

Association of Sleep Duration, Midday Napping with Atrial Fibrillation in Patients with Hypertension

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Objective: This study aimed to assess the associations of sleep duration, midday napping and the risk of atrial fibrillation (AF) in patients with hypertension.

Methods: We conducted a cross-sectional study enrolling 11,524 hypertensive participants from the Chinese Hypertension Registry Study. Information on sleep duration and midday napping were obtained by a self-administered questionnaire. Multivariate logistic regression analyses were performed to estimate odds ratio (OR) and 95% confidence intervals (CIs) for the risk of AF.

Results: Compared with sleeping 6 to ≤ 8 hours/night, those reporting shorter sleep duration (≤ 5 hours/night) had a greater risk of AF (OR 1.95; 95% CI 1.28–2.95) in the fully adjusted model, while longer sleep (≥ 9 hours/night) was not significantly associated with the risk of AF. Compared with nonhabitual nappers, nappers had a higher risk of AF (OR 1.28; 95% CI 1.03–1.60) in the fully adjusted model. Moreover, we observed significant joint effects of sleeping ≤ 5 hours/night and nap (OR 2.13; 95% CI 1.09–4.14) on the risk of AF after adjusting for confounding factors.

Conclusion: Short sleep duration and midday napping were independently and jointly associated with higher risks of AF in patients with hypertension.

Keywords: sleep medicine, midday napping, atrial fibrillation, hypertension

Introduction

Atrial fibrillation (AF) is the most common sustained heart arrhythmia in adults and affects an estimated 43.6 million individuals worldwide.¹ In China, the prevalence of AF increased 20-fold over 11 years, and the currently estimated prevalence of AF is 0.2% in individuals aged ≥ 20 years.² AF is a major societal health problem related to the risk of stroke,³ heart failure,⁴ coronary artery disease,⁵ and chronic kidney disease,⁶ as well as morbidity and mortality.⁷ Identifying all the risk factors for AF is important for creating population-based strategies to reduce the risk of AF-related morbidity.⁸ However, currently known risk factors for AF explain only about one-half of the population attributable risk, emphasizing the need to identify the novel risk factors for AF, especially modifiable risk factors.^{9–11}

Up to now, the relationship between sleep duration and AF remains in dispute. Several studies have indicated that longer sleep duration was associated with incident AF.^{12,13} However, recent data from Pittsburgh Medical Center sleep laboratories showed that short sleep duration is an independent risk factor for incident AF.¹⁴ Moreover, a Mendelian randomization (MR) study has demonstrated that too little sleep may be associated with the risk of AF.¹⁵ Evidence from prior studies indicated that both short and long sleep duration as the risk factors for various health problems including metabolic syndrome,¹⁶ diabetes,¹⁷ obesity,¹⁸ hypertension,¹⁹ dyslipidemia²⁰ and myocardial infarction.²¹ Moreover, all these diseases were related with AF. In addition, most of the studies have assessed the relationship of sleep duration with incident AF regardless of midday napping,

which is common throughout China. In the meantime, little is known, about the effect of sleep duration on the incident AF among hypertensive patients. Hypertension as a risk factor that leads to AF,²² a better understanding of the sleep duration in hypertensive patients may reduce the huge burden of AF. Therefore, we designed this study to explore the association between sleep duration, midday napping and incident AF in patients with hypertension.

Methods

Participants and Study Design

The patients in this study came from China H-type Hypertension Registry Study that was reported previously.^{23,24} Briefly, this is a real-world, multicenter, observational registry study. From March 2018 to August 2018, a total of 14,234 hypertensive subjects were enrolled in this study in Wuyuan, Jiangxi province of China. Eligible participants were adults aged 18 years and older who had hypertension. Hypertension was determined as seated, resting systolic blood pressure (SBP) ≥ 140 mmHg or diastolic blood pressure (DBP) ≥ 90 mmHg at the screening, self-report history of hypertension, or the use of antihypertensive drug(s). The exclusion criteria included neurological abnormalities, inability to follow up according to the study protocol, or plans to relocate shortly, and the patients, who were not suitable for inclusion or for long-term follow-up assessed by study physicians. All participants provided written informed consent. The protocol was approved by the Ethics Committee of Institute of BioMedicine, Anhui Medical University (Registration number: ChiCTR1800017274).

2710 patients were excluded for missing data on sleep duration or midday napping. Thus, a final sample of 11,524 subjects was included in the analyses ([Figure S1](#)).

Assessment of Covariates

The information on demographic characteristics (age and gender), lifestyle factors (smoking, drinking, and physical activity, sleep quality), and medical history (diabetes mellitus, coronary heart disease (CHD), hyperlipidemia, stroke, duration of hypertension, and use of medications) were collected by trained researchers. The body mass index (BMI) was calculated as weight (kg)/height (m²). Blood pressure (BP) was measured by the automated electronic device (Omron; Dalian, China). After 10 minutes of rest, we calculated SBP and DBP as the average of the three BP measurements.

Fasting venous blood samples were obtained from all study patients. Next, the blood samples would be frozen and analyzed Biaojia Biotechnology Laboratory, Shenzhen, China. Fasting plasma glucose (FPG) and fasting lipids (total cholesterol, high-density lipoprotein-cholesterol (HDL-C), low-density lipoprotein cholesterol (LDL-C), and triglycerides) were determined using automatic clinical analyzers (Beckman Coulter). The formula for estimated glomerular filtration rate (eGFR) was used for the Chronic Kidney Disease Epidemiology Collaboration (CKD-EPI) equation.

Assessment of Sleep Behavior

Sleep duration and midday napping were assessed by a self-administered questionnaire. Sleep duration was assessed by the question: "How many hours on average do you sleep per night?" Mean sleep duration was categorized into 3 groups: ≤ 5 , $6 \leq 8$ and ≥ 9 hours, based on previous studies.^{25,26} Midday napping was assessed by asking "Did you have midday napping habit?" with the following responses: yes, no. Sleep quality was ascertained by the question: "How was your sleep quality at night?" with the following responses: good, fair, poor. Snoring status was obtained by asking "Do you snore when you sleep?" with the following responses: "no," "sometimes," and "frequently."

Assessment of Atrial Fibrillation

AF was defined as AF or atrial flutter captured on standard 12-lead electrocardiogram. Subjects who were not found to have AF according to the ECG test, but had previous medical records or any prior ECG record for AF episode were also defined as having AF. The diagnosis of AF based on the ECG was first conducted by a physician and then verified by the senior physician.⁸

Statistical Analyses

Continuous variables were presented as mean \pm standard deviation (SD) or median (interquartile range (IQR)) and compared using the *t*-test. Categorical variables were summarized as count (%) and analyzed by chi-square tests. Multivariate logistic regression analysis was used to evaluate the risk of AF by calculating the odds ratio (OR) and 95% confidence interval (CI) with adjustment for major covariates in four models. Model 1: crude model; Model 2: adjusted for age, gender, BMI, physical activity, smoking, drinking; Model 3: added SBP, DBP, duration of hypertension, diabetes mellitus, hyperlipidemia, CHD, antihypertensive agents, antidiabetes agents, lipid-lowering agents, FPG, triglyceride, HDL-C, LDL-C, and eGFR to model 2. Model 4: added snoring status, sleep quality, sleep duration and midday napping to model 3. In addition, possible modifications on the associations between sleep duration, midday napping and AF were also evaluated by stratified analyses and interaction testing. Moreover, we estimated the joint effects of sleep duration and midday napping on the risk of AF, and using the moderate sleep duration (6–8 hours) and nonhabitual nap as the reference group.

All data analyzed were using the statistical package R (<http://www.r-project.org>) and Empower (R) (www.empowerstats.com; X&Y Solutions, Inc., Boston, MA). A 2-tailed $P < 0.05$ was considered to be statistically significant.

Results

Characteristics of the Subjects

A total of 11,524 participants with hypertension were included in the final analysis (5542 men and 5982 women, aged 64.1 ± 9.8 years). The prevalence of AF was 3.4% (388/11,524). Baseline characteristics are shown in Table 1. Of the 11,524 subjects, 4.3%, 51.0% and 44.7% reported they sleep ≤ 5 , 6–8 and ≥ 9 hours/night, respectively and 53.1% reported midday napping (Table 1). Compare with participants reporting 6 to ≤ 8 hours/night, those reporting sleep duration ≤ 5 hours/night had higher values of age, stroke, hyperlipidemia, CHD, lower values of male, physical activity, smoking, drinking and eGFR. In addition, compare with nonhabitual nappers, nappers had higher values of male, age, smoking, drinking, stroke, hyperlipidemia, diabetes mellitus, hyperlipidemia and antidiabetes agents.

Association Between Sleep Duration, Midday Napping and AF

Table 2 shows the associations of sleep duration and midday napping with AF. Compared with sleeping 6 to ≤ 8 hours/night, those reporting shorter sleep duration (≤ 5 hours/night) had a greater risk of AF (aOR 1.95; 95% CI 1.28, 2.95) in the fully adjusted model, while longer sleep (≥ 9 hours/night) had no significant associated on the risk of AF (aOR 0.89; 95% CI 0.71, 1.12). We further investigated the association between midday napping with incident AF. Compare with nonhabitual nappers, nappers had a higher risk of AF (aOR 1.28; 95% CI 1.03, 1.60). Moreover, we explored the joint effects of sleep duration and midday napping on the risks of AF. Compared with those reporting moderate sleep duration (6 to ≤ 8 hours) and no napping, subjects who short sleep duration ≤ 5 hours combined with midday napping showed a significantly increased risk of AF (aOR 2.16, 95% CI 1.11–4.23), whereas no significant joint effect of sleeping ≥ 9 hours and no napping on AF was observed (Figure 1).

Subgroup Analysis

A predefined subgroup analysis on the associations between sleep duration, midday napping and the risk of AF was done in the following variables: sex, age, BMI, smoking, drinking, CHD, hyperlipidemia, and diabetes mellitus. The association of the risk of AF and short sleep duration was consistent in all subgroups except for CHD (P for interaction = 0.02) (Figure 2). Moreover, the risk of AF and midday napping was consistent in all stratification subgroups (all P for interaction >0.05) (Figure 3).

Discussion

In this large cross-sectional study of the middle-aged and older hypertensive population, we examined the association between sleep duration, midday napping and the risk of AF. The main findings were that both short sleep duration (≤ 5 hours) and midday napping were significantly associated with higher risk of AF among patients with hypertension.

Table 1 Baseline Characteristics of the Study Participants According to Sleep Duration and Midday Napping

	Sleep Duration, Hours/Night			Midday Napping	
	≤5	6 ≤ 8	≥9	No	Yes
Number (%)	496 (4.3)	5872 (51.0)	5156 (44.7)	5400 (46.9)	6124 (53.1)
Males, n (%)	198 (39.9)	2648 (45.1)	2696 (52.3)	2258 (41.8)	3284 (53.6)
Age, y	66.3 ± 9.0	62.8 ± 9.6	65.2 ± 9.8	63.8 ± 9.9	64.3 ± 9.6
BMI, kg/m ²	23.2 ± 3.5	23.7 ± 3.6	23.3 ± 3.5	23.4 ± 3.5	23.6 ± 3.5
Physical activity, n (%)					
Mild	299 (60.3)	3256 (55.4)	2959 (57.4)	3040 (56.3)	3474 (56.7)
Moderate	97 (19.6)	1412 (24.0)	1132 (22.0)	1236 (22.9)	1405 (22.9)
Vigorous	100 (20.2)	1204 (20.5)	1065 (20.7)	1124 (20.8)	1245 (20.3)
Smoking, n (%)	115 (23.2)	1524 (26.0)	1424 (27.6)	1291 (23.9)	1772 (28.9)
Drinking, n (%)	89 (17.9)	1197 (20.4)	1152 (22.3)	1000 (18.5)	1438 (23.5)
Duration of hypertension, y	6.0 (3.0–10.0)	6.0 (2.0–10.0)	6.0 (3.0–10.0)	5.5 (2.0–9.0)	6.0 (3.0–10.0)
SBP, mmHg	147.7 ± 17.1	146.5 ± 17.4	147.7 ± 17.9	147.4 ± 17.6	146.8 ± 17.6
DBP, mmHg	88.4 ± 11.0	88.9 ± 10.6	88.4 ± 11.2	88.8 ± 10.9	88.5 ± 10.8
Stroke, n (%)	40 (8.1)	393 (6.7)	430 (8.3)	375 (6.9)	488 (8.0)
Diabetes mellitus, n (%)	89 (17.9)	1046 (17.8)	994 (19.3)	915 (16.9)	1214 (19.8)
Hyperlipidemia, n (%)	94 (19.0)	1089 (18.5)	816 (15.8)	853 (15.8)	1146 (18.7)
CHD, n (%)	49 (9.9)	323 (5.5)	317 (6.1)	333 (6.2)	356 (5.8)
Antihypertensive agents, n (%)	301 (60.7)	3625 (61.7)	3236 (62.8)	3315 (61.4)	3847 (62.8)
Antidiabetes agents, n (%)	22 (4.4)	276 (4.7)	269 (5.2)	229 (4.2)	338 (5.5)
Lipid-lowering agents, n (%)	17 (3.4)	195 (3.3)	205 (4.0)	181 (3.4)	236 (3.9)
eGFR, mL/min/1.73 m ²	83.5 ± 19.5	87.4 ± 18.9	84.2 ± 20.5	86.2 ± 19.8	85.4 ± 19.7
FPG, mmol/L	6.1 ± 1.4	6.1 ± 1.5	6.2 ± 1.6	6.1 ± 1.4	6.2 ± 1.7
TG, mmol/L	1.4 (1.0–2.0)	1.5 (1.1–2.2)	1.4 (1.0–2.1)	1.5 (1.0–2.1)	1.5 (1.0–2.2)
HDL-C, mmol/L	1.6 ± 0.4	1.5 ± 0.4	1.5 ± 0.4	1.5 ± 0.4	1.5 ± 0.4
LDL-C, mmol/L	2.9 ± 0.7	3.0 ± 0.8	2.9 ± 0.8	2.9 ± 0.8	2.9 ± 0.8
Sleep quality, n (%)					
Good	41 (8.3)	3034 (51.7)	3628 (70.4)	334 (6.2)	162 (2.6)
Fair	41 (8.3)	1455 (24.8)	814 (15.8)	3339 (61.8)	2533 (41.4)
Poor	414 (83.5)	1383 (23.6)	714 (13.8)	1727 (32.0)	3429 (56.0)
Snoring status, n (%)					
No	312 (62.9)	3201 (54.5)	2819 (54.7)	3119 (57.8)	3213 (52.5)
Sometimes	129 (26.0)	2001 (34.1)	1709 (33.1)	1667 (30.9)	2172 (35.5)
Frequently	55 (11.1)	670 (11.4)	628 (12.2)	614 (11.4)	739 (12.1)

Moreover, we observed significant joint effects of sleep duration and midday napping on the risk of incident AF. Furthermore, the relationships remained statistically significant after adjusting for potential confounding variables.

Several studies have reported that short sleep duration was significantly associated with higher risk of AF. Genuardi et al conducted a large observational study referred for polysomnography consisted of 31,206 subjects found that short sleep duration was associated with a 2.01-fold increased risk for current AF and a 1.44-fold increased risk for future AF.¹⁴ A Mendelian Randomization Analysis also showed that short sleep duration was associated with the higher risk of incident AF.¹⁵ However, the present findings were inconsistent with that of the study by Song et al on 87,693 participants (mean age, 50.24 years) in China.¹³ They found that long sleep duration (≥8 hours) was associated with incident AF during median follow-up of 7.89 years; Nevertheless, the total number of incident AF (n=322) during follow-up was comparatively low. Moreover, the Multi-Ethnic Study of Atherosclerosis indicated that sleep duration was not associated with AF risk; however, the small number of events may have limited power.²⁷ Moreover, a longitudinal cohort study of 14 millions California residents found that insomnia was associated with a 30% to 42% higher risk of AF, supporting the concept that sleep disruption was an independent risk factor of incident AF.²⁸ It is of note that the risk of AF with short sleep duration appeared to be more obvious in patients with CHD. It is well known that coronary artery disease was an important risk

Table 2 Association of Sleep Duration and Midday Napping with Atrial Fibrillation

	N	Events	OR for Prevalent Atrial Fibrillation			
			Model 1	Model 2	Model 3	Model 4
Sleep duration (hours/night)						
≤5	460	35	2.65 (1.85, 3.82)	2.22 (1.54, 3.21)	1.95 (1.32, 2.87)	1.95 (1.28, 2.95)
6 ≥ ≤ 8	5668	157	1.00	1.00	1.00	1.00
≥9	4854	146	1.10 (0.89, 1.37)	0.95 (0.76, 1.18)	0.95 (0.76, 1.19)	0.89 (0.71, 1.12)
Midday napping						
No	5128	141	1.00	1.00	1.00	1.00
Yes	5854	197	1.19 (0.97, 1.46)	1.18 (0.96, 1.46)	1.28 (1.03, 1.48)	1.28 (1.03, 1.60)

Notes: Model 1: crude model. Model 2: adjusted for age, sex, BMI, physical activity, smoking, drinking. Model 3: adjusted for age, sex, BMI, physical activity, smoking, drinking, SBP, DBP, duration of hypertension, diabetes mellitus, hyperlipidemia, CHD, antihypertensive agents, antidiabetes agents, lipid-lowering agents, FPG, TG, HDL-C, LDL-C, and eGFR. Model 4: adjusted for age, sex, BMI, physical activity, smoking, drinking, SBP, DBP, duration of hypertension, diabetes mellitus, hyperlipidemia, CHD, antihypertensive agents, antidiabetes agents, lipid-lowering agents, FPG, TG, HDL-C, LDL-C, eGFR, snoring status, sleep duration, sleep quality and midday napping. Each group adjusted for the other covariates except itself.

factor of incident AF.⁸ Thus, it is reasonable to assume that CHD might amplify the detrimental effect of short sleep duration on AF.

Several possible mechanisms may explain the relationship between short sleep duration and AF. One important biological pathway is through oxidative stress and inflammation, as previous studies have demonstrated that sleep deprivation could lead to the increase of oxidative stress and the release of inflammatory factors, which could promote the occurrence of AF.^{29,30} Moreover, short sleep duration and low sleep efficiency were associated with an elevated level of biomarkers of autonomic tone, additional pathways predisposing to AF.³¹ A cross-sectional study showed that sleep deprivation can increase QT interval and electromechanical delay in young adults, which are linked to AF.³² Short sleep duration has been demonstrated to be associated with obesity, diabetes, metabolic syndrome and myocardial infarction, all these diseases are contributed to AF incidence.^{16,18,19,21}

We also found that midday napping was an independent risk factor for AF. To date, this is the first study to investigate the relationship between daytime napping and AF in hypertensive patients. Although the underlying mechanisms

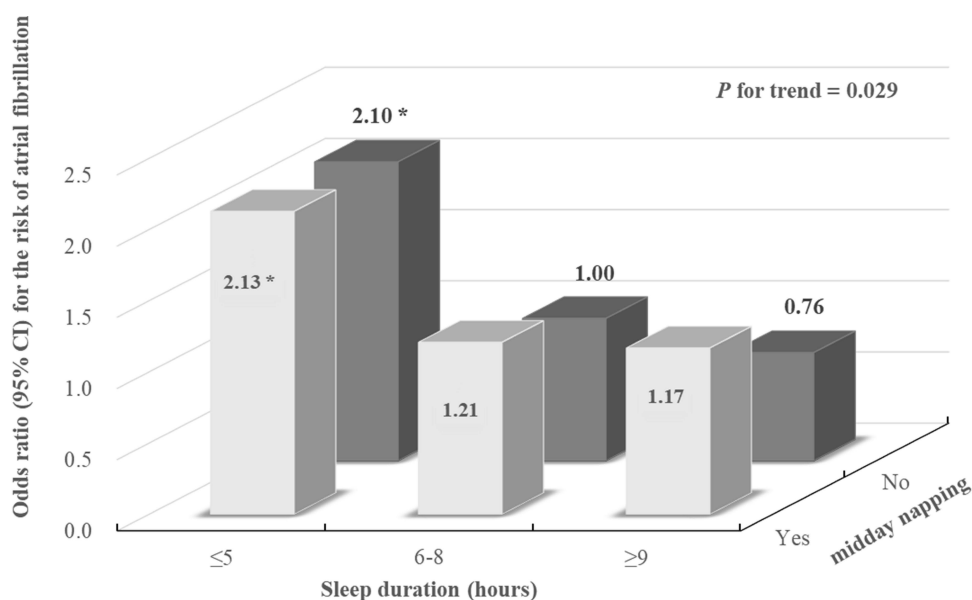


Figure 1 Joint effects of sleep duration and midday napping on incident AF risk. Each subgroup analysis adjusted, if not stratified, for age, sex, BMI, physical activity, smoking, drinking, SBP, DBP, duration of hypertension, diabetes mellitus, hyperlipidemia, CHD, antihypertensive agents, antidiabetes agents, lipid-lowering agents, FPG, TG, HDL-C, LDL-C, eGFR, snoring status and sleep quality (* $P < 0.05$).

Sleep duration (≤5 hours/night)

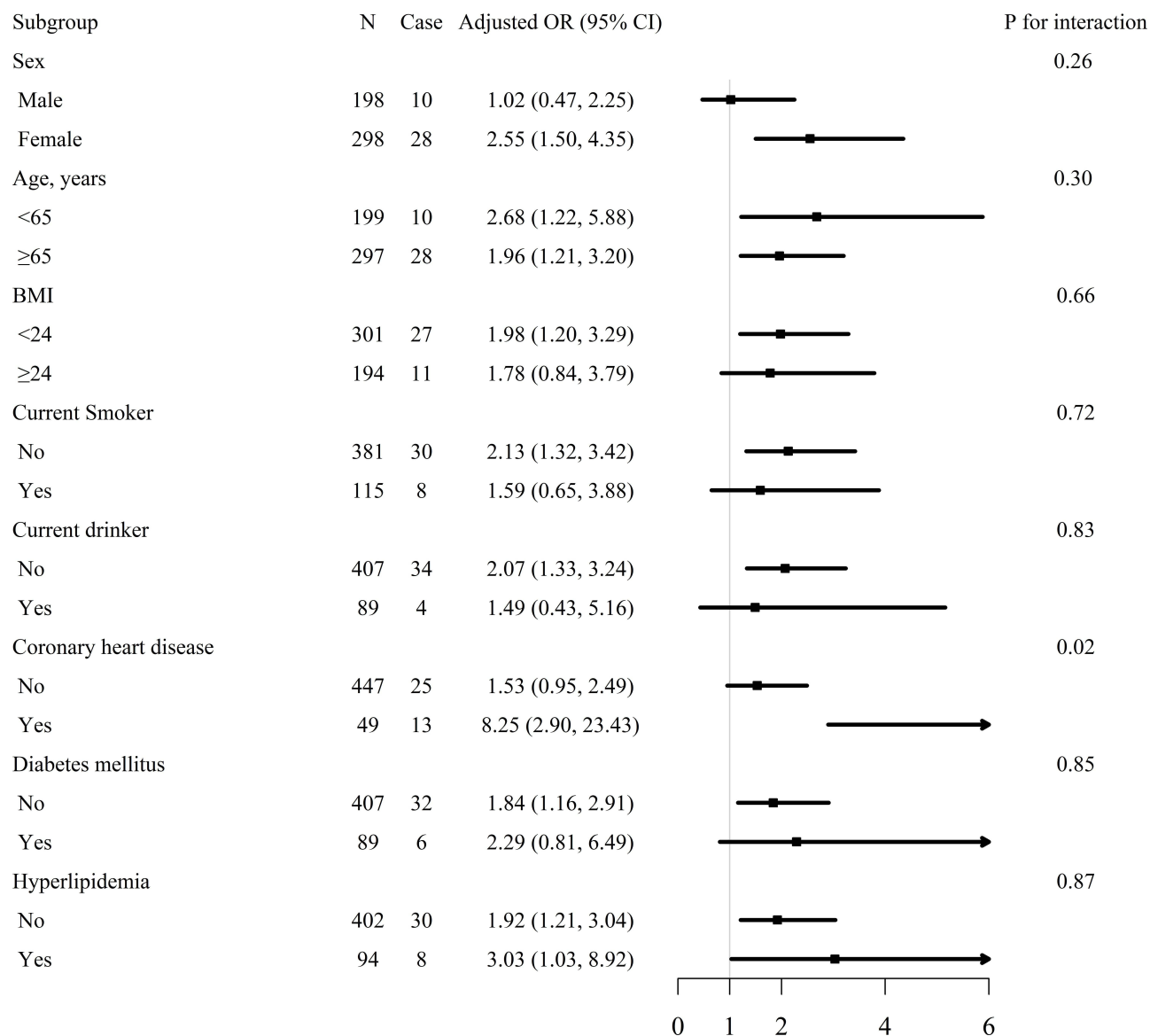


Figure 2 Subgroup analyses of the effect of sleep duration on incident AF risk. Each subgroup analysis adjusted, if not stratified, for age, sex, BMI, physical activity, smoking, drinking, SBP, DBP, duration of hypertension, diabetes mellitus, hyperlipidemia, CHD, antihypertensive agents, antidiabetes agents, lipid-lowering agents, FPG, TG, HDL-C, LDL-C, eGFR, snoring status, sleep quality and midday napping.

between daytime napping and AF are unclear, some epidemiologic and physiologic evidences might explain it. Midday napping is regarded as a risk factor for diabetes, CHD and metabolic syndrome,³⁴⁻³⁶ which are associated with the risk of AF. Midday napping, especially longer napping, might reflect an overall sedentary lifestyle, which is link to incident AF.³⁶

The strength of this study was the first to evaluate the relationship between sleep duration, midday napping and AF in a large hypertension cohort. Moreover, this study stressed the importance of moderate sleep duration for AF prevention in hypertensive patients. However, several potential limitations need to be addressed in this study. First, this study was a cross-sectional design, our results cannot provide a cause and effect association between short sleep duration, midday napping and the risk of AF. Although multivariate correction, it was difficult to exclude any potential confounding effect. Second, information on sleep duration and midday napping was collected from the questionnaire, which might overestimated actual sleep duration. However, it was unrealistic to obtain objective

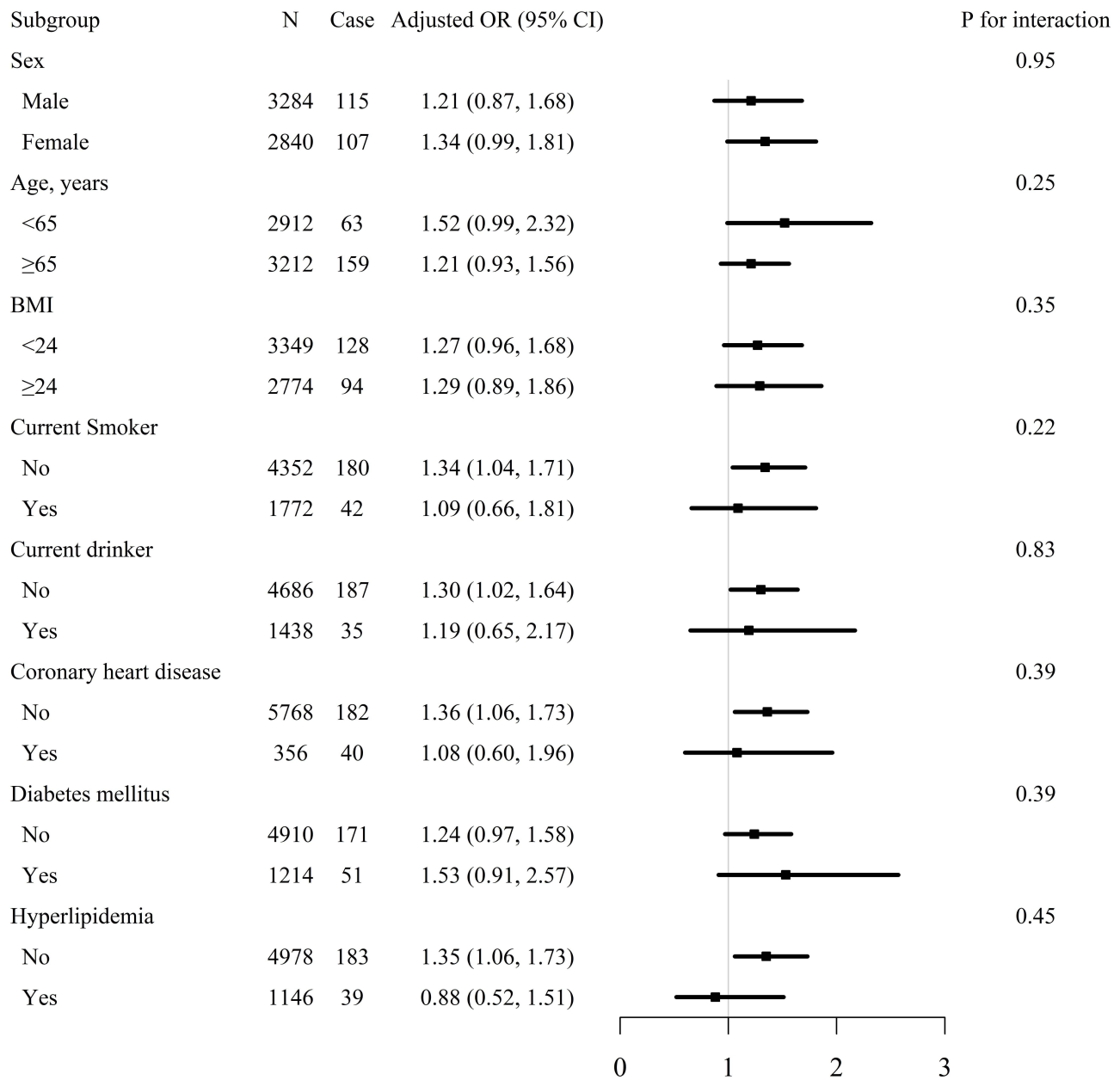
Midday napping (Yes)

Figure 3 Subgroup analyses of the effect of midday napping on incident AF risk. Each subgroup analysis adjusted, if not stratified, for age, sex, BMI, physical activity, smoking, drinking, SBP, DBP, duration of hypertension, diabetes mellitus, hyperlipidemia, CHD, antihypertensive agents, antidiabetes agents, lipid-lowering agents, FPG, TG, HDL-C, LDL-C, eGFR, snoring status and sleep duration.

measurements of sleep duration in a large sample population. And the method of obtaining sleep time and nap information through questionnaire has been reported in numerous studies.^{37,38} Third, we did not collect detailed information about nap time. Therefore, we could not perform regression analysis to clarify the association between nap duration and the risk of AF. Moreover, data about sleep apnoea was not collected, which may limit further exploration of whether this relationship could be modified by this factor. However, we collect the data about snoring, which is the the main symptom of sleep apnea. And we adjusted snoring status as a confounder in the statistical analysis. Fourth, a single assessment of sleep patterns at baseline was not satisfactory to reflect the dynamic change of sleep behaviors, which could lead to non-differential misclassification of sleep behaviors. Finally, AF was diagnosed based on electrocardiogram and medical record in this study, without ambulatory electrocardiogram

monitoring. Therefore, AF incidence can be underestimated due to electrocardiogram could not capture paroxysmal AF. However, inaccuracy of ascertainment of AF would also tend to bias findings toward the null hypothesis.

Conclusion

In this large cross-sectional study, we found that short sleep duration and midday napping were independently and jointly associated with higher risks of AF in patients with hypertension. Further research is needed to confirm our results in external populations and identify the potential mechanisms of the relationship.

Data Sharing Statement

The data are not publicly available due to privacy or ethical restrictions.

Ethics Approval and Consent to Participate

The study was conducted in accordance with the Declaration of Helsinki and was approved by the Institute of Biomedicine, Anhui Medical University, Hefei, China. Informed written consent was obtained from all patients before their enrollment in this study.

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Disclosure

The authors report no conflicts of interest in this work.

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