

■ TRAUMA

The impact of complications on quality of life and mortality after hip fracture

E. L. Goh,
M. E. Png,
D. Metcalfe,
J. Achten,
D. Appelbe,
X. L. Griffin,
J. A. Cook,
M. L. Costa,
on behalf of the
WHiTE Investigators

From University of
Oxford, Oxford, UK

Aims

There is limited information on how the development of complications influences quality of life (QoL) after hip fracture. The aim of this study was to investigate the relationship between complications, QoL, and mortality after hip fracture.

Methods

The World Hip Trauma Evaluation (WHiTE) study is a multicentre, prospective cohort study that collected data from patients aged ≥ 60 years who received surgical treatment for their hip fracture. Patients were followed up for 120 days after surgery. The primary and secondary outcomes were health-related QoL (EuroQoL five-dimension five-level questionnaire; EQ-5D-5L) and mortality, respectively. Linear and logistic regression models were fitted to assess the relationship between complications, EQ-5D-5L, and mortality.

Results

Among 24,523 patients with a hip fracture, the mean differences in EQ-5D-5L in patients who had surgery-specific complications were: prosthesis dislocation -0.14 (95% CI -0.20 to -0.08); fixation failure 0.00 (95% CI -0.15 to 0.14); periprosthetic or peri-implant fracture -0.08 (95% CI -0.18 to 0.02); reoperation for any indication -0.09 (95% CI -0.14 to -0.05); surgical site infection (SSI) -0.06 (95% CI -0.10 to -0.01); and deep SSI -0.13 (95% CI -0.20 to -0.07). The mean differences in EQ-5D-5L for the general complications were: acute kidney injury -0.05 (95% CI -0.07 to -0.02); blood transfusion -0.01 (95% CI -0.03 to 0.01); lower respiratory tract infection -0.07 (95% CI -0.09 to -0.05); urinary tract infection 0.01 (95% CI -0.01 to 0.03); cerebrovascular accident (CVA) -0.17 (95% CI -0.25 to -0.09); myocardial infarction (MI) -0.14 (95% CI -0.20 to -0.08); and venous thromboembolism 0.03 (95% CI -0.02 to 0.08).

Conclusion

We observed worse health-related QoL in patients who had a complication after hip fracture. Those who underwent revision surgery or had a prosthesis dislocation or deep SSI experienced similar levels of disability at 120 days to those with a CVA or MI.

Cite this article: *Bone Joint J* 2025;107-B(10):1118–1124.

Introduction

A hip fracture is the most common serious injury in older adults, with over ten million hip fractures occurring worldwide every year.¹ There is substantial disability and mortality that follows the injury, for which the burden of disease is estimated at 27 disability-adjusted life years per 1,000 individuals, equivalent to an average loss of 2.7% of healthy life expectancy.²⁻⁴ The majority of people experience a reduction in mobility and independence after their injury that they never fully regain.⁵ Mortality has been the focus of previous

hip fracture research, but it is increasingly evident that quality of life (QoL) after hip fracture is a more meaningful outcome.^{6,7}

There is a significant decrease in QoL that follows a hip fracture.^{8,9} Although a substantial recovery occurs after hip fracture surgery, patients do not tend to fully regain their pre-injury QoL.⁸⁻¹⁰ The same factors that influence mortality after hip fracture have also been shown to be associated with QoL, with previous work indicating that higher mortality and lower QoL is observed in patients who develop complications.^{8,11} Given

Correspondence should be sent to E. L. Goh; email: enlin.goh@ndorms.ox.ac.uk

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doi:10.1302/0301-620X.107B10.
BJJ-2024-1448.R1 \$2.00

Bone Joint J
2025;107-B(10):1118–1124.

Table I. Baseline characteristics of the World Hip Trauma Evaluation cohort.

Characteristic	Overall
Patients, n	24,523
Mean age, yrs (SD)	82.9 (8.4)
Sex, n (%)	
Male	7,337 (29.9)
Female	17,186 (70.1)
Regular smoker, n (%)	
Yes	2,139 (8.7)
No	20,958 (85.5)
Missing	1,426 (5.8)
Weekly alcohol consumption, n (%)	
0 to 7 units	20,453 (83.4)
8 to 14 units	1,458 (5.9)
15 to 21 units	507 (2.1)
> 21 units	604 (2.5)
Missing	2,001 (8.2)
Diabetic, n (%)	
Yes	3,603 (14.7)
No	19,566 (79.8)
Missing	1,354 (5.5)
Renal failure, n (%)	
Yes	1,807 (7.4)
No	21,335 (87.0)
Missing	1,381 (5.6)
Cognitive impairment, n (%)	
Yes	7,761 (31.6)
No	15,541 (63.4)
Missing	1,221 (5.0)
Residential status, n (%)	
Own home	13,971 (57.0)
Residential or nursing care	2,463 (10.0)
Rehabilitation unit	86 (0.4)
Acute hospital	85 (0.3)
Other	69 (0.3)
Missing	7,849 (32.0)
ASA grade, n (%)	
I	453 (1.8)
II	5,527 (22.5)
III	13,545 (55.2)
IV	3,546 (14.5)
V	72 (0.3)
Missing	1,380 (5.6)
Fracture type, n (%)	
Femoral neck – undisplaced (B1)	1,647 (6.7)
Femoral neck – displaced (B3)	15,103 (61.6)
Trochanteric – simple (A1)	2,393 (9.8)
Trochanteric – unstable (A2)	3,408 (13.9)
Trochanteric – transtrochanteric (A3)	817 (3.3)
Subtrochanteric	929 (3.8)
Missing	226 (0.9)
Operation type, n (%)	
Sliding hip screw	5,546 (22.6)
Cephalomedullary nail	2,645 (10.8)
Cannulated screws	644 (2.6)
Hip hemiarthroplasty	11,283 (46.0)
Total hip arthroplasty	2,076 (8.5)
Missing	2,329 (9.5)

ASA, American Society of Anesthesiologists.

that the risk of complications is high, preventing these complications is of great importance as this may mitigate the burden of disability.^{6,12,13} However, there are few data available on QoL after hip fracture, and how this may be affected by the complications occurring during recovery.¹⁴

The aim of this study was to investigate the relationship between complications, health-related QoL, and mortality after hip fracture.

Methods

Study design, setting, and participants. The World Hip Trauma Evaluation (WHiTE) study was a multicentre, prospective observational cohort study that collected data on the assessment, treatment, and recovery of a comprehensive hip fracture cohort in England, Wales, and Northern Ireland.¹⁵ Participants were eligible for inclusion if they were aged 60 years or older and received operative treatment for their hip fracture at any of the 77 participating NHS hospitals. On enrolment, patients received treatment under a standardized care pathway based on the National Institute for Health and Care Excellence Hip Fracture Guidelines (CG124).¹⁶ All patients were followed up for 120 days after injury by telephone interviews and postal questionnaires.

Ethics approval. Ethics approval was granted by the London-Camberwell St Giles Research Ethics Committee.¹⁵ This study was registered with the National Institute of Health and Care Research Portfolio (UKCRN ID12351) and the ISRCTN registry (ISRCTN63982700). Written consent to participate in the study was obtained from all patients. Patients who lacked capacity to consent to participate were still included in consultation with their carers.

Data sources. The WHiTE dataset contains data on a core outcome set of patient-reported outcome measures in addition to variables that are routinely measured by the UK National Hip Fracture Database.¹⁷ The full list of variables and outcomes collected as part of the study has been described previously.¹⁵ Data were stored on the OpenClinica v. 3.7 data collection system (OpenClinica, USA).

Variables. The primary variables of interest were the development of surgery-specific and general complications within 120 days of hip fracture. These complications were prespecified and are listed in Appendix A in the Supplementary Material.^{12,15} The process of how the data on complications were collected is described in Appendix B. The covariates selected for inclusion in the statistical models were specified a priori and are re-described in detail in Appendix C. The variables considered included patient-related factors such as the patient's demographics, pre-injury health-related quality of life (QoL), comorbidities, and injury, which have previously been shown to explain the variation in mortality and QoL after hip fracture.^{8,18,19}

Population. There were 24,523 patients enrolled in the WHiTE study, with complete follow-up available for 22,228 patients (90.6%). The mean age of the overall cohort was 82.9 years (SD 8.4) and 17,186 (70.1%) were female. Surgery-specific and general complications occurred in 1,052 (4.3%) and 4,983 (20.3%) patients, respectively. The baseline characteristics are described in Table I.

Table II. Number and proportion of patients with and without a complication and their respective mean pre-injury EuroQol five-dimension five-level questionnaire (EQ-5D-5L).

Complication	Cohort			Mean pre-injury EQ-5D-5L		
	Yes	No	Proportion (%)	Yes	No	p-value*
Surgery-specific complication						
Prosthesis dislocation	148	13,211	1.1	0.74	0.74	0.670
Fixation failure	78	8,757	0.9	0.79	0.75	0.074
Periprosthetic or peri-implant fracture	77	24,446	0.3	0.56	0.53	0.149
Reoperation for any indication	458	24,065	1.9	0.72	0.74	0.047
Revision surgery	210	24,313	0.9	0.73	0.74	0.426
Reoperation for infection	181	24,342	0.7	0.70	0.74	0.023
Surgical site infection (all)	767	23,756	3.1	0.72	0.74	0.004
Surgical site infection (deep or organ space)	305	24,218	1.2	0.72	0.74	0.059
General complication						
Acute kidney injury	810	23,713	3.3	0.68	0.74	< 0.001
Blood transfusion	1,425	23,098	5.8	0.71	0.75	< 0.001
Lower respiratory tract infection	2,132	22,391	8.7	0.67	0.75	< 0.001
Urinary tract infection	1,609	22,914	6.6	0.69	0.75	< 0.001
Cerebrovascular accident	155	24,368	0.6	0.73	0.74	0.605
Myocardial infarction	171	24,352	0.7	0.70	0.74	0.023
Venous thromboembolism	414	24,109	1.7	0.74	0.74	0.934
Deep vein thrombosis	304	24,219	1.2	0.77	0.74	0.072
Pulmonary embolism	157	24,366	0.6	0.69	0.74	0.020

*Linear regression model.

Outcomes. The primary outcome of interest was health-related QoL, as measured with the EuroQol five-dimension five-level questionnaire (EQ-5D-5L) at 120 days after hip fracture.²⁰ This is a validated self-administered patient-reported outcome measure that is based on a five-dimension health status classification system and a separate visual analogue scale (EQ-VAS) for overall health state (0 to 100 scale). The responses are converted into an overall score using a published utility algorithm for the UK population.²¹ The minimal clinically important difference in EQ-5D-5L is 0.05.²² The secondary outcome of interest was all-cause mortality during the same period.

Statistical analysis. The baseline characteristics of the study cohort were reported as mean (SD) for continuous measures, and proportions (binary) as appropriate. The cohort was divided into groups according to whether they had a complication or not, for each surgery-specific and general complication. The independent-samples *t*-test was used to compare the pre-injury EQ-5D-5L between these groups. Linear and logistic regression models were used to examine the relationship between each complication, EQ-5D-5L, and mortality, respectively. All models were adjusted for the aforementioned covariates. Death-adjusted EQ-5D-5L was used; patients who died before completing the postinjury EQ-5D-5L were assigned a value of zero.²³ Analyses were conducted on the complete case data (without imputation); only data from patients who completed the baseline and 120-day follow-up for EQ-5D-5L were analyzed. The results are presented as mean differences for the linear regression models and odds ratios (ORs) for the logistic regression models with corresponding 95% CI and p-values. A nominal two-sided 5% significance level was applied throughout. In the primary analysis, the models evaluated the interaction

between EQ-5D-5L at the 120-day follow-up or mortality during the 120-day follow-up, and complications within 30 days of follow-up, to minimize distortion of the effect size of the complication by events that may have occurred close to the 120-day follow-up timepoint. In the sensitivity analyses, the interaction between EQ-5D-5L at 120 days of follow-up or mortality during the 120 days of follow-up, and complications occurring at any point during the 120 days of follow-up, was compared. Statistical analyses were performed with R v. 4.2.3 (R Foundation for Statistical Computing, Austria). The EQ-5D-5L utility score was mapped from the five dimensions using the 'eq5d' R package.²⁴ Linear and logistic regression models were fitted using the 'lme4' R package.²⁵

Results

The mean pre-injury EQ-5D-5L was 0.73 (95% CI 0.73 to 0.74) and the mean postinjury EQ-5D-5L at 120 days of follow-up was 0.53 (95% CI 0.53 to 0.54), representing a change of -0.20 (95% CI -0.20 to -0.19). The number and proportion of patients with and without a complication and their respective pre-injury EQ-5D-5L are reported in Table II.

Surgery-specific complications. The adjusted mean difference in EQ-5D-5L in patients who had a surgery-specific complication within 30 days postoperatively is reported in Table III. The change in EQ-5D-5L for a surgery-specific complication was -0.07 (95% CI -0.10 to -0.04; $p < 0.001$); prosthesis dislocation -0.14 (95% CI -0.20 to -0.08; $p < 0.001$); fixation failure 0.00 (95% CI -0.15 to 0.14; $p = 0.969$); periprosthetic or peri-implant fracture -0.08 (95% CI -0.18 to 0.02; $p = 0.111$); reoperation for any indication -0.09 (95% CI -0.14 to -0.05; $p < 0.001$); revision surgery -0.14 (95% CI -0.21 to -0.06; $p < 0.001$); reoperation for infection -0.10 (95% CI -0.16 to -0.03; $p = 0.002$); surgical

Table III. Adjusted mean difference in pre- and post-injury EuroQol five-dimension five-level questionnaire (EQ-5D-5L) for a complication occurring within 30 days postoperatively. Symbols denote the clinical effect size of the change in pre- and post-injury EQ-5D-5L relative to the minimal clinically important difference of 0.05.

Complication	Mean difference (95% CI)	p-value
Surgery-specific complication		
Any surgery-specific complication*	-0.07 (-0.10 to -0.04)	< 0.001
Prosthesis dislocation†	-0.13 (-0.20 to -0.05)	< 0.001
Fixation failure	0.00 (-0.15 to 0.14)	0.969
Periprosthetic or peri-implant fracture	-0.08 (-0.18 to 0.02)	0.111
Reoperation for any indication†	-0.09 (-0.14 to -0.05)	< 0.001
Revision surgery†	-0.14 (-0.21 to -0.06)	< 0.001
Reoperation for infection*	-0.10 (-0.16 to -0.03)	0.002
Surgical site infection (all)*	-0.06 (-0.10 to -0.01)	0.007
Surgical site infection (deep or organ space)†	-0.13 (-0.20 to -0.07)	< 0.001
General complication		
Any general complication*	-0.03 (-0.04 to -0.02)	< 0.001
Acute kidney injury*	-0.05 (-0.07 to -0.02)	0.001
Blood transfusion	-0.01 (-0.03 to 0.01)	0.175
Lower respiratory tract infection†	-0.07 (-0.09 to -0.05)	< 0.001
Urinary tract infection	0.01 (-0.01 to 0.03)	0.454
Cerebrovascular accident†	-0.17 (-0.25 to -0.09)	< 0.001
Myocardial infarction†	-0.14 (-0.20 to -0.08)	< 0.001
Venous thromboembolism	0.03 (-0.02 to 0.08)	0.299
Deep vein thrombosis	0.06 (-0.01 to 0.12)	0.097
Pulmonary embolism	-0.01 (-0.08 to 0.06)	0.773

* Δ EQ-5D-5L (95% CI) < 0.05 | $p < 0.05$.

† Δ EQ-5D-5L (95% CI) ≥ 0.05 | $p < 0.05$.

site infection (SSI) -0.06 (95% CI -0.10 to -0.01; $p = 0.007$); and deep SSI -0.13 (95% CI -0.20 to -0.07; $p < 0.001$).

The risk of mortality in patients who had a surgery-specific complication within 30 days postoperatively is reported in Table IV. The risk of mortality associated with a surgery-specific complication was 1.51 (95% CI 1.11 to 2.01; $p < 0.001$); prosthesis dislocation 2.01 (95% CI 1.03 to 2.68; $p = 0.030$); periprosthetic or peri-implant fracture 0.33 (95% CI 0.05 to 1.11; $p = 0.134$); reoperation for any indication 1.57 (95% CI 1.04 to 2.31; $p = 0.028$); revision surgery 1.25 (95% CI 0.56 to 2.51; $p = 0.552$); reoperation for infection 1.78 (95% CI 1.01 to 3.03; $p = 0.044$), SSI 1.77 (95% CI 1.22 to 2.52; $p = 0.002$); and deep or organ space SSI 2.12 (95% CI 1.20 to 3.57; $p = 0.007$).

General complications. The adjusted mean difference in EQ-5D-5L in patients who had a general complication within 30 days postoperatively is reported in Table III. The change in EQ-5D-5L for a general complication was -0.03 (95% CI -0.04 to -0.02; $p < 0.001$); acute kidney injury (AKI) -0.05 (95% CI -0.07 to -0.02; $p = 0.001$); blood transfusion -0.01 (95% CI -0.03 to 0.01; $p = 0.175$); lower respiratory tract infection (LRTI) -0.07 (95% CI -0.09 to -0.05; $p < 0.001$); urinary tract infection (UTI) 0.01 (95% CI -0.01 to 0.03; $p = 0.454$); cerebrovascular accident (CVA) -0.17 (95% CI -0.25 to -0.09; $p < 0.001$); myocardial infarction (MI) -0.14 (95% CI -0.20 to -0.08; $p < 0.001$); venous thromboembolism (VTE) 0.03 (95%

Table IV. Adjusted odds ratio for mortality for a complication occurring within 30 days postoperatively.

Complication	OR (95% CI)	p-value
Surgery-specific complication		
Any surgery-specific complication	1.51 (1.11 to 2.01)	0.007
Prosthesis dislocation	2.01 (1.03 to 3.68)	0.030
Fixation failure*	< 0.01 (N/A)	0.948
Periprosthetic or peri-implant fracture	0.33 (0.05 to 1.11)	0.134
Reoperation for any indication	1.57 (1.04 to 2.31)	0.028
Revision surgery	1.25 (0.56 to 2.51)	0.552
Reoperation for infection	1.78 (1.00 to 3.03)	0.044
Surgical site infection (all)	1.77 (1.22 to 2.52)	0.002
Surgical site infection (deep or organ space)	2.12 (1.20 to 3.57)	0.007
General complication		
Any general complication	1.51 (1.35 to 1.69)	< 0.001
Acute kidney injury	1.76 (1.40 to 2.20)	< 0.001
Blood transfusion	1.05 (0.86 to 1.27)	0.623
Lower respiratory tract infection	2.46 (2.12 to 2.84)	< 0.001
Urinary tract infection	0.82 (0.65 to 1.02)	0.081
Cerebrovascular accident	2.81 (1.51 to 5.02)	< 0.001
Myocardial infarction	5.48 (3.50 to 8.53)	< 0.001
Venous thromboembolism	1.18 (0.67 to 1.96)	0.544
Deep vein thrombosis	0.48 (0.14 to 1.19)	0.164
Pulmonary embolism	1.53 (0.76 to 2.86)	0.201

*Unable to estimate coefficient due to the small number of events across the groups.

N/A, not applicable.

CI -0.02 to 0.08; $p = 0.299$); deep vein thrombosis (DVT) 0.06 (95% CI -0.01 to 0.12; $p = 0.097$); and pulmonary embolism (PE) -0.01 (95% CI -0.08 to 0.06; $p = 0.773$).

The risk of mortality in patients who had a general complication within 30 days post-operation is reported in Table IV. The risk of mortality associated with a general complication was 1.51 (95% CI 1.35 to 1.69; $p < 0.001$); AKI 1.76 (95% CI 1.40 to 2.20; $p < 0.001$); blood transfusion 1.05 (95% CI 0.86 to 1.27; $p = 0.623$); LRTI 2.46 (95% CI 2.12 to 2.84; $p < 0.001$); UTI 0.82 (95% CI 0.65 to 1.02; $p = 0.081$); CVA 2.81 (95% CI 1.51 to 5.02; $p < 0.001$); MI 5.48 (95% CI 3.50 to 8.53; $p < 0.001$); VTE 1.18 (95% CI 0.67 to 1.96; $p = 0.544$); DVT 0.48 (95% CI 0.14 to 1.19; $p = 0.164$); and PE 1.53 (95% CI 0.76 to 2.86; $p = 0.201$).

Sensitivity analysis. The results of the sensitivity analysis for EQ-5D-5L and mortality associated with surgery-specific and general complications are presented in Appendix D of the Supplementary Material. The mean differences for EQ-5D-5L and ORs for mortality were similar in both analyses.

Discussion

We observed a clinically meaningful reduction in health-related QoL after hip fracture in this study, equivalent to a decrease in EQ-5D-5L of 0.20. This decline is consistent with reports from previous studies in smaller hip fracture cohorts, where the residual deficit following injury has been found to range between 0.19 and 0.22.^{8,26-28} Our findings underline the profound impact that a hip fracture has on mobility, function, and independence in an already frail population. Patients who developed

a complication after hip fracture had a larger decrease in QoL compared with those who did not. Surgery-specific complications had a twofold greater impact on EQ-5D-5L compared with general complications. Much of the decline in QoL we observed in the cohort as a whole could be attributed to one of a small number of complications. In particular, patients who had a prosthesis dislocation, deep SSI, or revision surgery experienced a similar reduction in QoL at 120 days as those who had a CVA or MI.

Mortality during the follow-up period was 12.4%. It was significantly higher in patients who developed a complication than those who did not. Our analysis identified several complications that were independently associated with excess mortality after hip fracture. The largest effect sizes were observed with general complications; mortality was five times higher in patients with a MI, three times higher in those with a CVA, and two and a half times higher in those with a LRTI. For comparison, mortality was two times higher in patients with a prosthesis dislocation or deep SSI, and one and a half times higher in those who required further surgery for any indication compared with patients with no complication. Studies that have investigated the interaction between mortality and surgery-specific complications such as prosthesis dislocation, SSI, and reoperation have made similar observations, albeit with weaker associations, which can be attributed to the retrospective design, small population size, and incomplete representation.^{29–31}

Our findings indicate that the development of complications explains some of the variation observed between patients in QoL and mortality after hip fracture. We have previously observed that the same patient-related factors that influence QoL and mortality also influence the development of complications.¹¹ As such, these complications are likely to be an important mediating factor in the relationship between patient-, healthcare system-, and treatment-related factors and clinical outcome.^{11,19} A number of the complications that have been found to have the largest impact on outcome are potentially preventable.¹³ Notably, the risks of revision surgery and prosthesis dislocation are affected by operation type, whereas the risk of MI is influenced by the timing of surgery.^{11,13} Preventing such complications will be important in improving outcome after hip fracture.

A key strength of this study is large sample size and the high rate of complete follow-up (> 90%), which allows for examination of relationship between variables. The WHiTE cohort has been shown to be representative of the wider population of hip fractures in the UK and comparable with other populations worldwide.^{12,32} We accounted for mortality as part of the reporting of health-related QoL (using death-adjusted EQ-5D-5L); this is because excluding patients who died during follow-up overestimates the true effect size of EQ-5D-5L.²³ There are important limitations that should be considered. The associations identified between complications, QoL, and mortality cannot be taken as proof of a causal pathway per se. While we adjusted for known factors linked to outcomes after hip fracture, it is possible that there are unknown factors that were not accounted for, which might provide a link between hip fractures and QoL and mortality not mediated by complications.

In conclusion, patients who had a complication after hip fracture had worse health-related QoL and mortality than those who

did not. Those who had a prosthesis dislocation, deep SSI, or revision surgery experienced similar levels of disability to those with a CVA or MI. Interventions aimed at preventing these complications may improve QoL and reduce mortality after hip fracture and should be the focus of future research.



Take home message

- In the WHiTE study, patients with a complication after their hip fracture had worse quality of life and higher risk of death.
- Patients who had revision surgery, prosthesis dislocation or deep surgical site infection experienced similar levels of disability to those with a stroke or heart attack.

Social media

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Supplementary material



- List of prespecified complications of interest and of covariates included in the regression models, description of the data collection process for complications, and sensitivity analyses.

References

- Dong Y, Zhang Y, Song K, Kang H, Ye D, Li F.** What was the epidemiology and global burden of disease of hip fractures from 1990 to 2019? Results from an additional analysis of the global burden of disease study 2019. *Clin Orthop Relat Res.* 2023;481(6):1209–1220.
- Johnell O, Kanis JA.** An estimate of the worldwide prevalence and disability associated with osteoporotic fractures. *Osteoporos Int.* 2006;17(12):1726–1733.
- Shah A, Judge A, Griffin XL.** Incidence and quality of care for open fractures in England between 2008 and 2019: a cohort study using data collected by the trauma audit and research network. *Bone Joint J.* 2022;104-B(6):736–746.
- Papadimitriou N, Tsilidis KK, Orfanos P, et al.** Burden of hip fracture using disability-adjusted life-years: a pooled analysis of prospective cohorts in the chances consortium. *Lancet Public Health.* 2017;2(5):e239–e246.
- Dyer SM, Crotty M, Fairhall N, et al.** A critical review of the long-term disability outcomes following hip fracture. *BMC Geriatr.* 2016;16(1):158.
- Goh EL, Khatri A, Costa AB, et al.** Prevalence of complications in older adults after hip fracture surgery. *Bone Joint J.* 2025;107-B(2):139–148.
- Griffiths F, Mason V, Boardman F, et al.** Evaluating recovery following hip fracture: a qualitative interview study of what is important to patients. *BMJ Open.* 2015;5(1):e005406.
- Griffin XL, Parsons N, Achten J, Fernandez M, Costa ML.** Recovery of health-related quality of life in a United Kingdom hip fracture population. The Warwick Hip Trauma Evaluation: a prospective cohort study. *Bone Joint J.* 2015;97-B(3):372–382.
- Gjertsen JE, Baste V, Fevang JM, Furnes O, Engesaeter LB.** Quality of life following hip fractures: results from the Norwegian hip fracture register. *BMC Musculoskelet Disord.* 2016;17(1):265.
- Campenfeldt P, Ekström W, Al-Ani AN, Weibust E, Greve K, Hedström M.** Health related quality of life and mortality 10 years after a femoral neck fracture in patients younger than 70 years. *Injury.* 2020;51(10):2283–2288.
- Goh EL, Png ME, Metcalfe D, et al.** Risk factors associated with the development of complications after a hip fracture. *Bone Joint J.* 2025;107-B(9):950–956.
- Goh EL, Lerner RG, Achten J, Parsons N, Griffin XL, Costa PML.** Complications following hip fracture: results from the World Hip Trauma Evaluation cohort study. *Injury.* 2020;51(6):1331–1336.
- Goh EL, Png ME, Metcalfe D, et al.** The risk of complications after hip fracture. *Bone Joint J.* 2025;107-B(3):362–367.
- Walsh ME, Kristensen PK, Hjelholt TJ, et al.** Systematic review of multivariable prognostic models for outcomes at least 30 days after hip fracture finds 18 mortality

- models but no nonmortality models warranting validation. *J Clin Epidemiol*. 2024;173:111439.
15. **Costa ML, Griffin XL, Achten J, et al.** World hip trauma evaluation (WHiTE): framework for embedded comprehensive cohort studies. *BMJ Open*. 2016;6(10):e011679.
 16. **No authors listed.** Hip fracture: management: clinical guideline (CG124). National Institute for Health and Care Excellence. 2023. <https://www.nice.org.uk/guidance/cg124> (date last accessed 28 July 2025).
 17. **No authors listed.** 15 Years of Quality Improvement: The 2023 National Hip Fracture Database Report on 2022. National Hip Fracture Database, 2023. <https://www.nhfd.co.uk/2023report> (date last accessed 28 July 2025).
 18. **Schilling PL, Bozic KJ.** Development and validation of perioperative risk-adjustment models for hip fracture repair, total hip arthroplasty, and total knee arthroplasty. *J Bone Joint Surg Am*. 2016;98-A(1):e2.
 19. **Kjærøvik C, Gjertsen JE, Stensland E, Saltyte-Benth J, Soereide O.** Modifiable and non-modifiable risk factors in hip fracture mortality in Norway, 2014 to 2018: a linked multiregistry study. *Bone Joint J*. 2022;104-B(7):884–893.
 20. **Brooks R, De CF.** EuroQol: the current state of play. *Health Policy*. 1996;37(1): 53–72.
 21. **Dolan P.** Modeling valuations for EuroQol health states. *Med Care*. 1997;35(11):1095–1108.
 22. **McClure NS, Sayah FA, Xie F, Luo N, Johnson JA.** Instrument-defined estimates of the minimally important difference for EQ-5D-5L index scores. *Value Health*. 2017;20(4):644–650.
 23. **Parsons N, Griffin XL, Achten J, Chesser TJ, Lamb SE, Costa ML.** Modelling and estimation of health-related quality of life after hip fracture: a re-analysis of data from a prospective cohort study. *Bone Joint Res*. 2018;7(1):1–5.
 24. **Morton F, Singh Nijjar J.** Methods for analysing 'EQ-5D' data and calculating 'EQ-5D' index scores version 0.15.7. R-Project. <https://cran.r-project.org/web/packages/eq5d/eq5d.pdf> (date last accessed 28 July 2025).
 25. **Bates D, Mächler M, Bolker B, Walker S.** Fitting linear mixed-effects models using lme4. *J Stat Soft*. 2015;67(1):1–48.
 26. **Gjertsen J-E, Vinje T, Lie SA, et al.** Patient satisfaction, pain, and quality of life 4 months after displaced femoral neck fractures: a comparison of 663 fractures treated with internal fixation and 906 with bipolar hemiarthroplasty reported to the Norwegian hip fracture register. *Acta Orthop*. 2008;79(5):594–601.
 27. **Beaupre LA, Jones CA, Johnston DWC, Wilson DM, Majumdar SR.** Recovery of function following a hip fracture in geriatric ambulatory persons living in nursing homes: prospective cohort study. *J Am Geriatr Soc*. 2012;60(7):1268–1273.
 28. **Ekström W, Al-Ani AN, Sääf M, Cederholm T, Ponzer S, Hedström M.** Health related quality of life, reoperation rate and function in patients with diabetes mellitus and hip fracture—a 2 year follow-up study. *Injury*. 2013;44(6):769–775.
 29. **Blanco JF, da Casa C, Fidalgo H, et al.** Effect of hip hemiarthroplasty dislocation on mortality after hip fracture surgery. *Rev Esp Cir Ortop Traumatol*. 2023;67(1):3–11.
 30. **Pollard TCB, Newman JE, Barlow NJ, Price JD, Willett KM.** Deep wound infection after proximal femoral fracture: consequences and costs. *J Hosp Infect*. 2006;63(2):133–139.
 31. **Thakar C, Alsousou J, Hamilton TW, Willett K.** The cost and consequences of proximal femoral fractures which require further surgery following initial fixation. *J Bone Joint Surg Br*. 2010;92-B(12):1669–1677.
 32. **Metcalfe D, Costa ML, Parsons NR, et al.** Validation of a prospective cohort study of older adults with hip fractures. *Bone Joint J*. 2019;101-B(6):708–714.

Author information:

E. L. Goh, PhD, MRCS, Clinical Research Fellow in Musculoskeletal Trauma
D. Metcalfe, PhD, FRCEM, Kadoorie Associate Professor of Emergency
Medicine

J. Achten, PhD, Research Manager

D. Appelbe, PhD, Senior Research Information Specialist

M. L. Costa, PhD, FRCS (Orth), Professor of Orthopaedic Trauma Surgery
Oxford Trauma and Emergency Care, Nuffield Department of Orthopaedics,
Rheumatology and Musculoskeletal Sciences, Kadoorie Research Centre,
University of Oxford, Oxford, UK.

M. E. Png, PhD, Senior Researcher in Health Economics, Nuffield
Department of Primary Care Health Sciences, University of Oxford, Oxford,
UK.

X. L. Griffin, PhD, FRCS (Orth), Professor of Orthopaedic Trauma Surgery,
Bone and Joint Health, Blizzard Institute, Queen Mary University London,
London, UK.

J. A. Cook, PhD, Professor of Clinical Trials and Medical Statistics, Oxford
Clinical Trials Research Unit, Nuffield Department of Orthopaedics,
Rheumatology and Musculoskeletal Sciences, University of Oxford, Oxford,
UK.

Author contributions:

E. L. Goh: Conceptualization, Data curation, Formal analysis, Investigation,
Methodology, Software, Visualization, Writing – original draft, Writing –
review & editing.

M. E. Png: Conceptualization, Data curation, Investigation, Methodology,
Resources, Software, Supervision, Validation, Writing – original draft,
Writing – review & editing.

D. Metcalfe: Methodology, Resources, Supervision, Writing – original draft,
Writing – review & editing.

J. Achten: Funding acquisition, Investigation, Methodology, Project
administration, Resources, Writing – original draft, Writing – review &
editing.

D. Appelbe: Funding acquisition, Investigation, Methodology, Project
administration, Resources, Writing – original draft, Writing – review &
editing.

X. L. Griffin: Conceptualization, Funding acquisition, Methodology, Writing –
original draft, Writing – review & editing.

J. A. Cook: Conceptualization, Formal analysis, Methodology, Resources,
Software, Supervision, Visualization, Writing – original draft, Writing –
review & editing.

M. L. Costa: Conceptualization, Funding acquisition, Investigation,
Methodology, Resources, Supervision, Validation, Writing – original draft,
Writing – review & editing.

Funding statement:

The authors disclose receipt of the following financial or material support
for the research, authorship, and/or publication of this article: this study
was supported by National Institute for Health and Care Research (NIHR)
Oxford Biomedical Research Centre and NIHR Barts and St Georges
Biomedical Research Centre.

ICMJE COI statement:

E. L. Goh reports a doctoral research fellowship from the National Institute
for Health and Care Research (NIHR), which enabled research for this
study. J. Achten reports funding from the NIHR Oxford Biomedical
Research Centre, related to this study as well as grants or contracts
from the NIHR, unrelated to this study. D. Appelbe reports funding from
the NIHR Health Technology Assessment (HTA), related to this study.
D. Metcalfe reports an Advanced Fellowship from the NIHR, and an
infrastructure grant from the Kadoorie Charitable Foundation, both paid
to University of Oxford and not related to this study. X. L. Griffin reports
funding from the NIHR Research for Patient Benefit (RfPB), related to this
study, as well as multiple grants from UK Research and Innovation (UKRI)
and charity, unrelated to this study. J. A. Cook reports funding from the
NIHR, related to this study. M. L. Costa reports funding from the NIHR
and the Wellcome Trust, unrelated to this study.

Data sharing:

The datasets generated and analyzed in the current study are not publicly
available due to data protection regulations. Access to data is limited to the
researchers who have obtained permission for data processing. Further
inquiries can be made to the corresponding author.

Acknowledgements:

This study was funded by the National Institute of Health and Care
Research (NIHR) Oxford Biomedical Research Center (BRC) and supported
by the NIHR Barts and St George's BRC. The views expressed are those
of the authors and not necessarily those of the NIHR or the Department
of Health and Social Care. The authors thank the members of the World
Hip Trauma Evaluation (WHiTE) Oversight Committee (Tim Chesser, Iain
Moppett, Antony Johansen, Alwin McGibbon, Karen May, and Richard
Grant) and the WHiTE Scientific Committee (Stu White, Tim Chesser,
Jenny Gould, Josephine Rowling, Mark Baxter, Philip Bell, Rafa Pinedo-
Villanueva, Sallie Lamb, Andy Judge, and Chris Boulton).

Ethical review statement:

Ethics approval was granted by the London-Camberwell St Giles Research
Ethics Committee. Written consent to participate in the study was obtained

from all patients. Patients that lacked capacity to consent to participate were still included, in consultation with their carers.

Open access funding:

The open access fee for this article was funded by the National Institute for Health and Care Research (NIHR) Oxford Biomedical Research Centre.

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4.0/), which permits unrestricted use, distribution, and reproduction in any medium or format, provided the original author and source are credited.

Trial registration number:

This study was registered with the National Institute for Health and Care Research (NIHR) Portfolio (UKCRN ID12351) and the ISRCTN registry (ISRCTN63982700).

This article was primary edited by A. D. Liddle.