Admission Resting Heart Rate as an Independent Predictor of All-Cause Mortality in Elderly Patients with Hip Fracture

Zhicong Wang*

Xi Chen*

Yuxuan Wu*

Wei Jiang

Ling Yang

Hong Wang

Shuping Liu

Yuehong Liu

Orthopaedic Center of Deyang City,
Department of Orthopedics, Deyang
People's Hospital, Deyang, Sichuan,
People's Republic of China

*These authors contributed equally to
this work

Purpose: To investigate the relationship between admission resting heart rate (RHR) and all-cause mortality in elderly patients with hip fracture.

Methods: A retrospective cohort study with 837 patients based on the established hip fracture database was conducted. Admission RHR was measured via electrocardiogram, and patients were grouped by the median RHR value (beats per minute, bpm). The main outcomes were 1-year and total all-cause mortality. Cox proportional hazard models and restricted cubic spline were used to assess the relationship between RHR and mortality. Sensitivity analyses were further performed to determine whether the results were stable.

Results: The mean and median RHR were 82.3 and 80.0 bpm, respectively. After a median follow-up of 31.8 months, the 1-year and total all-cause mortality were 17.6% and 31.2%. Multivariable Cox analyses showed that high RHR was an independent risk factor for 1-year mortality ($HR = 1.51$; 95% CI : 1.08–2.13; $p = 0.016$), and total mortality ($HR = 1.44$; 95% CI : 1.12–1.85; $p = 0.005$). For each 10 bpm increase in RHR, the risk of 1-year death increased by 23.0% ($HR = 1.23$; 95% CI : 1.09–1.39; $p = 0.001$), and total death increased by 21.0% ($HR = 1.21$; 95% CI : 1.09–1.34; $p < 0.001$). A typical J-shaped curve was observed in the restricted cubic spline for the association between RHR and 1-year mortality, with the lowest mortality risk at 70 bpm. Sensitivity analyses yielded similar findings.

Conclusion: An increase in RHR was independently associated with all-cause mortality, and may be a useful prognostic predictor for elderly patients with hip fracture.

Keywords: elderly, hip fracture, resting heart rate, prognosis, mortality

Introduction

Resting heart rate (RHR) is an easily accessible physiological indicator which reflects the activity of the autonomic nervous system. Regarding its predictive value, elevated RHR was found to be associated with a higher risk of all-cause mortality in various cardiovascular diseases (CVD), such as hypertension,¹ heart failure,² myocardial infarction³ and chronic aortic regurgitation.⁴ Recently, this association has also been observed in patients with chronic obstructive pulmonary disease (COPD),⁵ diabetes,⁶ and even in the general population.⁷ Notably, Li et al⁸ found that high RHR was an important risk factor for both all-cause mortality and CVD events in older subjects but not in younger people.

Hip fracture is a common fracture in elderly patients, and frequently comorbid with the aforementioned diseases.⁹ Therefore, we hypothesized that an increase in RHR may have an important impact on prognosis in elderly patients with hip

Correspondence: Shuping Liu;
Yuehong Liu
Orthopaedic Center of Deyang City,
Department of Orthopedics, Deyang
People's Hospital, No. 173 Taishan North
Road, Jingyang District, Deyang, 618000,
People's Republic of China
Tel +86 838 241 8102
Fax +86 838 241 8102
Email doctorliushuping@126.com;
doctorliuyuehong@163.com

fracture. However, only one previous study has reported that older women with rapid RHR had an increased risk of mortality in hip fracture patients.¹⁰ In the present study, by utilizing the established hip fracture database,¹¹ we were able to investigate the relationship between admission RHR and all-cause mortality.

Methods

Study Design and Patients

This was a retrospective cohort design. As we previously described,¹¹ hip fracture patients admitted to the Orthopaedic Center of Deyang City from January 2014 to December 2020 were enrolled consecutively. Briefly, the inclusion criteria for this database were as follows: (1) diagnosis of hip fracture; (2) age ≥ 60 years; (3) fresh fracture less than 3 weeks. Any pathological fractures or fractures resulting from a high energy trauma were excluded from the database. In the current study, patients with atrial fibrillation or missing mortality data were then excluded. The study was performed according to the Declaration of Helsinki, and approved by the Institutional Ethics Committee of Deyang People's Hospital. As a retrospective study, the Ethics Committee agreed to waive the requirement for the informed consent, and all patient-related data were anonymized to ensure confidentiality.

Data Collection and Definition

The following characteristics were extracted from the clinical database: age at admission (years), sex (male or female), height (m) and weight (kg), current smoking and drinking status (yes or no), hip fracture type (femur neck fracture or intertrochanteric fracture), hypertension (yes or no) and treatment modality (surgical and non-surgical). Patient comorbidity was assessed with the Charlson comorbidity index (CCI), which is one of the most widely used comorbidity models.¹² Moreover, admission laboratory data related to the prognosis of hip fracture patients were also obtained, including lymphocyte [reference range: $(1.1-3.2) \times 10^9/L$], hemoglobin (reference range: 130.0–175.0 g/L), and albumin (reference range: 35.0–55.0 g/L).

For a better understanding of the results, age was grouped into four categories: 60–69 years, 70–79 years, 80–89 years and ≥ 90 years. Body mass index (BMI) was calculated as $\text{weight (kg)}/[\text{height (m)}]^2$, and grouped as follows: underweight ($\text{BMI} < 18.5 \text{ kg/m}^2$), normal weight

($\text{BMI} = 18.5-24.9 \text{ kg/m}^2$), and overweight ($\text{BMI} \geq 25.0 \text{ kg/m}^2$). The CCI score was calculated based on 17 diseases, and categorized as none ($\text{CCI} = 0$), low ($\text{CCI} = 1$) and high ($\text{CCI} \geq 2$).¹² Hypertension was defined as systolic blood pressure $\geq 140 \text{ mmHg}$ or diastolic blood pressure $\geq 90 \text{ mmHg}$ or taking any antihypertensive drugs. According to the lower limit of reference range, lymphopenia and hypoalbuminemia were defined as an absolute lymphocyte count $< 1.1 \times 10^9/L$ and serum albumin $< 35.0 \text{ g/L}$, respectively. Based on the World Health Organization (WHO) criteria, anemia was defined as hemoglobin level $< 130 \text{ g/L}$ for men and $< 120 \text{ g/L}$ for women.

RHR Measurement

On admission, all patients routinely underwent a 12-lead electrocardiography (aECG-18U; Nalong Technologies, China) after 5 minutes of bed rest. RHR (beats per minute, bpm) was measured by assessing the time between each R wave on the ECG record. According to the median value, patients were divided into 2 groups: low RHR group and high RHR group.

Follow-Up and Outcome

All surviving patients in the database were telephonically followed up after hospital discharge, and the follow-up contents included survival status (alive or dead) and the date of death. For patients who died in hospital, the date of death was obtained from medical record. Survival time was defined as the time interval from the date of hospital admission to the date of death from any cause or the last follow-up date (December 31, 2020), whichever occurred first. In this study, the main outcomes were 1-year all-cause mortality and total all-cause mortality at the last follow-up.

Statistical Analysis

Prior to analysis, missing values were checked and imputed using a multivariate normal imputation method.¹³ Continuous data were assessed for normality using the Shapiro–Wilk test. Normally distributed variables were described as means $\pm$ standard deviation (SD), and analyzed by independent Student's *t*-test, while non-normally distributed variables were expressed as median (Q1, Q3), and performed using Wilcoxon rank-sum test. Categorical data were presented as frequencies (percentages), and compared by chi-square test. Survival probability among the RHR groups was estimated by the Kaplan-Meier and Log rank test, and the median follow-

up time was calculated by the reverse Kaplan-Meier method. Univariable Cox proportional hazard models were used to identify potential predictor variables, and factors with $p < 0.10$ were then entered into the multi-variable Cox analyses to investigate the association between RHR and mortality. Model 1 was adjusted for age, sex and BMI. Model 2 was additionally adjusted for hip fracture type, CCI, treatment modality, lymphopenia, anemia and hypoalbuminemia. In addition, Cox models with the same covariates were also performed with RHR as a continuous variable (10 bpm increase). To better investigate the potential non-linear association between RHR and 1-year mortality, adjusted Cox model using restricted cubic spline with 5 knots was further constructed. Considering that RHR may be influenced by the use of antihypertensive drugs, sensitivity analyses were further performed to determine whether the results persisted in patients with or without hypertension. A two-tailed $p < 0.05$ was considered statistically significant. All the statistical analyses were performed with JMP Pro software (version 13.0.0; SAS Institute Inc., Cary, NC, USA).

Results

Baseline Characteristics

Out of 1044 patients in the original database, 207 patients were excluded according to the exclusion criteria. Ultimately, a total of 837 patients were included in the final analysis (Figure 1). The patient characteristics are listed in Table 1. Overall, the mean age was 78.8 ± 8.6 years, 552 (65.9%) were female, and the mean and median RHR were 82.3 and 80.0 bpm, respectively. Among these patients, 423 (50.5%) had below median RHR (≤ 80 bpm), and 414 (49.5%) had above median RHR (> 80 bpm). Patients with a higher RHR were more likely to receive non-surgical treatment (32.1% vs 23.2%, $p = 0.004$), while no significant differences were found with regard to demographics and laboratory data.

RHR and All-Cause Mortality

After a median follow-up of 31.8 months, 147 (17.6%) and 261 (31.2%) patients died within 1 year and the last follow-up. Compared with low RHR group, high RHR group suffered a higher 1-year mortality (22.3% vs 13.1%, $p = 0.001$) and total mortality (36.0% vs 26.5%, $p = 0.003$). Also, Kaplan-Meier analysis showed that high RHR patients had significantly worse survival than low RHR patients (log rank

Figure 1 Flow chart of the study.

$\chi^2 = 11.67$, $p = 0.001$; Figure 2). Consistent with the results, each 10 bpm increase in RHR was associated with a higher risk of 1-year and total mortality, with unadjusted hazard ratios (HR) of 1.44 (95% CI: 1.26–1.63) and 1.30 (95% CI: 1.18–1.44), respectively (Table 2). In addition, age, sex, BMI, hip fracture type, CCI score, treatment modality, lymphopenia, anemia and hypoalbuminemia were influencing factors for both 1-year and total mortality (all $p < 0.05$).

After further adjusting for the above potential confounding variables (model 2, Table 3), high RHR remained an independent risk factor for 1-year mortality (HR = 1.51; 95% CI: 1.08–2.13; $p = 0.016$), and total mortality (HR = 1.44; 95% CI: 1.12–1.85; $p = 0.005$). For each 10 bpm increase in RHR, the risk of 1-year death increased by 23.0% (HR = 1.23; 95% CI: 1.09–1.39; $p = 0.001$), and total death at the last follow-up increased by 21.0% (HR = 1.21; 95% CI: 1.09–1.34; $p < 0.001$). To further explore non-linear relationship, a typical J-shaped curve was observed in the restricted cubic spline for the association between RHR and 1-year mortality, with the lowest mortality risk at 70 bpm (Figure 3).

Sensitivity Analysis

To further identify whether hypertension might influence the relationship between RHR and mortality, sensitivity analyses were conducted (Table 4). The results were

Table 1 Patient Characteristics According to Admission Resting Heart Rate

Characteristics	Total (n = 837)	Low RHR ≤ 80 bpm (n = 423)	High RHR > 80 bpm (n = 414)	p value
Age (years)	78.8 ± 8.6	78.5 ± 8.2	79.2 ± 9.0	0.281
60–69 years, n (%)	145 (17.3)	74 (17.5)	71 (17.1)	0.229
70–79 years, n (%)	264 (31.5)	137 (32.4)	127 (30.7)	
80–89 years, n (%)	345 (41.2)	179 (42.3)	166 (40.10)	
≥ 90 years, n (%)	83 (9.9)	33 (7.8)	50 (12.1)	
Female, n (%)	552 (65.9)	273 (64.5)	279 (67.4)	0.384
BMI (kg/m ²)	21.64 ± 2.7	21.67 ± 2.6	21.61 ± 2.8	0.732
Underweight, n (%)	79 (9.4)	38 (9.0)	41 (9.9)	0.876
Normal weight, n (%)	689 (82.3)	349 (82.5)	340 (82.1)	
Overweight, n (%)	69 (8.2)	36 (8.5)	33 (8.0)	
Current smoker, n (%)	137 (16.4)	75 (17.7)	62 (15.0)	0.282
Current drinker, n (%)	83 (9.9)	48 (11.3)	35 (8.5)	0.161
Admission RHR (bpm)	82.3 ± 12.3	72.9 ± 6.6	91.9 ± 8.9	<0.001
Hip fracture type, n (%)				0.426
Femoral neck	416 (49.7)	216 (51.1)	200 (48.3)	
Intertrochanteric	421 (50.3)	207 (48.9)	214 (51.7)	
CCI, n (%)				0.695
None	428 (51.1)	222 (52.5)	206 (49.7)	
Low	247 (29.5)	123 (29.1)	124 (30.0)	
High	162 (19.4)	78 (18.4)	84 (20.3)	
Hypertension, n (%)	290 (34.6)	153 (36.2)	137 (33.1)	0.349
Surgical treatment, n (%)	606 (72.4)	325 (76.8)	281 (67.9)	0.004
Lymphocyte (×10 ⁹ /L)	0.96 (0.67, 1.25)	0.95 (0.68, 1.24)	0.97 (0.67, 1.25)	0.985
Lymphopenia, n (%)	535 (63.9)	269 (63.6)	266 (64.3)	0.843
Hemoglobin (g/L)	110.3 ± 20.9	111.3 ± 19.8	109.3 ± 21.9	0.183
Anemia, n (%)	594 (71.0)	299 (70.7)	295 (71.3)	0.856
Albumin (g/dL)	38.7 ± 4.7	38.9 ± 4.5	38.4 ± 4.9	0.151
Hypoalbuminemia, n (%)	176 (21.0)	88 (20.8)	88 (21.3)	0.873

Abbreviations: RHR, resting heart rate; bpm, beats per minute; BMI, body mass index; CCI, Charlson comorbidity index.

consistent with the main findings, and no significant hypertension × RHR interactions were observed.

Discussion

In this study, we consecutively included hip fracture patients from the established database, and found that admission RHR was significantly associated with 1-year mortality and total mortality at the last follow-up, even after controlling for potential confounding factors. Moreover, a typical J-shaped curve was observed in restricted cubic spline for the association between RHR and 1-year mortality, suggesting that RHR may be a useful prognostic predictor for elderly patients with hip fracture.

RHR is an important indicator of the health of the cardiovascular system, and easy to obtain clinically. Recently, increasing evidences have revealed that RHR may be a marker of healthy aging.¹⁴ Hartaigh et al¹⁵ analyzed 5691 older adults aged 65 years and over, and measured RHR annually for six consecutive years. The results showed that each 10 bpm increase in RHR had 33% greater hazard of all-cause mortality, which was consistent with the study by Floyd et al.¹⁶ Another study also found a similar risk relationship in the elderly population, but not in young adults.⁸ As is well known, hip fractures occur commonly in the elderly, and frequently combined with cardiovascular diseases, such as heart failure, coronary artery disease and atrial

Figure 2 Kaplan-Meier survival analysis for overall survival according to high and low resting heart rate (RHR).

fibrillation, which have been identified as independent risk factors for hip fracture survival.^{9,17,18} However, to our knowledge, this is the first study to clearly

demonstrate that an elevated RHR is associated with a higher all-cause mortality in elderly patients with hip fracture. Moreover, with each 10 bpm increase in RHR, the risk of 1-year death increased by 23.0%, and total death at the last follow-up increased by 21.0%.

Moreover, some studies utilized restricted cubic splines to explore the potential non-linear relationship between RHR and mortality. Cui et al¹⁹ found a U-shaped association between time-updated RHR and all-cause or cardiovascular mortality in a random middle-aged male cohort, and 60 bpm was associated with the lowest mortality, suggesting that both high and low RHR correlated with increased mortality in the general population. Also, the U-shaped curve was seen in critically ill patients, providing a new insight for optimizing HR control strategies.²⁰ Yet, other studies observed a J-shaped association in acute ischemic stroke patients,²¹ myocardial infarction patients,²² and

Table 2 Univariable Cox Regression Analysis for the Risk Factors Associated with 1-Year and Total All-Cause Mortality

Variables	1-Year Mortality		Total Mortality	
	HR (95% CI)	p value	HR (95% CI)	p value
Age (years)				
60–69	1.0 (reference)	-	1.0 (reference)	-
70–79	3.25 (1.56–7.93)	0.001	2.60 (1.53–4.70)	<0.001
80–89	4.83 (2.39–11.54)	<0.001	3.89 (2.35–6.92)	<0.001
≥ 90	6.58 (2.99–16.53)	<0.001	6.46 (3.68–11.99)	<0.001
Sex (male vs female)	1.68 (1.21–2.32)	0.002	1.28 (0.99–1.65)	0.056
BMI				
Underweight	1.10 (0.62–1.82)	0.722	0.92 (0.53–1.48)	0.739
Normal weight	1.0 (reference)	-	1.0 (reference)	-
Overweight	0.39 (0.14–0.86)	0.017	0.49 (0.22–0.93)	0.027
Current smoker	1.30 (0.85–1.93)	0.212	1.30 (0.94–1.77)	0.112
Current drinker	1.07 (0.60–1.76)	0.815	1.08 (0.70–1.59)	0.728
Admission RHR (High vs Low)	1.80 (1.29–2.52)	0.001	1.54 (1.20–1.97)	0.001
HR (Each 10 bpm increase)	1.44 (1.26–1.63)	<0.001	1.30 (1.18–1.44)	<0.001
Hip fracture type (intertrochanteric vs neck)	1.35 (0.97–1.87)	0.072	1.64 (1.27–2.11)	<0.001
CCI				
None	1.0 (reference)	-	1.0 (reference)	-
Low	1.96 (1.29–2.95)	0.002	1.83 (1.36–2.46)	<0.001
High	4.11 (2.78–6.11)	<0.001	3.48 (2.57–4.69)	<0.001
Surgical treatment (No vs Yes)	8.36 (5.90–12.05)	<0.001	4.82 (3.76–6.18)	<0.001
Lymphopenia (Yes vs No)	1.74 (1.21–2.55)	0.002	1.49 (1.15–1.94)	0.003
Anemia (Yes vs No)	1.82 (1.22–2.80)	0.003	1.71 (1.27–2.35)	<0.001
Hypoalbuminemia (Yes vs No)	2.13 (1.51–2.97)	<0.001	1.77 (1.35–2.29)	<0.001

Abbreviations: HR, hazard ratio; CI, confidence interval; BMI, body mass index; RHR, resting heart rate; bpm, beats per minute; CCI, Charlson comorbidity index.

Table 3 Multivariable Cox Regression Analysis for Admission Resting Heart Rate Associated with 1-Year and Total All-Cause Mortality

	1-Year Mortality		Total Mortality	
	HR (95% CI)	p value	HR (95% CI)	p value
Admission RHR (High vs Low)				
Model 1	1.79 (1.28–2.51)	0.001	1.56 (1.22–2.00)	0.001
Model 2	1.51 (1.08–2.13)	0.016	1.44 (1.12–1.85)	0.005
Admission RHR (Each 10 bpm increase)				
Model 1	1.43 (1.26–1.62)	<0.001	1.32 (1.19–1.46)	<0.001
Model 2	1.23 (1.09–1.39)	0.001	1.21 (1.09–1.34)	<0.001

Notes: Model 1 adjusted for age, sex and body mass index. Model 2 adjusted for model 1 plus hip fracture type, Charlson comorbidity index, treatment modality, lymphopenia, anemia and hypoalbuminemia.

Abbreviations: HR, hazard ratio; CI, confidence interval; RHR, resting heart rate.

even in the general population.²³ In agreement with the latter observations, we also observed a typical J-shaped curve for the association between RHR and 1-year mortality, and 70 bpm exhibited the lowest mortality risk.

Indeed, RHR may be easily affected by the use of antihypertensive medications, such as beta-blocker treatment. Recently, a meta-analysis that included 189,385 patients revealed no association between beta-blockers use and all-cause mortality in acute myocardial infarction.²⁴ Nevertheless, a beneficial effect of beta-blocker treatment has been previously reported in elderly patients with heart failure.^{25,26} In addition,

Figure 3 Restricted cubic spline for the association between resting heart rate (RHR) and 1-year all-cause mortality.

2 large cohort studies with more than 120,000 hip fracture patients also found that beta-blocker therapy was independently associated with a reduction in mortality.^{27,28} Due to the lack of detailed and reliable medication history, we were unable to conduct an analysis among patients with or without beta-blocker treatment. However, we attempted to address this limitation with sensitivity analysis in patients with or without hypertension. After multivariable adjustments, this relationship remained rather stable across hypertension status, and no significant hypertension $\times$ RHR interactions were observed, which was consistent with our main findings and previous study.¹

However, this study had several limitations. First, its retrospective nature reduce the statistical power of this study. Meanwhile, the sample size was relatively small, which limited our ability to assess the short-term prognosis due to the small number of death events. Thus, prospective, large sample size studies are required to validate the results of this study. Second, we could not obtain any medication information that may directly affect the RHR. This may be an important confounder, but we were unable to incorporate into the multivariable models. Third, only admission RHR was included in the analysis, which could not fully reflect the entire period. Finally, although we adjusted for potential confounders, many unknown variables might not have been included in the analyses.

In conclusion, an increase in RHR was independently associated with all-cause mortality, and may be a useful prognostic predictor for elderly patients with hip fracture.

Table 4 Sensitivity Analysis by Hypertension for 1-Year and Total All-Cause Mortality

	1-Year Mortality			Total Mortality		
	HR (95% CI)	p value	Interaction p value	HR (95% CI)	p value	Interaction p value
Hypertension						
Yes	1.40 (1.09–1.79)	0.008	0.144	1.30 (1.07–1.57)	0.007	0.875
No	1.17 (1.01–1.35)	0.031		1.16 (1.03–1.31)	0.017	

Notes: Adjusted for age, sex, body mass index, hip fracture type, Charlson comorbidity index, treatment modality, lymphopenia, anemia and hypoalbuminemia.

Abbreviations: HR, hazard ratio; CI, confidence interval.

Data Sharing Statement

The datasets used and/or analysed during the current study are available from the corresponding author on reasonable request.

Ethics Approval and Consent to Participate

The study was performed according to the Declaration of Helsinki, and approved by the Institutional Ethics Committee of Deyang People's Hospital. As a retrospective study, the Ethics Committee agreed to waive the requirement for the informed consent, and all patient-related data were anonymized to ensure confidentiality.

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Disclosure

The authors declare that they have no competing interests.

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