control group
group could not be found in our study. However, as we had a limited number of participants in our study, the power was probably insufficient to investigate this issue.

Lower BMI was found in the fracture group. Higher BMI has been argued to protect against a hip fracture, but probably not in this age group [38]. The lower BMI might also be associated with the larger number of smokers in the fracture group. Smoking is associated with negative influence on nutrition, which may result in lower BMI among smokers [33].

Our study suggests that the presence of osteoporosis and smoking were the strongest differences between the fracture group and the control group with a seven times higher OR for patients with displaced FNF when adjusting for smoking, comorbidity and BMD.

The main strength of our study is that the DXA measurements in the fracture group were performed at the time of injury, documenting the current bone status. Furthermore, we present results from a representative and relatively homogeneous gender- and age-matched population.

However, this study has limitations, as case-control studies are prone to bias, especially selection and recall bias. The number of fracture patients was limited, as the incidence of hip fractures in this age group is low [19]. The sample size was small and included both genders, recognizing the differences in risk factors and epidemiology of hip fracture in women and men [15, 18, 38, 39]. Exact power calculations were not performed, but tripling the number of controls was used to improve statistical strength [26]. We used a standard logistic regression model for a loose-matching strategy within pair correlation, approving that conditional logistic regression is the appropriate analytical method for matched case-control studies. We did not record several well-known risk factors for hip fractures and osteoporosis, such as alcohol consumption, calcium and vitamin D levels, differences in bone-associated comorbidities (e.g. malabsorption), hormonal deficits, or the use of specific medication influencing bone quality (steroids, anti-epileptic medication) [20, 34, 35, 39–41].

Conclusions

Patients between 55 and 70 years with a displaced femoral neck fracture were more osteoporotic and have more comorbidities, than a comparable gender- and age- matched group from the general population without a femoral neck fracture. Our data indicate that these patients are frailer than expected and should probably not be compared with their peers of the same age. From a clinical perspective, patients aged 55-70 years may benefit from a similar treatment as those over 70 years, within a treatment algorithm considering biological age, individual factors, and medical challenges.

Abbreviations

95% CI: 95% confidence interval; ASA: American Society of Anesthesiologists; BMD: Bone mineral density; BMI: Body Mass Index; CCI: Charlson comorbidity index; DXA: Dual-energy x-ray absorptiometry; FNF: Femoral neck fracture; HHS: Harris Hip Score; HOOS: Hip Dysfunction and Osteoarthritis Outcome Score; IF: Internal fixation; OR: Odds ratio

Acknowledgements

We thank Jurate Saltyte-Benth from the Department of Data and Analytics at Akershus University Hospital for statistical assistance.

Author’s contributions

SB contributed to study conception and design, data collection, performed the statistical analysis with support from the Department of Data and Analytics at Akershus University Hospital and wrote the manuscript. JEG contributed to study conception and design, data collection, interpretation of data, and critical revision of the manuscript. FF contributed to study conception and design, data collection, interpretation of data, and critical revision of the manuscript. SEU contributed to study conception and design, data collection, interpretation of data, and critical revision of the manuscript. CR contributed to study conception and design, data collection, interpretation of data, and critical revision of the manuscript. All authors read and approved the final manuscript.

Funding

The first author received funding from Strategic Research Funding Akershus University Hospital and from Sophies Minde Ortopedi AS, a subsidiary of Oslo University Hospital and Akershus University Hospital. The funding body had no role in the design of the study, data collection, data analysis, data interpretation or in writing the manuscript.

Availability of data and materials

Due to regulations from the Norwegian Data Inspectorate and according to Norwegian personal protections laws, publication of the complete dataset is not legal or appropriate.

Table 4 Logistic regression with Odd ratio obtaining displaced FNF for multivariate analyses and with adjustments

|                          | Unadjusted | Adjusted a |
|--------------------------|------------|------------|
|                          | OR 95%CI p-values | OR 95%CI p-values |
| Smoking                  | 8.3 3.8-17.9 < 0.001 | 6.8 2.8-16.1 < 0.001 |
| Charlson Comorbidity Index Score | 2.6 1.6-4.1 < 0.001 | 2.0 1.2-3.6 0.011 |
| DXA femoral neck T-score > -1 normal bone | 1 1 < 0.001 | 1 1 < 0.001 |
| DXA femoral neck T-score -1 -2.5 osteopenia | 3.1 1.0-9.6 0.049 | 2.1 0.6-6.9 0.236 |
| DXA femoral neck T-score ≤ -2.5 osteoporosis | 13.6 4.1-45.3 < 0.001 | 7.6 2.1-27.9 < 0.001 |

*Adjusted for all other variables in the table in a multiple regression analysis model
Ethics approval and consent to participate
The study was performed in accordance with the Helsinki Declaration, and the protocol was approved by the Regional Ethics Committee South-East Norway (reference number REK Ser-ØS Ø 2013/1023) and the institutional ethics committee of the Akershus University Hospital, Lørenskog, Norway. Informed consent was obtained from all individual participants included in the study.

Consent for publication
Not applicable.

Competing interests
The authors declare that they have no competing interests.

Author details
1Department of Orthopedic Surgery, Akershus University Hospital, Lørenskog, Norway. 2Norwegian Hip Fracture Register, Department of Orthopedic Surgery, Haukeland University Hospital, Bergen, Norway. 3Department of Clinical Medicine, University of Bergen, Bergen, Norway. 4Department of Orthopedic Surgery, Oslo University Hospital, Oslo, Norway. 5Department of Orthopedics, Skåne University Hospital, Lund University, Malmö, Sweden. 6Institute of Clinical Medicine, University of Oslo, Oslo, Norway.

Received: 25 March 2019 Accepted: 19 July 2019
Published online: 14 August 2019

References
1. WHO. Prevention and management of osteoporosis. World Health Organ Tech Rep Ser. 2003;921:1–164.
2. Cooper C, Cole ZA, Harvey NC, Dennison EM, et al. Secular trends in the incidence of hip and other osteoporotic fractures. Osteoporos Int. 2011;22(5):1277–88.
3. Johnell O, Kanis JA. An estimate of the worldwide prevalence and disability associated with osteoporotic fractures. Osteoporos Int. 2006;17(12):1726–33.
4. Stoen RO, Nordsetten L, Meyer HE, Frihagen F, Falch JA, Loftus CM. Hip fracture incidence is decreasing in the high incidence area of Oslo. Norway Osteoporos Int. 2012;23(10):2527–34.
5. Omsland TK, Holvik K, Meyer HE, Center JR, Emaus N, Tell GS, et al. Fractureincidence in Norway 1999–2008: time trends in total incidence and hip fracture incidence: a NOREPOS study. Eur J Epidemiol. 2012;27(10):807–14.
6. Ahlborg HG, Rosengren BE, Linvenin TL, Rogmark C, Nilsson JA, Sembo J, et al. Prevalence of osteoporosis and incidence of hip fractures in women—several trends over 30 years. BMC Musculoskelet Disord. 2010;11:48.
7. Leslie WD, O’Donnell S, Jean S, Lagace C, Walsh F, Bancej C, et al. Trends in hip fracture rates in Canada. Jama. 2009;302(8):885–9.
8. Gjertsen JE, Virje T, Engesaeter LB, Lie SA, Havelin LI, Furnes O, et al. Internal screw fixation compared with bipolar hemiarthroplasty for treatment of displaced femoral neck fractures in elderly patients. J Bone Joint Surg Am. 2010;92(3):619–28.
9. Gao H, Liu Z, Xing D, Gong M. Which is the best alternative for displaced femoral neck fractures in the elderly?: a meta-analysis. Clin Orthop Relat ResClin Orthop Relat Res. 2012;470(6):1782–91.
10. Xu D, Li X, Bi F, Ma C, Lu L, Gao J. Hemiarthroplasty compared with total hip arthroplasty for displaced fractures of femoral neck in the elderly: a systematic review and meta-analysis of fourteen randomized clinical trials. Int J Clin Exp Med. 2018;11(6):5430–43.
11. Rogmark C, Kristensen MT, Vilberg B, Ronquist SS, Overgaard S, Palm H. Hip fractures in the non-elderly-who, why and whither? Injury. 2018;49(8):1465–50.
12. National Institute for Health and Care Excellence. Hip fracture management clinical guideline 124. https://www.nice.org.uk/guidance/cg124/chapter/Recommendations-for-research2017.
13. AAOS. Management of hip fractures in the elderly: evidence-based clinical practice guideline 2015. https://www.orthoguidelines.org/hipfractureguideline.
14. Frihagen F, Nordsetten L, Madsen JJE. Hemiarthroplasty or internal fixation for intracapsular displaced femoral neck fractures: randomised controlled trial. BMJ (Clinical research ed). 2007;335(7632):1251–4.
15. Sogaard AJ, Holvik K, Meyer HE, Tell GS, Gjersdal CG, Emaus N, et al. Continued decline in hip fracture incidence in Norway: a NOREPOS study. Osteoporos Int. 2016;27(7):2127–22.
16. National Hip Fracture Register, Haukeland University Hospital Bergen, annual report 2017, http://nhfwebhelse.net/.
17. Statistics Norway SSB. Available 2019, www.ssb.no
18. Lofthus CM, Oines EK, Meyer HE, Kristiansen IS, Nordsetten L, Falch JA. Young patients with hip fracture: a population-based study of bone mass analysis and risk factors for osteoporosis. Osteoporos Int. 2006;17(11):1666–72.
19. Karantana A, Boulton C, Bouloumis G, Shu KS, Scammell BE, Moran CG. Epidemiology and outcome of fracture of the hip in women aged 65 years and under: a cohort study. J Bone Joint Surg (Br). 2011;93(5):658–64.
20. Al-Ani AN, Neander G, Samuelsson B, Blomfeldt R, Ekstrom W, Hedstrom M. Risk factors for osteoporosis are common in young and middle-aged patients with femoral neck fractures regardless of trauma mechanism. Acta Orthop. 2013;84(1):54–9.
21. Heeveld MJ, Raaymakers EL, Luitse JS, Nijhof M, Gouma DJ. Femoral neck fractures: can physiologic status determine treatment choice? Clin Orthop Relat ResClin Orthop Relat Res. 2007;461:203–12.
22. Slobogean GP, Sprague SA, Scott T, Bhandari M. Complications following young femoral neck fractures. Injury. 2015;46(3):848–91.
23. Upadhyay A, Jain P, Mishra P, Manni L, Gautum VK, Dhaon BK. Delayed internal fixation of fractures of the neck of the femur in young adults. A prospective, randomised study comparing closed and open reduction. J Bone Joint Surg (Br). 2004;86(7):1035–40.
24. Bartels S, Gjertsen JE, Frihagen F, Rogmark C, Utvag SE. High failure rate after internal fixation and beneficial outcome after arthroplasty in treatment of displaced femoral neck fractures in patients between 55 and 70 years. Acta Orthop. 2018;89(1):53–8.
25. Kuo CL, Duan Y, Grady J. Unconditional or conditional logistic regression model for age-matched case-control data? Front Public Health. 2016;4:57.
26. Hodge SE, Subaran RL, Weissman MM, Fyer AJ. Designing case-control studies: decisions about the controls. Am J Psychiatry. 2012;169(8):785–9.
27. Charlson ME, Pompei P, Ales KL, Mackenzie CR. A new method of classifying prognostic comorbidity in longitudinal studies: development and validation. J Chronic Dis. 1987;40(5):373–83.
28. American Society of Anesthesiologists. New classification of physical status. Anesthesiology. 1963;24:111.
29. Harris WH. Traumatic arthritis of the hip after dislocation and acetabular fractures: treatment by mold arthroplasty. An end-result study using a new method of result evaluation. J Bone Joint Surg Am. 1969;51(4):737–55.
30. Dawson J, Fitzpatrick R, Carr A, Murray D. Questionnaire on the perceptions of patients about total hip replacement. J Bone Joint Surg (Br). 1996;78(2):185–90.
31. Nilsson AK, Klassbo M, Roos EM. Hip disability and osteoarthritis outcome score (HOOS)–validity and responsiveness in total hip replacement. BMC Musculoskelet Disord. 2003;4:10.
32. Kanis JA, Adachi JD, Cooper C, Clark P, Cummings SR, Diaz-Curiel M, et al. Standardising the descriptive epidemiology of osteoporosis: recommendations from the epidemiology and quality of life working group of IOF. Osteoporos Int. 2013;24(11):2763–4.
33. Kanis JA, Johnell O, Oden A, Johansson H, De Laet C, Esim LA, et al. Smoking and fracture risk: a meta-analysis. Osteoporos Int. 2005;16(2):155–62.
34. Holtmberg AH, Johnell O, Nilsson PM, Nilsson J, Berglund G, Akesson K. Risk factors for fragility fracture in middle age: A prospective population-based study of 33,000 men and women. Osteoporos Int. 2006;17(1):1065–77.
35. Pasoto SG, Yoshihara LA, Maeda LC, Bernik MM, Lotufo PA, Bonfa E, et al. Osteoporotic hip fractures in non-elderly patients: relevance of associated co-morbidities. Rheumatol Int. 2012;32(10):3149–53.
36. Fan Y, Wei F, Lang Y, Liu Y. Diabetes mellitus and risk of hip fractures: a meta-analysis. Osteoporos Int. 2016;27(12):2129–28.
37. Jia P, Bao L, Chen H, Yuan J, Liu W, Feng F, et al. Risk of low-energy fracture in type 2 diabetes patients: a meta-analysis of observational studies. Osteoporos Int. 2017;28(11):3113–21.
38. Sogaard AJ, Holvik K, Omsland TK, Tell GS, Dahl C, Schiøtt B, et al. Age and sex differences in body mass index as a predictor of hip fracture: a NOREPOS study. Am J Epidemiol. 2016;184(7):510–9.
39. Sogaard AJ, Ranhoff AH, Meyer HE, Omsland TK, Nystad W, Tell GS, et al. The association between alcohol consumption and risk of hip fracture
differs by age and gender in cohort of Norway: a NOREPOS study. Osteoporos Int. 2018;29(1):2457–67.

40. Stearns AT, Jaberoo MC, Ashraf R, Wheelwright EF, Maclean AD. Displaced intracapsular hip fractures in the working age alcohol-abusing patient. Scott Med J. 2009;54(1):16–20.

41. Kanis JA, Johanson H, Johnell O, Odén A, De Laet C, Eisman JA, et al. Alcohol intake as a risk factor for fracture. Osteoporos Int. 2005;16(7):737–42.

Publisher’s Note
Springer Nature remains neutral with regard to jurisdictional claims in published maps and institutional affiliations.