



## Diabetes mellitus and gender have a negative impact on the outcome of hip fracture surgery – a pilot study

Title	Diabetes mellitus and gender have a negative impact on the outcome of hip fracture surgery – a pilot study
Author(s)	Galbraith, Adam Samuel;Sanz-Nogués, Clara;Glynn, Sharon;Coleman, Cynthia M.;Murphy, Colin
Publication Date	2019-11-07
Publisher	Wiley

1 **Title:**

2 Diabetes Mellitus and Gender have a Negative Impact on the Outcome of Hip Fracture  
3 Surgery – a Pilot Study

4 **Running Title:**

5 Hio Fracture Mortality and Diabetes

6 **Authors:**

7 1. **Name:** Adam Samuel Galbraith (Corresponding Author)

8 **Affiliation address:** Department of Trauma & Orthopaedics, Galway University  
9 Hospital, Newcastle Road, Galway, Ireland

10 **Telephone:** 0858136764

11 **Email:** adamgalbraith@rcsi.com

12

13 2. **Name:** Clara Sanz-Nogués

14 **Affiliation address:** Regenerative Medicine Institute, National University of  
15 Ireland Galway, Biomedical Science 1<sup>st</sup> Floor South, Upper Newcastle, Galway,  
16 Ireland

17 **Email:** clara.sanznogues@nuigalway.ie

18

19 3. **Name:** Sharon Glynn

20 **Affiliation address:** Discipline of Pathology, Lambe Institute for Translational  
21 Medicine, School of Medicine, National University of Ireland Galway, Costello  
22 Road, Galway, Ireland

23 **Email:** sharon.glynn@nuigalway.ie

24

25 4. **Name:** Cynthia M. Coleman

26 **Affiliation address:** Regenerative Medicine Institute, National University of  
27 Ireland Galway, Biomedical Science 1<sup>st</sup> Floor South, Upper Newcastle, Galway,  
28 Ireland

29 **Email:** cynthia.coleman@nuigalway.ie

30

31           **5. Name:** Colin Murphy

32           **Affiliation address:** Department of Trauma & Orthopaedics, Galway University  
33           Hospital, Newcastle Road, Galway, Ireland

34           **Email:** cmurphy@rcsi.ie

35

36   **Author Contributions Statement:**

37   Adam Galbraith performed data collection, database organisation, assisted data analysis,  
38   constructed and authored the main manuscript.

39

40   Clara Sanz-Nogués executed data analysis and interpretation, created the resultant tables,  
41   figures and contributed to the associated narrative, reviewed and edited drafts of the  
42   manuscript.

43

44   Sharon Glynn contributed to data analysis and interpretation, performed survival analysis,  
45   reviewed and edited drafts of the manuscript.

46

47   Cynthia Coleman co-conceived the concept of this study, wrote and obtained ethical  
48   permission to conduct the investigation, secured collaborations to execute the investigation,  
49   contributed to database organisation and data interpretation, reviewed and edited drafts of the  
50   manuscript.

51

52   Colin Murphy co-conceived concept of study, secured collaboration to execute research,  
53   contributed to data collection and database organisation, reviewed and edited draft  
54   manuscript.

55

56   All authors have read and approved the final, submitted manuscript.

57

58

59

60

61 **Abstract**

62 Diabetes mellitus (DM) is associated with an elevated risk of post-operative complications.  
63 The impact it has on patients living with DM following hip fracture surgery (HFS) is not  
64 completely understood, and may represent a predictor of increased mortality. This study  
65 investigates the impact of DM, gender, American Society of Anaesthesiologists (ASA) grade  
66 and fracture location, on outcome of HFS in Ireland. The Hospital Inpatient Enquiry (HIPE)  
67 database records all fragility hip fractures within Galway University Hospital. Retrospective  
68 data collection was performed over a three-year period. Data collected included patient age,  
69 gender, date of HFS, anatomical fracture location, type of operation, ASA grade, DM status  
70 and mortality. A database of 650 individuals was created including 461 females and 189  
71 males, with an average group age of  $80.2 \pm 9.3$  years. Results showed a significantly higher  
72 incidence of hip fractures in males with DM (19.57%) than females with DM (12.36%) ( $\chi^2$   
73 test,  $p = 0.020$ ). Cox regression survival analysis indicated that DM status and ASA grade  
74 were the two main independent predictors of patient survival following HFS. Nevertheless,  
75 when examining the combined impact of gender and DM status on survival after HFS, results  
76 showed that survival post HFS differed significantly with gender and presence of DM (log-  
77 rank test,  $p < 0.001$ ), with males with DM performing worse than females with DM  
78 ( $p=0.021$ ) or males without DM ( $p=0.001$ ). This gender and disease-associated outcome  
79 should prompt early multi-disciplinary team approach to the management of hip fractures in  
80 patients with DM.

81 **Keywords:**

82 Hip Fracture Surgery, Diabetes Mellitus, Mortality, ASA grade, Diabetic Osteopathy

83

84

## 85 **Introduction**

86 In line with improvements in global well-being and healthcare delivery, a vast growth in the  
87 proportion of the population aged 65 years and over has occurred and is expected to double  
88 by 2060 <sup>1</sup>. The 2018 European Ageing Report projects that the old-age dependency ratio will  
89 increase significantly from 25% in 2010 to 51.2% in 2070 <sup>2</sup>. In keeping with this growth in  
90 the elderly populations, an equivalent increase in the incidence of hip fractures is expected <sup>2</sup>.  
91 <sup>3</sup>. The absolute number of all fragility fracture admissions increased by 30% between the  
92 years of 2002 to 2014 in Europe <sup>2</sup>. In Ireland, the Health Service Executive (HSE) has  
93 identified hip fractures as “one of the most serious illness pertaining to long-term hospital  
94 admission” <sup>4</sup>. The Irish Hip Fracture Database (IHFD) reported over 3,000 hip fractures  
95 annually in a total population of 4.7 million people since it was established nationally in 2015  
96 <sup>5</sup>.

97 A better understanding of factors impacting upon hip fracture surgery (HFS) outcomes is  
98 becoming apparent, following the introduction of hip fracture care pathways and  
99 collaborative review of national data. The blue book standards outlined by the British  
100 Orthopaedic Association (BOA) have been referenced globally <sup>6</sup>. Their adoption in Ireland, in  
101 the form of the Irish Hip Fracture Standards (IHFS) has led to a transformation in the delivery  
102 of hip fracture care. However, internationally the reported one-year mortality following hip  
103 fractures remains persistently high ranging from 8.4% to 34% <sup>7-9</sup>. Hip fracture patients  
104 represent a high-risk surgical group, yet the individual influence of any single comorbidity  
105 remains unclear <sup>7, 10, 11</sup>.

106 The impact of diabetes mellitus (DM) in HFS is unclear <sup>12-14</sup>. The global prevalence of DM  
107 among adults over 18 years of age has risen from 4.7% in 1980 to 8.5% in 2014 with the  
108 WHO projecting that DM will be the seventh leading cause of death in 2030 <sup>15</sup>. DM has been

109 reported an independent risk factor for fragility fractures, with research reporting an  
110 incidence of hip fracture in DM patients up to 70% higher than patients without DM<sup>16</sup>. DM  
111 is associated with higher level of osteoporosis and osteopenia, increased osteoblast apoptosis  
112 and osteoclast mediated bone resorption resulting in poorer bone healing and regenerative  
113 capacity following injury<sup>17-20</sup>. The impact of DM on HFS rehabilitation and long-term post-  
114 operative outcomes remains unclear, as individuals living with DM are reported to be at an  
115 increased risk of post-operative complications and mortality following HFS<sup>12, 13</sup>, while others  
116 studies have reported no significant difference between patients living with and without DM  
117 regarding HFS<sup>21</sup>.

118 The aim of this study was to evaluate the impact of DM on HFS outcomes, with particular  
119 interest regarding any associations between the presence of DM, gender, anatomical fracture  
120 location, type of fixation, American Society of Anaesthesiologists (ASA) grade and early  
121 post-operative mortality rates following HFS.

## 122 **Methodology**

123 This was a Level 3 retrospective cohort study. All patients admitted to Galway University  
124 Hospital with fragility hip fractures were recorded in the hospital in-patient enquiry (HIPE)  
125 database and included in the study. Data was collected retrospectively from 1<sup>st</sup> January 2014  
126 to 31<sup>st</sup> December 2016 in adherence with the STrengthening the Reporting of OBServational  
127 studies in Epidemiology (STROBE) guidelines<sup>22</sup>. In line with the criteria utilised by the  
128 National Hip Fracture Database and IHFD, all patients over 60 years old with hip fractures  
129 (intracapsular, intertrochanteric and subtrochanteric) other than periprosthetic fractures were  
130 included, regardless of cause<sup>5</sup>. A total of 650 patients were included in the analysis. Data  
131 collected included patient age, gender, DM status, anatomical neck of femur fracture location,  
132 date of primary HFS, type of fixation, ASA grade and patient mortality<sup>23</sup>. ASA grade is

133 recorded by the IHFD as a surrogate marker for co-morbidities. Registry measured endpoints  
134 were followed and therefore additional specific patient co-morbidities have not been recorded  
135 <sup>5</sup>. Time to surgery was under 48 hours for 75% of the patients, as per the IHFD annual reports  
136 <sup>5</sup>. Diabetic status was confirmed by a consultant endocrinologist and cross-referencing with  
137 the hospital laboratory system provided identification of patients' HbA1c level.

138 Participants' mortality was checked up to the 1<sup>st</sup> November 2017. The electronic patient  
139 demographic system (PAS system) was used to identify deceased patients. A patient database  
140 linked the computerised PAS system to the HIPE database. HIPE is an Irish national database  
141 of coded discharge summaries from acute public hospitals. Ireland has used the International  
142 Classification of Diseases, Ninth Revision, Clinical Modification (ICD-9-CM) for both  
143 diagnosis and procedure coding from 1990 to 2005 and ICD-10-CM thereafter <sup>24</sup>. Following  
144 discharge from hospital, the hospital administrators update the PAS data when they are made  
145 aware of a patient's death. In addition, an online public database of death notifications  
146 (RIP.ie) was utilised to cross-check all notifications. Other countries link inpatient  
147 admissions with national births/deaths registries via a unique identifier, allowing for real-time  
148 accuracy. Lacking such a system in Ireland, mortality rates are likely underestimated.

149 Cross-referencing with the hospital radiology system was also performed to confirm the  
150 anatomical fracture location and definitive surgical intervention performed. Ethical approval  
151 for this study was granted by the Galway University Hospitals Research Ethics Committee  
152 (CA1783).

153

154

155

## 156 **Statistical Analysis**

157 Statistical analyses were performed using the Minitab17 software package for Windows.  
158 Survival curves and cox regression analysis was performed using Stata/SE 14 statistical  
159 software (Stata Corp) as previously described <sup>25</sup>. All statistical tests were 2-sided, and an  
160 association was considered statistically significant with *p*-values less than 0.05. A Student's *t*-  
161 test was used to analyse differences in mean age and follow-up time between the DM and  
162 non-DM groups. Pearson's Chi Squared ( $\chi^2$ ) analysis was used to determine the association  
163 between explanatory variables such as DM status, gender, fracture location, type of operation  
164 and ASA grade. Survival was calculated for the period from the date of primary HFS to the  
165 date of last completed search for death entries (1<sup>st</sup> November 2017) for the 650 case patients  
166 with hip fracture. The Kaplan-Meier method and the log-rank test were used for univariable  
167 survival analysis. Multivariable cox regression survival analysis was used to calculate an  
168 adjusted hazard ratio (HR). The following covariates were included in the analysis: age at  
169 diagnosis (as a continuous variable), gender (male *vs.* female), DM (yes *vs.* no), fracture  
170 location (subtrochanteric *vs.* intracapsular *vs.* intertrochanteric), and ASA grade (2 *vs.* 3 *vs.*  
171 4). A statistical test for interaction was performed in Stata to determine the association  
172 between gender and DM with survival after primary HFS.

## 173 **Results**

### 174 **Demographics:**

175 A database of 650 individuals was collated including 461 (70.9%) females and 189 (29.1%)  
176 males, with an average patient age of 80.2±9.3 years ranging between 60 and 101 years.  
177 Those categorized as living with DM (n=79, n=15 missing data) had an average HbA1c of  
178 52.76±18.97 mmol/mol ranging from 26 to 128 mmol/mol. There were no statistically  
179 significant differences among the age of participants living with DM (79.9±8.4 years) and



180 those living without DM ( $80.3 \pm 9.5$  years) (2-sided  $t$ -test,  $p = 0.739$ ) or male ( $79.4 \pm 9.6$  years)  
181 vs female ( $80.6 \pm 9.2$  years) (2-sided  $t$ -test,  $p=0.137$ ). When looking at the percentage of  
182 females and males with hip fractures that had DM, there were significantly more males with  
183 DM (19.57%) than females with DM (12.36%) ( $\chi^2$  test;  $p=0.020$ ) (Table 1). DM was not  
184 found to affect the anatomical location of neck of femur fractures, with a similar distribution  
185 of fracture location among individuals with DM and without DM ( $\chi^2$  test;  $p=0.864$ ).  
186 Furthermore, the type of fixation did not significantly differ among individuals living with  
187 DM and those without DM ( $\chi^2$  test;  $p=0.434$ ) (Table 1).

188 Pearson's  $\chi^2$  analysis revealed no association between fracture location and gender: male  
189 intracapsular (59.8%), intertrochanteric (31.2%), subtrochanteric (9.0%) compared with  
190 female intracapsular (56.6%), intertrochanteric (34.1%), subtrochanteric (9.33%) ( $\chi^2$  test;  $p=$   
191 0.750) (Table 2). We identified an association with increasing ASA grade in patients living  
192 with DM (ASA  $\geq 3$ , 72.63%,  $n = 61$ ) compared with non-DM patients (ASA  $\geq 3$ ,  $n=247$ ) ( $p=$   
193 0.001) (Table 1). Pearson's  $\chi^2$  analysis identified a significant association between ASA  
194 grade and gender: male ASA  $\geq 3$ , 65.1% ( $n=110$ ) compared with females 48.41% ( $n=198$ )  
195 (Table 2).

### 196 **Survival Analysis:**

197 The effect of age, gender, DM status, anatomical fracture location and ASA grade on  
198 predicting patient survival was examined using the Kaplan-Meier method (Figure S-1) and  
199 univariate and multivariable cox regression survival analysis (Table 3). The Kaplan-Meier  
200 analysis and univariate cox regression analysis showed that only the presence of DM (log-  
201 rank test,  $p=0.002$ ,) and ASA grade (log-rank test,  $p=<0.01$ ) were statistically significant  
202 predictors of patient survival, which remained significant in the multivariable cox regression  
203 analysis after adjusting for age at diagnosis. Other factors such as gender and fracture

204 location were not found to be statistically significant. In the multivariable cox regression  
205 analysis adjusting for age at diagnosis, gender became significantly associated with patient  
206 outcome ( $p=0.035$ ). Nevertheless, in a combined multivariable cox regression survival  
207 analysis only DM status and ASA grade were shown to be good independent predictors of  
208 patient outcome.

209 We then examined the combined impact of gender and DM status on patient outcome after  
210 HFS as shown in Figure S-2. These results demonstrate that patient survival differed  
211 significantly depending on gender and DM status (log-rank test,  $p<0.001$ ). Univariate cox  
212 regression survival analysis showed no significant differences between males and females  
213 without DM (HR 1.09, 95% CI 0.70-1.70,  $p=0.706$ ) or females without DM versus females  
214 with DM (HR 1.36, 95% CI 0.75-2.45,  $p=0.314$ ). Interestingly, there was a significant  
215 difference between males without DM versus males with DM (HR 2.72, 95% CI 1.46-5.06,  
216  $p=0.002$ ), and females with DM versus males with DM (HR 2.32, 95% CI 1.11-4.85,  
217  $p=0.025$ ), which remained significant in the multivariable cox regression analysis after  
218 adjusting for age at diagnosis, and in the combined multivariable cox regression analysis  
219 (Table 4). A test for statistical interaction between gender and DM status on patients' survival  
220 showed a near significant interaction ( $p=0.092$ ), corroborating these findings.

## 221 **Discussion:**

222 Patients with hip fractures represent a high-risk surgical group. However, the individual  
223 influence of any single comorbidity remains unclear<sup>7, 10, 11</sup>. Furthermore, Franklin et al, when  
224 investigating patient characteristics and pre-operative co-morbidities between European and  
225 American patients undergoing elective surgery found significant differences between the two  
226 groups with respect to pre-operative characteristics and co-morbidities<sup>26</sup>. Evidence has  
227 shown that patients living with DM are at increased risk of sustaining fragility fractures,  
228 however the overall impact of DM patients undergoing HFS is unknown<sup>12, 13, 27</sup>. Gulcelik et

229 al, reported a significant increase in the probability of one year survival following HFS in  
230 patients living without DM (87.3%) compared to patients living with DM (68.0%)<sup>14</sup>. Our  
231 research is the first to describe the association of DM and mortality following HFS in an Irish  
232 population. The demographic represented in this study is similar in characteristics to data  
233 published by hip fracture databases nationally and internationally with respect to gender,  
234 ASA grade and hip fracture location<sup>5, 28</sup>. We have identified a significant increase in  
235 mortality of individuals living with DM undergoing HFS with a particular gender effect on  
236 patient's outcome. Furthermore, we have identified an association between hip fracture  
237 patients living with DM and higher ASA grade which may be a contributing factor to the  
238 increased post-operative mortality observed. This research brings into focus the need for  
239 early multi-disciplinary team management in patients requiring HFS and those who are living  
240 with co-morbid DM.

#### 241 **DM and overall mortality of patients living with DM following HFS**

242 This study has demonstrated that patients with DM had a significantly greater post-operative  
243 mortality following HFS when compared to patients without DM. Two previously published  
244 studies have reported that the presence of DM does not negatively impact survival following  
245 HFS<sup>12, 13</sup>. Norris et al, reported that patients living with DM had an increased inpatient length  
246 of stay and were more likely to develop post-operative complications including pressure sores  
247 and cardiovascular issues, however the one-year post-operative mortality between patients  
248 living with DM and non-DM patients was not affected<sup>12</sup>. Ekstrom et al have similarly  
249 reported that although individuals living with DM may have increased post-operative pain  
250 and risk of overall post-operative complications, the presence of DM did not impact upon the  
251 long-term rehabilitative capacity of this patient cohort<sup>12, 13, 21</sup>. Although the research  
252 presented here has not examined the impact of DM specifically on post-operative

253 rehabilitation, it does indicate a negative impact of patients living with DM on long-term  
254 survival and overall mortality. The increase in overall mortality identified here is likely  
255 multifactorial, and may be explicable due to differing ASA grade and overall incidence of  
256 DM in our cohort when compared to Norris and Ekstrom<sup>12, 13</sup>.

### 257 **ASA grade and overall mortality of patients living with DM following HFS**

258 The ASA classification first described in 1941 is a highly effective grading system which  
259 identifies patient risk of post-operative morbidity and mortality. It forms an integral  
260 component of the WHO pre-operative checklist (Haynes, 2009), a check-list that following  
261 implementation has shown an overall reduction in post-operative mortality at one year by up  
262 to 50%<sup>29</sup>. The ASA classification continues as a widely utilised and effective assessment  
263 tool, recently updated by the American Society of Anaesthesiologists in 2014<sup>30-32</sup>. The ASA  
264 classification is a subjective assessment performed by anaesthesiologists in which patients are  
265 assigned as grade one to five based upon increasing risk of post-operative morbidity<sup>32</sup>. The  
266 system also carries a subclassification “E” which is added to a patients’ baseline grade in the  
267 event that their surgery is emergency, in the setting of HFS by virtue of their pathology hip  
268 fracture patients are immediately assigned to this subclassification. ASA grade is recorded by  
269 the IFHD as a surrogate marker for co-morbidities and previous research has indicated its  
270 importance in the setting of HFS<sup>5, 33, 34</sup>.

271 As previously discussed, all hip fracture patients represent a high-risk surgical patient cohort  
272 in which, up to 70% of patients are classified as ASA grade  $\geq 3$ <sup>35</sup>. Overall in, this study  
273 (n=650), almost 53% of hip fracture patients had ASA grade  $\geq 3$ . Nevertheless, we found that  
274 a higher percentage of patients with DM had ASA grade  $\geq 3$  (72.62%) compared to those  
275 without DM (49.79%), which could be explained by the higher presence of co-morbidities in  
276 DM patients.

277 In this investigation, we identified that presence of DM and higher ASA grade were both  
278 independent predictors of patient mortality after primary HFS ( $p=0.025$  and  $p<0.001$   
279 respectively). Similarly, Hu et al, conducted a systematic review and meta-analysis assessing  
280 the pre-operative predictors of mortality following HFS in which they also identified higher  
281 ASA grade and the presence of DM as strong predictors of overall mortality<sup>10</sup>.

282 The use of the validated assessments of patient frailty, such as the Frailty Index (FI), may  
283 represent a more encompassing assessment tool for predicting adverse outcomes and  
284 mortality following HFS<sup>27, 36</sup>. The FI assesses patient health based on accumulation of  
285 disease, including the presence or absence of DM. The incidence of a greater number of  
286 patient co-morbidities indicates increased frailty<sup>36</sup>. When assessing patient frailty with  
287 respect to fracture risk, Li et al, identified that patients living with DM were significantly  
288 frailer than those living without DM, with a significant relationship between the FI and the  
289 risk of incident fragility fracture, “a hazard ratio (HR) of 1.02 (95% CI 1.01-1.03) and 1.19  
290 (95% CI 1.10-1.33) for per-0.01 and per-0.10 FI increase, respectively ( $p=0.018$ )”<sup>37</sup>.

291 Further research is required to understand this association. In addition, research has shown  
292 that increased FI is associated with increased overall mortality<sup>37, 38</sup>. Improved management of  
293 DM as a disease entity has the potential to improve both FI and post-operative outcomes for  
294 this high-risk group<sup>38</sup>.

### 295 **Gender and overall mortality of patients living with DM following HFS**

296 Several studies have identified an association between male gender and increased mortality  
297 following HFS, the aetiology of which is not fully understood<sup>39</sup>. Endo et al, established that  
298 overall ASA grade was higher in males and the incidence of post-operatively complications  
299 were more common in men<sup>40</sup>. In this study we also found that ASA grade was higher in  
300 males (57.4% ASA grade 3 and 7.69% ASA grade 4) than in females (44.99% ASA grade 3

301 and 2.93% ASA grade 4) ( $\chi^2$  test;  $p=0.001$ ). When we look at the combined impact of gender  
302 and DM status on patient outcome after HFS, our results showed that males with DM  
303 performed poorer than males without DM ( $p=0.001$ ) and females with DM ( $p=0.025$ ).  
304 Multiple studies have shown that overall, irrespective of DM status that male patients may  
305 have up to a 10% overall increase in one-year mortality following HFS than their female  
306 counterparts<sup>41-43</sup>. Multiple factors may be responsible for this difference. Diagnosis of  
307 osteoporosis is an important preventative measure which has been shown to reduce the  
308 incidence of hip fractures when treated appropriately. It has been suggested that osteoporosis  
309 may be underdiagnosed in the male population. Diagnosis of osteoporosis currently relies  
310 upon bone mineral density (BMD) based on a non-gender specific reference value. Cawthon  
311 et al, argue that specific gender reference values of BMD should be created. As a result, more  
312 men would be diagnosed with osteoporosis, despite having an overall higher BMD reference  
313 value of their counterpart females<sup>44</sup>. Gregg et al suggest that the presence of DM contributes  
314 to a substantial reduction in overall life expectancy of both sexes, but the impact is greater in  
315 females<sup>45</sup>. The development of DM-related cardiovascular disease may represent a  
316 significant aetiological factor, where the risk is as high as a six-fold increase in females  
317 compared with a two to three-fold increase in males<sup>46</sup>. It is postulated that these gender  
318 difference may in part be due to physiological differences between males and females and the  
319 impact of diabetic nephropathy upon oestrogen regulation<sup>47</sup>.

320 Identifying the aetiological explanation regarding our finding of reduced long-term survival  
321 of men living with DM versus their female counterparts following HFS goes beyond the  
322 scope of this research and highlights another area in which national databases could extend  
323 their data collection profile to include co-morbidities and other previously identified  
324 predictors of increased mortality.

325

326 **Strengths and Limitations**

327 Galway University Hospital has a population census of greater than 300,000 individuals, of  
328 which this study cohort included a total of 650 patients, in which there was a higher presence  
329 of females with hip fractures (n=461) compared to males (n=189). Nevertheless, this study is  
330 representative of a significant percentage of the Irish population improving the application of  
331 our findings to the national population.

332 The principle limitation of this study is its retrospective nature. The study was performed in a  
333 single tertiary orthopaedic facility, which may reduce generalisability of the study. However,  
334 the clinical protocols followed in this facility are homogenous both nationally and  
335 internationally and this dataset is representative of the rest of the nation. In addition, no data  
336 was obtained regarding additional comorbidities, modes of treatment or duration of control  
337 due to ethical restrictions. It was therefore not possible to delineate whether patients were  
338 T1DM or T2DM, but paucity of T1DM cases in this cohort of patients has been reported at  
339 0.12 – 0.18% <sup>48</sup>. In addition, in this study, ASA grade was used as a surrogate marker for co-  
340 morbidities, as recorded by the IFHD. There is scope for a prospective study or national hip  
341 fracture registry data which should aim to capture all potential confounding factors  
342 influencing long-term mortality following HFS such as BMI, cardiovascular disease and  
343 lifestyle factors.

344 The absolute mortality rate is likely to be underestimated. HIPE data has the limitation of  
345 recording inpatient mortality only and is not linked with national Central Statistics Office  
346 (CSO) data. Our research utilised publicly available databases to crosscheck and minimise  
347 this potential deficiency error, and our mortality trend is consistent with that of a previous  
348 research that based upon CSO data <sup>3, 49</sup>.

349

350 **Conclusion**

351 Patients living with DM are at an increased risk of hip fracture <sup>16, 50</sup>. Internationally,  
352 improvement measures of hip fracture patients are focused on a myriad of factors including  
353 admission time to appropriate orthopaedic units, timely access to surgery and integrated peri-  
354 and post-operative multi-disciplinary team input. This study is the first in Ireland to note a  
355 deleterious effect of DM and gender on post-operative mortality following HFS and indicates  
356 the necessity of an early multi-disciplinary approach for the management of hip fracture  
357 patients living with DM. Increased cross specialty awareness is required to appreciate the  
358 increased rate of fragility fractures in patients living with DM and the associated increase in  
359 post-operative mortality as outlined in this research. Further studies are recommended to  
360 consider the systemic physiological impact of DM and gender and its role in the context of  
361 the biochemical and biomechanical impact of DM on bone morphology.

362 **Acknowledgments:**

363 Cynthia Coleman acknowledges granted funding support from the Diabetes Ireland Research  
364 Alliance/Medical Research Charity Group/Health Research Board Joint Funding Scheme  
365 (HRB-MRCG-2016-2). The authors have no professional or financial affiliations that relate to  
366 or influence the contents of this publication.

367

368

369

370

371



372 **Bibliography**

- 373 1. Rechel B, Grundy E, Robine JM, et al. Ageing in the European Union. *Lancet (London,*  
374 *England)*. 2013; 381: 1312-22.
- 375 2. The 2018 Ageing Report Underlying Assumptions and Projection Methodologies European  
376 Economy2017.
- 377 3. Elbattah M and Molloy O. *The Economic Burden of Hip Fractures among Elderly Patients in*  
378 *Ireland: A Combined Perspective of System Dynamics and Machine Learning*. 2016.
- 379 4. Laffoy M. Strategy to prevent falls and fractures in Ireland's ageing population summary,  
380 conclusions and recommendations. Health Service Executive (HSE) Strategic Health Planning (Dublin  
381 8.);National Council on Ageing and Older People (NCAOP);Department of Health and Children  
382 (DOHC)
- 383 2008.
- 384 5. Audit NOoC. Irish Hip Fracture Database National Report 2017.
- 385 6. *The Care of Patients with Fragility Fracture*. British Orthopaedic Association, 2007.
- 386 7. Smith T, Pelpola K, Ball M, Ong A and Myint PK. Pre-operative indicators for mortality  
387 following hip fracture surgery: a systematic review and meta-analysis. *Age and ageing*. 2014; 43:  
388 464-71.
- 389 8. Lee YK, Lee YJ, Ha YC and Koo KH. Five-year relative survival of patients with osteoporotic hip  
390 fracture. *The Journal of clinical endocrinology and metabolism*. 2014; 99: 97-100.
- 391 9. Tran T, Bliuc D, Hansen L, et al. Persistence of Excess Mortality Following Individual Nonhip  
392 Fractures: A Relative Survival Analysis. *The Journal of clinical endocrinology and metabolism*. 2018;  
393 103: 3205-14.
- 394 10. Hu F, Jiang C, Shen J, Tang P and Wang Y. Preoperative predictors for mortality following hip  
395 fracture surgery: a systematic review and meta-analysis. *Injury*. 2012; 43: 676-85.
- 396 11. Le Manach Y, Collins G, Bhandari M, et al. Outcomes After Hip Fracture Surgery Compared  
397 With Elective Total Hip Replacement. *Jama*. 2015; 314: 1159-66.
- 398 12. Norris R and Parker M. Diabetes mellitus and hip fracture: a study of 5966 cases. *Injury*.  
399 2011; 42: 1313-6.
- 400 13. Ekstrom W, Al-Ani AN, Saaf M, Cederholm T, Ponzer S and Hedstrom M. Health related  
401 quality of life, reoperation rate and function in patients with diabetes mellitus and hip fracture--a 2  
402 year follow-up study. *Injury*. 2013; 44: 769-75.
- 403 14. Gulcelik NE, Bayraktar M, Caglar O, Alpaslan M and Karakaya J. Mortality after hip fracture in  
404 diabetic patients. *Experimental and clinical endocrinology & diabetes : official journal, German*  
405 *Society of Endocrinology [and] German Diabetes Association*. 2011; 119: 414-8.
- 406 15. Mathers CD and Loncar D. Projections of global mortality and burden of disease from 2002  
407 to 2030. *PLoS medicine*. 2006; 3: e442.
- 408 16. Janghorbani M, Van Dam RM, Willett WC and Hu FB. Systematic review of type 1 and type 2  
409 diabetes mellitus and risk of fracture. *American journal of epidemiology*. 2007; 166: 495-505.
- 410 17. Gandhi A, Beam HA, O'Connor JP, Parsons JR and Lin SS. The effects of local insulin delivery  
411 on diabetic fracture healing. *Bone*. 2005; 37: 482-90.
- 412 18. Kagel EM, Majeska RJ and Einhorn TA. Effects of diabetes and steroids on fracture healing.  
413 *Current opinion in orthopaedics*. 1995; 6: 7-13.
- 414 19. Wongdee K and Charoenphandhu N. Update on type 2 diabetes-related osteoporosis. *World*  
415 *journal of diabetes*. 2015; 6: 673-8.
- 416 20. Hamann C, Goettsch C, Mettelsiefen J, et al. Delayed bone regeneration and low bone mass  
417 in a rat model of insulin-resistant type 2 diabetes mellitus is due to impaired osteoblast function.  
418 *American journal of physiology Endocrinology and metabolism*. 2011; 301: E1220-8.
- 419 21. Mizrahi EH, Fleissig Y, Arad M and Adunsky A. Functional outcome of elderly hip fracture  
420 patients: does diabetes matter? *Arch Gerontol Geriatr*. 2006; 43: 165-73.

- 421 22. von Elm E, Altman DG, Egger M, Pocock SJ, Gotsche PC and Vandembroucke JP.  
422 Strengthening the Reporting of Observational Studies in Epidemiology (STROBE) statement:  
423 guidelines for reporting observational studies. *BMJ (Clinical research ed)*. 2007; 335: 806-8.
- 424 23. Daabiss M. American Society of Anaesthesiologists physical status classification. *Indian*  
425 *journal of anaesthesia*. 2011; 55: 111-5.
- 426 24. O'Neill B. The hospital in-patient enquiry scheme-a study of data accuracy. *Irish medical*  
427 *journal*. 1982; 75: 238-9.
- 428 25. Glynn SA, Boersma BJ, Dorsey TH, et al. Increased NOS2 predicts poor survival in estrogen  
429 receptor-negative breast cancer patients. *The Journal of clinical investigation*. 2010; 120: 3843-54.
- 430 26. Franklin PD, Miozzari H, Christofilopoulos P, Hoffmeyer P, Ayers DC and Lubbeke A.  
431 Important patient characteristics differ prior to total knee arthroplasty and total hip arthroplasty  
432 between Switzerland and the United States. *BMC musculoskeletal disorders*. 2017; 18: 14.
- 433 27. Sternberg SA, Wershof Schwartz A, Karunanathan S, Bergman H and Mark Clarfield A. The  
434 identification of frailty: a systematic literature review. *Journal of the American Geriatrics Society*.  
435 2011; 59: 2129-38.
- 436 28. Registry ANZHF. Bi-national Annual Report of Hip Fracture Care. 2018.
- 437 29. Haynes AB, Weiser TG, Berry WR, et al. A Surgical Safety Checklist to Reduce Morbidity and  
438 Mortality in a Global Population. *New England Journal of Medicine*. 2009; 360: 491-9.
- 439 30. Hopkins TJ, Raghunathan K, Barbeito A, et al. Associations between ASA Physical Status and  
440 postoperative mortality at 48 h: a contemporary dataset analysis compared to a historical cohort.  
441 *Perioperative medicine (London, England)*. 2016; 5: 29-.
- 442 31. Saklad M, M.D. GRADING OF PATIENTS FOR SURGICAL PROCEDURES. *Anesthesiology: The*  
443 *Journal of the American Society of Anesthesiologists*. 1941; 2: 281-4.
- 444 32. Anaesthesiologists ASo. ASA Physical Status Classification System. 2014.
- 445 33. Hamlet WP, Lieberman JR, Freedman EL, Dorey FJ, Fletcher A and Johnson EE. Influence of  
446 health status and the timing of surgery on mortality in hip fracture patients. *American journal of*  
447 *orthopedics (Belle Mead, NJ)*. 1997; 26: 621-7.
- 448 34. Richmond J, Aharonoff GB, Zuckerman JD and Koval KJ. Mortality Risk After Hip Fracture.  
449 *Journal of orthopaedic trauma*. 2003; 17: S2-S5.
- 450 35. Griffiths R, Alper J, Beckingsale A, et al. Management of proximal femoral fractures 2011:  
451 Association of Anaesthetists of Great Britain and Ireland. *Anaesthesia*. 2012; 67: 85-98.
- 452 36. Mitnitski AB, Mogilner AJ and Rockwood K. Accumulation of deficits as a proxy measure of  
453 aging. *TheScientificWorldJournal*. 2001; 1: 323-36.
- 454 37. Li G, Prior JC, Leslie WD, et al. Frailty and Risk of Fractures in Patients With Type 2 Diabetes.  
455 *Diabetes care*. 2019.
- 456 38. Hoogendijk EO, Theou O, Rockwood K, Onwuteaka-Philipsen BD, Deeg DJH and Huisman M.  
457 Development and validation of a frailty index in the Longitudinal Aging Study Amsterdam. *Aging*  
458 *clinical and experimental research*. 2017; 29: 927-33.
- 459 39. Penrod JD, Litke A, Hawkes WG, et al. The association of race, gender, and comorbidity with  
460 mortality and function after hip fracture. *The journals of gerontology Series A, Biological sciences*  
461 *and medical sciences*. 2008; 63: 867-72.
- 462 40. Endo Y, Aharonoff GB, Zuckerman JD, Egol KA and Koval KJ. Gender differences in patients  
463 with hip fracture: a greater risk of morbidity and mortality in men. *Journal of orthopaedic trauma*.  
464 2005; 19: 29-35.
- 465 41. Elliott J, Beringer T, Kee F, Marsh D, Willis C and Stevenson M. Predicting survival after  
466 treatment for fracture of the proximal femur and the effect of delays to surgery. *Journal of clinical*  
467 *epidemiology*. 2003; 56: 788-95.
- 468 42. Jamal Sepah Y, Umer M, Khan A and Ullah Khan Niazi A. Functional outcome, mortality and  
469 in-hospital complications of operative treatment in elderly patients with hip fractures in the  
470 developing world. *International orthopaedics*. 2010; 34: 431-5.

471 43. Roche JJ, Wenn RT, Sahota O and Moran CG. Effect of comorbidities and postoperative  
472 complications on mortality after hip fracture in elderly people: prospective observational cohort  
473 study. *BMJ (Clinical research ed)*. 2005; 331: 1374.

474 44. Cawthon PM. Gender differences in osteoporosis and fractures. *Clinical orthopaedics and  
475 related research*. 2011; 469: 1900-5.

476 45. Gregg EW, Gu Q, Cheng YJ, Narayan KM and Cowie CC. Mortality trends in men and women  
477 with diabetes, 1971 to 2000. *Annals of internal medicine*. 2007; 147: 149-55.

478 46. Legato MJ, Gelzer A, Golland R, et al. Gender-specific care of the patient with diabetes:  
479 review and recommendations. *Gender medicine*. 2006; 3: 131-58.

480 47. Maric C and Sullivan S. Estrogens and the diabetic kidney. *Gender medicine*. 2008; 5 Suppl A:  
481 S103-13.

482 48. Forsen L, Meyer HE, Midthjell K and Edna TH. Diabetes mellitus and the incidence of hip  
483 fracture: results from the Nord-Trondelag Health Survey. *Diabetologia*. 1999; 42: 920-5.

484 49. Elbattah M and Molloy O. The Economic Burden of Hip Fractures among Elderly Patients in  
485 Ireland: A Combined Perspective of System Dynamics and Machine Learning. 2016.

486 50. Tebe C, Martinez-Laguna D, Moreno V, et al. Differential Mortality and the Excess Rates of  
487 Hip Fracture Associated With Type 2 Diabetes: Accounting for Competing Risks in Fracture Prediction  
488 Matters. *Journal of bone and mineral research : the official journal of the American Society for Bone  
489 and Mineral Research*. 2018.

490

491

492

493

494

495

496

497

498

499

500

501

502

503

504 **Figure Legends**

505 **Figure S-1. Kaplan-Meier survival estimates following primary HFS.** A) Kaplan-Meier  
506 cumulative survival curves of hip fracture patients by DM status ( $n = 644$ ). The survival of  
507 patients with DM ( $n = 93$ ) was significantly poorer than the survival of patients without DM  
508 ( $n=551$ , log-rank test,  $p=0.002$ ). B) Kaplan-Meier cumulative survival curves of hip fracture  
509 patients by gender ( $n = 644$ ). No significant difference in survival was observed between male  
510 ( $n=186$ ) versus female ( $n = 458$ ) hip fracture patients (log-rank test,  $p=0.089$ ). C) Kaplan-  
511 Meier cumulative survival curves of hip fracture patients by fracture location ( $n=644$ ). No  
512 significant difference in survival was observed depending on fracture location (log-rank  
513 test,  $p=0.620$ ). D) Kaplan-Meier cumulative survival curves of hip fracture patients by ASA  
514 grade 2, 3 and 4 ( $n=588$ ). An increase in poor outcome was seen in patients with ASA 2  
515 versus ASA 3 (log-rank test,  $p<0.001$ ), ASA 3 versus ASA 4 (log-rank test,  $p=0.001$ ), and  
516 ASA 2 versus ASA 4 (log-rank test,  $p<0.001$ ).

517

518 **Figure S-2. Impact of gender and diabetes status on patient survival after HFS.** Kaplan-  
519 Meier cumulative survival curves of hip fracture patients depending on gender and presence  
520 of DM ( $n=644$ ). The survival of patients differed significantly depending on gender and DM  
521 status (log-rank test,  $p<0.001$ ). While there was no significant difference between males and  
522 female without DM ( $p=0.705$ ) or females without DM versus DM ( $p=0.311$ ), there was a  
523 significant difference between males without DM versus DM ( $p=0.001$ ), and females with  
524 DM versus males with DM patients ( $p=0.021$ ). (Note – within figure M=male; F=female;  
525 D=diabetes; ND=no diabetes.)

526