Association between sleep duration on workdays and blood pressure in non-overweight/obese population in NHANES: a public database research
Yingjie Su, Changluo Li, Yong Long, Liudang He & Ning Ding

This study aimed to explore the association between sleep duration on workdays and blood pressure (BP) including systolic blood pressure (SBP) and diastolic blood pressure (DBP) in non-overweight/obese population. A cross-sectional study composed of 2887 individuals from NHANES was conducted. Subjective sleep duration on workdays were evaluated by the questionnaire. Multiple linear regression analyses were done to explore the relationship between sleep duration and BP. Compared with sleep duration of 6–8 h, both sleep duration < 6 h and ≥ 8 h on workdays were significantly associated with increased SBP (β, 3.58 [95% CI 1.60, 5.56] and 1.70 [95% CI 0.76, 2.64], respectively). However, the significant association was not founded in DBP. The stratified analyses showed that in females, sleep duration (< 6 h or ≥ 8 h) on workdays were associated with SBP (β = 5.99 and 2.41, respectively, both \( P < 0.0005 \)). In addition, the SBP levels were much higher among participants aged (≥ 60) with sleep duration < 6 h. The effect size was 7.23 (\( P = 0.0217 \)). In the subgroup classified by race, a significantly positive association between sleep duration (< 6 h, ≥ 8 h) and SBP can be seen in the White population (β = 6.64 (\( P = 0.0007 \)) and 1.91 (\( P = 0.0215 \)), respectively). In non-overweight/obese population, both short sleep duration (< 6 h) and long sleep duration (≥ 8 h) on workdays were correlated with higher level of SBP.

Abbreviations
SBP  Systolic blood pressure
DBP  Diastolic blood pressure
BP  Blood pressure
TC  Total cholesterol
BMI  Body mass index
AST  Aspartate aminotransferase
HDL  High-density lipoprotein
CI  Confidence interval
OR  Odds ratio

Hypertension is a global health challenge, caused by the integration and accumulation of environmental and genetic risk factors. In the United States, approximately one-third of adults suffer from hypertension. Despite the fact that the public’s attention and treatment of hypertension has improved, hypertension no longer remains well-managed. According to recent literature, it has been calculated that during 2015, systolic blood pressure (SBP) of at least 110–115 mmHg was correlated with greater than 10 million deaths and over 211 million disability adjusted life year (DALYS).

In recent years, the adjustment of lifestyle as the main means to prevent hypertension has attracted people’s attention. Sleep, as a critical role in cardiovascular fitness, has been identified as a critical life-style risk element for cardiovascular disease. Sleep duration, particularly lack of sleep, could affect blood pressure (BP) through...
disturbances in autonomic and hormone balances, and also lead to increased obesity and metabolic disorders, and circadian rhythm disorders. Meta-analysis showed that short sleep time was associated with an increased risk of hypertension. Growing evidence has showed that BP is influenced by sleep duration, which is based on different age, genders, and races to explore the association.

However, few studies have been performed to investigate the relationship between sleep duration and BP in people with different body mass indexes (BMI). Previous research has identified that people who are overweight and obesity were both strong influence factors on BP. So far, few studies have been done to explore the possible relationship between sleep duration and elements of BP in people who are not overweight or obese. Hence, this study aimed to determine the association between sleep duration on workdays and BP in non-overweight/obese population.

Methods

Study population. The National Health and Nutrition Examination Survey (NHANES) is the only national survey that offers a cross-sectional view of nutrition and health in the United States population. It collects information about demographics, health and health behaviors. Data researchers and users can use the survey data of the NHANES on the Internet. Details statistics of NHANES can be found on www.cdc.gov/nchs/nhanes/. All methods in our research were performed in accordance with the Declaration of Helsinki.

According to WHO guidelines, BMI is divided into underweight (< 18.5 kg/m²), normal weight (18.5–24.99 kg/m²), overweight (25–29.99 kg/m²) and obesity (≥ 30 kg/m²). Non-overweight/obese is defined as people with BMI < 25. This research combined 2015–2018 data for analysis. A total of 19,225 potentially participants were enrolled, 16,338 participants were excluded for the following reasons: missing sleep duration data (n = 6818), missing BP data (n = 1055), taking antihypertensive medications (n = 2944), missing BMI and BMI ≥ 25 (n = 5521). Finally, 2887 participants were included in the study (Fig. 1).

Definition. Sleep duration on workdays was evaluated by the questionnaire with the following questions: “Number of hours usually sleep on weekdays or workdays.” Sleep duration was divided into three groups, which were < 6 h, 6–8 h, ≥ 8 h respectively, of which 6–8 h was used as the reference group.

The trained and certified examiners used the standardized protocols and calibrated equipment to get the blood pressure readings. Three consecutive BP readings were acquired via auscultatory means. If a BP measurement was not successfully completed, a fourth measurement was implemented. The average of all available measures was used.
Covariates. Race was divided into four groups: Mexican American, white, black and other race. Alcohol consumption was defined as the response to the question: “In the past 12 months, how often did you drink any type of alcoholic beverage?”, the responses was classified into three groups: drinking, no drinking, not recorded. Smoking was defined as the response to the question: “Do you now smoke cigarettes?”, the responses was classified into three groups: smoking, not smoking, not recorded. Diabetes was defined as the responses to the question: “Have you ever been told by a doctor or health professional that you have diabetes or sugar diabetes?”, the responses was classified into four group: yes, no, borderline, not recorded. Hypertension was defined as the response to the question: “Have you ever been told by a doctor or other health professional that you had hypertension, also called high blood pressure?”. The response was classified into three group: yes, no, and not recorded. The snort or stop breathing was defined as the response to the question: “In the past 12 months, how often did you snort, gasp, or stop breathing while you were asleep?”. The answers were classified into three group: yes, no, and not recorded. The method of obtaining other covariates, include gender, age, albumin, creatinine, hemoglobin, total cholesterol (TC), aspartate aminotransferase (AST), high-density lipoprotein (HDL), body mass index (BMI), can be found at www.cdc.gov/nchs/nhanes/. Among this covariates, age, albumin, creatinine, hemoglobin, TC, AST, HDL, BMI as continuous variables. Gender, alcohol consumption, diabetes, smoking, race, hypertension, snort or stop breathing as categorical variables.

Statistical analysis. All estimates were calculated accounting for NHANES sample weights. A weighted multiple linear regression model was used to assess the correlation between sleep duration on workdays and BP including systolic blood pressure(SBP) and diastolic blood pressure(DBP). The covariates mentioned above were adjusted as potential effect modifiers. The mean ± S.D and percentage were used to represent continuous and categorical variables, respectively. To calculate the differences between males and females, weighted linear regression models were used for continuous variables or weighted chi-square tests for categorical variables. The values of missing continuous covariates were indicated by dummy variables, including albumin, hemoglobin, creatinine and TC, AST, HDL and the missing ratios were 7.8%, 6.3%, 7.8%, 7.7%, 8.0%, 7.7% respectively. The missing categorical variables were included in the analysis as a single group. The statistical software packages R (http://www.R-project.org) and EmpowerStats (http://www.empowerstats.com) were used for the data analyses. When the P value was < 0.05, it was considered statistically significant.

Ethics approval and consent to participate. The ethics review board of the National Center for Health Statistics approved all NHANES protocols. Informed consent was obtained from all subjects and/or their legal guardian(s).

Results

Participant characteristics and univariate analysis. Table 1 showed the description of weighted sociodemographic and baseline characteristics. In the study, 2887 participants were subclassified based on gender. Among the participants, the proportion of males and females were 44.81% (n = 1378) and 55.19% (n = 1509), respectively. With ethnicity, the proportion of Mexican American, White and Black were 6.27%, 64.62% and 9.94%, respectively. Overall, the mean (SD) values for age, albumin, creatinine, SBP, DBP, hemoglobin, TC, AST, HDL, BMI were 38.54 (17.72) years, 43.48 (3.58) g/L, 73.21 (19.73) umol/L, 115.33 (14.56) mmHg, 68.47 (10.08) mmHg, 14.07 (1.44)/dL, 4.67 (1.01) mmol/L, 23.18 (13.01) IU/L, 1.59 (0.43) mmol/L, and 21.97 (2.06) kg/m², respectively. Among the participants, 66.01% were consumed alcohol drinkers, 2.61% were diabetics, 3.16% were hypertension, 6.72% have snort or stop breathing. 17.99% were smokers. Sleep duration was divided into three groups, which were < 6 h, 6–8 h, ≥ 8 h, each with a proportion of 6.26%, 40.53%, 53.21%, respectively. The univariate analysis of potential confounding factors is shown in Table 2.

Association between sleep duration on workdays and blood pressure. The results of multiple linear regression analysis used to explore the relationship between sleep duration and SBP were shown in Table 3. In the crude model, the sleep duration with 6–8 h was compared as the control group. Sleep duration < 6 h was significantly positively correlated with SBP (β, 6.15 [95% CI 3.88, 8.42]). However, a significant relationship was not found between sleep duration ≥ 8 h and SBP. After adjustment for gender, age, race (model I), we can observe a significantly positive association between sleep duration and SBP, the effect size of the group < 6 h and ≥ 8 h were (β, 4.17 [95% CI 2.19, 6.15]), (β, 1.55 [95% CI 0.60, 2.51]), respectively. Similarity, after controlling for all the potential confounding factors (model II), the relationship between the two was still present. The effect size of the group < 6 h and ≥ 8 h was (β, 3.58 [95% CI 1.60, 5.56]), (β, 1.70 [95% CI 0.76, 2.64]), respectively. In terms of DBP, the results of multiple linear regression analysis were illuminated in supplementary table 1. After controlling all the potential confounding factors, the significant association was not found in sleep duration < 6 h (β, 0.28 [95% CI −1.25, 1.82]) and ≥ 8 h (β, −0.41 [95% CI −1.34, 0.52]).

Subgroup analyses of factors influencing the association between sleep duration and SBP. In the subgroup analysis stratified by gender, age and race, the association between sleep duration and SBP was explored in Table 4. The positive effect was evident in most subgroups. All the potential confounding factors except the subgroup variable were adjusted. It showed that in females, sleep duration (6 h, ≥ 8 h) on workdays was associated with SBP (β = 5.99, 2.41, respectively, all P < 0.0005). Moreover, the association was much more obvious among participants aged (≥ 60) with sleep duration < 6 h. The effect size was 7.23 (P = 0.0217). In the subgroup classified by race, a significantly positive association was found in White whose sleep duration < 6 h or ≥ 8 h (β = 6.64 (P = 0.0007), and 1.91 (P = 0.0215), respectively). In others race, sleep duration (≥ 8 h) was associated with SBP (β = 2.06, P = 0.0097).
Discussion
The sleep duration of the general population has been affected by modern life, which also has been an important public health issue that has attracted the attention of researchers. Previous studies proved that sleep duration might also contribute to the increase in blood pressure. We found sleep duration in non-overweight/obese people positively correlated with SBP. In females, sleep duration < 6 h or ≥ 8 h on workdays were associated with SBP. In middle and old age, insufficient sleep duration (< 6 h) can lead to higher levels of SBP. In comparison with other ethnic groups, sleep duration < 6 h or ≥ 8 h was also associated with higher SBP among the White population.

A lot of cross-sectional and longitudinal epidemiological studies were used to explore the connection among sleep duration and hypertension. In the 2007–2009 National Healthy Interview Surveys (NHIS), compared with the 8-h group, adults who slept for less than 6 h or 6 h were more likely to develop hypertension (odds ratio (OR): 1.49 (1.34–1.64) and 1.15 (1.08–1.23), respectively). Several meta-analysis researches

| Total(n = 2887) | Male(n = 1378) | Female(n = 1509) | P value |
|----------------|---------------|-----------------|---------|
| Age(years)     | 38.54 ± 17.72 | 37.15 ± 17.89   | 39.68 ± 17.50 | 0.0001 |
| Albumin(g/L)   | 43.48 ± 3.58  | 44.57 ± 3.55    | 42.60 ± 3.35 | < 0.0001 |
| Creatinine(umol/L) | 73.21 ± 19.73 | 83.91 ± 15.74 | 64.56 ± 18.34 | < 0.0001 |
| SBP(mmHg)      | 115.33 ± 14.56 | 118.08 ± 13.48 | 113.10 ± 15.01 | < 0.0001 |
| DBP(mmHg)      | 68.47 ± 10.08  | 69.46 ± 10.68   | 67.66 ± 9.49 | < 0.0001 |
| Hemoglobin (g/dL) | 14.07 ± 1.44   | 15.04 ± 1.16    | 13.29 ± 1.14 | < 0.0001 |
| TC (mmol/L)    | 4.67 ± 1.01    | 4.51 ± 0.96     | 4.80 ± 1.03 | < 0.0001 |
| AST (IU/L)     | 23.18 ± 13.01  | 25.30 ± 15.67   | 21.47 ± 10.05 | < 0.0001 |
| HDL (mmol/L)   | 1.59 ± 0.43    | 1.43 ± 0.37     | 1.73 ± 0.43 | < 0.0001 |
| BMI (kg/m²)    | 21.97 ± 2.06   | 22.13 ± 2.06    | 21.85 ± 2.06 | 0.0004 |

Table 1. General characteristics of 2887 participants included in the present study. Mean +/- SD for: continuous variables. P value was calculated by weighted linear regression model. % for: categorical variables P value was calculated by weighted chi-square test. SBP systolic blood pressure, DBP diastolic blood pressure, TC total cholesterol, BMI body mass index, AST aspartate aminotransferase, HDL high-density lipoprotein.
Table 2. Univariate analysis for SBP and DBP. CI confidence interval, Ref reference, TC total cholesterol, BMI body mass index, AST aspartate aminotransferase, HDL high-density lipoprotein, SBP systolic blood pressure, DBP diastolic blood pressure.

| Covariate                | SBP (β, 95% CI, P) | DBP (β, 95% CI, P) |
|--------------------------|--------------------|--------------------|
| **Gender**               |                    |                    |
| Male                     | Ref                | Ref                |
| Female                   | −4.99 (−6.04, −3.93) < 0.0001 | −1.80 (−2.53, −1.06) < 0.0001 |
| Age(years)               | 0.37 (0.34, 0.40) < 0.0001 | 0.13 (0.11, 0.15) < 0.0001 |
| **Race**                 |                    |                    |
| Mexican American         | Ref                | Ref                |
| White                    | 2.47 (0.25, 4.68) 0.0291 | 1.93 (0.39, 3.46) 0.0142 |
| Black                    | 5.77 (3.07, 8.47) < 0.0001 | 2.07 (0.20, 3.95) 0.0303 |
| Others                   | 1.46 (−0.98, 3.89) 0.2415 | 2.40 (0.71, 4.09) 0.0055 |
| Albumin(g/L)             | −0.28 (−0.44, −0.13) 0.0002 | −0.11 (−0.22, −0.01) 0.0344 |
| Creatinine(umol/L)       | 0.12 (0.09, 0.14) < 0.0001 | 0.04 (0.02, 0.06) < 0.0001 |
| Hemoglobin (g/dL)        | 0.74 (0.36, 1.12) 0.0001 | 0.86 (0.60, 1.12) < 0.0001 |
| TC(mmol/L)               | 3.21 (2.68, 3.74) < 0.0001 | 1.71 (1.34, 2.08) < 0.0001 |
| AST(IU/L)                | 0.10 (0.06, 0.14) < 0.0001 | 0.04 (0.01, 0.07) 0.0060 |
| HDL(mmol/L)              | 1.81 (0.55, 3.07) 0.0048 | 1.03 (0.16, 1.91) 0.0209 |
| BMI (kg/m²)              | 1.03 (0.77, 1.28) < 0.0001 | 0.39 (0.22, 0.57) < 0.0001 |
| **Alcohol**              |                    |                    |
| No drinking              | Ref                | Ref                |
| Drinking                 | −5.00 (−6.85, −3.15) < 0.0001 | −1.24 (−2.52, 0.05) 0.0592 |
| Not recorded              | −7.89 (−9.92, −5.86) < 0.0001 | −4.66 (−6.06, −3.25) < 0.0001 |
| **Diabetes**             |                    |                    |
| Yes                      | Ref                | Ref                |
| No                       | −8.75 (−12.06, −5.43) < 0.0001 | −0.91 (−3.23, 1.40) 0.4833 |
| Borderline               | −3.99 (−9.51, 1.52) 0.1559 | 0.55 (−3.29, 4.39) 0.7787 |
| **Smoking**              |                    |                    |
| No smoking               | Ref                | Ref                |
| Not recorded              | −5.13 (−6.52, −3.75) < 0.0001 | −3.03 (−4.00, −2.06) < 0.0001 |
| **Hypertension**         |                    |                    |
| Yes                      | Ref                | Ref                |
| No                       | −16.43 (−19.46, −13.40) < 0.0001 | −10.14 (−12.24, −8.04) < 0.0001 |
| Not recorded              | −15.47 (−18.49, −12.44) < 0.0001 | −10.79 (−12.89, −8.69) < 0.0001 |
| **Snort or stop breathing** |        |                    |
| No                       | Ref                | Ref                |
| Yes                      | 4.61 (2.40, 6.81) < 0.0001 | 1.00 (−0.53, 2.52) 0.1998 |
| Not recorded              | 0.67 (−0.43, 1.77) 0.2345 | −1.21 (−1.98, −0.45) 0.0019 |

Table 3. Relationship between sleep duration and SBP in different models. Crude model adjust for: None; Model I adjust for: Gender; Age; Race; Model II adjust for: Gender; Age; Race; alcohol; Albumin; Creatinine; Hemoglobin; diabetes; hypertension; snort or stop breathing; smoke; TC; BMI; AST; HDL. CI confidence interval, Ref reference, TC total cholesterol, BMI body mass index, AST aspartate aminotransferase, HDL high-density lipoprotein, SBP systolic blood pressure.

| Exposure              | Crude model (β, 95% CI, P) | Model I (β, 95% CI, P) | Model II (β, 95% CI, P) |
|-----------------------|---------------------------|------------------------|-------------------------|
| **Sleep duration**    |                           |                        |                         |
| 6–8 h                 | Ref                       | Ref                    | Ref                     |
| < 6 h                 | 6.15 (3.88, 8.42) < 0.0001 | 4.17 (2.19, 6.15) < 0.0001 | 3.58 (1.66, 5.56) 0.0004 |
| ≥ 8 h                 | 0.11 (−1.00, 1.21) 0.8513  | 1.55 (0.60, 2.51) 0.0015 | 1.70 (0.76, 2.64) 0.0004 |
confirmed that short sleep duration (≤ 5 h or ≤ 6 h) was associated with hypertension, but there was no evidence of heterogeneity. However, few studies supported that short sleep duration had no impact on hypertension. A rhythm regulation study showed that short sleep duration will disrupt the 24-h sleep–wake cycle, which is an integrated process forming function light information to the brain through retinal ganglion cells, such as sleep–wake and circadian rhythm. This change will inevitably have a certain impact on blood pressure. Research have shown that when a person's sleep–wake cycle is inconsistent with the external environment, the average arterial pressure will increase by 3%. The inflammatory process may also play a vital role in the pathogenesis and pathophysiology of the relationship between short or long sleep duration and BP. Inflammatory factors such as c-reactive protein and interleukin-6 are increased with prolonged sleep duration, which can cause drowsiness and fatigue, and may also increase the risk of hypertension in people with long sleep duration. Another study showed that short sleep duration was associated with elevated C-reactive protein level. Long sleep duration was also associated with an increased risk of obesity, metabolic syndrome, and type 2 diabetes. The underlying mechanisms were that long sleep duration could disrupt circadian clocks and decrease insulin sensitivity, leading to unhealthy eating habits, decreased calorie consumption, and elevated systemic inflammation. Additionally, long sleep duration has been identified to be related to sleep fragments, which could activate sympathetic nervous system and lead to increased BP.

Our research can bring some inspiration to clinical work. For example, for non-overweight/obese female patients with borderline hypertension or poor blood pressure control, we can improve SBP by adjusting sleep time. We need to pay more attention to the impact of sleep duration on SBP among non-overweight/obese people, especially in regards to females and the elderly.

### Table 4. Effect size of sleep duration on SBP in each subgroup. Adjusted for: Gender; Age; Race; alcohol; Albumin; Creatinine; Hemoglobin; diabetes; hypertension; snort or stop breathing; smoke; TC; BMI; AST; HDL except the subgroup variable. CI confidence interval, Ref reference, TC total cholesterol, BMI body mass index, AST aspartate aminotransferase, HDL high-density lipoprotein.

| Characteristic | N  | 6–8 h (β, 95% CI, P) | <6 h (β, 95% CI, P) | ≥8 h (β, 95% CI, P) |
|---------------|----|---------------------|---------------------|---------------------|
| Gender        |    |                     |                     |                     |
| Male          | 1378 Ref | 1.22 (−1.31, 3.76) 0.3440 | 0.92 (−0.39, 2.33) 0.1685 |
| Female        | 1509 Ref | 5.99 (2.93, 9.04) 0.0001 | 2.41 (1.07, 3.75) 0.0004 |
| Age           |    |                     |                     |                     |
| <45           | 1884 Ref | 1.18 (−0.85, 3.20) 0.2543 | 0.74 (−0.16, 1.65) 0.1082 |
| ≥45, <60      | 495 Ref | 4.80 (−0.72, 10.32) 0.0890 | 0.21 (−2.44, 2.86) 0.8777 |
| ≥60           | 508 Ref | 7.23 (1.08, 13.38) 0.0217 | 1.86 (−1.60, 5.32) 0.2920 |
| Race          |    |                     |                     |                     |
| Mexican American | 302 Ref | 2.09 (−4.11, 8.30) 0.5096 | −0.38 (−2.87, 2.11) 0.7649 |
| White         | 968 Ref | 6.64 (2.82, 10.45) 0.0007 | 1.91 (0.28, 3.54) 0.0215 |
| Black         | 550 Ref | −0.76 (−4.12, 2.60) 0.6587 | 1.04 (−1.41, 3.49) 0.4063 |
| Others        | 1067 Ref | 2.95 (−0.24, 6.13) 0.0699 | 2.06 (0.50, 3.61) 0.0097 |

### Conclusions

In non-overweight/obese population, especially in females, both short sleep duration (<6 h) and long sleep duration (≥ 8 h) on workdays were correlated with higher levels of SBP. In old age populations, insufficient sleep duration (<6 h) was associated with higher levels of SBP. Compared with other ethnic groups, sleep duration (<6 h or ≥ 8 h) was also associated with higher SBP in Whites.
Data availability

The datasets used and/or analyzed during the present study were availed by the corresponding author on reasonable request.

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Author contributions

Conception and design: Y.S., N.D.; Administrative support: N.D.; Provision of study materials or patients: C.L., Y.L., L.H.; Collection and assembly of data: Y.S., N.D.; Data analysis and interpretation: Y.S., L.H.; Final approval of manuscript: All authors.
Competing interests
The authors declare no competing interests.

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