

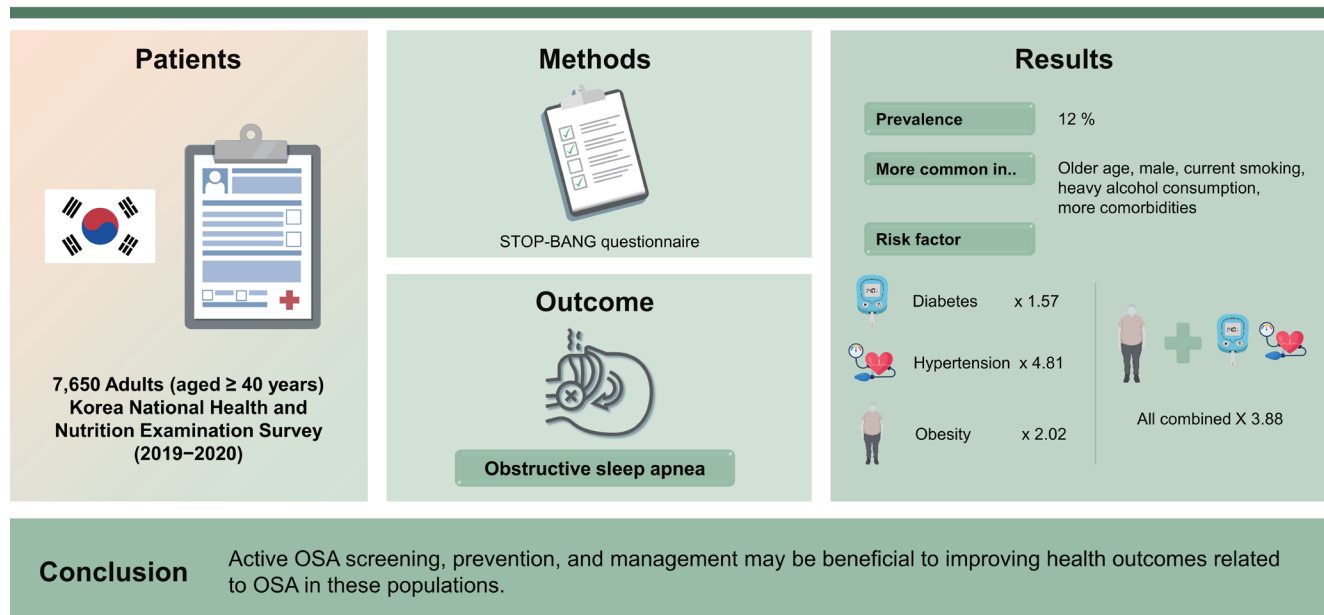


Comorbidities associated with high-risk obstructive sleep apnea based on the STOP-BANG questionnaire: a nationwide population-based study

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Background/Aims: We investigated the prevalence, associated factors, and comorbidities of high-risk obstructive sleep apnea (OSA) as determined by the STOP-BANG questionnaire in a Korean adult population.

Methods: Data were obtained from 7,650 adults (aged ≥ 40 years) who participated in the Korea National Health and Nutrition Examination Survey (2019–2020). Multiple logistic regression analyses were used to evaluate the association of socioeconomic status, lifestyle related factors, and comorbidities with high-risk OSA (STOP-BANG score, 5–8).

Results: The prevalence of high-risk OSA according to the STOP-BANG questionnaire was 12.0 %. Older age, male, current smoking, heavy alcohol consumption, and more comorbidities were associated with higher STOP-BANG scores. In multivariable adjusted analysis, diabetes mellitus (DM) (odds ratio [OR], 1.57; 95% confidence interval [CI], 1.25 to 1.97), hypertension (OR, 4.81; 95% CI, 3.88 to 5.97), and obesity (OR, 2.02; 95% CI, 1.60 to 2.56) were associated with high risk of OSA, and when hypertension, DM, and obesity were combined, the risk increased synergistically (OR, 3.88; 95% CI, 2.94 to 5.11).

Conclusions: According to the STOP-BANG questionnaire, the high risk of OSA was more common in males, and was associated with chronic metabolic disease, particularly in those with DM, hypertension, and obesity combined. Active OSA screening, prevention, and management may be beneficial to improving health outcomes related to OSA in these populations.

Keywords: Sleep apnea, obstructive; STOP-BANG; Hypertension; Diabetes mellitus; Obesity

INTRODUCTION

Obstructive sleep apnea (OSA) is characterized as a condition in which the upper airway repetitively collapses during sleep, completely or partially, resulting in frequent arousal and oxygen desaturation. According to previous reports, an estimated 425 million people aged 30 to 69 years have moderate to severe OSA, and up to 90% of OSA in the general adult population is misdiagnosed, and consequently untreated [1]. OSA is a growing health concern that has been related to a variety of comorbid conditions, including metabolic, cardiovascular (CV), renal, pulmonary, and neuropsychiatric disorders [2]. The lack of oxygen caused by OSA leads to increased health risks, such as CV disease, aortic aneurysm, high blood pressure, stroke, diabetes mellitus (DM), depressive illness, cognitive impairment, and even mortality, if not treated [3,4]. Given that undiagnosed OSA has major consequences in terms of public safety and all-cause mortality, focusing on the early detection and treatment of OSA is crucial.

Despite the crucial implications of untreated OSA, prompt diagnosis is often difficult. The gold standard for diagnosing OSA requires overnight polysomnography, which is time-consuming, expensive, and labor-intensive [5]. Furthermore, assessing OSA necessitates specialized personnel and equipment, which are rarely found in primary care clinics. As a result, to triage patients for early detection of OSA, a

simple and reliable screening method is necessary. Several screening methods have previously been developed, and their efficacy has been validated. However, the limitations of these methods include the duration, complications associated with the method, and the requirement of upper airway evaluation [6,7].

The STOP-BANG questionnaire is an OSA screening tool that includes four self-reportable (STOP: snoring, weariness, witnessed apnea, and elevated blood pressure) and four demographic (BANG: body mass index [BMI], age, neck circumference, and gender) questions [8]. In the original validation trial, the STOP-BANG questionnaire with a score of ≥ 3 had a sensitivity of (84, 93, and 100) % for detecting mild, moderate, and severe OSA, respectively. The STOP-BANG questionnaire is frequently employed, given its high diagnostic accuracy, ease of use, and clear risk stratification criteria. However, BMI and neck circumference, two STOP-BANG questionnaire questions, are impacted by region-specific body features, which may alter the STOP-BANG questionnaire performance in different geographic locations [9].

The objective of this study was to investigate the prevalence of high-risk OSA as determined by the STOP-BANG questionnaire using nationwide data collected from the 2019 to 2020 Korean National Health and Nutrition Survey (KNHANES). We further intended to investigate the clinical features and comorbidities associated with high-risk OSA.

METHODS

Study population

We used data from the KNHANES, a nationwide population-based cross-sectional survey. This survey was conducted in eight phases: KNHANES phases I (1998), II (2001), III (2005), IV (2007–2009), V (2010–2012), VI (2013–2015), VII (2016–2018), and VIII (2019–2021). Data from KNHANES VIII–1 and VIII–2 (2019–2020) conducted by the Korea Centers for Disease Control and Prevention were assessed in this study. Written informed consent was obtained from all participants prior to survey administration. The present study was exempted from review by the Institutional Review Board of the Catholic University of Korea (VC22ZISI0106), because it employs deidentified and publicly available data.

The KNHANES consists of several surveys about general health and nutritional status, health examinations, and laboratory investigations [10]. This survey used a stratified multistage probability sampling method to represent the Korean adult population, and the method considered geographic area, sex, and age group by referring to household registries. There were 15,469 participants from KNHANES VIII–1 and VIII–2. We excluded those under the age of 40 years (n = 6,173). Among them, 8,061 participants properly answered OSA-related questionnaires, such as the STOP-BANG questionnaire. We also excluded participants without information on other confounding factors, such as DM, hypertension, smoking, drinking, income, or education level. Finally, the data of 7,650 participants were included in the

analysis (Fig. 1).

Survey for OSA

Patients aged ≥ 40 years were asked about the risk factors associated with OSA using the STOP-BANG questionnaire, which contains four questions and four objective measures with yes/no answers: (1) Snoring: Do you snore loudly (louder than talking or loud enough to be heard through closed doors)?; (2) Tired: Do you often feel tired, fatigued, or sleepy during daytime?; (3) Observed: Has anyone observed you stop breathing during your sleep?; (4) Pressure: Do you have or are you being treated for high blood pressure?; (5) BMI > 30 kg/m²; (6) Age > 50 years old; (7) Neck circumference > 36.3 cm; and (8) Gender (if male) [11]. The total sum of “yes” responses was counted.

Clinical and laboratory measurements

Anthropometric, socioeconomic, health-related variables, and biochemical measurements were included in the analysis. BMI was calculated as weight in kilograms divided by height in square meters. BMI was categorized into five groups: underweight (< 18.5 kg/m²), normal weight (≥ 18.5, < 23 kg/m²), overweight (≥ 23, < 25 kg/m²), obese (≥ 25, < 30 kg/m²), and severely obese (≥ 30 kg/m²) according to the World Health Organization guidelines for the Asian-Pacific region [12]. Obesity was defined as subjects with BMI ≥ 25 kg/m². Waist circumference (WC) in the KNHANES was assessed by measuring at the midpoint between the lowest rib and the anterior iliac crest in the standing position. Ab-

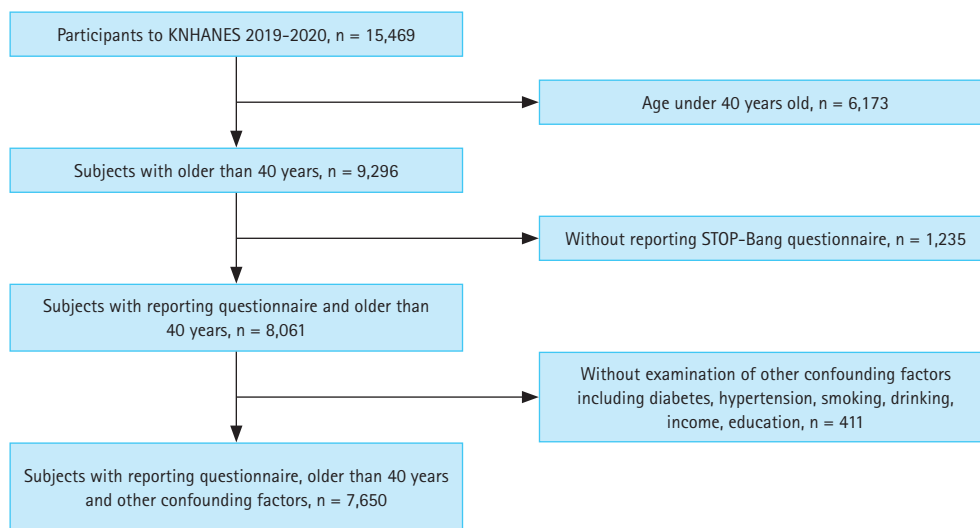


Figure 1. Flow chart presenting the selection of study participants. KNHANES, Korean National Health and Nutrition Survey.

dominal obesity was defined as WC \geq 90 cm in men, and \geq 85 cm in women, according to the Korean Society for the Study of Obesity [13].

Socioeconomic variables included gender, age, household income, educational level occupation (employed or unemployed), and residency area (urban or rural). Educational attainment was classified as elementary school or less, middle school, high school, and college or more. Residence area was defined as urban residence when residing in Seoul, Gyeonggi-do, or five metropolitan cities in Korea. The information on household income was obtained through the questionnaire, and stratified into five groups for each quintile. Household income was calculated as an equivalent income by dividing monthly income into the square root of family size.

Health-related variables included smoking and alcohol drinking status, physical activity, fasting plasma glucose (FPG), renal function, and lipid profiles. Smoking status was classified into three groups: never smoker, former smoker, and current smoker. Former smokers were distinguished from current smokers based on their present smoking status. Subjects who ceased smoking at the time of the survey were considered former smokers, regardless of the duration of smoking cessation. Alcohol drinking status was classified into three groups: never, mild-to-moderate, and heavy drinkers, according to daily alcohol consumption at the time of the survey. Never drinker was defined as individuals consuming on $<$ 1 occasion/month. Mild-to-moderate drinking was defined as individuals drinking $<$ 30 g of alcohol/day. A heavy drinker was defined as an individual drinking \geq 30 g of alcohol/day [14]. Physical activity was defined as performing moderate-intensity physical activity for at least 30 min/day at least 5 times/week or strenuous-intensity physical activity for at least 20 min/day at least 3 times/week [15].

For biochemical measurements, FPG and lipid profile levels were measured after overnight fasting using a Hitachi Autonomic Analyzer 7600-210 (Hitachi, Tokyo, Japan). Glycosylated hemoglobin (HbA1c) levels were measured by high-performance liquid chromatography on a Tosoh G8 device (Tosoh, Tokyo, Japan) [16]. The estimated glomerular filtration rate (eGFR) was calculated using the Modification of Renal Diet equation from baseline serum creatinine [17].

Primary outcome and assessment of covariates

The primary outcome was the prevalence of high risk of

OSA, and OSA risk was assessed using the STOP-BANG score as follows: (1) high risk of OSA: yes to 5–8 items; (2) intermediate risk of OSA: yes to 3–4 items; and (3) low risk of OSA: yes to 0–2 items [18].

The presence of comorbidities was determined by asking respondents if they were ever diagnosed with, or treated for, the following conditions: DM, hypertension, hypercholesterolemia, or chronic kidney disease (CKD). The diagnosis of DM was indicated if any of the following were present: FPG \geq 126 mg/dL, current use of antidiabetic medication, a previous history of DM, or HbA1c \geq 6.5% [19]. Hypertension was diagnosed if any of the following were present: systolic blood pressure \geq 140 mmHg or diastolic blood pressure \geq 90 mmHg, the current use of antihypertensive medications, or a self-reported physician diagnosis of hypertension [20]. Total cholesterol \geq 240 mg/dL, or use of cholesterol-lowering medications, were all used to diagnose hypercholesterolemia [19]. CKD was defined as eGFR $<$ 60 mL/min/1.73 m² [17].

Statistical analysis

General characteristics are presented as percentages (standard errors [SE]), and as the means and SE for quantitative variables. To perform group comparisons, the Rao-Scott chi-square test and analysis of variance (ANOVA) were used for categorical and continuous data, respectively. Both univariate and multiple logistic regression analyses were used to calculate odds ratios (ORs) and 95% confidence intervals (CIs) for evaluating factors associated with the OSA high-risk group (STOP-BANG score 5–8). In addition to the unadjusted model, the following confounding factors were considered in the multivariable regression model: age and sex for model 2; and age, sex, education level, household income quintile, smoking status, alcohol consumption, physical activity, DM, hypertension, hypercholesterolemia, CKD, obesity, and abdominal obesity for model 3. All statistical analyses were conducted using SAS version 9.4 for Windows (SAS Institute Inc., Cary, NC, USA). The *p* values provided are two-sided, with a level of significance of 0.05.

RESULTS

Clinical characteristics according to STOP-BANG score category

Table 1 demonstrates the clinical characteristics of the study

Table 1. Clinical characteristics of the study population according to the STOP-BANG score

Characteristic	STOP-BANG score			p value
	0–2	3–4	5–8	
No. of patients	3,863 (50.5)	2,868 (37.5)	919 (12.0)	
Age groups, % (SE)				< 0.001
40–49	40.4 (1.2)	22.2 (1.2)	16.7 (1.5)	
50–59	27.5 (0.9)	32.8 (1.2)	43.2 (2.1)	
60–69	18.6 (0.8)	25.5 (1.0)	26.4 (1.8)	
70–79	10.0 (0.6)	15.0 (0.8)	11.0 (1.0)	
≥ 80	3.5 (0.4)	4.5 (0.4)	2.7 (0.5)	
Male sex, % (SE)	17.0 (0.7)	74.6 (0.9)	92.0 (1.0)	< 0.001
Educational level, % (SE)				< 0.001
Elementary or lower	16.9 (0.8)	19.6 (1.0)	11.7 (1.1)	
Middle school	9.9 (0.6)	12.7 (0.8)	12.4 (1.3)	
High school	36.2 (1.1)	34.3 (1.2)	36.0 (1.9)	
College or higher	37.0 (1.3)	33.5 (1.5)	39.9 (2.1)	
Household income, % (SE)				< 0.001
Quintile 1 (lowest)	11.8 (0.7)	14.2 (1.0)	12.5 (1.2)	
Quintile 2	16.4 (0.8)	18.6 (1.0)	17.9 (1.5)	
Quintile 3	20.4 (0.9)	20.3 (1.0)	19.9 (1.6)	
Quintile 4	25.8 (0.9)	24.2 (1.2)	20.8 (1.5)	
Quintile 5 (highest)	25.6 (1.2)	22.8 (1.3)	29.0 (2.0)	
Region, urban, % (SE)	71.7 (1.8)	68.3 (1.9)	67.6 (2.4)	0.019
Occupation, yes, % (SE)	59.1 (1.1)	68.1 (1.1)	74.2 (1.7)	< 0.001
Smoking, % (SE)				< 0.001
Non	81.3 (0.7)	38.1 (1.1)	23.9 (1.6)	
Former	9.6 (0.5)	37.3 (1.1)	47.7 (1.9)	
Current	9.2 (0.6)	24.6 (1.0)	28.4 (1.7)	
Drinking, % (SE)				< 0.001
Non	35.6 (1.0)	27.0 (1.1)	20.8 (1.6)	
Mild to moderate	61.2 (1.0)	61.0 (1.1)	60.2 (1.9)	
Heavy	3.3 (0.4)	12.0 (0.8)	19.1 (1.5)	
Physical activity, % (SE)	39.5 (0.9)	41.0 (1.1)	40.5 (1.9)	0.543
BMI groups, kg/m ² , % (SE)				< 0.001
< 18.5	4.0 (0.4)	1.2 (0.2)	0.15 (0.1)	
≥ 18.5–< 23	47.8 (1.0)	24.5 (1.0)	10.3 (1.1)	
≥ 23–< 25	23.5 (0.8)	28.9 (1.1)	23.4 (1.6)	
≥ 25–< 30	23.5 (0.8)	39.5 (1.1)	46.7 (2.0)	
≥ 30	1.1 (0.2)	5.9 (0.5)	19.4 (1.5)	
Abdominal obesity, % (SE)	27.3 (0.8)	48.7 (1.0)	67.1 (1.7)	< 0.001
Diabetes mellitus, % (SE)	10.8 (0.6)	22.3 (1.0)	32.7 (2.0)	< 0.001
Hypertension, % (SE)	19.6 (0.8)	49.7 (1.2)	73.3 (1.8)	< 0.001
Hypercholesterolemia, % (SE)	27.3 (0.9)	32.6 (1.0)	38.8 (1.7)	< 0.001

Table 1. Continued

Characteristic	STOP-BANG score			p value
	0–2	3–4	5–8	
CKD, % (SE)	1.7 (0.2)	4.3 (0.4)	4.5 (0.8)	< 0.001
Age, yr	55.1 ± 0.3	58.9 ± 0.3	58.1 ± 0.4	< 0.001
Height, cm	159.6 ± 0.2	166.2 ± 0.2	169.0 ± 0.3	< 0.001
Weight, kg	58.9 ± 0.2	69.0 ± 0.3	76.8 ± 0.4	< 0.001
BMI, kg/m ²	23.1 ± 0.1	24.9 ± 0.2	26.9 ± 0.1	< 0.001
Waist circumference, cm	80.7 ± 0.2	88.8 ± 0.3	94.4 ± 0.3	< 0.001
Fasting glucose, mg/dL	99.3 ± 0.4	107.1 ± 0.6	112.1 ± 1.2	< 0.001
Systolic BP, mmHg	117.2 ± 0.4	124.7 ± 0.4	127.5 ± 0.6	< 0.001
Diastolic BP, mmHg	74.7 ± 0.2	78.5 ± 0.2	81.9 ± 0.4	< 0.001
Total cholesterol, mg/dL	198.8 ± 0.8	190.7 ± 0.9	189.5 ± 1.7	< 0.001
LDL-C, mg/dL	120.6 ± 0.7	113.3 ± 0.8	111.2 ± 1.5	< 0.001
HDL-C, mg/dL	54.9 ± 0.3	48.2 ± 0.3	46.5 ± 0.4	< 0.001
eGFR, mL/min/1.73 m ²	97.0 ± 0.4	90.8 ± 0.5	90.3 ± 0.8	< 0.001
Triglyceride, mg/dL	102.6 (100.6–104.7)	131.9 (128.2–135.7)	147.4 (140.9–154.2)	< 0.001

Values are presented as number (%), percentage (standard error), mean ± standard error, or geometric mean (95% confidence interval). Abdominal obesity was defined as waist circumference ≥ 90 cm in men, and ≥ 85 cm in women.

BMI, body mass index; CKD, chronic kidney disease; BP, blood pressure; LDL-C, low-density lipoprotein cholesterol; HDL-C, high-density lipoprotein cholesterol; eGFR, estimated glomerular filtration rate.

population according to the STOP-BANG score category. Among 7,650 total individuals, 919 subjects (12.0%) had a high risk of OSA (score 5–8). Compared to the intermediate- or low-risk group, the high-risk OSA group showed higher proportions of men, greater BMI, increased rates of smoking, heavy alcohol consumption, and abdominal obesity, and a higher proportion of employed responders. Regarding health-related factors, the high-risk OSA group had higher FPG, blood pressure, and triglyceride levels, and lower total cholesterol, high-density lipoprotein cholesterol, and eGFR levels. The high-risk STOP-BANG score group showed a significantly higher comorbidity prevalence. In the high-risk group, 67.1% had abdominal obesity, 32.7% DM, 73.3% hypertension, and 66.1% obesity, which was significantly higher, compared to the intermediate and low risk group. The clinical characteristics for males and females were both similar to the general population when they were analyzed separately (Supplementary Table 1).

In the STOP questionnaire, the prevalence of each item decreased in the following order: hypertension (38.0%), tired feeling (30.9%), snoring (20.3%), and observed stopping breathing during sleep (10.0%) (Table 2). In the BANG

questionnaire, the prevalence of each item decreased in the following order: age > 50 years (69.6 %), male sex (48.5 %), neck circumference > 36.6 cm (41.9 %), and BMI > 30 kg/m² (5.3 %).

Fig. 2 shows trends in the prevalence of a high risk of OSA categorized by the STOP-BANG scoring system by age and sex groups. When classified according to age group, the prevalence of high risk of OSA was highest in patients in their 50s (17.8%) and 60s (15.5%), and lowest in the patients in their 40s (7.2%) in the total (*p* for trend < 0.001) and male populations (*p* for trend < 0.001). However, in the female group, the prevalence of a high risk of OSA was highest in the patients in their 60s (3.1%) and 50s (2.9%), and lowest in those in their 40s (0.1%) (*p* for trend < 0.001). The prevalence was increased approximately 10-fold in males compared with females across all age categories.

Combined comorbidities and high risk of OSA

Those with high-risk OSA showed significantly higher proportions of individuals with comorbidities (DM, hypertension, hypercholesterolemia, CKD, obesity, and abdominal obesity) than the low- or intermediate-risk group (*p* < 0.001

Table 2. STOP-BANG score distribution according to questionnaire items

Variable	Total	STOP-BANG score										Score by risk group ^a		
		0	1	2	3	4	5	6	7	8	0-2	3-4	5-8	
No. of patients	7,650	551	1,444	1,868	1,634	1,234	608	235	67	9	3,863	2,868	919	
Percentage		7.2	18.9	24.4	21.4	16.1	7.9	3.1	0.9	0.1	50.5	37.5	12.0	
STOP														
Snore	20.3 (0.6)	0.0 (0.0)	1.9 (0.4)	6.7 (0.7)	19.5 (1.2)	30.4 (1.5)	61.4 (2.3)	87.6 (2.4)	100.0 (0.0)	100.0 (0.0)	3.8 (0.4)	24.2 (1.0)	71.5 (1.8)	
Tired	30.9 (0.7)	0.0 (0.0)	20.4 (1.3)	25.6 (1.3)	35.7 (1.4)	35.1 (1.7)	55.3 (2.4)	64.4 (3.6)	90.2 (3.8)	100.0 (0.0)	19.5 (0.8)	35.4 (1.1)	60.6 (1.9)	
Observed	10.0 (0.4)	0.0 (0.0)	0.6 (0.2)	1.4 (0.3)	5.2 (0.7)	12.6 (1.0)	35.5 (2.4)	69.2 (3.4)	96.7 (2.0)	100.0 (0.0)	0.8 (0.2)	8.4 (0.6)	49.4 (2.0)	
Pressure	38.0 (0.8)	0.0 (0.0)	3.8 (0.5)	39.3 (1.3)	39.7 (1.5)	63.0 (1.8)	68.9 (2.4)	77.2 (3.0)	95.4 (2.8)	100.0 (0.0)	19.6 (0.8)	49.7 (1.2)	73.3 (1.8)	
BANG														
BMI	5.3 (0.3)	0.0 (0.0)	0.5 (0.2)	2.0 (0.4)	4.3 (0.6)	8.1 (0.9)	15.1 (1.7)	23.2 (3.2)	34.9 (6.9)	100.0 (0.0)	1.1 (0.2)	5.9 (0.5)	19.4 (1.5)	
Age	69.6 (0.9)	0.0 (0.0)	65.8 (1.6)	75.2 (1.3)	73.9 (1.6)	83.1 (1.4)	82.2 (1.9)	84.4 (2.9)	86.5 (4.8)	100.0 (0.0)	59.6 (1.2)	77.9 (1.2)	83.3 (1.5)	
Neck	41.9 (0.6)	0.0 (0.0)	0.8 (0.3)	18.5 (1.2)	55.3 (1.5)	82.7 (1.2)	91.3 (1.3)	99.0 (0.7)	100.0 (0.0)	100.0 (0.0)	8.9 (0.6)	67.1 (1.1)	94.1 (0.9)	
Gender male	48.5 (0.5)	0.0 (0.0)	6.3 (0.8)	31.5 (1.4)	66.6 (1.3)	85.2 (1.1)	90.2 (1.4)	95.0 (1.6)	96.4 (2.2)	100.0 (0.0)	17.0 (0.7)	74.6 (0.9)	92.0 (1.0)	

Values are presented as percentage (standard error).

Key words represent following questions. Snore: Do you snore loudly (louder than talking or loud enough to be heard through closed doors)?; Tired: Do you often feel tired, fatigued, or sleepy during daytime?; Observed: Has anyone observed you stop breathing during your sleep?; Pressure: Do you have or are you being treated for high blood pressure?; BMI: Body mass index > 30 kg/m²?; Age: Age over 50 years old?; Neck: Neck circumference > 36.6 cm?; Gender: Gender male?;

^aRisk levels: 0-2, low risk of OSA; 3-4, intermediate risk of OSA; 5-8, high risk of OSA.

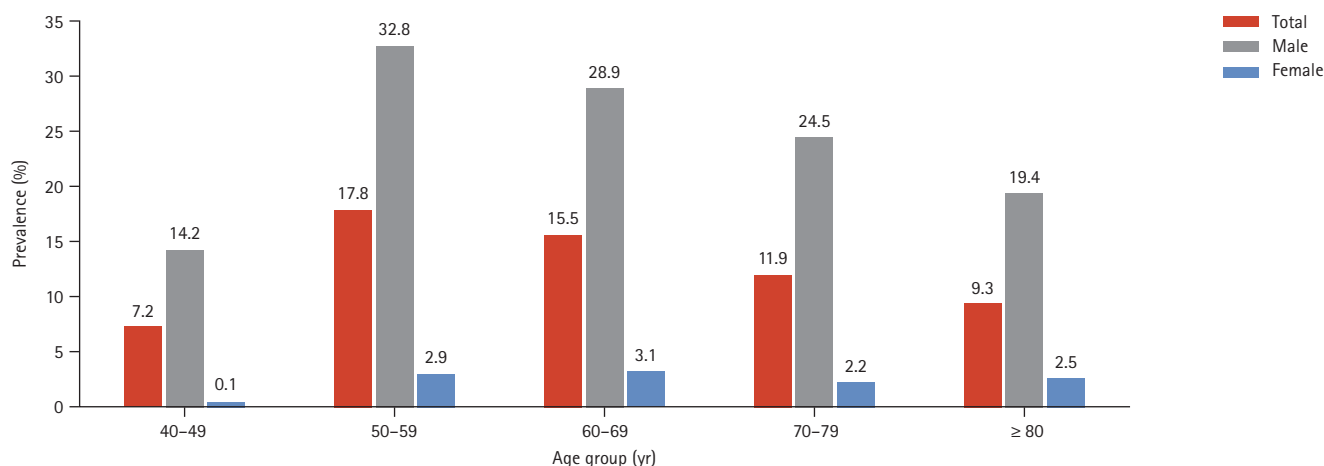


Figure 2. Prevalence of high risk of obstructive sleep apnea (OSA) using the STOP-BANG scoring system according to age and sex group. In both the total and male populations, the prevalence of high risk of OSA was highest in patients in their 50s, and lowest in patients in their 40s. Patients in their 60s had the largest female group, while those in their 40s had the lowest. The prevalence increased approximately 10-fold in males compared with females across all age categories. The chi-square test *p* for trend: total and male, < 0.001; female, < 0.001.

for all, except for CKD, $p = 0.264$) (Fig. 3). In particular, if the subjects had DM, hypertension, and obesity at the same time, the prevalence of high risk in the OSA population increased up to approximately four times, compared to those who did not have any of the comorbidities (42.3% vs. 11.1%, $p < 0.001$). The association was also apparent when the prevalence of high-risk OSA was obtained based on the measures of hypertension, diabetes, and obesity according to decile group (Supplementary Table 2). The prevalence tended to increase with higher measures of systolic/diastolic blood pressure, FPG, HbA1c, BMI, and WC.

In multivariable logistic regression analysis, the high risk of OSA was significantly associated with increasing age (OR, 1.01; 95% CI, 1.00 to 1.02), males (OR, 13.24; 95% CI, 9.34 to 18.77), and higher education (OR, 1.93; 95% CI, 1.34 to 2.77), after adjusting for multiple demographic, biochemical, and comorbidity confounders (Table 3). However, no association was noted between household income status, employment, smoking, alcohol consumption status, or physical activity, and higher STOP-BANG scores. The presence of comorbidities, such as DM (OR, 1.57; 95% CI, 1.25 to 1.97), hypertension (OR, 4.81; 95% CI, 3.88 to 5.97), obesity (OR, 2.02; 95% CI, 1.60 to 2.56), and abdominal obesity (OR, 1.61; 95% CI, 1.28 to 2.02) were significantly associated with a high risk of OSA. Participants who had DM, hypertension, and obesity altogether had almost 4-fold

high risk of OSA, compared to those with no comorbidities (OR, 3.88; 95% CI, 2.94 to 5.11). In terms of gender, the high risk of OSA was significantly associated with age, education level, and abdominal obesity only in males. In contrast, DM, hypertension, and obesity were all associated with the high risk of OSA in both males and females (Supplementary Table 3).

DISCUSSION

In this recently investigated, large nationwide cross-sectional study, we found that comorbidities, such as DM, hypertension, and obesity, were significantly associated with a high risk of OSA, based on the STOP-BANG questionnaire in a Korean adult population. Notably, these three comorbidities had a synergistic impact in elevating OSA risk. The STOP-BANG scoring questionnaire was first introduced in the KNHANES VIII in 2019. Hence, this is the first nationwide study on the association of chronic metabolic disease with a high risk of OSA in the representative Korean population utilizing the STOP-BANG questionnaire. According to the results of our study, screening for OSA should be more actively recommended for patients with DM, hypertension, and obesity, particularly males.

The STOP-BANG questionnaire is well recognized as a reli-

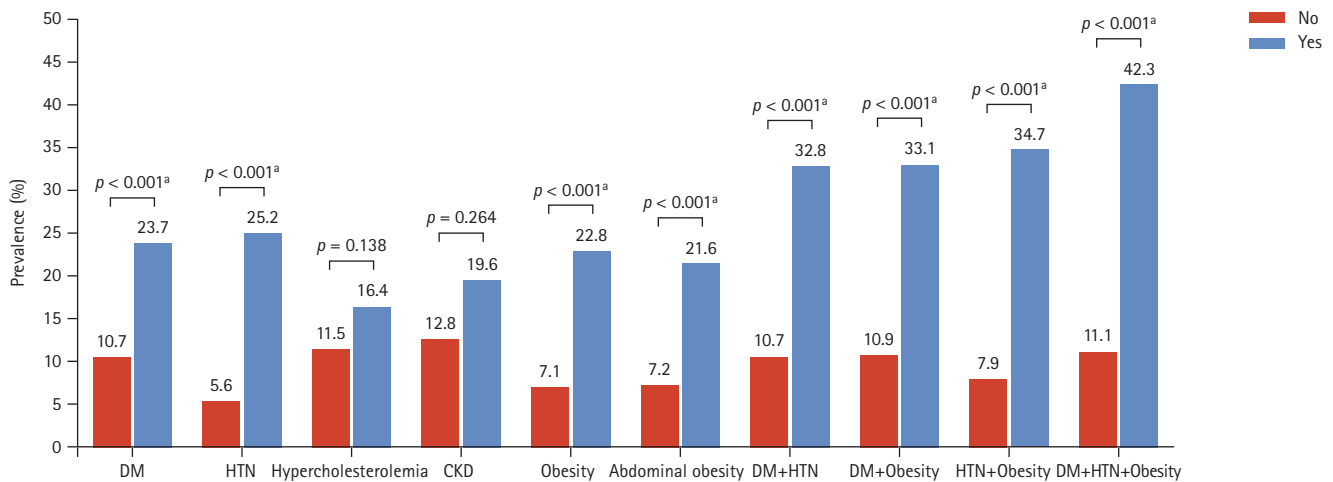


Figure 3. Prevalence of high risk of obstructive sleep apnea (OSA) using the STOP-BANG scoring system according to comorbidities. Diabetes mellitus (DM), hypertension (HTN), obesity, and abdominal obesity were significantly associated with a high risk of OSA. Diabetes, hypertension, and obesity show a synergistic effect on the prevalence of high-risk OSA. When the three comorbidities were combined, the odds of having a high risk of OSA increased nearly 4-fold, compared to people who had no comorbidities. Abdominal obesity was defined as waist circumference ≥ 90 cm in men, and ≥ 85 cm in women. Obesity was defined as subjects with body mass index ≥ 25 kg/m². Red bar: prevalence without comorbidity, blue bar: prevalence with comorbidity. CKD, chronic kidney disease. ^a $p < 0.05$.

Table 3. Multivariate logistic regression analysis for the high-risk group of obstructive sleep apnea

Variable	Model 1		Model 2		Model 3	
	OR (95% CI)	p value	OR (95% CI)	p value	OR (95% CI)	p value
Age	1.01 (1.00–1.02)	< 0.001	1.02 (1.01–1.03)	< 0.001	1.01 (1.00–1.02)	0.012
Male sex	15.94 (12.04–21.08)	< 0.001	16.42 (12.43–21.70)	< 0.001	13.24 (9.34–18.77)	< 0.001
Education		< 0.001		0.077		0.003
Elementary or lower	1 (ref)		1 (ref)		1 (ref)	
Middle	1.72 (1.27–2.32)		1.38 (0.99–1.93)		1.41 (0.98–2.03)	
High	1.57 (1.23–2.00)		1.45 (1.08–1.95)		1.82 (1.30–2.55)	
College	1.73 (1.37–2.20)		1.45 (1.07–1.96)		1.93 (1.34–2.77)	
Household income		0.037		0.126		0.208
Q1	1 (ref)		1 (ref)		1 (ref)	
Q2	1.05 (0.81–1.37)		1.04 (0.76–1.40)		0.93 (0.66–1.33)	
Q3	1.00 (0.76–1.30)		0.99 (0.73–1.35)		0.84 (0.59–1.20)	
Q4	0.85 (0.66–1.09)		0.83 (0.61–1.12)		0.75 (0.52–1.09)	
Q5	1.21 (0.95–1.55)		1.16 (0.85–1.59)		1.02 (0.69–1.51)	
Occupation, no	1 (ref)	< 0.001	1 (ref)	0.334	1 (ref)	0.330
Yes	1.69 (1.42–2.00)		1.11 (0.90–1.39)		1.13 (0.88–1.46)	
Smoking		< 0.001		0.042		0.540
Non	1 (ref)		1 (ref)		1 (ref)	
Former	5.80 (4.79–7.02)		1.33 (1.07–1.67)		1.16 (0.89–1.50)	
Current	4.69 (3.79–5.81)		1.19 (0.93–1.52)		1.13 (0.86–1.49)	
Alcohol consumption				< 0.001		0.272
Non	1 (ref)	< 0.001	1 (ref)		1 (ref)	
Mild to moderate	1.51 (1.24–1.84)		1.07 (0.86–1.340)		1.06 (0.80–1.39)	
Heavy	4.15 (3.19–5.41)		1.67 (1.247–2.24)		1.27 (0.91–1.76)	
Physical activity, no	1 (ref)	0.841	1 (ref)	0.423	1 (ref)	0.481
Yes	1.02 (0.86–1.20)		0.93 (0.78–1.11)		0.93 (0.76–1.14)	
DM, no	1 (ref)	< 0.001	1 (ref)	< 0.001	1 (ref)	< 0.001
Yes	2.60 (2.14–3.15)		2.27 (1.83–2.81)		1.57 (1.25–1.97)	
HTN, no	1 (ref)	< 0.001	1 (ref)	< 0.001	1 (ref)	< 0.001
Yes	5.67 (4.71–6.82)		6.13 (5.01–7.49)		4.81 (3.88–5.97)	
Hypercholesterolemia, no	1 (ref)	< 0.001	1 (ref)	< 0.001	1 (ref)	0.138
Yes	1.51 (1.29–1.76)		1.86 (1.56–2.21)		1.18 (0.95–1.45)	
CKD, no	1 (ref)	0.011	1 (ref)	0.313	1 (ref)	0.264
Yes	1.65 (1.12–2.43)		1.26 (0.81–1.96)		0.77 (0.49–1.22)	
Obesity, no	1 (ref)	< 0.001	1 (ref)	< 0.001	1 (ref)	< 0.001
Yes	3.86 (3.27–4.56)		3.66 (3.06–4.38)		2.02 (1.60–2.56)	
Abdominal obesity, no	1 (ref)	< 0.001	1 (ref)	< 0.001	1 (ref)	< 0.001
Yes	3.55 (3.02–4.17)		3.35 (2.82–3.99)		1.61 (1.28–2.02)	
BMI, kg/m ²		< 0.001		< 0.001		< 0.001
< 18.5	0.20 (0.05–0.85)		0.17 (0.04–0.72)		0.17 (0.04–0.78)	
≥ 18.5–< 23	1 (ref)		1 (ref)			

Table 3. Continued

Variable	Model 1		Model 2		Model 3	
	OR (95% CI)	<i>p</i> value	OR (95% CI)	<i>p</i> value	OR (95% CI)	<i>p</i> value
≥ 23–< 25	3.30 (2.50–4.36)		2.59 (1.94–3.47)		2.19 (1.60–3.00)	
≥ 25–< 30	5.61 (4.32–7.27)		4.49 (3.42–5.90)		3.20 (2.37–4.33)	
≥ 30–< 35	22.15 (15.66–31.31)		47.53 (30.06–75.14)		31.33 (19.40–50.58)	
≥ 35	23.47 (10.87–50.66)		74.16 (21.81–252.23)		38.44 (12.26–120.51)	
Obesity group		< 0.001		< 0.001		< 0.001
Low BMI, low WC	1 (ref)		1 (ref)		1 (ref)	
Low BMI, high WC	1.79 (1.29–2.47)		1.51 (1.08–2.11)		1.21 (0.86–1.72)	
High BMI, low WC	2.29 (1.62–3.25)		1.86 (1.30–2.66)		1.48 (1.01–2.17)	
High BMI, high WC	4.74 (3.95–5.68)		4.48 (3.67–5.46)		3.19 (2.56–3.97)	
DM + HTN + obesity	1 (ref)	< 0.001	1 (ref)	< 0.001	1 (ref)	< 0.001
Yes	5.86 (4.67–7.35)		6.26 (4.83–8.10)		3.88 (2.94–5.11)	

Model 1 was unadjusted; Model 2 was adjusted for age and sex; Model 3 was adjusted for age, sex, education, household income, occupation, smoking, drinking, physical activity, DM, hypertension, hypercholesterolemia, CKD, obesity, and abdominal obesity (weight circumference: men ≥ 90 cm/women 85 cm). Abdominal obesity was defined as WC ≥ 90 cm in men, and ≥ 85 cm in women. Low and high WC was determined by this standard. Obesity was defined as subjects with BMI ≥ 25 kg/m².

OR, odds ratio; CI, confidence interval; DM, diabetes mellitus; HTN, hypertension; CKD, chronic kidney disease; BMI, body mass index; WC, waist circumference.

able OSA screening tool that is both simple and self-reportable [8]. The validity of this method is evidenced in several prior systemic reviews. A recent meta-analysis confirmed that the STOP-BANG questionnaire can be used as a screening tool to help triage individuals with suspected OSA [9]. Another previous meta-analysis validated the strong performance of the STOP-BANG questionnaire for OSA screening [21]. The correlation between STOP-BANG scores and the probability and severity of OSA is also well indicated. A STOP-BANG score of (5–8) indicated individuals with a high chance of moderate/severe OSA in recent prospective research using a surgical population [18]. Another previous study also classified patients with a STOP-BANG score of (0–2) as low risk for moderate to severe OSA, whereas those with a score of (5–8) were classified as high risk for moderate to severe OSA [22]. This classification was utilized in our study to evaluate clinical features according to the severity of OSA.

Several studies employing the STOP-BANG questionnaire, including the Asian population, have been reported. According to earlier systematic reviews, the prevalence of moderate-to-severe OSA was 21.3%, whereas that of severe OSA was 7.8% [23]. In a study of the Singapore population, the prevalence of moderate-to-severe OSA was

28.1%, and that of severe OSA was 10.7% [24]. Other questionnaire-based studies reported that the prevalence of high-risk OSA ranged (12.4 to 15.8) % in the Korean adult population; however, those studies of the Korean population utilized the Berlin questionnaire [25,26]. Altogether, the results of our study seem to be consistent with previous data regarding the adult population.

Our study presents a remarkably higher prevalence of high-risk OSA in males, compared with females. The male predominance in high-risk OSA was frequently presented in previous epidemiologic studies, and male sex is a widely accepted risk factor for OSA [27]. The 10-fold predominance of high-risk OSA in males correlates with an old renowned study conducted in 1979 [28]. However, recent studies show a rather mild male predominance, compared to our study [26]. This discrepancy may be explained by a more recent study that indicated an unacceptably low sensitivity of the STOP-BANG questionnaire in females, compared to males [29]. Due to the bias toward males, sex-specific cutoffs for STOP-BANG should be carefully investigated. Although the predominance of male in high-risk OSA is apparent in our study, the clinical characteristics among males and females were not largely different, and the tendency was comparable to the general population.

Our study also presents that socioeconomic factors, such as education level, household income, region of residence, and occupation status, were associated with high risk of OSA. The hypothesis that lower socioeconomic status is linked to poor sleep quality and increased prevalence of OSA is supported by various research [30,31], and a significant higher risk for OSA was observed in association with less education and lower economic income [32]. Common comprehensive measures of socioeconomic status include household income, location of residence, occupation status, and basic education or education-derived variables [25]. Although there is no conclusive evidence on the relationship between socioeconomic status and the risk of OSA, it is widely accepted that socioeconomic status has an impact on a range of health practices, biomarkers, and chronic diseases [30]. Therefore, our results support the notion that composite indicators of socioeconomic status, such as household income, area of residence, occupation status, and education attainment, are associated with the risk of OSA.

The most significant finding of our study is the association of a higher STOP-BANG score with a higher comorbidity prevalence, as well as a synergistic effect, when all three comorbidities are present. According to our results, hypertension, DM, and obesity were significantly associated with high-risk OSA, and the prevalence of high-risk OSA consistently increased with higher measures of hypertension, DM, and obesity. OSA has long been understood to play a role in CV disorders, particularly hypertension, heart failure, and stroke [3]. OSA is further exacerbated by an increase in body weight. According to studies, gaining 10% of one's body weight increases the risk of OSA by six times [33]. Obesity-related fat deposition in the neck can obstruct the pharyngeal lumen, causing it to collapse when sleeping [34]. OSA is also known to cause arterial stiffness, which can lead to a CV disorder, and OSA therapy can help to reduce arterial stiffness [35]. Furthermore, sleep breathing disorders may increase the risk of DM through affecting plasma insulin levels and glycemia [36]. Thus, it is possible to deduce that OSA and comorbidities, such as hypertension, DM, and obesity, interact, and have an impact on one another. According to our results, OSA should be screened and detected early in patients with DM, hypertension, or obesity, especially if all three are present. The health impact of OSA in certain chronic metabolic illnesses could be decreased by using this validated short questionnaire, and the importance

of lifestyle modification and active treatment of coexisting diseases should be emphasized for those individuals.

Our study has several limitations. First, the subjects were not clinically confirmed to have OSA using polysomnography. The risk stratification was exclusively based on the participant's self-reporting questionnaire. The fact that our study's prevalence of high-risk OSA was comparable to that of prior population-based studies implies that our sampling approach was rather reliable. Second, given that our sample excluded patients aged less than 40 years, the biased sample may have affected the true clinical significance of our data. However, previous data show that utilization of health care is only substantial in OSA patients over the age of 40 years, which supports the KHANES policy of only administering the STOP-BANG questionnaire to those over the age of 40 years [37,38]. As a result, in terms of social welfare, populations above the age of 40 years should be prioritized for the early identification of OSA. Finally, this study was inevitably performed with a cross-sectional design, given the retrospective analysis of the established dataset. Thus, causal associations between high-risk OSA and other comorbidities, or the use of antihyperglycemic, antihyperlipidemic, and antihypertensive medications, cannot be completely clarified. However, this research has advantages in retrieving data from a nationally representative survey sample with a high response rate; therefore, a large amount of data was provided to overcome potential confounding issues. Furthermore, our data were obtained from a large sample, which improved the precision of our findings, and allowed for numerous statistical adjustments. This study is the first to report the prevalence of high-risk groups of OSA based on the STOP-BANG questionnaire in a Korean population, and provides the clinical implication of a high-risk OSA group.

In conclusion, we present for the first time that based on the STOP-BANG questionnaire, the prevalence of high-risk OSA groups in the general Korean population was 12.0%. OSA must be screened and detected early in patients with DM, hypertension, and obesity. The health burden of OSA in chronic metabolic disease could be reduced using this simple and validated short questionnaire. In the future, the exact prevalence of OSA needs to be diagnosed by a confirmed method in this high-risk group of OSA in the Korean population.

KEY MESSAGE

1. The STOP-BANG questionnaire revealed that the prevalence of a high risk of obstructive sleep apnea (OSA) was 12.0% in the Korean population over age 40 years.
2. A high risk of OSA was associated with socioeconomic (older age, male gender, current smoker, heavy alcohol drinker) and chronic metabolic disease, particularly in those with diabetes mellitus, hypertension, or obesity.
3. Active OSA screening, prevention, and management may improve health outcomes related to OSA in patients with diabetes mellitus, hypertension, or obesity.

Conflict of interest

No potential conflict of interest relevant to this article was reported.

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REFERENCES

1. Benjafield AV, Ayas NT, Eastwood PR, et al. Estimation of the global prevalence and burden of obstructive sleep apnoea: a literature-based analysis. *Lancet Respir Med* 2019;7:687-698.
2. McNicholas WT. Obstructive sleep apnoea and comorbidity: an overview of the association and impact of continuous positive airway pressure therapy. *Expert Rev Respir Med* 2019;13:251-261.
3. Bradley TD, Floras JS. Obstructive sleep apnoea and its cardiovascular consequences. *Lancet* 2009;373:82-93.
4. Kerner NA, Roose SP. Obstructive sleep apnea is linked to depression and cognitive impairment: evidence and potential mechanisms. *Am J Geriatr Psychiatry* 2016;24:496-508.
5. Phua CQ, Jang IJ, Tan KB, et al. Reducing cost and time to diagnosis and treatment of obstructive sleep apnea using ambulatory sleep study: a Singapore sleep centre experience. *Sleep Breath* 2021;25:281-288.
6. Patil SP, Schneider H, Schwartz AR, Smith PL. Adult obstructive sleep apnea: pathophysiology and diagnosis. *Chest* 2007;132:325-337.
7. Amra B, Rahmati B, Soltaninejad F, Feizi A. Screening questionnaires for obstructive sleep apnea: an updated systematic review. *Oman Med J* 2018;33:184-192.
8. Chung F, Yegneswaran B, Liao P, et al. STOP questionnaire: a tool to screen patients for obstructive sleep apnea. *Anesthesiology* 2008;108:812-821.
9. Pivetta B, Chen L, Nagappa M, et al. Use and performance of the STOP-Bang questionnaire for obstructive sleep apnea screening across geographic regions: a systematic review and meta-analysis. *JAMA Netw Open* 2021;4:e211009.
10. Kweon S, Kim Y, Jang MJ, et al. Data resource profile: the Korea National Health and Nutrition Examination Survey (KNHANES). *Int J Epidemiol* 2014;43:69-77.
11. Byun JI, Kim DH, Kim JS, Shin WC. Usefulness of using alternative body-mass index and neck circumference criteria for STOP-Bang questionnaire in screening South Korean obstructive sleep apnea patients. *Sleep Med Res* 2020;11:38-43.
12. World Health Organization. *The Asia-Pacific Perspective: Redefining Obesity and Its Treatment*. Sydney (AU): WHO, 2000.
13. Seo MH, Lee WY, Kim SS, et al. 2018 Korean Society for the Study of Obesity guideline for the management of obesity in Korea. *J Obes Metab Syndr* 2019;28:40-45.
14. Park S, Ahn J, Lee BK. Very-low-fat diets may be associated with increased risk of metabolic syndrome in the adult population. *Clin Nutr* 2016;35:1159-1167.
15. Rosenberg DE, Bull FC, Marshall AL, Sallis JF, Bauman AE. Assessment of sedentary behavior with the International Physical Activity Questionnaire. *J Phys Act Health* 2008;5 Suppl 1: S30-S44.
16. Jung CH, Son JW, Kang S, et al. Diabetes fact sheets in Korea, 2020: an appraisal of current status. *Diabetes Metab J* 2021;45:1-10.
17. Lamb EJ, Tomson CR, Roderick PJ; Clinical Sciences Reviews Committee of the Association for Clinical Biochemistry. Estimating kidney function in adults using formulae. *Ann Clin Biochem* 2005;42(Pt 5):321-345.
18. Chung F, Subramanyam R, Liao P, Sasaki E, Shapiro C, Sun Y. High STOP-Bang score indicates a high probability of obstructive sleep apnoea. *Br J Anaesth* 2012;108:768-775.
19. Ko YM, Ko SH, Han K, et al. Importance of awareness and treatment for diabetes in influenza vaccination coverage of diabetic patients under 65 years: a population-based study. *Diabetes Metab J* 2021;45:55-66.

20. Whelton PK, Carey RM, Aronow WS, et al. 2017 ACC/AHA/AAPA/ABC/ACPM/AGS/APhA/ASH/ASPC/NMA/PCNA guideline for the prevention, detection, evaluation, and management of high blood pressure in adults: a report of the American College of Cardiology/American Heart Association task force on clinical practice guidelines. *J Am Coll Cardiol* 2018;71:e127-e248.
21. Nagappa M, Liao P, Wong J, et al. Validation of the STOP-Bang questionnaire as a screening tool for obstructive sleep apnea among different populations: a systematic review and meta-analysis. *PLoS One* 2015;10:e0143697.
22. Chung F, Abdullah HR, Liao P. STOP-Bang questionnaire: a practical approach to screen for obstructive sleep apnea. *Chest* 2016;149:631-638.
23. Chen L, Pivetta B, Nagappa M, et al. Validation of the STOP-Bang questionnaire for screening of obstructive sleep apnea in the general population and commercial drivers: a systematic review and meta-analysis. *Sleep Breath* 2021;25:1741-1751.
24. Tan A, Yin JD, Tan LW, van Dam RM, Cheung YY, Lee CH. Predicting obstructive sleep apnea using the STOP-Bang Questionnaire in the general population. *Sleep Med* 2016;27-28:66-71.
25. Kang K, Seo JG, Seo SH, Park KS, Lee HW. Prevalence and related factors for high-risk of obstructive sleep apnea in a large Korean population: results of a questionnaire-based study. *J Clin Neurol* 2014;10:42-49.
26. Sunwoo JS, Hwangbo Y, Kim WJ, Chu MK, Yun CH, Yang KI. Prevalence, sleep characteristics, and comorbidities in a population at high risk for obstructive sleep apnea: a nationwide questionnaire study in South Korea. *PLoS One* 2018;13:e0193549.
27. Huang T, Lin BM, Markt SC, et al. Sex differences in the associations of obstructive sleep apnoea with epidemiological factors. *Eur Respir J* 2018;51:1702421.
28. Block AJ, Boysen PG, Wynne JW, Hunt LA. Sleep apnea, hypopnea and oxygen desaturation in normal subjects: a strong male predominance. *N Engl J Med* 1979;300:513-517.
29. Bauters FA, Loof S, Hertegonne KB, Chirinos JA, De Buyzere ML, Rietzschel ER. Sex-specific sleep apnea screening questionnaires: closing the performance gap in women. *Sleep Med* 2020;67:91-98.
30. Etindele Sosso FA, Matos E. Socioeconomic disparities in obstructive sleep apnea: a systematic review of empirical research. *Sleep Breath* 2021;25:1729-1739.
31. Li X, Sundquist K, Sundquist J. Socioeconomic status and occupation as risk factors for obstructive sleep apnea in Sweden: a population-based study. *Sleep Med* 2008;9:129-136.
32. Adams RJ, Piantadosi C, Appleton SL, et al. Investigating obstructive sleep apnoea: will the health system have the capacity to cope?: a population study. *Aust Health Rev* 2012;36:424-429.
33. Peppard PE, Young T, Palta M, Dempsey J, Skatrud J. Longitudinal study of moderate weight change and sleep-disordered breathing. *JAMA* 2000;284:3015-3021.
34. Ryan CM, Bradley TD. Pathogenesis of obstructive sleep apnea. *J Appl Physiol* (1985) 2005;99:2440-2450.
35. Buchner NJ, Quack I, Stegbauer J, Woznowski M, Kaufmann A, Rump LC. Treatment of obstructive sleep apnea reduces arterial stiffness. *Sleep Breath* 2012;16:123-133.
36. Elmasry A, Lindberg E, Berne C, et al. Sleep-disordered breathing and glucose metabolism in hypertensive men: a population-based study. *J Intern Med* 2001;249:153-161.
37. Reuveni H, Greenberg-Dotan S, Simon-Tuval T, Oksenberg A, Tarasiuk A. Elevated healthcare utilisation in young adult males with obstructive sleep apnoea. *Eur Respir J* 2008;31:273-279.
38. Kao LT, Lee HC, Lin HC, Tsai MC, Chung SD. Healthcare service utilization by patients with obstructive sleep apnea: a population-based study. *PLoS One* 2015;10:e0137459.

Supplementary Table 1. Clinical characteristics compared by gender of the study population according to the STOP-BANG score

Characteristic	STOP-BANG score								
	Male				Female				
	0–2	3–4	5–8	<i>p</i> value	0–2	3–4	5–8	<i>p</i> value	
Number	516	1,969	836		3,347	899	83		
Age groups, yr								< 0.001	< 0.001
40–49	66.5 (2.4)	26.8 (1.4)	18.0 (1.6)		35.1 (1.2)	8.5 (1.2)	2.1 (1.5)		
50–59	12.7 (1.6)	33.9 (1.4)	43.1 (2.1)		30.6 (0.9)	29.7 (2.1)	43.6 (6.9)		
60–69	12.2 (1.6)	23.3 (1.1)	25.7 (1.8)		19.9 (0.8)	32.0 (1.8)	34.7 (6.6)		
70–79	6.0 (1.0)	12.3 (0.8)	10.7 (1.0)		10.8 (0.6)	22.8 (1.7)	14.1 (4.0)		
≥ 80	2.6 (0.7)	3.6 (0.4)	2.5 (0.5)		3.7 (0.4)	7.1 (1.0)	5.5 (2.8)		
Educational level								< 0.001	< 0.001
Elementary school or lower	7.4 (1.2)	12.7 (1.0)	9.8 (1.0)		18.8 (0.9)	39.7 (2.0)	33.8 (5.9)		
Middle school	6.8 (1.1)	11.9 (0.9)	12.2 (1.4)		10.6 (0.6)	14.9 (1.4)	14.6 (4.9)		
High school	30.9 (2.4)	35.6 (1.4)	36.5 (2.0)		37.3 (1.1)	30.5 (2.1)	30.3 (5.9)		
College or higher	54.8 (2.6)	39.8 (1.7)	41.5 (2.1)		33.4 (1.3)	14.9 (1.5)	21.3 (5.9)		
Household income								0.016	< 0.001
Quintile 1 (lowest)	8.8 (1.5)	11.1 (0.9)	11.3 (1.2)		12.4 (0.8)	23.3 (1.8)	27.2 (5.5)		
Quintile 2	13.5 (1.7)	16.6 (1.0)	17.2 (1.5)		17.0 (0.8)	24.2 (1.8)	25.6 (6.0)		
Quintile 3	19.4 (2.1)	20.6 (1.1)	20.3 (1.7)		20.6 (0.9)	19.2 (1.6)	14.2 (3.6)		
Quintile 4	31.8 (2.6)	25.3 (1.2)	20.8 (1.6)		24.6 (0.9)	20.8 (1.6)	20.1 (5.8)		
Quintile 5 (highest)	26.6 (2.6)	26.4 (1.5)	30.4 (2.0)		25.4 (1.2)	12.4 (1.3)	12.9 (4.1)		
Region, urban	70.1 (2.9)	70.0 (2.0)	67.1 (2.5)	0.456	72.0 (1.7)	63.4 (2.6)	73.5 (5.9)	< 0.001	
Occupation, yes	85.9 (1.7)	75.8 (1.1)	77.3 (1.7)	< 0.001	53.6 (1.2)	45.4 (2.2)	37.9 (6.4)	< 0.001	
Smoking								< 0.001	0.244
Non	26.7 (2.3)	20.3 (1.0)	18.1 (1.5)		92.4 (0.5)	90.5 (1.2)	90.9 (4.3)		
Former	36.3 (2.4)	48.5 (1.3)	51.3 (2.0)		4.1 (0.4)	4.4 (0.8)	7.4 (4.1)		
Current	37.0 (2.7)	31.2 (1.3)	30.7 (1.8)		3.5 (0.4)	5.1 (0.9)	1.7 (1.7)		
Drinking								0.002	0.046
Non	16.7 (1.8)	20.5 (1.1)	18.9 (1.6)		39.4 (1.0)	46.0 (2.0)	42.3 (6.6)		
Mild to moderate	70.7 (2.2)	63.9 (1.4)	60.5 (2.0)		59.2 (1.0)	52.6 (2.0)	55.9 (6.7)		
Heavy	12.7 (1.7)	15.6 (1.0)	20.6 (1.6)		1.4 (0.2)	1.4 (0.5)	1.8 (1.3)		
Physical activity	39.9 (2.4)	43.9 (1.3)	42.0 (2.0)	0.301	39.4 (1.0)	32.6 (2.0)	24.0 (5.2)	< 0.001	
BMI groups, kg/m ²								< 0.001	< 0.001
< 18.5	6.1 (1.2)	1.6 (0.3)	0.2 (0.1)		3.6 (0.4)	0.2 (0.1)	0.0 (0.0)		
≥ 18.5–< 23	50.5 (2.4)	24.8 (1.2)	10.6 (1.2)		47.3 (1.1)	23.6 (1.7)	8.0 (3.2)		
≥ 23–< 25	22.8 (2.1)	31.4 (1.3)	25.3 (1.7)		23.7 (0.8)	21.6 (1.7)	1.3 (0.8)		
≥ 25–< 30	20.6 (2.0)	39.9 (1.3)	48.2 (2.0)		24.1 (0.9)	38.3 (1.8)	30.2 (6.1)		
≥ 30	0.0 (0.0)	2.3 (0.4)	15.8 (1.5)		1.3 (0.2)	16.4 (1.4)	60.5 (6.5)		
Abdominal obesity	17.8 (1.9)	44.3 (1.2)	65.2 (1.9)	< 0.001	29.3 (1.0)	61.5 (1.9)	89.7 (3.6)	< 0.001	
Diabetes mellitus	7.2 (1.3)	20.4 (1.1)	32.0 (2.2)	< 0.001	11.5 (0.7)	27.9 (1.7)	40.7 (6.1)	< 0.001	
Hypertension	2.5 (0.8)	40.0 (1.3)	72.0 (1.9)	< 0.001	23.2 (0.9)	78.3 (1.7)	88.8 (4.4)	< 0.001	
Hypercholesterolemia	15.8 (1.9)	26.8 (1.2)	37.9 (1.8)	< 0.001	29.7 (1.0)	49.5 (2.0)	49.5 (6.5)	< 0.001	
CKD	0.5 (0.2)	3.9 (0.5)	4.1 (0.8)	< 0.001	1.9 (0.3)	5.3 (0.9)	9.7 (3.8)	< 0.001	

Supplementary Table 1. Continued

Characteristic	STOP-BANG score							
	Male				Female			
	0-2	3-4	5-8	<i>p</i> value	0-2	3-4	5-8	<i>p</i> value
Age, yr	50.9 ± 0.5	57.4 ± 0.3	57.8 ± 0.4	< 0.001	56.0 ± 0.3	63.2 ± 0.5	61.5 ± 1.2	< 0.001
Height, cm	170.8 ± 0.3	170.0 ± 0.2	170.0 ± 0.2	0.065	157.4 ± 0.1	155.2 ± 0.2	157.4 ± 0.7	< 0.001
Weight, kg	82.6 ± 0.4	89.0 ± 0.2	94.0 ± 0.4	< 0.001	80.4 ± 0.2	88.3 ± 0.4	97.9 ± 1.3	< 0.001
BMI, kg/m ²	22.7 ± 0.1	24.6 ± 0.1	26.6 ± 0.1	< 0.001	23.2 ± 0.1	26.0 ± 0.2	30.1 ± 0.5	< 0.001
Waist circumference, cm	82.6 ± 0.4	89.0 ± 0.2	94.0 ± 0.4	< 0.001	80.4 ± 0.2	88.3 ± 0.4	97.9 ± 1.3	< 0.001
Fasting glucose, mg/dL	100.1 ± 0.9	107.5 ± 0.7	112.2 ± 1.3	< 0.001	99.1 ± 0.4	106.0 ± 0.8	111.0 ± 3.6	< 0.001
Systolic BP, mmHg	115.1 ± 0.6	122.4 ± 0.4	127.3 ± 0.6	< 0.001	117.6 ± 0.4	131.3 ± 0.8	130.6 ± 1.8	< 0.001
Diastolic BP, mmHg	75.8 ± 0.4	78.8 ± 0.3	81.9 ± 0.5	< 0.001	74.5 ± 0.2	77.6 ± 0.4	81.0 ± 1.2	< 0.001
Total cholesterol, mg/dL	195.1 ± 1.8	191.1 ± 1.0	189.4 ± 1.8	0.065	199.6 ± 0.9	189.4 ± 1.5	190.8 ± 5.3	< 0.001
LDL-C, mg/dL	117.7 ± 1.7	113.6 ± 0.9	111.2 ± 1.6	0.016	121.2 ± 0.8	112.4 ± 1.3	111.9 ± 4.9	< 0.001
HDL-C, mg/dL	49.7 ± 0.6	46.9 ± 0.3	46.4 ± 0.4	< 0.001	56.0 ± 0.3	52.0 ± 0.5	48.3 ± 1.2	< 0.001
eGFR, mL/min/1.73 m ²	96.2 ± 0.8	90.4 ± 0.5	90.0 ± 0.8	< 0.001	97.2 ± 0.5	91.9 ± 0.9	92.7 ± 2.9	< 0.001

Values are presented as percentage (standard error) or mean ± standard error.

BMI, body mass index; CKD, chronic kidney disease; BP, blood pressure; LDL-C, low-density lipoprotein cholesterol; HDL-C, high-density lipoprotein cholesterol; eGFR, estimated glomerular filtration rate.

Supplementary Table 2. The prevalence of high-risk OSA for measures of hypertension, diabetes, and obesity, according to decile group

Decile	Prevalence of high-risk OSA according to decile group											
	SBP		DBP		FPG		HbA1c		BMI		WC	
	%	SE	%	SE	%	SE	%	SE	%	SE	%	SE
1	4.2	1.0	7.7	1.1	6.8	1.2	7.1	1.2	1.3	0.4	1.7	0.6
2	7.2	1.0	6.8	1.0	5.1	0.8	5.4	0.8	2.9	0.8	5.1	1.0
3	7.5	1.2	6.8	1.0	7.0	1.0	9.3	1.3	4.8	0.9	9.8	1.2
4	9.8	1.3	9.2	1.3	11.8	1.8	13.9	1.6	7.5	1.2	9.9	1.3
5	13.8	1.5	11.3	1.6	12.2	1.4	10.3	1.3	12.4	1.5	9.9	1.2
6	17.2	1.8	12.1	1.3	13.5	1.6	12.7	1.5	10.2	1.2	12.8	1.4
7	14.7	1.5	11.5	1.4	13.8	1.5	14.0	1.8	18.3	1.6	12.2	1.3
8	18.9	1.8	14.7	1.4	16.1	1.7	16.3	1.3	12.6	1.5	13.9	1.6
9	22.0	1.7	15.8	1.6	20.8	1.8	21.7	2.1	22.4	2.0	20.4	1.9
10	19.0	1.6	30.9	1.9	25.7	1.9	24.4	1.8	35.8	2.2	37.2	2.1

OSA, obstructive sleep apnea; SBP, systolic blood pressure; DBP, diastolic blood pressure; FPG, fasting plasma glucose; HbA1c, glycosylated hemoglobin; BMI, body mass index; WC, waist circumference; SE, standard error.

Supplementary Table 3. Multivariate logistic regression analysis for the high-risk group of obstructive sleep apnea, compared by gender of the study population

Variable	Male						Female					
	Model 1		Model 2		Model 3		Model 1		Model 2		Model 3	
	OR (95% CI)	p value	OR (95% CI)	p value	OR (95% CI)	p value	OR (95% CI)	p value	OR (95% CI)	p value	OR (95% CI)	p value
Age	1.02 (1.01-1.02)	<0.001	1.02 (1.01-1.02)	<0.001	1.02 (1.01-1.03)	<0.001	1.03 (1.01-1.05)	<0.001	1.03 (1.01-1.05)	<0.001	0.98 (0.95-1.00)	0.071
Education		0.336		0.013		0.003		0.160		0.959		0.447
Elementary	1 (ref)		1 (ref)		1 (ref)		1 (ref)		1 (ref)		1 (ref)	
Middle	1.34 (0.93-1.93)		1.53 (1.06-2.22)		1.46 (0.97-2.18)		0.86 (0.38-1.95)		1.03 (0.43-2.44)		1.31 (0.55-3.09)	
High	1.25 (0.94-1.66)		1.65 (1.21-2.25)		1.90 (1.33-2.71)		0.57 (0.31-1.05)		0.84 (0.38-1.85)		1.77 (0.73-4.31)	
College	1.13 (0.85-1.51)		1.64 (1.19-2.25)		1.96 (1.34-2.85)		0.48 (0.23-1.01)		0.78 (0.28-2.30)		2.42 (0.83-7.09)	
Household income		0.044		0.056		0.215		0.015		0.189		0.100
Q1	1 (ref)		1 (ref)		1 (ref)		1 (ref)		1 (ref)		1 (ref)	
Q2	1.01 (0.73-1.41)		1.10 (0.79-1.53)		0.95 (0.65-1.39)		0.74 (0.36-1.52)		0.85 (0.42-1.70)		0.92 (0.45-1.91)	
Q3	0.94 (0.68-1.30)		1.12 (0.80-1.57)		0.90 (0.61-1.33)		0.37 (0.19-0.74)		0.47 (0.22-0.99)		0.37 (0.17-0.81)	
Q4	0.73 (0.53-1.00)		0.90 (0.64-1.27)		0.78 (0.52-1.18)		0.45 (0.20-1.02)		0.58 (0.25-1.36)		0.50 (0.19-1.37)	
Q5	1.08 (0.79-1.47)		1.33 (0.95-1.87)		1.10 (0.72-1.67)		0.30 (0.13-0.67)		0.39 (0.17-0.90)		0.41 (0.17-1.01)	
Occupation, no	1 (ref)		1 (ref)		1 (ref)		1 (ref)		1 (ref)		1 (ref)	
Yes	0.96 (0.77-1.18)		1.21 (0.95-1.56)		1.26 (0.95-1.67)		0.56 (0.33-0.96)		0.67 (0.37-1.20)		0.56 (0.31-1.02)	
Smoking		0.027		0.049		0.515		0.445		0.394		0.482
Non	1 (ref)		1 (ref)		1 (ref)		1 (ref)		1 (ref)		1 (ref)	
Former	1.35 (1.07-1.70)		1.33 (1.06-1.67)		1.15 (0.89-1.49)		1.82 (0.56-5.86)		2.05 (0.64-6.58)		1.61 (0.55-4.70)	
Current	1.14 (0.89-1.46)		1.19 (0.93-1.53)		1.15 (0.88-1.51)		0.46 (0.06-3.43)		0.54 (0.07-4.02)		0.44 (0.06-3.48)	
Alcohol consumption		0.004		<0.001		0.242		0.908		0.642		0.510
Non	1 (ref)		1 (ref)		1 (ref)		1 (ref)		1 (ref)		1 (ref)	
Mild to moderate	0.96 (0.76-1.23)		1.06 (0.83-1.36)		1.01 (0.74-1.37)		0.93 (0.54-1.59)		1.22 (0.66-2.23)		1.10 (0.58-2.09)	
Heavy	1.43 (1.06-1.93)		1.65 (1.21-2.25)		1.24 (0.87-1.76)		1.25 (0.29-5.33)		1.92 (0.42-8.72)		2.70 (0.50-14.44)	
Physical activity, no	1 (ref)		1 (ref)		1 (ref)		1 (ref)		1 (ref)		1 (ref)	
Yes	0.96 (0.80-1.16)		0.98 (0.81-1.18)		0.96 (0.78-1.19)		0.51 (0.29-0.90)		0.56 (0.32-0.98)		0.59 (0.33-1.08)	
DM, no	1 (ref)		1 (ref)		1 (ref)		1 (ref)		1 (ref)		1 (ref)	
Yes	2.24 (1.78-2.82)		2.14 (1.69-2.71)		1.54 (1.19-1.99)		4.00 (2.41-6.63)		3.49 (1.99-6.10)		2.01 (1.15-3.50)	
HTN, no	1 (ref)		1 (ref)		1 (ref)		1 (ref)		1 (ref)		1 (ref)	
Yes	5.61 (4.62-6.83)		5.65 (4.59-6.96)		4.63 (3.71-5.78)		15.61 (6.54-37.28)		18.54 (7.27-47.31)		11.53 (4.31-30.80)	
Hypercholesterolemia, no	1 (ref)		1 (ref)		1 (ref)		1 (ref)		1 (ref)		1 (ref)	
Yes	1.90 (1.59-2.28)		1.87 (1.56-2.25)		1.24 (0.99-1.54)		1.96 (1.17-3.28)		1.66 (0.99-2.80)		1.00 (0.58-1.74)	
CKD, no	1 (ref)		1 (ref)		1 (ref)		1 (ref)		1 (ref)		1 (ref)	
Yes	1.32 (0.84-2.06)		1.08 (0.68-1.72)		0.67 (0.42-1.08)		4.09 (1.73-9.67)		2.86 (1.16-7.09)		1.46 (0.64-3.34)	

Supplementary Table 3. Continued

Variable	Male						Female					
	Model 1		Model 2		Model 3		Model 1		Model 2		Model 3	
	OR (95% CI)	p value	OR (95% CI)	p value	OR (95% CI)	p value	OR (95% CI)	p value	OR (95% CI)	p value	OR (95% CI)	p value
Obesity, no	1 (ref)	<0.001	1 (ref)	<0.001	1 (ref)	<0.001	1 (ref)	<0.001	1 (ref)	<0.001	1 (ref)	<0.001
Yes	3.00 (2.47–3.63)		3.18 (2.63–3.85)		2.35 (1.91–2.91)		21.90 (10.16–47.23)		20.89 (9.55–45.73)		14.74 (6.70–32.43)	
Abdominal obesity, no	1 (ref)	<0.001	1 (ref)	<0.001	1 (ref)	<0.001	1 (ref)	<0.001	1 (ref)	<0.001	1 (ref)	0.106
Yes	3.02 (2.52–3.63)		3.00 (2.50–3.61)		1.57 (1.24–2.00)		15.94 (7.38–34.42)		15.41 (6.67–35.58)		2.14 (0.85–5.37)	
BMI level, kg/m ²		<0.001		<0.001		<0.001		<0.001		<0.001		<0.001
< 18.5	0.19 (0.04–0.82)		0.18 (0.04–0.76)		0.18 (0.04–0.81)		-		-		-	
< 23	1 (ref)		1 (ref)		1 (ref)		1 (ref)		1 (ref)		1 (ref)	
< 25	2.51 (1.86–3.38)		2.71 (1.99–3.68)		2.29 (1.65–3.18)		0.29 (0.07–1.23)		0.26 (0.06–1.10)		0.20 (0.05–0.88)	
< 30	3.96 (2.99–5.25)		4.40 (3.30–5.86)		3.11 (2.26–4.27)		6.05 (2.36–15.55)		5.26 (2.00–13.81)		3.47 (1.26–9.57)	
< 35	26.45 (15.71–44.52)		35.94 (21.58–59.86)		23.16 (13.55–39.60)		72.17 (29.15–178.68)		71.73 (28.97–177.60)		46.45 (18.53–116.43)	
≥ 35	20.26 (5.70–72.03)		29.09 (8.42–100.48)		16.02 (4.76–53.94)		129.79 (35.54–473.98)		165.09 (46.17–590.26)		65.53 (17.23–249.29)	
Obesity group		<0.001		<0.001		<0.001		<0.001		<0.001		<0.001
Low BMI, low WC	1 (ref)		1 (ref)		1 (ref)		1 (ref)		1 (ref)		1 (ref)	
Low BMI, high WC	1.72 (1.22–2.41)		1.54 (1.09–2.18)		1.25 (0.87–1.81)		1.54 (0.36–6.65)		1.29 (0.29–5.92)		0.92 (0.19–4.38)	
High BMI, low WC	1.65 (1.14–2.41)		1.76 (1.21–2.55)		1.37 (0.92–2.04)		4.60 (0.86–24.56)		4.75 (0.89–25.19)		4.64 (0.82–26.24)	
High BMI, high WC	3.76 (3.05–4.65)		3.87 (3.14–4.78)		2.74 (2.17–3.47)		27.10 (11.19–65.62)		25.07 (9.90–63.48)		15.79 (6.10–40.90)	
DM + HTN + obesity, no	1 (ref)	<0.001	1 (ref)	<0.001	1 (ref)	<0.001	1 (ref)	<0.001	1 (ref)	<0.001	1 (ref)	<0.001
Yes	5.73 (4.31–7.61)		5.58 (4.18–7.44)		3.56 (2.61–4.86)		11.22 (6.56–19.18)		9.98 (5.60–17.79)		4.71 (2.62–8.48)	

Model 1 was unadjusted; Model 2 was adjusted for age and sex; Model 3 was adjusted for age, sex, education, household income, occupation, smoking, drinking, physical activity, DM, hypertension, hypercholesterolemia, CKD, obesity, and abdominal obesity (weight circumference: men ≥ 90/women 85 cm). Abdominal obesity was defined as WC ≥ 90 cm in men, and ≥ 85 cm in women. Low and high WC was determined by this standard. Obesity was defined as subjects with BMI ≥ 25 kg/m². OR, odds ratio; CI, confidence interval; DM, diabetes mellitus; HTN, hypertension; CKD, chronic kidney disease; BMI, body mass index; WC, waist circumference.