

Mortality After Hip Fracture in Austria 2008–2011

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Abstract Osteoporosis-related hip fractures represent a substantial cause of mortality and morbidity in industrialized countries like Austria. Identification of groups at high risk for mortality after hip fracture is crucial for health policy decisions. To determine in-hospital, long-term, and excess mortality after osteoporosis-related hip fracture in Austrian patients, we conducted a retrospective cohort analysis of pseudonymized invoice data from Austrian social insurance authorities covering roughly 98 % of the entire population. The data set included 31,668 subjects aged 50 years and above sustaining a hip fracture between July 2008 and December 2010 with follow-up until June

2011, and an age-, gender-, and regionally matched control population without hip fractures (56,320 subjects). Kaplan–Meier and Cox hazard regression analyses served to determine unadjusted and adjusted mortality rates: Unadjusted all-cause 1-year mortality amounted to 20.2 % (95 % CI: 19.7–20.7 %). Males had significantly higher long-term, in-hospital, and excess mortality rates than females, but younger males exhibited lower excess mortality than their female counterparts. Advanced age correlated with increased long-term and in-hospital mortality, but lower excess mortality. Excess mortality, particularly in males, was highest in the first 6 months after hip fracture, but remained statistically significantly elevated throughout the observation period of 3 years. Longer hospital stay per fracture was correlated with mortality reduction in older patients and in patients with more subsequent fractures. In conclusion, more efforts are needed to identify causes and effectively prevent excess mortality especially in male osteoporosis patients.

The authors declare that they have no conflict of interest.

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Introduction

Being a source of substantial morbidity and mortality, in particular among the elderly, hip fractures represent a significant public health concern [1, 2]. Osteoporosis-related bone loss and deterioration of bone quality with advancing age are the main cause of non-traumatic hip fractures [3, 4]. Whereas a decline in hip fracture incidence rates has been reported for a number of countries during the past years including Austria [5–8], mortality after fracture has remained consistently high [2, 9]. Both incidence and

Table 1 Baseline characteristics of the hip fracture study population

	Women	Men	<i>P</i>	All
Median age ^a (IQR)	83.56 (10.44)	78.32 (16.37)	<0.001	82.45 (12.41)
Age group (years) ^a				
All	23192 (73.2 %)	8476 (26.8 %)	<0.001	31668
50–54	288 (41.3 %)	409 (58.7 %)	<0.001	697
55–59	492 (49.1 %)	511 (50.9 %)	0.40	1003
60–64	752 (53.8 %)	646 (46.2 %)	<0.001	1398
65–69	1341 (59.0 %)	932 (41.0 %)	<0.001	2273
70–74	1706 (64.8 %)	927 (35.2 %)	<0.001	2633
75–79	3227 (70.9 %)	1327 (29.1 %)	<0.001	4554
80–84	5693 (77.6 %)	1645 (22.4 %)	<0.001	7338
85–89	6552 (82.2 %)	1422 (17.8 %)	<0.001	7974
90–94	2240 (82.5 %)	476 (17.5 %)	<0.001	2716
95+	901 (83.3 %)	181 (16.7 %)	<0.001	1082
Number of fractures				
1	18056 (72.5 %)	6846 (27.5 %)	<0.001	24902
2	4304 (76.3 %)	1336 (23.7 %)	<0.001	5640
3	596 (73.2 %)	218 (26.8 %)	<0.001	814
4	168 (76.0 %)	53 (24.0 %)	<0.001	221
≥5	68 (74.7 %)	23 (25.3 %)	<0.001	91

^a At date of discharge from hospital or time of death in hospital following index fracture

mortality rates follow a geographical pattern suggesting a north-south gradient with the highest risk reported from Northern Europe and North America [2, 10]. Thus, 1-year mortality rates have recently been described to range from 14 to 36 % [11]. Excess mortality rates, i.e., mortality among hip fracture patients as compared to that among the matched general population, were demonstrated to decrease dramatically during the first half year following fracture and remain elevated thereafter even after 10 years in some studies [12, 13]. Other investigations focusing on institutionalized patients, however, found no significant excess mortality already after 6 months [14, 15]. In addition, excess mortality following hip fracture was found to be inversely correlated with age [16].

Apart from advanced age and male gender, several comorbidities such as end-stage kidney disease, shock due to the fracture event, cardiac disease, congestive heart failure, and chronic liver disease have recently been determined as risk factors [16–18]. Consistently, a recent meta-analysis found a Charlson comorbidity index score of at least one, an American Society of Anesthesiologists score of at least three, and dementia predicted elevated mortality risk following hip fracture [19]. On the other hand, hip fracture itself was described to predict excess mortality independently from comorbidities and known hip fracture risk factors [20].

Previous epidemiologic studies in Austria on hip fracture reported short-term in-hospital mortality only [21, 22]. Therefore, we performed a comprehensive, retrospective

nation-wide study, based on personalized data retrieved from social insurance registries from more than 31,000 elderly hip fracture patients and 56,000 controls to determine in-hospital mortality as well as long-term absolute and excess mortality.

Methods

Study Design and Patients

In this retrospective nation-wide study, pseudonymized patient data were retrieved from thirteen Austrian social insurance authorities covering roughly 98 % of the entire population, using as database program SAS, version 9.3 (SAS Institute Inc., Cary, North Carolina, USA). We identified 31,668 inpatients aged 50 years and above having sustained a hip fracture between July 2008 and December 2010. Follow-up data on survival were available until June 2011. Baseline characteristics of the study population are shown in Table 1. Mean follow-up time after release from hospital upon first fracture recorded in the study period (the index fracture) was 528 days (95 % CI: 524–531 days). Diagnosis of hip fractures was in agreement with the ICD-10 definition S72 [23]. Information on sex, age at hospital discharge from index fracture, residence, subsequent fractures, date of death, and hip fracture-related length of hospital stay was available. Furthermore, 56,320 individuals without hip fractures matched

for age, sex, and residency served as control population, whose mean follow-up time was 964 days (95 % CI: 962–966 days). Recruitment of control subjects was performed using the “ranuni” function which is a tool implemented in the SAS statistical software, version 9.3 (SAS Institute Inc., Cary, North Carolina, USA). This function generated a uniform distribution on the interval (0, 1) covering all persons in our database who were registered by Austrian social insurance authorities, with the exclusion of individuals without a history of hip fracture.

Data Analysis

Kaplan–Meier survival analysis identified crude all-cause mortality rates, and Cox hazard regression modeling yielded mortality rates adjusted for age, gender, number of fractures, and hip fracture-related hospital days per fracture. Validity of the proportional hazard assumption was verified in all instances by log-minus-log representations of hazards over time. The χ^2 , Fisher’s exact as well as Mann–Whitney U tests, where appropriate, informed on differences that, at a confidence level of 95 %, were considered statistically significant. All analyses were conducted in SPSS, version 19 (SPSS Inc., Chicago, Illinois, USA).

Crude and excess all-cause mortality rates were calculated for two time points of reference: Embarking on directly obtainable dates of hospital discharge after first fracture in our study disregarded those patients who died in hospital upon first fracture. On the other side, hospital admission dates, equaling dates of first fracture, were ascertainable from hip fracture-related hospital days only for those patients who sustained no more than one fracture within the study interval (24,902 of 31,668, corresponding to 78.6 %; Table 1), because records of hip fracture-related hospital days were available in total only. To allow also for patients with ≥ 2 fractures in the study interval (maximum: 22 fractures), median hospital days contingent upon patient age and gender were calculated for each number of fractures (Online Resource, supplemental Table 1), which served to assess the proportion of hospital days assigned to each fracture. Rounded hospital days pertaining to the first fracture were thus subtracted from the known date of hospital discharge, resulting in an estimate of the date of first fracture for patients with ≥ 2 fractures.

Results

Table 1 lists baseline characteristics of the fracture study population. Most female hip fracture patients were aged 85–89 years, whereas most males were 80–84 years. With respect to number of fractures, gender proportions were similar regardless of number of fractures. However, χ^2

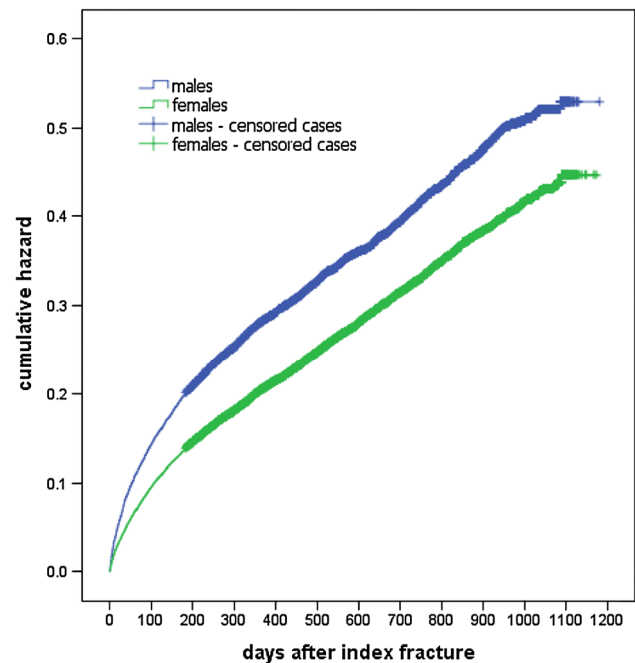


Fig. 1 Kaplan–Meier survival diagram depicting cumulative absolute mortality hazards over time for female and male hip fracture patients upon admission to hospital following index fracture

analysis revealed a statistically significant difference ($P < 0.001$) between patients with only one fracture and those with more fractures (not shown). Patients sustaining a hip fracture stayed a mean of 22.0 (SD: 16.4) days in hospital (median 16 days (IQR: 17); Online Resource, supplemental Table 1). Women stayed significantly longer in hospital than men overall (median 17 days (IQR: 17) vs 16 days (IQR: 16), $P < 0.001$), which was particularly reflected by patients between 65 and 89 years at hospital discharge after index fracture. Overall and for both genders, peak age for hospital days was between 75 and 84 years, i.e., up to a median of 18 days (IQR: 18) corresponding to a mean of 23.8 (SD: 17.4) days among female patients aged 75–79.

Cumulative hazards for mortality following hospital admission upon index fracture are shown in Fig. 1: Overall, 1-year mortality rates in women and men were 18.6 % (95 % CI: 18.1–19.1 %) and 24.5 % (95 % CI: 23.6–25.5 %), respectively (Fig. 1; Online Resource, supplemental Table 2). Online Resource, supplemental Table 2 provides detailed information on cumulative all-cause mortality rates of various time intervals from 30 days to two and a half years after both first hospital discharge and date of first admission (estimated for ≥ 2 fractures from hip fracture-related hospital days). As expected, cumulative mortality increases in both women and men over time, and at all time points, men are more prone to death than women (Fig. 1; Online Resource, supplemental Table 2).

Table 2 Cumulative all-cause one-year mortality age group and gender-wise after (i) first hospital discharge (upper row in each age group) and (ii) first admission (estimated for ≥ 2 fractures; lower row in each age group) in the study

Age group (years) at 1st		Women	Men	<i>P</i>	All
Discharge	Admission	Mortality (95 % CI)	Mortality (95 % CI)		Mortality (95 % CI)
All		16.5 % (16.0–17.0 %)	20.8 % (19.9–21.8 %)	<0.001	17.6 % (17.2–18.1 %)
	All	18.6 % (18.1–19.1 %)	24.5 % (23.6–25.5 %)	<0.001	20.2 % (19.7–20.7 %)
50–54		5.8 % (2.9–8.8 %)	6.2 % (3.7–8.7 %)	0.84	5.9 % (4.1–7.8 %)
	50–54	6.9 % (3.7–10.0 %)	7.2 % (4.5–9.8 %)	0.91	7.0 % (5.0–9.0 %)
55–59		5.3 % (3.1–7.4 %)	6.2 % (3.9–8.4 %)	0.51	5.6 % (4.1–7.2 %)
	55–59	5.1 % (3.1–7.2 %)	8.4 % (5.9–10.9 %)	0.036	6.7 % (5.1–8.4 %)
60–64		6.3 % (4.4–8.1 %)	8.6 % (6.3–10.9 %)	0.123	7.3 % (5.8–8.8 %)
	60–64	6.7 % (4.8–8.6 %)	9.4 % (7.0–11.8 %)	0.101	7.9 % (6.4–9.4 %)
65–69		7.9 % (6.4–9.5 %)	10.8 % (8.7–12.9 %)	0.033	9.1 % (7.8–10.3 %)
	65–69	9.1 % (7.5–10.7 %)	12.8 % (10.6–15.1 %)	0.001	10.6 % (9.3–11.9 %)
70–74		7.2 % (5.9–8.5 %)	13.8 % (11.5–16.2 %)	<0.001	9.5 % (8.3–10.6 %)
	70–74	8.2 % (6.8–9.5 %)	17.0 % (14.5–19.5 %)	<0.001	11.3 % (10.0–12.5 %)
75–79		9.1 % (8.0–10.1 %)	20.3 % (18.0–22.6 %)	<0.001	12.3 % (11.3–13.4 %)
	75–79	10.3 % (9.2–11.4 %)	23.0 % (20.7–25.3 %)	<0.001	14.0 % (13.0–15.1 %)
80–84		15.4 % (14.4–16.4 %)	27.6 % (25.3–30.0 %)	<0.001	18.1 % (17.1–19.0 %)
	80–84	17.4 % (16.4–18.4 %)	31.5 % (29.2–33.8 %)	<0.001	20.6 % (19.6–21.5 %)
85–89		20.9 % (19.9–22.0 %)	31.2 % (28.6–33.8 %)	<0.001	22.7 % (21.7–23.7 %)
	85–89	23.3 % (22.2–24.2 %)	36.4 % (33.8–39.0 %)	<0.001	25.7 % (24.7–26.7 %)
90–94		28.6 % (26.6–30.6 %)	40.0 % (35.0–44.9 %)	<0.001	30.5 % (28.6–32.3 %)
	90–94	32.3 % (30.2–34.3 %)	47.0 % (42.3–51.7 %)	<0.001	34.9 % (33.0–36.8 %)
95+		38.9 % (35.4–42.3 %)	51.6 % (43.6–59.7 %)	<0.001	40.9 % (37.7–44.1 %)
	95+	43.1 % (39.7–46.5 %)	57.8 % (50.2–65.3 %)	<0.001	45.5 % (42.4–48.6 %)

Furthermore, mortality sharply increases above 80 years in women and 75 years in men (Table 2).

In-hospital mortality was analyzed for all fractures occurring during the study period (Table 3), amounting to overall 3.6 % (95 % CI: 3.4–3.8 %) and 4.0 % (95 % CI: 3.8–4.3 %) at first and last fracture, respectively. Male patients had an elevated mortality rate ($P < 0.001$) at both their first (5.1 %, 95 % CI: 4.6–5.6 %) and last fracture (5.7 %, 95 % CI: 5.2–6.2 %). Reported by age groups, overall in-hospital mortality was lowest among patients aged 60–64 years both for first and last fractures. In contrast, in-hospital mortality was as high as 8.9 % (95 % CI: 7.2–10.6 %) among individuals aged 95 years and above. At all ages except for patients aged 50–54 and 60–64 years, men had significantly higher in-hospital mortality than women.

Unadjusted and adjusted hazard ratios (HRs) were calculated for mortality of the entire study interval based on first admission dates (estimated for ≥ 2 fractures) including variables gender, age group, hospital days, and number of fractures (Table 4). Men's hazard of shorter time to death relative to female patients rose from 1.3 in the unadjusted analysis to approximately 1.75 in any of the adjusted

models, and advanced age entailed a mortality risk of more than 12-fold for the oldest in relation to the youngest patients. Moreover, an effect of hospital days per fracture on risk reduction was evident in the adjusted cases, resulting in a 0.3–0.4 % hazard reduction of each additional hospital day in the adjusted models. Finally, sustaining more than one fracture was associated with improved survival. An interaction model comprising all variables (Online Resource, supplemental Table 3) demonstrated that more advanced age increased mortality hazard, likewise more hospital days per fracture were weakly but significantly associated with higher risk. In detail, male patients at age 70–79 years were at elevated mortality risk, whereas increased hospitalization time reduced risk in patients aged 75 years and over and in patients sustaining three fractures.

Next, we determined hip fracture patients' mortality risk relative to a control population represented by individuals without a history of hip fracture. Indeed, we found increased risk at all time points up to 3 years as compared to controls (Table 5): Excess mortality expressed as relative hazard (RH) was highest within 30 days after hip fracture, being twice as high in men as compared to women

Table 3 In-hospital mortality age group and gender-wise, accounting for each patient's (i) only first fracture (upper row in each age group) or (ii) last fracture (lower row in each age group) recorded within study interval

Age group (years) at		Women	Men	<i>P</i>	All
First fracture	Last fracture	Mortality (95 % CI)	Mortality (95 % CI)		Mortality (95 % CI)
All		3.0 % (2.8–3.3 %)	5.1 % (4.6–5.6 %)	<0.001	3.6 % (3.4–3.8 %)
	All	3.4 % (3.2–3.7 %)	5.7 % (5.2–6.2 %)	<0.001	4.0 % (3.8–4.3 %)
50–54		0.8 % (0.0–1.8 %)	1.3 % (0.2–2.4 %)	0.71	1.1 % (0.3–1.9 %)
	50–54	0.8 % (0.0–1.8 %)	1.3 % (0.2–2.4 %)	0.71	1.1 % (0.3–1.9 %)
55–59		0.4 % (0.0–1.0 %)	2.7 % (1.3–4.2 %)	0.004	1.6 % (0.8–2.4 %)
	55–59	0.4 % (0.0–1.1 %)	2.9 % (1.4–4.5 %)	0.003	1.7 % (0.9–2.5 %)
60–64		0.4 % (0.0–0.9 %)	1.2 % (0.3–2.0 %)	0.137	0.8 % (0.3–1.2 %)
	60–64	0.7 % (0.1–1.4 %)	1.2 % (0.3–2.0 %)	0.42	0.9 % (0.4–1.5 %)
65–69		1.3 % (0.6–1.9 %)	2.6 % (1.6–3.6 %)	0.019	1.8 % (1.3–2.4 %)
	65–69	1.4 % (0.7–2.0 %)	2.8 % (1.7–3.9 %)	0.015	2.0 % (1.4–2.6 %)
70–74		1.1 % (0.6–1.6 %)	3.8 % (2.5–5.0 %)	<0.001	2.1 % (1.5–2.6 %)
	70–74	1.3 % (0.8–1.9 %)	4.0 % (2.7–5.3 %)	<0.001	2.2 % (1.7–2.8 %)
75–79		1.7 % (1.2–2.1 %)	3.6 % (2.6–4.6 %)	<0.001	2.3 % (1.8–2.7 %)
	75–79	1.9 % (1.4–2.4 %)	4.2 % (3.1–5.3 %)	<0.001	2.5 % (2.1–3.0 %)
80–84		2.8 % (2.4–3.3 %)	6.3 % (5.1–7.5 %)	<0.001	3.6 % (3.2–4.0 %)
	80–84	3.2 % (2.7–3.7 %)	6.9 % (5.7–8.2 %)	<0.001	4.0 % (3.6–4.5 %)
85–89		3.7 % (3.2–4.2 %)	8.0 % (6.5–9.4 %)	<0.001	4.5 % (4.0–4.9 %)
	85–89	4.3 % (3.8–4.8 %)	8.7 % (7.2–10.2 %)	<0.001	5.1 % (4.6–5.6 %)
90–94		5.8 % (4.8–6.8 %)	12.6 % (9.5–15.6 %)	<0.001	7.0 % (6.0–8.0 %)
	90–94	6.4 % (5.4–7.5 %)	14.9 % (11.6–18.1 %)	<0.001	7.9 % (6.9–8.9 %)
95+		7.3 % (5.5–9.0 %)	12.6 % (7.7–17.6 %)	0.017	8.2 % (6.5–9.9 %)
	95+	7.9 % (6.1–9.7 %)	13.7 % (8.6–18.8 %)	0.013	8.9 % (7.2–10.6 %)

(RH 16.28 (95 % CI: 12.47–21.26) versus 8.29 (95 % CI: 7.09–9.70), respectively, first admission as reference time point). RHs decreased considerably within 6 months in both men and women, but remained statistically significant for the whole observation period. Furthermore, supplemental Tables 4 and 5 in the Online Resource show age-group dependent RHs for the first and second year, respectively. RHs decreased with advancing age, with the exception of males aged 95 years and over where RH increased again. Interestingly, whereas women displayed RHs higher than their age-matched male counterparts up to 65–69 years both within the first and during the second year, and RHs within the second year were roughly equal among genders between 70 and 80 years of age, it was men's RHs that exceeded those of women at ages above.

Discussion

Herein, we present a large-scale up-to-date analysis of mortality rates up to 3 years after hip fractures in Austrian patients in the period of 2008–2011. As expected and also described previously for the same population, hip fractures

occur much more frequently in women than in men (ratio roughly 3:1), comparable to other nation-wide studies [8, 12, 24–26]. Interestingly, the proportion of men decreased at subsequent fractures (Table 1, “Results” section) probably being partially related to higher male mortality after first hip fracture.

We also analyzed length of hospital stay due to hip fracture. As compared to a previous publication from the same country analyzing data from 1995, we found very similar hospitalization time in both investigations (mean 22 days) [21]. Also, peak mean hospital days were between 75 and 84 years in both studies. Advanced age seemed to be associated with longer hospitalization among women as compared with men in the same report [21], though no *P* values are listed. Overall, our results on age- and gender-adjusted length of hospital stay after hip fracture suggest no substantial changes from 1995–2011 [21].

Considering all-cause mortality, we found death rates of 4.8 and 20.2 % after 30 days and 1 year, respectively. These numbers are among the lowest compared to investigations of similar size from other countries. A large retrospective cohort study from Denmark observed 13 and 29 % deaths after 30 days and 1 year, respectively, accounting for all

Table 4 Unadjusted and adjusted hazard ratios (HRs) for risk of shorter survival, based on first admission date (estimated for ≥ 2 fractures)

Variable	Unadjusted HR (95 % CI)	<i>P</i>	Adjusted HR (95 % CI)	<i>P</i>	Adjusted HR (95 % CI)	<i>P</i>	Adjusted HR (95 % CI)	<i>P</i>
Gender								
Women	Reference		Reference		Reference		Reference	
Men	1.30 (1.24–1.36)	<0.001	1.76 (1.67–1.84)	<0.001	1.76 (1.67–1.84)	<0.001	1.74 (1.66–1.83)	<0.001
Age group (years)								
50–54	Reference		Reference		Reference		Reference	
55–59	1.14 (0.81–1.59)	0.45	1.20 (0.86–1.68)	0.29	1.21 (0.86–1.69)	0.27	1.20 (0.86–1.68)	0.29
60–64	1.33 (0.97–1.81)	0.076	1.43 (1.04–1.95)	0.026	1.44 (1.05–1.97)	0.023	1.44 (1.05–1.97)	0.022
65–69	1.64 (1.23–2.20)	<0.001	1.83 (1.37–2.44)	<0.001	1.84 (1.38–2.46)	<0.001	1.84 (1.38–2.46)	<0.001
70–74	1.86 (1.40–2.48)	<0.001	2.16 (1.62–2.87)	<0.001	2.19 (1.65–2.91)	<0.001	2.21 (1.66–2.94)	<0.001
75–79	2.53 (1.92–3.33)	<0.001	3.05 (2.32–4.01)	<0.001	3.10 (2.36–4.09)	<0.001	3.16 (2.40–4.17)	<0.001
80–84	3.70 (2.83–4.85)	<0.001	4.67 (3.56–6.12)	<0.001	4.76 (3.63–6.24)	<0.001	4.84 (3.69–6.35)	<0.001
85–89	4.83 (3.69–6.32)	<0.001	6.25 (4.77–8.19)	<0.001	6.36 (4.85–8.34)	<0.001	6.46 (4.92–8.46)	<0.001
90–94	7.01 (5.34–9.21)	<0.001	9.18 (6.98–12.08)	<0.001	9.33 (7.09–12.27)	<0.001	9.38 (7.13–12.33)	<0.001
95+	9.73 (7.36–12.86)	<0.001	12.78 (9.66–16.92)	<0.001	12.89 (9.74–17.07)	<0.001	12.76 (9.64–16.89)	<0.001
Hospital days per fracture	0.999 (0.997–1.000)	0.131			0.997 (0.995–0.999)	<0.001	0.996 (0.994–0.998)	<0.001
Number of fractures								
1	Reference						Reference	
2	0.68 (0.64–0.72)	<0.001					0.68 (0.64–0.73)	<0.001
3	0.62 (0.53–0.72)	<0.001					0.63 (0.54–0.74)	<0.001
4	0.70 (0.52–0.93)	0.015					0.70 (0.52–0.93)	0.015
≥ 5	0.41 (0.23–0.71)	0.002					0.39 (0.22–0.69)	0.001

subjects irrespective of age; first patients were, however, recruited already in 1977 [12]. A more recent Danish nationwide cohort study including all hip fracture patients between 1999 and 2002 that were born up to 1945 revealed a cumulative 1-year mortality rate of 26.4 % for female and 37.1 % for male patients [24]. Bass et al. [27] retrospectively analyzed survival in veterans aged 65 and above registered in the US Medicare database, reporting unadjusted mortality rates of 9.7 and 32.2 % for 30 days and 1 year, respectively. A retrospective cohort study of Canadian hip fracture patients aged over 50 found all-cause mortality at 1 year post-fracture to be 36.3 % [28]. Along the same lines, in-hospital mortality of overall 4.0 % (95 % CI: 3.8–4.3 %) reported herein (Table 3) is low compared with previous reports from Austria including all age groups that stated 6.8 % in 1995 [21], and 3.8 and 3.2 % for male and female patients, respectively, in 2004 [22]. Reports from other countries incorporating patients aged 65 and over stated 4.5 % in a nationwide Danish study [29]), 5.1 % in New York state hospitals over a period of 12 years [30], and as high a figure as 14.3 % in an extensive investigation from England [31]. Conversely, nationwide investigations from South Korea including hip fracture patients aged 50 and older stated

overall 1-year mortality of 16.55 % in 2003 [25] and ranging from 17.8 to 19.0 % between 2005 and 2007 [32]. Recently, a nationwide study from Taiwan on patients aged 60 years and over reported declining annual hip fracture-related mortality from overall 18.10 % in 1999 to 13.98 % in 2009, corresponding to an average 1-year mortality of 16.32 % for the whole time interval, and 1-month mortality was even found as low as 2.49 % [33]. Thus, Austrian hip fracture patients seem to be at lower mortality risk as compared to most other countries except Eastern Asian populations.

Our results underscore the role of advanced age and male gender in hip fracture patients' mortality risk. Moreover, our analyses show that both longer hospitalization time per fracture and more fractures are in association with a significantly reduced risk. Well in line, longer medical care in hospital after hip fracture correlated with lower mortality in Japan as compared with the United States, even upon adjustment for confounding factors [34]. Alternatively, an Australian study claimed longer waiting times for healthier patients destined to enter a rehabilitation center than for more co-morbid patients who are transferred to nursing homes usually rather quickly [35]. However, variable interaction confers a statistically

Table 5 Excess mortality expressed as relative hazard (RH) of various time intervals after (i) first hospital discharge (upper row in each interval) and (ii) first admission (estimated for ≥ 2 fractures; lower row in each interval) in the study, both adjusted for gender

Interval after first		Women	<i>P</i>	Men	<i>P</i>	All	<i>P</i>
Discharge	Admission	RH (95 % CI)		RH (95 % CI)		RH (95 % CI)	
0–30 days		5.40 (4.59–6.36)	<0.001	10.41 (7.91–13.71)	<0.001	6.60 (5.73–7.59)	<0.001
	0–30 days	8.29 (7.09–9.70)	<0.001	16.28 (12.47–21.27)	<0.001	10.21 (8.93–11.68)	<0.001
31–90 days		3.77 (3.36–4.24)	<0.001	6.05 (4.97–7.38)	<0.001	4.31 (3.90–4.77)	<0.001
	31–90 days	4.15 (3.70–4.66)	<0.001	7.25 (5.98–8.80)	<0.001	4.89 (4.43–5.40)	<0.001
91 days–6 months		2.34 (2.12–2.58)	<0.001	3.00 (2.57–3.49)	<0.001	2.52 (2.32–2.74)	<0.001
	91 days–6 months	2.55 (2.32–2.80)	<0.001	3.39 (2.92–3.93)	<0.001	2.78 (2.57–3.02)	<0.001
6 months–1 year		1.63 (1.50–1.76)	<0.001	2.50 (2.19–2.85)	<0.001	1.83 (1.71–1.96)	<0.001
	6 months–1 year	1.70 (1.57–1.84)	<0.001	2.60 (2.28–2.96)	<0.001	1.908 (1.78–2.04)	<0.001
1–1½ years		1.61 (1.47–1.77)	<0.001	2.10 (1.80–2.45)	<0.001	1.73 (1.60–1.88)	<0.001
	1–1½ years	1.59 (1.45–1.74)	<0.001	2.03 (1.74–2.37)	<0.001	1.70 (1.57–1.84)	<0.001
1½–2 years		1.75 (1.57–1.95)	<0.001	2.45 (2.04–2.95)	<0.001	1.91 (1.74–2.09)	<0.001
	1½–2 years	1.79 (1.61–1.98)	<0.001	2.49 (2.08–2.98)	<0.001	1.94 (1.77–2.13)	<0.001
2–2½ years		1.66 (1.46–1.89)	<0.001	2.68 (2.17–3.31)	<0.001	1.88 (1.68–2.09)	<0.001
	2–2½ years	1.65 (1.46–1.87)	<0.001	2.71 (2.21–3.32)	<0.001	1.87 (1.68–2.08)	<0.001
2½–3 years		1.44 (1.15–1.80)	0.002	1.71 (1.10–2.64)	0.016	1.50 (1.23–1.84)	<0.001
	2½–3 years	1.48 (1.20–1.82)	<0.001	1.90 (1.29–2.78)	0.001	1.57 (1.31–1.88)	<0.001
0–90 days		4.28 (3.90–4.71)	<0.001	7.44 (6.34–8.72)	<0.001	5.03 (4.64–5.46)	<0.001
	0–90 days	5.45 (4.97–5.97)	<0.001	10.12 (8.67–11.81)	<0.001	6.57 (6.07–7.10)	<0.001
0–6 months		3.23 (3.02–3.46)	<0.001	4.82 (4.33–5.37)	<0.001	3.64 (3.44–3.86)	<0.001
	0–6 months	3.88 (3.64–4.14)	<0.001	6.15 (5.54–6.82)	<0.001	4.47 (4.23–4.72)	<0.001
0–1 year		2.44 (2.32–2.57)	<0.001	3.75 (3.45–4.07)	<0.001	2.76 (2.65–2.89)	<0.001
	0–1 year	2.80 (2.67–2.94)	<0.001	4.50 (4.16–4.87)	<0.001	3.22 (3.09–3.36)	<0.001
1–2 years		1.67 (1.56–1.79)	<0.001	2.24 (1.99–2.52)	<0.001	1.80 (1.70–1.92)	<0.001
	1–2 years	1.67 (1.56–1.79)	<0.001	2.21 (1.96–2.48)	<0.001	1.80 (1.69–1.91)	<0.001
2–3 years		1.60 (1.43–1.79)	<0.001	2.43 (2.01–2.93)	<0.001	1.78 (1.62–1.96)	<0.001
	2–3 years	1.60 (1.44–1.78)	<0.001	2.48 (2.07–2.97)	<0.001	1.79 (1.63–1.96)	<0.001

significant effect of longer hospitalization at age 75 and over, associated with decreased mortality risk. This implies that aged patients who need more time to recover benefit in particular from longer in-hospital care. Thus following the explanation offered by Williams et al. [35], one would have to assume either low co-morbidity or particularly long waiting times until rehabilitation among older patients; both scenarios are unlikely in Austrian patients. Concerning subsequent fractures as protective factor, this could be due to the fact that surviving patients are relatively healthy but prone to the next fracture as a consequence of osteoporosis.

Notably, the impact of gender and consecutive fractures as such vanished in the interaction model unless in conjunction with other variables (Online Resource, supplemental Table 3): Though hazards rose with age independently of gender, interaction terms revealed a significantly elevated HR for men aged 70–74 compared to their female counterparts, implying particular mortality risk for male patients of that age.

Relevance for fracture number was observed only in interaction with hospital days per fracture entailing risk reduction for (at least) three fractures (no statistical significance for ≥ 4 fractures is probably due to low patient numbers, HRs for 4 and ≥ 5 fractures are below that for 3 fractures); therefore it is patients with more subsequent fractures who benefit the most from a longer stay in hospital in terms of survival.

Relative hazards (RHs) for all-cause excess mortality during various time intervals from hospital admission upon index fracture in our patients are comparable with those of a previous meta-analysis [13] although male mortality rates are slightly higher among our patients where RHs dropped to below 2 after 6 months and remained in this range up to 3 years. Our results are in line with others reporting a rapid decline within the first year post-fracture, yet a persistent elevation for an observation period of as long as 10 years [12, 13]. In contrast, studies investigating institutionalized patients reported no significant excess mortality among patients surviving the first 6 months after fracture [14, 15].

Another observation in our study is that during both first and second year post-fracture, excess mortality up to 69 years of age was higher among women than men. However, these findings should be interpreted cautiously in view of the large-sized 95 % CIs at these age groups that are due to fewer study subjects and more censored cases at those ages as compared with more advanced age groups. On the other hand, studies from South Korea [32] and Taiwan [33] found higher female excess mortality rates up to 79 years in the first and second year post-fracture, and higher male excess mortality only after 80 years of age. These findings are at odds with another study claiming excess mortality to be higher in men than women at all ages [36] but are consistent with results of a meta-analysis that reported higher excess mortality for men only at 70 years and above [13].

Enhanced administration of anti-resorptive drugs such as bisphosphonates in conjunction with effective orthogeriatric care measures could be causative for the low mortality rates following hip fracture reported herein in relation to data from other countries. Also, we found reduced in-hospital mortality relative to one previous national study that analyzed data from 1995 [21]. In recent years in Austria, a number of orthogeriatric care services were initiated before or within the study interval of the present investigation, however, on a regional and local basis rather than as part of an overarching, coordinated nationwide program [37–39]. These initiatives not only ensure due prescription of medication like bisphosphonates and adherence to therapy, they also implement and reinforce integrated measures of postoperative care [40, 41]. Mortality data from one such care center on hip fracture patients aged 80 years and above were reported to be 2.8 % in hospital and 23.2 % one year post-fracture, which is strikingly low for this age group [42]. Moreover, mortality outcomes of international orthogeriatric service programs suggest improved survival rates at least in hospital (summarized in [40]), therefore declining in-hospital and perhaps also lowered longer term mortality might in part be attributed to efficacious orthogeriatric care measures in Austria. As for bisphosphonates, an increase in their prescription during the 2000s was demonstrated to coincide with a reduction in hip fracture incidence in two studies [43, 44], and during the same period, a decreasing trend in hip fracture incidence was also observed in Austria [8]. Further, there is evidence for an impact of bisphosphonate treatment on prolonged survival after hip fractures [45–47]. The fact that male excess mortality rates are above those seen in other studies might be based on suboptimal osteoporosis treatment in these patients, as previously described for Denmark [47]. Collectively, disparate study results during the last 10–15 years on short-term and long-term

mortality after hip fracture reported at different times and in different countries could mirror varying evolution stages in the implementation of orthogeriatric services and anti-resorptive treatment regimes alike.

Our study is subject to several limitations. First, no information on lifestyle factors and co-morbidities was available. Second, the date of first admission was not directly available for 21.4 % of patients sustaining more than one fracture during the study interval, for whom it had to be assessed from hip fracture-related hospital days. An alternative to this approach would have been exclusion of subsequent fractures like elsewhere [29], however, we believe that such a procedure would have conferred a substantial bias to our investigation. Third, mean follow-up time of 528 days for hip fracture patients did not permit conclusions on long-term mortality beyond 3 years, unlike other previous comparable studies (e.g., [12–14]). Finally, there was no criterion to ensure that first fractures recorded in the study period (index fractures for which hospital discharge dates were available) were first fractures that patients sustained: Exclusion of index fractures that were in fact not patients' first fractures might have tilted estimates towards slightly shorter survival times. On the other hand, we are convinced that substantive strengths of our work encompass (i) the participation of almost all social insurance authorities, corresponding to coverage of approximately 98 % of the total Austrian population, (ii) the access to personalized data thus ruling out multiple registrations, implying that all fracture events could be unambiguously assigned to study subjects, and (iii) the allowance for mortality rates beginning with hospital discharge upon first fracture in the study interval, which represents valuable information regarding survivors of (first) hospitalization.

Taken together, our results demonstrate in a nation-wide retrospective study that recent all-cause mortality rates including in-hospital mortality after hip fracture among elderly patients are low in Austria compared with many other countries, however, male excess mortality is elevated. Moreover, in a multivariate model accounting for variable interactions, we found that longer hospital care favors patients aged 75 and above and those sustaining (at least) three fractures the most, and that the highest gender difference in mortality risk after hip fracture is among patients aged 70–79 years with males at greater risk. Further research is necessary to identify causes of male excess mortality. Elderly male osteoporotic patients are therefore an important high-risk group after hip fracture in Austria.

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