

## Predicted Atherosclerotic Cardiovascular Disease Risk and Masked Hypertension Among Blacks in the Jackson Heart Study

D. Edmund Anstey, MD; John N. Booth III, MS; Marwah Abdalla, MD, MPH; Tanya M. Spruill, PhD; Yuan-I Min, PhD; Paul Muntner, PhD; Daichi Shimbo, MD

**Background**—Among individuals without hypertension based on clinic blood pressure (BP), it is unclear who should be screened for masked hypertension, defined as having hypertension based on out-of-clinic BP. We hypothesized that individuals with a higher 10-year predicted atherosclerotic cardiovascular disease (ASCVD) risk, calculated using the pooled cohort risk equations, have a higher prevalence of masked hypertension.

**Methods and Results**—We analyzed data from the Jackson Heart Study—a population-based cohort of blacks—to determine the association of predicted ASCVD risk with masked hypertension. The sample included 644 participants, 40 to 79 years of age, with clinic systolic/diastolic BP <140/90 mmHg, who completed ambulatory BP monitoring, were free of cardiovascular disease, and had data on factors needed to calculate ASCVD risk. Ten-year predicted ASCVD risk was calculated using the pooled cohort risk equations. Any masked hypertension was defined as masked daytime hypertension (mean daytime systolic/diastolic BP  $\geq$ 135/85 mmHg), masked nighttime hypertension (mean nighttime systolic/diastolic BP  $\geq$ 120/70 mmHg), or masked 24-hour hypertension (mean 24-hour systolic/diastolic BP  $\geq$ 130/80 mmHg). The prevalence of any masked hypertension was 54.0%. Compared with participants in the lowest (<5%) predicted ASCVD risk category, multivariable-adjusted prevalence ratios (95% confidence interval) for any masked hypertension were 1.36 (1.03–1.79), 1.62 (1.22–2.16), and 1.91 (1.47–2.48) for those with ASCVD risk of 5% to <7.5%, 7.5% to <10%, and  $\geq$ 10%, respectively. The C statistic for discriminating between participants with versus without any masked hypertension was 0.681 (95% confidence interval, 0.640–0.723) for ASCVD risk and 0.703 (95% confidence interval, 0.663–0.744) for clinic systolic BP and diastolic BP.

**Conclusions**—Higher ASCVD risk was associated with an increased prevalence of masked hypertension. Although the discrimination of ASCVD risk for masked hypertension was not superior to clinic BP, risk prediction equations may be useful for identifying the subgroup of individuals with both masked hypertension and high predicted ASCVD risk. (*Circ Cardiovasc Qual Outcomes*. 2017;10:e003421. DOI: 10.1161/CIRCOUTCOMES.116.003421.)

**Key Words:** blood pressure ■ cardiovascular diseases ■ humans ■ masked hypertension ■ prevalence

Blood pressure (BP) is traditionally measured in the clinic setting. However, studies have demonstrated that BP measured in the clinic differs substantially when measured outside of the clinic using ambulatory BP monitoring (ABPM).<sup>1,2</sup> Masked hypertension, defined as not having hypertension on clinic BP measurements but having hypertension based on daytime BP measurements from outside of the clinic setting, is a common phenotype with prevalence estimates ranging from 15% to 30% in population-based studies.<sup>1</sup> When the definition of masked hypertension includes having hypertension based on nighttime BP, the prevalence has been reported to be substantially higher.<sup>3</sup> Masked hypertension has been associated with an increased risk for

atherosclerotic cardiovascular disease (ASCVD) events in several prior studies.<sup>1–6</sup>

### See Editorial by Morris and Bisognano

It is unclear who should undergo ABPM to identify masked hypertension among adults without hypertension based on BP measured in the clinic. In prior studies, the prevalence of masked hypertension has increased with higher clinic systolic BP (SBP) and diastolic BP (DBP).<sup>1,2</sup> In addition to higher clinic BP, older age, male sex, smoking, diabetes mellitus, and antihypertensive medication use have each been associated with an increased prevalence of masked hypertension.<sup>1,2</sup> Therefore, a composite score based on multiple ASCVD risk

Received November 14, 2016; accepted May 26, 2017.

From the Department of Medicine, Columbia University Medical Center, New York, NY (D.E.A., M.A., D.S.); Department of Epidemiology, University of Alabama at Birmingham, Birmingham, AL (J.N.B., P.M.); Department of Population Health, New York University School of Medicine, New York, NY (T.M.S.); and School of Medicine, University of Mississippi Medical Center, Jackson, MS (Y.-I.M.).

The Data Supplement is available at <http://circoutcomes.ahajournals.org/lookup/suppl/doi:10.1161/CIRCOUTCOMES.116.003421/-/DC1>.

Correspondence to D. Edmund Anstey, MD, Columbia University Medical Center, 622 W 168th St, PH 9–310, New York, NY 10032. E-mail [dea2123@cumc.columbia.edu](mailto:dea2123@cumc.columbia.edu)

© 2017 American Heart Association, Inc.

*Circ Cardiovasc Qual Outcomes* is available at <http://circoutcomes.ahajournals.org>

DOI: 10.1161/CIRCOUTCOMES.116.003421

### WHAT IS KNOWN

- Blood pressure measured in the clinic can differ substantially from blood pressure measured outside of the clinic assessed using ambulatory blood pressure monitoring.
- Masked hypertension, defined as not having hypertension on clinic blood pressure measurements but having hypertension on ambulatory blood pressure monitoring, is a common phenotype and is associated with an increased risk of cardiovascular disease events.
- Current guidelines do not specify whom with non-elevated clinic blood pressure should be screened for masked hypertension with ambulatory blood pressure monitoring.

### WHAT THE STUDY ADDS

- Higher predicted atherosclerotic cardiovascular disease risk using the pooled cohort risk equations is associated with an increased prevalence of masked hypertension among blacks.
- Cardiovascular disease risk prediction equations may be useful in determining whom should be screened for masked hypertension using ABPM.

factors may be a useful tool for choosing who to screen with ABPM for identifying masked hypertension.

In the current study, we evaluated whether higher 10-year predicted ASCVD risk, assessed using the pooled cohort risk equations,<sup>7</sup> is associated with a higher prevalence of masked hypertension. Additionally, we compared 10-year predicted ASCVD risk to clinic BP for discriminating participants with and without masked hypertension. These analyses were conducted in the JHS (Jackson Heart Study)—a large population-based cohort of blacks. The results of this study may help identify individuals who should undergo ABPM for detecting masked hypertension.

## Methods

### Study Population

The JHS—a population-based prospective cohort study—was designed to evaluate cardiovascular disease risk among blacks.<sup>8</sup> Briefly, the JHS enrolled 5306 noninstitutionalized blacks, aged  $\geq 20$  years, between 2000 and 2004 from the Atherosclerosis Risk in the Community site in Jackson, MS—a representative sample of urban and rural Jackson—Mississippi metropolitan tricounty (Hinds, Madison, and Rankin counties) residents, volunteers, randomly contacted individuals, and secondary family members of participants.<sup>9,10</sup>

The current analysis was restricted to JHS participants who underwent ABPM after the baseline examination ( $n=1148$ ). Participants who did not meet the international database on ABPM in relation to cardiovascular outcomes<sup>11</sup> criteria for a complete ABPM recording ( $n=102$ ; described below) or were missing clinic BP or information on antihypertensive medication use ( $n=30$ ) were excluded from the current analysis. Because masked hypertension can only be present among individuals without hypertension based on clinic-measured BP (ie, clinic SBP  $< 140$  mmHg and clinic DBP  $< 90$  mmHg), participants who had clinic SBP  $\geq 140$  mmHg or clinic DBP  $\geq 90$  mmHg ( $n=196$ ) were also excluded. Also, because the pooled cohort risk

equations were developed and validated among adults 40 to 79 years of age, participants  $< 40$  years of age or  $> 79$  years of age ( $n=51$ ) were excluded.<sup>7</sup> Participants were excluded if they had missing data on ASCVD risk factors ( $n=64$ ), including age, sex, total cholesterol, high-density lipoprotein cholesterol, smoking status, and diabetes mellitus. Finally, because the pooled cohort risk equations were designed to evaluate predicted ASCVD risk among adults without cardiovascular disease,<sup>7</sup> participants with a history of myocardial infarction or stroke ( $n=61$ ) were excluded from the current study, leaving a final sample size of 644 participants. The institutional review board governing human subjects' research at the participating institutions approved the JHS protocol and all data collection procedures. All participants provided written informed consent. The current analysis was approved by the Institutional Review Board at Columbia University and University of Alabama at Birmingham.

### Data Collection

Detailed descriptions of data collection, methodology, specimen collection, and processing have been previously described.<sup>10,12</sup> Data for this analysis were collected during the baseline in-home interview and study visit and through ABPM.

### Baseline Characteristics

Age, sex, and education were obtained by self-report using standardized interviewer-administered questionnaires. Current smoking was defined by affirmative responses to the questions "Have you smoked more than 400 cigarettes in your lifetime?" and "Do you now smoke cigarettes?" Antihypertensive medication use in the 2 weeks before the study visit was self-reported. Height, weight, and clinic BP were measured, and blood samples were collected by trained staff during the study visit. Body mass index was calculated as weight in kilograms divided by height in meters squared. Total and high-density lipoprotein cholesterol were quantified by an oxidase method.<sup>12</sup> Serum glucose was measured using a glucose oxidase method on a Vitros 250 or 950, ortho-clinical diagnostics analyzer.<sup>12</sup> Hemoglobin A1c was measured using a TOSOH high-performance liquid chromatography system. Diabetes mellitus was defined as a fasting ( $\geq 8$  hours) serum glucose  $\geq 126$  mg/dL or hemoglobin A1c  $\geq 6.5\%$  or self-reported use of insulin or oral hypoglycemic medications within 2 weeks before the study visit. Serum creatinine was measured using a multipoint enzymatic spectrophotometric assay on a Vitros 950, ortho-clinical diagnostic analyzer. Estimated glomerular filtration rate was calculated using the chronic kidney disease epidemiology collaboration equation.<sup>13</sup> Reduced estimated glomerular filtration rate was defined as  $< 60$  mL/min/1.73 m<sup>2</sup>.

### Clinic BP Measurement

A standardized protocol was followed to measure clinic BP. Participants were asked to avoid caffeine, eating, heavy physical activity, smoking, and alcohol intake for 12 hours before their study visit. An appropriately sized cuff, determined from an arm circumference measurement, and a random-zero sphygmomanometer (Hawksley and Sons, Ltd) were used for the BP measurement.<sup>10,14</sup> Participants sat for at least 5 minutes in an upright position with their back and arms supported, feet flat on the floor, and legs uncrossed before trained staff conducted 2 BP measurements, separated by 1 minute, in the right arm. The JHS Coordinating Center conducted quality control through training and semiannual retraining of staff,<sup>15</sup> monitoring digit preference for each staff member, and by comparing mean BP measurements within and between trained technicians. As previously described,<sup>16</sup> a BP comparability substudy for which BP was assessed simultaneously, using a Y connector, by random-zero sphygmomanometer and an Omron HEM-907XL oscillometric device was conducted among 2115 JHS participants in 2005 to 2008. The random-zero BP measurements were calibrated to the oscillometric device using robust regression. The average of the 2 clinic BP measurements was used for analysis.

### Ambulatory BP Monitoring

After the baseline study visit, participants were fitted with an ABPM device (Spacelabs 90207) on their nondominant arm. Ambulatory BP was recorded every 20 minutes for 24 hours. Data were evaluated for quality and processed with Medifacts International Medicom software (Rockville, MD). International Database on ABPM in relation to Cardiovascular Outcomes criteria were used to define whether the ABPM measurement was complete.<sup>11</sup> Participants were considered to have a complete ABPM if they had  $\geq 10$  daytime (10:00 AM to 8:00 PM) and  $\geq 5$  nighttime (midnight to 6:00 AM) SBP and DBP measurements.<sup>11</sup>

### Hypertension Categories

Daytime hypertension was defined as a mean ambulatory SBP  $\geq 135$  mm Hg or a mean ambulatory DBP  $\geq 85$  mm Hg using BP measurements obtained between 10:00 AM and 8:00 PM.<sup>17</sup> Nighttime hypertension was defined by a mean ambulatory SBP  $\geq 120$  mm Hg or a mean ambulatory DBP  $\geq 70$  mm Hg using BP measurements obtained between midnight and 6:00 AM, and 24-hour hypertension was defined as a mean ambulatory SBP  $\geq 130$  mm Hg or a mean ambulatory DBP  $\geq 80$  mm Hg using all BP measurements obtained on ABPM.<sup>17</sup> As the current analysis was restricted to participants with clinic SBP  $< 140$  mm Hg and clinic DBP  $< 90$  mm Hg, those with daytime, nighttime, and 24-hour hypertension were considered to have masked daytime hypertension, masked nighttime hypertension, and masked 24-hour hypertension, respectively. Participants with masked daytime hypertension, masked nighttime hypertension, or masked 24-hour hypertension were categorized as having any masked hypertension. Participants were categorized as either having prehypertension, defined by a mean clinic SBP 120 to 139 mm Hg or a mean clinic DBP 80 to 89 mm Hg, or normal clinic BP, defined by a mean clinic SBP  $< 120$  mm Hg and a mean clinic DBP  $< 80$  mm Hg. For participants taking antihypertensive medication, on-treatment clinic SBP/DBP of 120 to 139/80 to 89 mm Hg and masked uncontrolled hypertension are corresponding terms for prehypertension and masked hypertension, respectively.<sup>18</sup> For simplicity in the presentation of the results, we use the terms prehypertension and masked hypertension for all participants, regardless of antihypertensive medication use.

### Ten-Year Predicted ASCVD Risk

In 2013, the American College of Cardiology and American Heart Association recommended the use of the pooled cohort risk equations for estimating 10-year ASCVD risk.<sup>7</sup> There are 4 race- and sex-specific pooled cohort risk equations, each of which uses age, total cholesterol, high-density lipoprotein cholesterol, clinic SBP, and status of antihypertensive medication use, current smoking, and diabetes mellitus to calculate 10-year predicted ASCVD risk.<sup>19</sup> For the primary analyses, participants were categorized into four 10-year predicted ASCVD risk categories:  $< 5\%$ ,  $5\%$  to  $< 7.5\%$ ,  $7.5\%$  to  $< 10\%$ , and  $\geq 10\%$ .<sup>7</sup>

### Statistical Analyses

Participant characteristics were calculated by 10-year predicted ASCVD risk category for the overall analytic sample and for those taking and not taking antihypertensive medication, separately. Herein, we describe the analysis for any masked hypertension. Identical analyses were performed for masked daytime hypertension, masked nighttime hypertension, and masked 24-hour hypertension. The prevalence of any masked hypertension was calculated by category of 10-year predicted ASCVD risk. Using the lowest 10-year predicted ASCVD risk category ( $< 5\%$ ) as the referent group, prevalence ratios (95% confidence interval [CI]) for any masked hypertension were calculated for participants with 10-year predicted ASCVD risk of  $5\%$  to  $< 7.5\%$ ,  $7.5\%$  to  $< 10\%$ , and  $\geq 10\%$ . Associations were examined in an unadjusted model and after adjustment for age, sex, body mass index, less than high school education, and reduced estimated glomerular filtration rate.

The *P*-trend across the categories was calculated by modeling category of 10-year predicted ASCVD risk as a continuous variable using a Cochran–Armitage test for trend. Analyses were repeated for participants taking and not taking antihypertensive medication, separately. Because of a limited sample size being available in some subgroups, participants were grouped into 3 categories of 10-year predicted ASCVD risk ( $< 5\%$ ,  $5\%$  to  $< 7.5\%$ , and  $\geq 7.5\%$ ) for these stratified analyses. The distribution of 10-year predicted ASCVD risk categories was then calculated for participants with and without any masked hypertension, separately.

As previous studies have demonstrated a substantial diagnostic overlap between prehypertension and masked hypertension,<sup>6,20,21</sup> the prevalence and prevalence ratios (95% CI) for any masked hypertension associated with 10-year predicted ASCVD risk were calculated for participants with prehypertension and normal clinic BP, separately. Also, adjusted prevalence ratios (95% CI) for any masked hypertension were calculated for each SD higher 10-year predicted ASCVD risk, clinic SBP, and clinic DBP.

The ability of the 10-year predicted ASCVD risk score and clinic SBP and DBP, clinic SBP alone, and clinic DBP alone to discriminate between participants with versus without any masked hypertension was determined by calculating C statistics. Differences in the C statistics were calculated comparing clinic SBP and clinic DBP together, clinic SBP alone, and clinic DBP alone versus 10-year predicted ASCVD risk. Also, differences in C statistics were calculated comparing clinic SBP and DBP versus clinic SBP alone and separately, clinic DBP alone. Differences in the C statistics were calculated overall and after stratifying participants into those not taking and taking antihypertensive medication. The 95% CIs for the C statistics and the differences in C statistics were calculated using 1000 iteration bootstraps with each iteration including the same sample size as in the main analyses.

*P* values  $< 0.05$  were considered statistically significant. All data analyses were conducted using SAS, version 9.4 (SAS Institute, Cary, NC), or Stata/IC, version 12.1 (Stata, Inc, College Station, TX).

## Results

### Participant Characteristics

Participants with higher 10-year predicted ASCVD risk were older, more likely to be men, have less than high school education, be current smokers, have diabetes mellitus and reduced estimated glomerular filtration rate, and to be taking antihypertensive medication (Table 1). Mean total and low-density lipoprotein cholesterol were higher and high-density lipoprotein cholesterol was lower among participants with higher 10-year predicted ASCVD risk. Also, those with higher ASCVD risk had higher clinic, daytime, nighttime, and 24-hour SBP, higher clinic and nighttime DBP, and a higher prevalence of prehypertension (Table I in the [Data Supplement](#)). Participant characteristics and the clinic and ambulatory BP levels among participants not taking and taking antihypertensive medication, separately, are reported in Table II in the [Data Supplement](#).

### ASCVD Risk Category and Masked Hypertension

The prevalence of any masked hypertension, masked daytime hypertension, masked nighttime hypertension, and masked 24-hour hypertension was 54.0%, 31.5%, 49.2%, and 34.8%, respectively. Participants with higher ASCVD risk had a higher prevalence of each type of masked hypertension (Figure 1). As compared with 10-year predicted ASCVD risk  $< 5\%$ , the adjusted prevalence ratio (95% CI) for having any masked hypertension was 1.36 (1.03–1.79), 1.62 (1.22–2.16), and 1.91 (1.47–2.48) for 10-year predicted ASCVD

**Table 1. Characteristics of Jackson Heart Study Participants by Category of 10-Year Predicted ASCVD Risk**

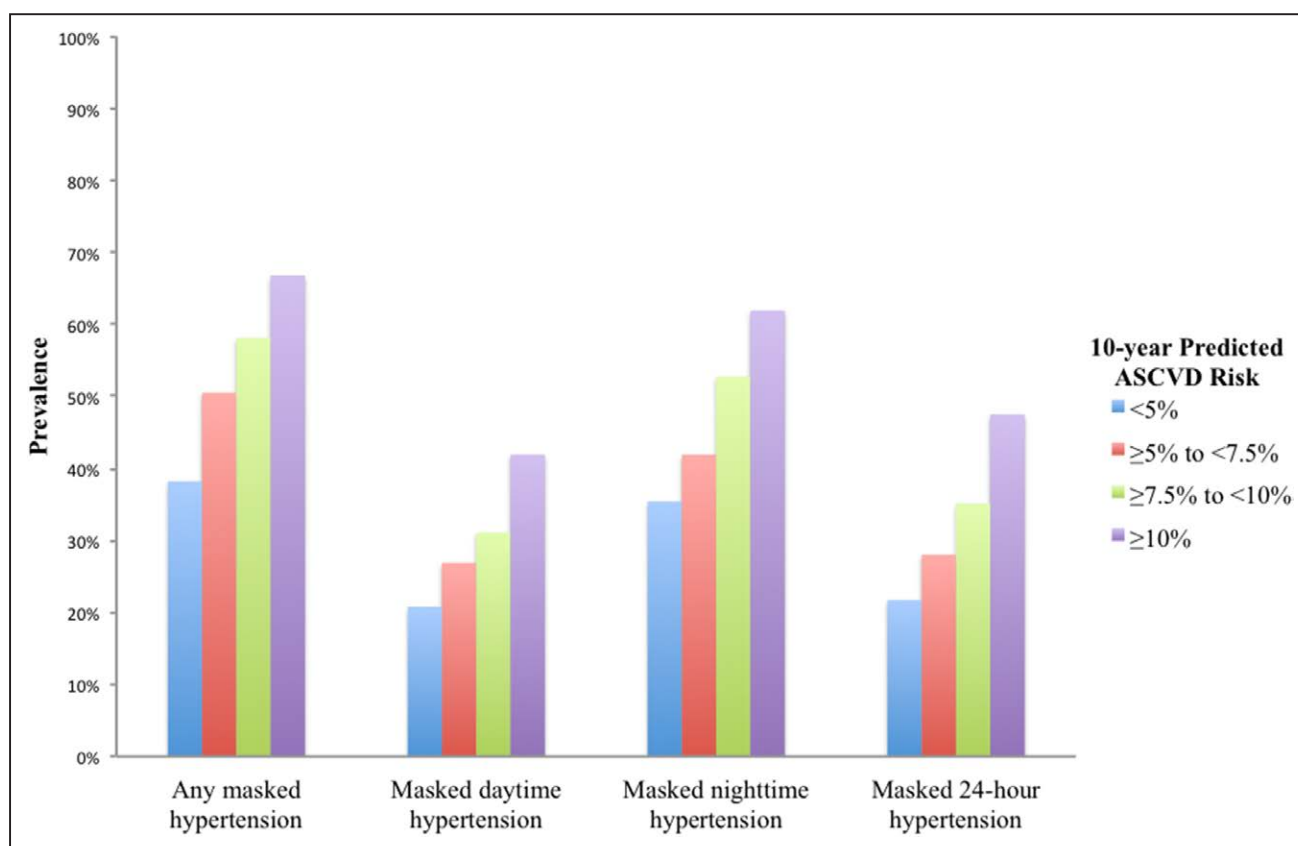
Characteristics	10-y Predicted ASCVD Risk				P-Trend
	<5% (n=212)	≥5% to <7.5% (n=93)	≥7.5% to <10% (n=74)	≥10% (n=265)	
Age, y	49.6±6.1	57.8±6.7	61.8±6.4	66.2±6.8	<0.001
Men, %	14.2	33.3	33.8	40.4	<0.001
Less than high school education, %	6.2	19.4	16.2	21.7	<0.001
Current smoking, %	5.7	7.5	5.4	10.9	0.045
Diabetes mellitus, %	1.9	6.5	13.5	40.4	<0.001
eGFR <60 mL/min/m <sup>2</sup> , %	0.0	5.4	10.8	14.7	<0.001
Body mass index, kg/m <sup>2</sup>	31.1±7.6	31.7±6.1	31.3±6.6	30.6±5.2	0.261
Total cholesterol, mg/dL	196.9±35.5	199.9±36.3	202.3±36.6	208.7±42.9	<0.001
LDL cholesterol, mg/dL	121.9±35.0	125.5±36.4	126.8±35.4	131.1±36.4	0.005
HDL cholesterol, mg/dL	56.7±15.1	52.9±14.4	54.2±12.4	54.4±15.0	0.135
Antihypertensive medication use, %	25.0	51.6	59.5	73.2	<0.001

The numbers in the table are mean±SD or percentages. ASCVD indicates atherosclerotic cardiovascular disease; eGFR, estimated glomerular filtration rate; HDL, high-density lipoprotein; and LDL, low-density lipoprotein.

risk ≥5% to <7.5%, ≥7.5% to <10%, and ≥10%, respectively ( $P$ -trend=<0.001; Table 2). Similar findings were observed for masked daytime hypertension, masked nighttime hypertension, and masked 24-hour hypertension, and among participants not taking and taking antihypertensive medication, separately (Table III in the [Data Supplement](#))

### Prevalence of ASCVD Risk Category by Masked Hypertension Status

The percentage of participants with 10-year predicted ASCVD risk ≥10% was higher among those with versus without any masked hypertension (50.9% versus 29.7%), masked daytime hypertension (54.7% versus 34.9%), masked nighttime



**Figure 1.** Prevalence of masked hypertension by category of 10-y predicted atherosclerotic cardiovascular disease (ASCVD) risk.



**Table 2. Prevalence and Prevalence Ratios for Masked Hypertension by Category of 10-Year Predicted ASCVD Disease Risk**

Type of Masked Hypertension	10-y Predicted ASCVD Risk				P-Trend
	<5%	≥5% to <7.5%	≥7.5% to <10%	≥10%	
	(n=212)	(n=93)	(n=74)	(n=265)	
Prevalence					
Any masked hypertension, %	38.2	50.5	58.1	66.8	<0.001
Masked daytime hypertension, %	20.8	26.9	31.1	41.9	<0.001
Masked nighttime hypertension, %	35.4	41.9	52.7	61.9	<0.001
Masked 24-h hypertension, %	21.7	28.0	35.1	47.5	<0.001
Unadjusted prevalence ratio (95% CI)					
Any masked hypertension	1 (ref)	1.32 (1.02–1.72)	1.52 (1.17–1.97)	1.75 (1.44–2.12)	<0.001
Masked daytime hypertension	1 (ref)	1.30 (0.85–1.98)	1.50 (0.97–2.30)	2.02 (1.50–2.72)	<0.001
Masked nighttime hypertension	1 (ref)	1.19 (0.88–1.60)	1.49 (1.12–1.98)	1.75 (1.43–2.15)	<0.001
Masked 24-h hypertension	1 (ref)	1.29 (0.85–1.95)	1.62 (1.08–2.42)	2.19 (1.65–2.91)	<0.001
Adjusted prevalence ratio (95% CI)*					
Any masked hypertension	1 (ref)	1.36 (1.03–1.79)	1.62 (1.22–2.16)	1.91 (1.47–2.48)	<0.001
Masked daytime hypertension	1 (ref)	1.52 (0.97–2.37)	1.82 (1.13–2.91)	2.64 (1.74–4.01)	<0.001
Masked nighttime hypertension	1 (ref)	1.13 (0.83–1.55)	1.43 (1.04–1.96)	1.66 (1.24–2.23)	<0.001
Masked 24-h hypertension	1 (ref)	1.34 (0.87–2.06)	1.69 (1.10–2.60)	2.36 (1.60–3.48)	<0.001

ASCVD indicates atherosclerotic cardiovascular disease; and CI, confidence interval.

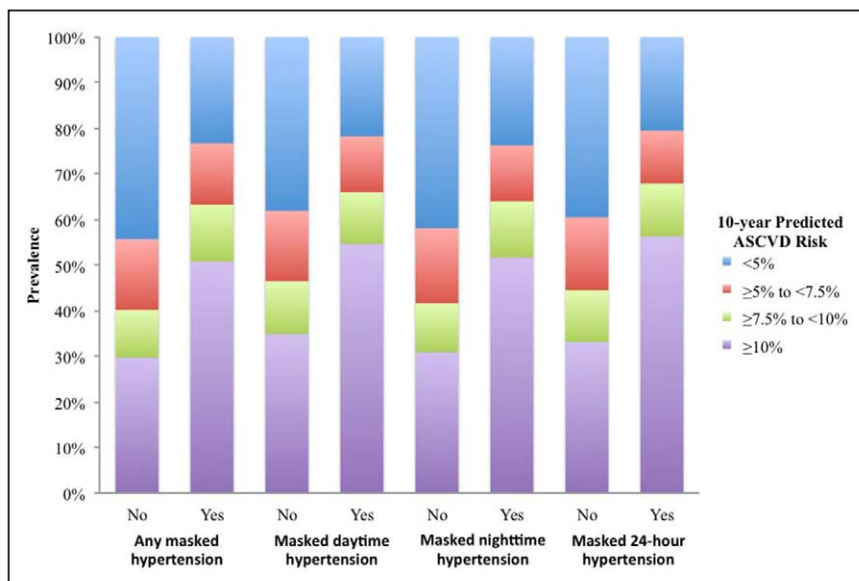
\*Adjusted for age, sex, body mass index, less than high school education, and estimated glomerular filtration rate <60 mL/min/m.

hypertension (51.7% versus 30.9%), and masked 24-hour hypertension (56.3% versus 33.1%; Figure 2; Table IV in the [Data Supplement](#)). Among participants not taking and taking antihypertensive medication, separately, the percentage in the ≥7.5% ASCVD risk category was higher among those with versus without each type of masked hypertension (Table V in the [Data Supplement](#); Figure I in the [Data Supplement](#)).

### ASCVD Risk Category and Masked Hypertension, Stratified by Prehypertension Status

The prevalence of 10-year predicted ASCVD risk ≥10% was 25.6% and 50.0% for participants with normal clinic

BP and prehypertension, respectively. Among participants with normal clinic BP, as compared with 10-year predicted ASCVD risk <5%, the adjusted prevalence ratio (95% CI) for having any masked hypertension was 1.73 (1.01–2.97), 1.88 (1.00–3.54), and 2.22 (1.24–3.99) for 10-year predicted ASCVD risk ≥5% to <7.5%, ≥7.5% to <10%, and ≥10%, respectively ( $P$ -trend=0.015; Table VI in the [Data Supplement](#)). Similar findings were observed for masked nighttime hypertension but not masked daytime hypertension or masked 24-hour hypertension. Among participants with prehypertension, the adjusted prevalence ratio (95% CI) for having any masked hypertension was 1.06 (0.77–1.44), 1.29 (0.94–1.75),



**Figure 2.** Distribution of 10-y predicted atherosclerotic cardiovascular disease (ASCVD) risk categories among Jackson Heart Study participants with and without masked hypertension. The prevalence data, depicted here, are presented in Table 4 in the [Data Supplement](#).

and 1.42 (1.07–1.89) for 10-year predicted ASCVD risk  $\geq 5\%$  to  $<7.5\%$ ,  $\geq 7.5\%$  to  $<10\%$ , and  $\geq 10\%$ , respectively ( $P$ -trend=0.007). Similar findings were observed for masked daytime hypertension and masked 24-hour hypertension but not masked nighttime hypertension.

### ASCVD Risk Category and Clinic BP for Detecting Masked Hypertension

The prevalence ratios for any masked hypertension were larger for each SD higher clinic SBP (1.31; 95% CI, 1.21–1.41) compared with each SD higher ASCVD risk score (1.23; 95% CI, 1.14–1.33) or clinic DBP (1.13; 95% CI, 1.05–1.22; Table 3). Similar findings were observed for masked daytime hypertension, masked nighttime hypertension, and masked 24-hour hypertension. The multivariable-adjusted C statistic for masked 24-hour hypertension was higher for clinic SBP and DBP versus ASCVD risk score (difference=0.036; 95% CI, 0.002–0.073) and lower for clinic DBP alone versus ASCVD risk score (difference=–0.033; 95% CI, –0.070 to –0.001; Table 4). There was no statistically significant difference in the C statistics for masked 24-hour hypertension associated with clinic SBP alone versus ASCVD risk score. Also, there were no statistically significant differences in the C statistic for any masked hypertension, masked daytime hypertension, and masked nighttime hypertension for clinic SBP and DBP, clinic SBP alone, and clinic DBP alone versus ASCVD risk score. There was no statistically significant difference in the C statistic for clinic SBP alone versus clinic SBP and DBP in detecting any masked hypertension, masked daytime hypertension, masked nighttime hypertension, or masked 24-hour hypertension. The C statistics for any masked hypertension, masked daytime hypertension, masked nighttime hypertension, and masked 24-hour hypertension were each lower for clinic DBP alone versus clinic SBP and DBP (difference=–0.050, 95% CI [–0.088 to –0.024]; –0.070, 95% CI [–0.111 to –0.036]; –0.043, 95% CI [–0.077 to –0.017]; and –0.069, 95% CI [–0.111 to –0.039], respectively).

The C statistics for masked hypertension among participants not taking and, separately, taking antihypertensive medication are shown in Table VII in the [Data Supplement](#). Among participants not taking antihypertensive medication, the C statistics for any masked hypertension and masked daytime hypertension were higher for clinic SBP and DBP versus ASCVD risk score (difference=0.085; 95% CI, 0.037–0.150 and 0.039; 95% CI, 0.000–0.030, respectively). The C statistic

for any masked hypertension was also higher for clinic SBP alone versus ASCVD risk score (difference=0.085; 95% CI, 0.038–0.154). Among participants taking antihypertensive medication, the C statistics for masked daytime hypertension, masked nighttime hypertension, and masked 24-hour hypertension were higher for both clinic SBP and DBP (difference=0.039, 95% CI [0.041–0.095]; 0.025, 95% CI [0.013–0.064]; and 0.036, 95% CI [0.031–0.076], respectively) and clinic SBP alone (difference=0.039, 95% CI [0.043–0.094]; 0.025, 95% CI [0.016–0.053]; and 0.035, 95% CI [0.034–0.071], respectively) versus the ASCVD risk score.

### Discussion

In this population-based sample of blacks, higher 10-year predicted ASCVD risk was associated with a higher prevalence of any masked hypertension, masked daytime hypertension, masked nighttime hypertension, and masked 24-hour hypertension. These associations were consistent for participants taking and not taking antihypertensive medication, and among those with prehypertension and normal clinic BP. Also, a majority of participants with each type of masked hypertension had a 10-year predicted ASCVD risk  $\geq 10\%$ . The C statistic for discriminating between participants with versus without masked hypertension was similar or lower for predicted ASCVD risk when compared with clinic SBP and clinic DBP.

Previous studies of European and Japanese cohorts have demonstrated a high prevalence of masked hypertension and a strong association between masked hypertension and ASCVD events.<sup>1,2,4–6</sup> Similar observations have been made in the JHS.<sup>3</sup> In the current study, among participants with clinic SBP/DBP  $<140/90$  mm Hg, there was a high prevalence of masked hypertension, ranging from 31.5% to 54.0% depending on the type of masked hypertension being evaluated. There may be a role for masked hypertension screening using ABPM among blacks given its high prevalence and strong association with ASCVD events in this population.

The optimal approach for identifying whom to screen for masked hypertension using ABPM is unknown. Several recent position papers, scientific statements, and guidelines recommend that ABPM should be used to assess out-of-clinic BP.<sup>17,22–25</sup> However, recommendations on who to screen with ABPM to identify masked hypertension were not provided. One approach may be to perform ABPM among all

**Table 3. Adjusted Prevalence Ratios for Masked Hypertension Associated With 1 SD Higher ASCVD Risk, Clinic Systolic Blood Pressure, and Clinic Diastolic Blood Pressure**

	Adjusted Prevalence Ratio (95% CI)*			
	Any Masked Hypertension	Masked Daytime Hypertension	Masked Nighttime Hypertension	Masked 24-h Hypertension
10-y ASCVD risk (SD=8.3%)	1.23 (1.14–1.33)	1.48 (1.32–1.67)	1.22 (1.12–1.33)	1.42 (1.27–1.58)
Clinic SBP (SD=10.4 mm Hg)	1.31 (1.21–1.41)	1.64 (1.44–1.87)	1.30 (1.19–1.42)	1.57 (1.39–1.77)
Clinic DBP (SD=7.4 mm Hg)	1.13 (1.05–1.22)	1.22 (1.08–1.39)	1.13 (1.04–1.23)	1.18 (1.05–1.33)

ASCVD indicates atherosclerotic cardiovascular disease; CI, confidence interval; DBP, diastolic blood pressure; and SBP, systolic blood pressure.

\*Adjusted for age, sex, body mass index, less than high school education, and estimated glomerular filtration rate  $<60$  mL/min/m<sup>2</sup>.

**Table 4. Adjusted C statistics\* for Masked Hypertension Associated With 10-Year Predicted ASCVD Risk, Clinic Systolic BP and Diastolic BP, Clinic Systolic BP Alone, and Clinic Diastolic BP Alone**

Outcome	Adjusted C Statistic (95% CI)*				Difference in C Statistic (95% CI)				
	ASCVD Risk Score	Clinic SBP and DBP	Clinic SBP	Clinic DBP	Clinic SBP and DBP vs ASCVD Risk Score	Clinic SBP vs ASCVD Risk Score	Clinic DBP vs ASCVD Risk Score	Clinic SBP vs Clinic SBP and DBP	Clinic DBP vs Clinic SBP and DBP
Any masked hypertension	0.681 (0.640–0.723)	0.703 (0.663–0.744)	0.703 (0.662–0.744)	0.653 (0.611–0.696)	0.022 (–0.014 to 0.056)	0.022 (–0.012 to 0.057)	–0.028 (–0.67 to 0.002)	0.000 (–0.003 to 0.002)	–0.050 (–0.088 to –0.024)†
Masked daytime hypertension	0.682 (0.636–0.728)	0.721 (0.680–0.763)	0.721 (0.679–0.762)	0.651 (0.606–0.697)	0.039 (–0.002 to 0.081)	0.039 (–0.001 to 0.080)	–0.031 (–0.074 to 0.004)	0.000 (–0.007 to 0.002)	–0.070 (–0.111 to –0.036)†
Masked nighttime hypertension	0.664 (0.621–0.706)	0.689 (0.648–0.730)	0.689 (0.648–0.730)	0.646 (0.603–0.689)	0.025 (–0.007 to 0.060)	0.025 (–0.005 to 0.060)	–0.018 (–0.051 to 0.017)	0.000 (–0.001 to 0.002)	–0.043 (–0.077 to –0.017)†
Masked 24-h hypertension	0.696 (0.652–0.739)	0.732 (0.691–0.772)	0.730 (0.690–0.771)	0.662 (0.618–0.707)	0.036 (0.002–0.073)†	0.035 (0.000–0.072)	–0.033 (–0.070 to –0.001)†	–0.001 (–0.011 to 0.002)	–0.069 (–0.111 to –0.039)†

ASCVD indicates atherosclerotic cardiovascular disease; CI, confidence interval; DBP, diastolic blood pressure; and SBP, systolic blood pressure.

\*Adjusted for age, sex, body mass index, less than high school education, and estimated glomerular filtration rate <60 mL/min/m<sup>2</sup>.

†Statistically significant differences vs ASCVD risk score.

individuals with clinic SBP/DBP <140/90 mm Hg. We previously estimated that 153 million US adults would have to be screened using this approach,<sup>26</sup> making it impractical. Another strategy is to perform ABPM in individuals who have risk factors for masked hypertension. Previous studies have shown that among individuals with clinic SBP/DBP <140/90 mm Hg, higher clinic BP, particularly in the prehypertension range, is a strong predictor of masked hypertension.<sup>1,6,20,27</sup> In the current analysis, higher 10-year predicted ASCVD risk, estimated with the pooled cohort risk equations, was associated with a higher prevalence of masked hypertension. However, the C statistic for discriminating between participants with versus without masked hypertension was similar or lower using predicted ASCVD risk when compared with clinic SBP and clinic DBP. Despite the prior evidence that several ASCVD risk factors are associated with masked hypertension,<sup>1,2,21,28,29</sup> clinic BP, particularly in the prehypertension range,<sup>1,6,20,26</sup> may be sufficient for identifying individuals with masked hypertension.

In the 2013 guidelines for the management of arterial hypertension, the European Society of Hypertension and European Society of Cardiology recommended that lifestyle measures and antihypertensive drug treatment should be considered for individuals with masked hypertension (class IIa recommendation, level C evidence).<sup>30</sup> However, there have been no randomized trials evaluating whether these interventions reduce the risk of ASCVD events among individuals with masked hypertension. There is increasing evidence that predicted ASCVD risk should be considered when deciding when to initiate or intensify hypertension treatment.<sup>31,32</sup> In the current study, the majority (50.9%) of participants with any masked hypertension were in the highest (≥10%) ASCVD risk category. Individuals with masked hypertension and elevated ASCVD risk might derive a greater absolute cardiovascular benefit from treatment compared with individuals with masked hypertension and low ASCVD risk.<sup>31</sup> Therefore, using elevated ASCVD risk as a screening criterion for conducting ABPM may identify the majority of individuals with masked hypertension and the majority of those likely to have ASCVD events.<sup>19</sup> We acknowledge that a large percentage (38.2%)

of participants with the lowest ASCVD risk (<5%) had any masked hypertension. It remains unknown whether masked hypertension is associated with an increased risk of ASCVD events among individuals with low predicted ASCVD risk. It is likely that antihypertensive medication treatment will have limited benefit in this group given their low predicted ASCVD risk. This issue has important implications for whether ABPM should be performed for individuals with low ASCVD risk to identify masked hypertension and underscores the need for further research in this area.

There are several strengths of the current study. We used data from JHS—a population-based cohort—comprised of blacks with a broad range of ASCVD risk. The JHS includes one of the largest samples of ABPM conducted among blacks, and ABPM and clinic BP were conducted following standardized protocols. The large sample size allowed us to evaluate the association between ASCVD risk categories and masked hypertension among participants with normal clinic BP and prehypertension, separately, and by antihypertensive medication use. There are also several potential limitations to the current study. Participants in the JHS underwent only 1 ABPM session, and, therefore, we cannot exclude the possibility that the results would have differed with the inclusion of additional 24-hour periods of ABPM. Without additional ABPM recordings before visit 1, it is also not possible to determine how long participants had masked hypertension. Further, clinic BP was measured using a manual device. However, for the current analysis, clinic BP values were calibrated with an oscillometric device. Finally, there was a relatively small sample size in some of the ASCVD risk categories in the subgroup analyses.

In conclusion, among blacks without BP in the hypertensive range based on clinic measurements, higher ASCVD risk was associated with a higher prevalence of any masked hypertension, masked daytime hypertension, masked nighttime hypertension, and masked 24-hour hypertension. Over 50% of participants with each type of masked hypertension had a 10-year predicted ASCVD risk ≥10%. The C statistic discriminating between individuals with and without masked hypertension associated with ASCVD risk was equivalent to or lower than clinic-measured BP. However, risk prediction equations may be useful



for identifying individuals with both masked hypertension and increased ASCVD risk who may derive the greatest absolute risk reduction from antihypertensive treatment.

### Acknowledgments

We wish to thank the staff and participants of the JHS (Jackson Heart Study).

### Sources of Funding

The JHS (Jackson Heart Study) is supported and conducted in collaboration with Jackson State University (HHSN268201300049C and HHSN268201300050C); University of Mississippi Medical Center (HHSN268201300046C and HHSN268201300047C); and Touglao College (HHSN268201300048C) contracts from the National Heart, Lung, and Blood Institute (NHLBI) and the National Center on Minority Health and Health Disparities at the National Institute of Health. The current study is also supported by R01 HL117323 from the NHLBI. Dr Anstey receives support through 2T32HL007854-21. J.N. Booth III receives support through F31 HL129701 from the NHLBI. Dr Abdalla receives support through HL117323-02S2. Dr Shimbo receives support through K24-HL125704 and P01-HL047540 from the NHLBI. Drs Muntner and Shimbo receive support through 15SFRN2390002 from the American Heart Association. The views expressed in this article are those of the authors and do not necessarily represent the views of the National Heart, Lung, and Blood Institute; the National Institutes of Health; or the US Department of Health and Human Services.

### Disclosures

P. Muntner received an institutional grant from Amgen, Inc, unrelated to the topic of the current article. D. Shimbo is a consultant for Abbott Vascular and Novartis Pharmaceuticals Corp. The other authors report no conflicts.

### References

- Peacock J, Diaz KM, Viera AJ, Schwartz J, Shimbo D. Unmasking masked hypertension: prevalence, clinical implications, diagnosis, correlates, and future directions. *J Hum Hypertens*. 2014;28:521–528. doi: 10.1038/jhh.2014.9.
- Angeli F, Reboldi G, Verdecchia P. Masked hypertension: evaluation, prognosis, and treatment. *Am J Hypertens*. 2010;23:941–948. doi: 10.1038/ajh.2010.112.
- Booth JN 3rd, Diaz KM, Seals SR, Sims M, Ravenell J, Muntner P, Shimbo D. Masked hypertension and cardiovascular disease events in a prospective cohort of blacks: the Jackson heart study. *Hypertension*. 2016;68:501–510. doi: 10.1161/HYPERTENSIONAHA.116.07553.
- Fagard RH, Cornelissen VA. Incidence of cardiovascular events in white-coat, masked and sustained hypertension versus true normotension: a meta-analysis. *J Hypertens*. 2007;25:2193–2198. doi: 10.1097/HJH.0b013e3282ef6185.
- Hansen TW, Kikuya M, Thijs L, Bjorklund-Bodegard K, Kuznetsova T, Ohkubo T, Richart T, Torp-Pedersen C, Lind L, Jeppesen J, Ibsen H, Imai Y, Staessen JA; IDACO Investigators. Prognostic superiority of daytime ambulatory over conventional blood pressure in four populations: a meta-analysis of 7,030 individuals. *J Hypertens*. 2007;25:1554–1564. doi: 10.1097/HJH.0b013e3281c49da5.
- Redmond N, Booth JN 3rd, Tanner RM, Diaz KM, Abdalla M, Sims M, Muntner P, Shimbo D. Prevalence of masked hypertension and its association with subclinical cardiovascular disease in African Americans: results from the Jackson heart study. *J Am Heart Assoc*. 2016;5:e002284. doi: 10.1161/JAHA.115.002284.
- Goff DC Jr, Lloyd-Jones DM, Bennett G, Coady S, D'Agostino RB Sr, Gibbons R, Greenland P, Lackland DT, Levy D, O'Donnell CJ, Robinson JG, Schwartz JS, Shero ST, Smith SC Jr, Sorlie P, Stone NJ, Wilson PW; American College of Cardiology/American Heart Association Task Force on Practice Guidelines. 2013 ACC/AHA guideline on the assessment of cardiovascular risk: a report of the American College of Cardiology/American Heart Association Task Force on Practice Guidelines. *J Am Coll Cardiol*. 2014;63:2935–2959. doi: 10.1016/j.jacc.2013.11.005.
- Taylor HA Jr. The Jackson heart study: an overview. *Ethn Dis*. 2005;15(4 suppl 6):S6-1–3.
- Wilson JG, Rotimi CN, Ekinwe L, Royal CD, Crump ME, Wyatt SB, Steffes MW, Adeyemo A, Zhou J, Taylor HA Jr, Jaquish C. Study design for genetic analysis in the Jackson heart study. *Ethn Dis*. 2005;15(4 suppl 6):S6-30–37.
- Taylor HA Jr, Wilson JG, Jones DW, Sarpong DF, Srinivasan A, Garrison RJ, Nelson C, Wyatt SB. Toward resolution of cardiovascular health disparities in African Americans: design and methods of the Jackson heart study. *Ethn Dis*. 2005;15(4 suppl 6):S6-4–17.
- Thijs L, Hansen TW, Kikuya M, Bjorklund-Bodegard K, Li Y, Dolan E, Tikhonoff V, Seidlerova J, Kuznetsova T, Stolarz K, Bianchi M, Richart T, Casiglia E, Malyutina S, Filipovsky J, Kawecka-Jaszcz K, Nikitin Y, Ohkubo T, Sandoya E, Wang J, Torp-Pedersen C, Lind L, Ibsen H, Imai Y, Staessen JA, O'Brien E; IDACO Investigators. The international database of ambulatory blood pressure in relation to cardiovascular outcome (IDACO): protocol and research perspectives. *Blood Press Monit*. 2007;12:255–262.
- Carpenter MA, Crow R, Steffes M, Rock W, Heilbraun J, Evans G, Skelton T, Jensen R, Sarpong D. Laboratory, reading center, and coordinating center data management methods in the Jackson Heart Study. *Am J Med Sci*. 2004;328:131–144.
- Levey AS, Stevens LA, Schmid CH, Zhang YL, Castro AF 3rd, Feldman HI, Kusek JW, Eggers P, Van Lente F, Greene T, Coresh J; CKD-EPI (Chronic Kidney Disease Epidemiology Collaboration). A new equation to estimate glomerular filtration rate. *Ann Intern Med*. 2009;150:604–612.
- Barker MH, Erlanger J, Meakins J, Schneider R, Scholz SB, Ungerleider H, White PD, Wiggers C, Wright I, Bramwell C, Cotton TF, Evans W, Gilchrist AR, Hay J, Campbell M, Pressure CSB, Pressure CSB. Standard method for taking and recording blood pressure readings. *J Amer Med Assoc*. 1939;113:294–297.
- Jackson Heart Study. Jackson heart study protocol, manual 4, blood pressure, visit 1. 2001. [https://www.jacksonheartstudy.org/jhsinfo/Portals/0/pdf/manuals1/Blood\\_pressure\\_manual4\\_02-18-2001\(1\).pdf](https://www.jacksonheartstudy.org/jhsinfo/Portals/0/pdf/manuals1/Blood_pressure_manual4_02-18-2001(1).pdf). Accessed July 13, 2016.
- Abdalla M, Booth JN 3rd, Seals SR, Spruill TM, Viera AJ, Diaz KM, Sims M, Muntner P, Shimbo D. Masked hypertension and incident clinic hypertension among blacks in the Jackson heart study. *Hypertension*. 2016;68:220–226. doi: 10.1161/HYPERTENSIONAHA.115.06904.
- O'Brien E, Parati G, Stergiou G, Asmar R, Beilin L, Bilo G, Clement D, de la Sierra A, de Leeuw P, Dolan E, Fagard R, Graves J, Head GA, Imai Y, Kario K, Lurbe E, Mallion JM, Mancia G, Mengden T, Myers M, Ogedegbe G, Ohkubo T, Omboni S, Palatini P, Redon J, Ruilope LM, Shennan A, Staessen JA, vanMontfrans G, Verdecchia P, Waerber B, Wang J, Zanchetti A, Zhang Y, European Society of Hypertension Working Group on Blood Pressure Monitoring. European society of hypertension position paper on ambulatory blood pressure monitoring. *J Hypertens*. 2013;31:1731–1768. doi: 10.1097/HJH.0b013e328363e964.
- Shimbo D, Abdalla M, Falzon L, Townsend RR, Muntner P. Role of ambulatory and home blood pressure monitoring in clinical practice: a narrative review. *Ann Intern Med*. 2015;163:691–700. doi: 10.7326/M15-1270.
- Muntner P, Colantonio LD, Cushman M, Goff DC Jr, Howard G, Howard VJ, Kissela B, Levitan EB, Lloyd-Jones DM, Safford MM. Validation of the atherosclerotic cardiovascular disease pooled cohort risk equations. *JAMA*. 2014;311:1406–1415. doi: 10.1001/jama.2014.2630.
- Shimbo D, Newman JD, Schwartz JE. Masked hypertension and prehypertension: diagnostic overlap and interrelationships with left ventricular mass: the masked hypertension study. *Am J Hypertens*. 2012;25:664–671. doi: 10.1038/ajh.2012.15.
- Diaz KM, Veerabhadrapa P, Brown MD, Whited MC, Dubbert PM, Hickson DA. Prevalence, determinants, and clinical significance of masked hypertension in a population-based sample of African Americans: the Jackson heart study. *Am J Hypertens*. 2015;28:900–908. doi: 10.1093/ajh/hpu241.
- Hackam DG, Quinn RR, Ravani P, Rabi DM, Dasgupta K, Daskalopoulou SS, Khan NA, Herman RJ, Bacon SL, Cloutier L, Dawes M, Rabkin SW, Gilbert RE, Ruzicka M, McKay DW, Campbell TS, Grover S, Honos G, Schiffrin EL, Bolli P, Wilson TW, Feldman RD, Lindsay P, Hill MD, Gelfer M, Burns KD, Vallee M, Prasad GV, Lebel M, McLean D, Arnold JM, Moe GW, Howlett JG, Boulanger JM, Laroche P, Leiter LA, Jones C, Ogilvie RI, Woo V, Kaczorowski J, Trudeau L, Petrella RJ, Milot A, Stone JA, Drouin D, Lavoie KL, Lamarre-Cliche M, Godwin M, Tremblay G, Hamet P, Fodor G, Carruthers SG, Pylypchuk GB, Burgess E, Lewanczuk R, Dresser GK, Penner SB, Hegele RA, McFarlane PA, Sharma M, Reid DJ, Tobe SW, Poirier L, Padwal RS; Canadian Hypertension Education Program. The 2013 Canadian hypertension education program



- recommendations for blood pressure measurement, diagnosis, assessment of risk, prevention, and treatment of hypertension. *Can J Cardiol*. 2013;29:528–542. doi: 10.1016/j.cjca.2013.01.005.
23. National Clinical Guideline Centre (UK). *Hypertension: The Clinical Management of Primary Hypertension in Adults: Update of Clinical Guidelines 18 and 34*. London, UK: Royal College of Physicians; 2011.
  24. Redon J, Lurbe E. Ambulatory blood pressure monitoring is ready to replace clinic blood pressure in the diagnosis of hypertension: con side of the argument. *Hypertension*. 2014;64:1169–1174; discussion 1174. doi: 10.1161/HYPERTENSIONAHA.114.03883.
  25. Siu AL; U.S. Preventive Services Task Force. Screening for high blood pressure in adults: U.S. preventive services task force recommendation statement. *Ann Intern Med*. 2015;163:778–786. doi: 10.7326/M15-2223.
  26. Booth JN 3rd, Muntner P, Diaz KM, Viera AJ, Bello NA, Schwartz JE, Shimbo D. Evaluation of criteria to detect masked hypertension. *J Clin Hypertens (Greenwich)*. 2016;18:1086–1094. doi: 10.1111/jch.12830.
  27. Alwan H, Pruijm M, Ponte B, Ackermann D, Guessous I, Ehret G, Staessen JA, Asayama K, Vuistiner P, Younes SE, Paccaud F, Wuerzner G, Pechere-Bertschi A, Mohaupt M, Vogt B, Martin PY, Burnier M, Bochud M. Epidemiology of masked and white-coat hypertension: the family-based SKIPOGH study. *PLoS One*. 2014;9:e92522. doi: 10.1371/journal.pone.0092522.
  28. Verberk WJ, Kessels AG, de Leeuw PW. Prevalence, causes, and consequences of masked hypertension: a meta-analysis. *Am J Hypertens*. 2008;21:969–975. doi: 10.1038/ajh.2008.221.
  29. Sobrino J, Domenech M, Camafort M, Vinyoles E, Coca A; ESTHEN group investigators. Prevalence of masked hypertension and associated factors in normotensive healthcare workers. *Blood Press Monit*. 2013;18:326–331. doi: 10.1097/MBP.0000000000000002.
  30. Mancia G, Fagard R, Narkiewicz K, Redon J, Zanchetti A, Böhm M, Christiaens T, Cifkova R, De Backer G, Dominiczak A, Galderisi M, Grobbee DE, Jaarsma T, Kirchhof P, Kjeldsen SE, Laurent S, Manolis AJ, Nilsson PM, Ruilope LM, Schmieder RE, Sirnes PA, Sleight P, Viigimaa M, Waeber B, Zannad F, Redon J, Dominiczak A, Narkiewicz K, Nilsson PM, Burnier M, Viigimaa M, Ambrosioni E, Caulfield M, Coca A, Olsen MH, Schmieder RE, Tsioufis C, van de Borne P, Zamorano JL, Achenbach S, Baumgartner H, Bax JJ, Bueno H, Dean V, Deaton C, Erol C, Fagard R, Ferrari R, Hasdai D, Hoes AW, Kirchhof P, Knuuti J, Kolh P, Lancellotti P, Linhart A, Nihoyannopoulos P, Piepoli MF, Ponikowski P, Sirnes PA, Tamargo JL, Tenders M, Torbicki A, Wijns W, Windecker S, Clement DL, Coca A, Gillebert TC, Tendera M, Rosei EA, Ambrosioni E, Anker SD, Bauersachs J, Hitij JB, Caulfield M, De Buyzere M, De Geest S, Derumeaux GA, Erdine S, Farsang C, Funck-Brentano C, Gerc V, Germano G, Gielen S, Haller H, Hoes AW, Jordan J, Kahan T, Komajda M, Lovic D, Mahrholdt H, Olsen MH, Ostergren J, Parati G, Perk J, Polonia J, Popescu BA, Reiner Z, Rydén L, Sirenko Y, Stanton A, Struijker-Boudier H, Tsioufis C, van de Borne P, Vlachopoulos C, Volpe M, Wood DA. 2013 ESH/ESC guidelines for the management of arterial hypertension: the task force for the management of arterial hypertension of the European society of hypertension (ESH) and of the European society of cardiology (ESC). *Eur Heart J*. 2013;34:2159–2219. doi: 10.1093/eurheartj/ehf151.
  31. Jackson R, Lawes CM, Bennett DA, Milne RJ, Rodgers A. Treatment with drugs to lower blood pressure and blood cholesterol based on an individual's absolute cardiovascular risk. *Lancet*. 2005;365:434–441. doi: 10.1016/S0140-6736(05)17833-7.
  32. Blood Pressure Lowering Treatment Trialists' Collaboration; Sundstrom J, Arima H, Woodward M, Jackson R, Karmali K, Lloyd-Jones D, Baigent C, Emberson J, Rahimi K, MacMahon S, Patel A, Perkovic V, Turnbull F, Neal B. Blood pressure-lowering treatment based on cardiovascular risk: a meta-analysis of individual patient data. *Lancet*. 2014;384:591–598. doi: 10.1016/S0140-6736(14)61212-5.

## Predicted Atherosclerotic Cardiovascular Disease Risk and Masked Hypertension Among Blacks in the Jackson Heart Study

D. Edmund Anstey, John N. Booth III, Marwah Abdalla, Tanya M. Spruill, Yuan-I Min, Paul Muntner and Daichi Shimbo

*Circ Cardiovasc Qual Outcomes.* 2017;10:

doi: 10.1161/CIRCOUTCOMES.116.003421

*Circulation: Cardiovascular Quality and Outcomes* is published by the American Heart Association, 7272 Greenville Avenue, Dallas, TX 75231

Copyright © 2017 American Heart Association, Inc. All rights reserved.

Print ISSN: 1941-7705. Online ISSN: 1941-7713

The online version of this article, along with updated information and services, is located on the World Wide Web at:

<http://circoutcomes.ahajournals.org/content/10/7/e003421>

Data Supplement (unedited) at:

<http://circoutcomes.ahajournals.org/content/suppl/2017/07/11/CIRCOUTCOMES.116.003421.DC1>

**Permissions:** Requests for permissions to reproduce figures, tables, or portions of articles originally published in *Circulation: Cardiovascular Quality and Outcomes* can be obtained via RightsLink, a service of the Copyright Clearance Center, not the Editorial Office. Once the online version of the published article for which permission is being requested is located, click Request Permissions in the middle column of the Web page under Services. Further information about this process is available in the [Permissions and Rights Question and Answer](#) document.

**Reprints:** Information about reprints can be found online at:

<http://www.lww.com/reprints>

**Subscriptions:** Information about subscribing to *Circulation: Cardiovascular Quality and Outcomes* is online at:

<http://circoutcomes.ahajournals.org/subscriptions/>

## **SUPPLEMENTAL MATERIAL**

### **Predicted Atherosclerotic Cardiovascular Disease Risk and Masked Hypertension Among African Americans in the Jackson Heart Study**

Short title: Cardiovascular disease risk and masked hypertension

D. Edmund Anstey, MD<sup>1</sup>, John N. Booth, III<sup>2</sup>, MS, Marwah Abdalla, MD, MPH<sup>1</sup>, Tanya M. Spruill, PhD<sup>3</sup>, Yuan-I Min, PhD<sup>4</sup>, Paul Muntner<sup>2</sup>, PhD, Daichi Shimbo, MD<sup>1</sup>

<sup>1</sup>Columbia University Medical Center, New York, New York

<sup>2</sup>University of Alabama at Birmingham, Birmingham, Alabama

<sup>3</sup>New York University School of Medicine, New York, New York

<sup>4</sup>School of Medicine, University of Mississippi Medical Center, Jackson, Mississippi

**SUPPLEMENTAL TABLES**

**Supplemental Table 1.** Clinic and ambulatory blood pressure levels among Jackson Heart Study participants by category of 10-year predicted atherosclerotic cardiovascular risk.

	10-year predicted ASCVD risk category				p-trend
	<5%	≥5% to <7.5%	≥7.5% to <10%	≥10%	
	(n=212)	(n=93)	(n=74)	(n=265)	
<b>SBP, mmHg</b>					
Clinic SBP	117.3 ± 10.2	121.2 ± 9.5	122.5 ± 10.6	125.9 ± 9.3	<0.001
Daytime SBP	121.9 ± 9.7	125.9 ± 10.9	126.7 ± 10.1	131.2 ± 12.4	<0.001
Nighttime SBP	111.7 ± 11.4	116.3 ± 13.2	118.0 ± 11.0	123.0 ± 15.1	<0.001
24-hour SBP	117.8 ± 9.7	122.3 ± 10.8	123.7 ± 9.9	128.0 ± 12.5	<0.001
<b>DBP, mmHg</b>					
Clinic DBP	74.2 ± 7.7	73.5 ± 7.5	72.5 ± 8.0	72.3 ± 6.9	0.003
Daytime DBP	77.7 ± 8.4	77.2 ± 7.7	76.6 ± 8.3	77.2 ± 9.5	0.473
Nighttime DBP	66.4 ± 8.7	66.2 ± 9.1	68.2 ± 8.5	68.3 ± 10.1	0.015
24-hour DBP	73.2 ± 8.0	72.7 ± 7.5	73.5 ± 7.9	73.5 ± 9.0	0.553
Prehypertension, %	47.2%	61.3%	64.9%	77.4%	<0.001

The numbers in the table are mean ± standard deviation or percentages.

ASCVD: Atherosclerotic cardiovascular disease

SBP: systolic blood pressure

DBP: diastolic blood pressure



**Supplemental Table 2.** Characteristics and clinic and ambulatory blood pressure among Jackson Heart Study participants by category of 10-year predicted atherosclerotic cardiovascular risk stratified by those not taking antihypertensive medication (upper panel) and those taking antihypertensive medication (lower panel).

	10-year predicted ASCVD risk category			p-trend
	<5%	≥5% to <7.5%	≥7.5%	
<b>Participants not taking antihypertensive medication</b>				
	(n=159)	(n=45)	(n=101)	
Age, years	49.3 ± 6.1	57.0 ± 6.1	66.0 ± 6.4	<0.001
Male, %	18.2%	62.2%	51.5%	<0.001
Less than high school education, %	4.4%	17.8%	21.8%	<0.001
Current smoking, %	6.9%	4.4%	13.9%	0.074
Diabetes, %	1.9%	4.4%	26.7%	<0.001
eGFR < 60 ml/min/m <sup>2</sup> , %	0.0%	2.2%	8.9%	0.014
Body mass index, kg/m <sup>2</sup>	30.7 ± 7.6	29.8 ± 6.0	29.8 ± 5.0	0.254
Total cholesterol, mg/dL	198.2 ± 36.5	206.5 ± 34.4	216.8 ± 42.0	<0.001
LDL cholesterol, mg/dL	124.3 ± 35.9	136.7 ± 33.2	139.8 ± 36.6	<0.001
HDL cholesterol, mg/dL	55.5 ± 13.2	49.4 ± 12.9	55.6 ± 16.2	0.832
<b>SBP, mmHg</b>				
Clinic SBP	116.9 ± 10.5	120.1 ± 9.6	126.1 ± 8.4	<0.001
Daytime SBP	121.5 ± 9.4	127.3 ± 10.2	130.4 ± 11.1	<0.001
Nighttime SBP	111.4 ± 10.7	117.0 ± 12.9	120.8 ± 13.5	<0.001
24-hour SBP	117.4 ± 9.4	123.5 ± 10.6	126.8 ± 11.4	<0.001
<b>DBP, mmHg</b>				
Clinic DBP	74.0 ± 7.6	73.4 ± 8.2	72.2 ± 6.0	0.060
Daytime DBP	77.3 ± 8.4	79.1 ± 6.9	77.4 ± 8.7	0.824
Nighttime DBP	66.1 ± 8.2	68.6 ± 9.1	68.6 ± 8.7	0.016
24-hour DBP	72.8 ± 7.9	74.8 ± 7.1	73.9 ± 7.9	0.226
Prehypertension, %	47.2%	57.8%	80.2%	<0.001
<b>Participants taking antihypertensive medication</b>				
	(n=53)	(n=48)	(n=238)	
Age, years	50.6 ± 5.9	58.5 ± 7.2	65.0 ± 7.2	<0.001
Male, %	1.9%	6.3%	33.6%	<0.001
Less than high school education, %	11.3%	20.8%	19.9%	0.209
Current smoking, %	1.9%	10.4%	8.0%	0.234
Diabetes, %	1.9%	8.3%	37.8%	<0.001
eGFR < 60 ml/min/m <sup>2</sup> , %	0.0%	8.3%	16.0%	0.004
Body mass index, kg/m <sup>2</sup>	32.6 ± 7.4	33.4 ± 5.7	31.1 ± 5.7	0.034
Total cholesterol, mg/dL	192.8 ± 32.2	193.7 ± 37.3	203.2 ± 40.9	0.041
LDL cholesterol, mg/dL	115.0 ± 31.5	115.4 ± 36.6	126.1 ± 35.2	0.015
HDL cholesterol, mg/dL	60.3 ± 19.5	56.1 ± 15.1	53.8 ± 13.6	0.004

SBP, mmHg				
Clinic SBP	118.4 ± 9.3	122.3 ± 9.4	124.8 ± 10.2	<0.001
Daytime SBP	123.0 ± 10.6	124.5 ± 11.4	130.1 ± 12.5	<0.001
Nighttime SBP	112.7 ± 13.3	115.7 ± 13.6	122.4 ± 14.9	<0.001
24-hour SBP	119.0 ± 10.6	121.2 ± 10.9	127.2 ± 12.4	<0.001
DBP, mmHg				
Clinic DBP	74.9 ± 8.0	73.7 ± 6.9	72.3 ± 7.6	0.018
Daytime DBP	79.1 ± 8.5	75.4 ± 8.0	76.9 ± 9.4	0.262
Nighttime DBP	67.3 ± 10.1	64.0 ± 8.6	68.1 ± 10.2	0.216
24-hour DBP	74.4 ± 8.2	70.7 ± 7.3	73.4 ± 9.1	0.911
Prehypertension, %	47.2%	64.6%	72.3%	<0.001

The numbers in the table are mean ± standard deviation or percentages.

ASCVD: Atherosclerotic cardiovascular disease

eGFR: estimated glomerular filtration rate

HDL: high-density lipoprotein

LDL: low-density lipoprotein

SBP: systolic blood pressure

DBP: diastolic blood pressure

**Supplemental Table 3.** Prevalence and prevalence ratios for masked hypertension by category of 10-year predicted atherosclerotic cardiovascular disease risk among participants not taking antihypertensive medication (upper panel) and taking antihypertensive medication (bottom panel).

	10-year predicted ASCVD risk category			
	<5%	≥5% to <7.5%	≥7.5%	p-trend
<b>Participants not taking antihypertensive medication</b>	(n=159)	(n=45)	(n=101)	
	Prevalence			
Any masked hypertension	35.2%	62.2%	63.4%	<0.001
Masked daytime hypertension	20.1%	31.1%	37.6%	0.002
Masked nighttime hypertension	33.3%	53.3%	61.4%	<0.001
Masked 24-hour hypertension	22.0%	40.0%	46.5%	<0.001
	Unadjusted prevalence ratio (95% CI)			
Any masked hypertension	1 (ref)	1.60 (1.13 - 2.27)	1.84 (1.41 - 2.41)	<0.001
Masked daytime hypertension	1 (ref)	1.55 (0.91 - 2.64)	1.87 (1.25 - 2.79)	0.002
Masked nighttime hypertension	1 (ref)	1.77 (1.30 - 2.41)	1.80 (1.39 - 2.33)	<0.001
Masked 24-hour hypertension	1 (ref)	1.82 (1.14 - 2.88)	2.11 (1.48 - 3.03)	<0.001
	Adjusted prevalence ratio (95% CI)*			
Any masked hypertension	1 (ref)	1.62 (1.07 - 2.46)	1.94 (1.22 - 3.10)	0.006
Masked daytime hypertension	1 (ref)	1.49 (0.79 - 2.80)	1.88 (0.97 - 3.67)	0.064
Masked nighttime hypertension	1 (ref)	1.85 (1.27 - 2.71)	2.09 (1.35 - 3.23)	0.001
Masked 24-hour hypertension	1 (ref)	2.12 (1.22 - 3.66)	2.65 (1.48 - 4.76)	0.001
<b>Participants taking antihypertensive medication</b>	(n=53)	(n=48)	(n=238)	
	Prevalence			
Any masked hypertension	47.2%	39.6%	65.5%	0.001
Masked daytime hypertension	22.6%	22.9%	40.3%	0.004
Masked nighttime hypertension	41.5%	31.3%	59.2%	0.002
Masked 24-hour hypertension	20.8%	16.7%	44.1%	<0.001
	Unadjusted prevalence ratio (95% CI)			
Any masked hypertension	1 (ref)	0.75 (0.44 - 1.28)	1.43 (1.02 - 2.00)	0.006

Masked daytime hypertension	1 (ref)	1.01 (0.49 - 2.08)	1.78 (1.06 - 3.00)	0.009
Masked nighttime hypertension	1 (ref)	0.84 (0.53 - 1.32)	1.39 (1.03 - 1.87)	0.005
Masked 24-hour hypertension	1 (ref)	0.80 (0.35 - 1.83)	2.13 (1.23 - 3.67)	<0.001
	Adjusted prevalence ratio (95% CI)*			
Any masked hypertension	1 (ref)	0.72 (0.43 - 1.22)	1.21 (0.81 - 1.81)	0.154
Masked daytime hypertension	1 (ref)	1.30 (0.63 - 2.67)	2.14 (1.17 - 3.94)	0.008
Masked nighttime hypertension	1 (ref)	0.89 (0.57 - 1.40)	1.38 (0.96 - 1.98)	0.031
Masked 24-hour hypertension	1 (ref)	0.84 (0.37 - 1.90)	1.86 (1.02 - 3.39)	0.015

ASCVD: Atherosclerotic cardiovascular disease

CI: Confidence interval

\*Adjusted for age, sex, body mass index, less than high school education, and estimated glomerular filtration rate <60 ml/min/m<sup>2</sup>



**Supplemental Table 4.** Distribution of 10-year predicted atherosclerotic cardiovascular disease risk categories among Jackson Heart Study participants with and without masked hypertension.

ASCVD risk category	Type of masked hypertension							
	Any		Daytime		Nighttime		24-hour	
	No	Yes	No	Yes	No	Yes	No	Yes
	n=441	n=203	n=327	n=317	n=420	n=224	n=296	n=348
<5%	44.3%	23.3%	38.1%	21.7%	41.9%	23.7%	39.5%	20.5%
≥5% to <7.5%	15.5%	13.5%	15.4%	12.3%	16.5%	12.3%	16.0%	11.6%
≥7.5% to <10%	10.5%	12.4%	11.6%	11.3%	10.7%	12.3%	11.4%	11.6%
≥10%	29.7%	50.9%	34.9%	54.7%	30.9%	51.7%	33.1%	56.3%

ASCVD: Atherosclerotic cardiovascular disease

**Supplemental Table 5.** Prevalence of 10-year predicted atherosclerotic cardiovascular disease risk categories among participants with and without masked hypertension stratified by those not taking antihypertensive medication (upper panel) and those taking antihypertensive medication (lower panel).

	Type of masked hypertension							
	Any		Daytime		Nighttime		24-hour	
	No	Yes	No	Yes	No	Yes	No	Yes
ASCVD risk category	Participants not taking antihypertensive medications							
	n=221	n=84	n=166	n=139	n=205	n=100	n=157	n=148
<5%	57.5%	38.1%	63.9%	38.1%	60.5%	35.0%	65.6%	37.8%
≥5% to <7.5%	14.0%	16.7%	12.7%	17.3%	13.2%	18.0%	10.8%	18.9%
≥7.5%	28.5%	45.2%	23.5%	44.6%	26.3%	47.0%	23.6%	43.2%
ASCVD risk category	Participants taking antihypertensive medications							
	n=220	n=119	n=161	n=178	n=215	n=124	n=139	n=200
<5%	18.6%	10.1%	19.3%	12.4%	19.5%	8.9%	20.1%	12.5%
≥5% to <7.5%	16.8%	9.2%	20.5%	8.4%	18.6%	6.5%	20.9%	9.5%
≥7.5%	64.5%	80.7%	60.2%	79.2%	61.9%	84.7%	59.0%	78.0%

ASCVD: Atherosclerotic cardiovascular disease

**Supplemental Table 6.** Prevalence and prevalence ratios for masked hypertension by category of atherosclerotic cardiovascular disease risk among Jackson Heart Study participants with normal clinic blood pressure (upper panel) and with prehypertension (upper panel).

	10-year predicted ASCVD risk category				p-trend
	<5%	≥5% to <7.5%	≥7.5% to <10%	≥10%	
<b>Among participants with normal clinic blood pressure</b>	n=112	n=36	n=26	n=60	
	Prevalence				
Any masked hypertension	25.9%	44.4%	46.2%	53.3%	<0.001
Masked daytime hypertension	8.9%	22.2%	23.1%	23.3%	0.001
Masked nighttime hypertension	23.2%	33.3%	38.5%	50.0%	<0.001
Masked 24-hour hypertension	9.8%	22.2%	23.1%	30.0%	0.001
	Unadjusted prevalence ratio (95% CI)				
Any masked hypertension	1 (ref)	1.72 (1.06 - 2.78)	1.78 (1.06 - 3.00)	2.06 (1.39 - 3.05)	<0.001
Masked daytime hypertension	1 (ref)	2.49 (1.06 - 5.83)	2.58 (1.03 - 6.47)	2.61 (1.24 - 5.52)	0.007
Masked nighttime hypertension	1 (ref)	1.44 (0.81 - 2.54)	1.66 (0.92 - 2.99)	2.15 (1.41 - 3.28)	<0.001
Masked 24-hour hypertension	1 (ref)	2.26 (0.99 - 5.19)	2.35 (0.96 - 5.77)	3.05 (1.55 - 6.04)	<0.001
	Adjusted prevalence ratio (95% CI)*				
Any masked hypertension	1 (ref)	1.73 (1.01 - 2.97)	1.88 (1.00 - 3.54)	2.22 (1.24 - 3.99)	0.015
Masked daytime hypertension	1 (ref)	2.73 (0.97 - 7.69)	2.72 (0.82 - 8.98)	3.35 (1.02 - 11.01)	0.109
Masked nighttime hypertension	1 (ref)	1.45 (0.78 - 2.70)	1.77 (0.87 - 3.58)	2.38 (1.27 - 4.45)	0.006
Masked 24-hour hypertension	1 (ref)	1.97 (0.80 - 4.86)	2.04 (0.68 - 6.09)	2.75 (1.01 - 7.45)	0.078
<b>Among participants with prehypertension</b>	n=100	n=57	n=48	n=205	
	Prevalence				
Any masked hypertension	52.0%	54.4%	64.6%	70.7%	0.001
Masked daytime hypertension	34.0%	29.8%	35.4%	47.3%	0.009
Masked nighttime hypertension	49.0%	47.4%	60.4%	65.4%	0.002
Masked 24-hour hypertension	35.0%	31.6%	41.7%	52.7%	0.001
	Unadjusted prevalence ratio (95% CI)				
Any masked hypertension	1 (ref)	1.05 (0.77 - 1.42)	1.24 (0.94 - 1.65)	1.36 (1.10 - 1.67)	0.001
Masked daytime hypertension	1 (ref)	0.88 (0.54 - 1.42)	1.04 (0.65 - 1.67)	1.39 (1.02 - 1.90)	0.012
Masked nighttime hypertension	1 (ref)	0.97 (0.69 - 1.36)	1.23 (0.91 - 1.67)	1.33 (1.07 - 1.67)	0.003

Masked 24-hour hypertension	1 (ref)	0.90 (0.57 - 1.44)	1.19 (0.78 - 1.83)	1.51 (1.12 - 2.03)	0.001
	Adjusted prevalence ratio (95% CI)*				
Any masked hypertension	1 (ref)	1.06 (0.77 - 1.44)	1.29 (0.94 - 1.75)	1.42 (1.07 - 1.89)	0.007
Masked daytime hypertension	1 (ref)	0.98 (0.60 - 1.61)	1.20 (0.73 - 1.97)	1.72 (1.11 - 2.66)	0.006
Masked nighttime hypertension	1 (ref)	0.89 (0.63 - 1.27)	1.11 (0.79 - 1.57)	1.13 (0.81 - 1.58)	0.282
Masked 24-hour hypertension	1 (ref)	0.94 (0.58 - 1.51)	1.22 (0.78 - 1.90)	1.55 (1.02 - 2.35)	0.013

ASCVD: Atherosclerotic cardiovascular disease

CI: Confidence interval

Normal clinic blood pressure is defined as mean clinic systolic blood pressure <120 mm Hg and mean clinic diastolic blood pressure <80 mm Hg. Prehypertension is defined as mean clinic systolic blood pressure 120 to 139 mm Hg or mean clinic diastolic blood pressure 80 to 89 mm Hg.

\*Adjusted for age, sex, body mass index, less than high school education, and estimated glomerular filtration rate <60 ml/min/m<sup>2</sup>



**Supplemental Table 7.** C-statistic for masked hypertension using 10-year predicted atherosclerotic cardiovascular disease (ASCVD) risk compared with clinic systolic blood pressure (SBP) and diastolic blood pressure (DBP), clinic SBP alone, and clinic DBP alone among Jackson Heart Study participants not taking and taking antihypertensive medication in adjusted models.

	Adjusted c-statistic*				Difference in c-statistic (95% CI)		
	10-year predicted ASCVD Risk Score	Clinic SBP and DBP	Clinic SBP	Clinic DBP	Clinic SBP and DBP versus ASCVD risk score†	Clinic SBP versus ASCVD risk score‡	Clinic DBP versus ASCVD risk score§
Participants not taking antihypertensive medication							
Any masked hypertension	0.655 (0.593 to 0.718)	0.740 (0.684 to 0.797)	0.740 (0.684 to 0.797)	0.656 (0.594 to 0.718)	<b>0.085</b> ( <b>0.037 to 0.150</b> )	<b>0.085</b> ( <b>0.038 to 0.154</b> )	0.001 (-0.054 to 0.056)
Masked daytime hypertension	0.682 (0.636 to 0.728)	0.721 (0.680 to 0.763)	0.721 (0.679 to 0.762)	0.651 (0.606 to 0.697)	<b>0.039</b> ( <b>0.000 to 0.030</b> )	0.039 (-0.006 to 0.031)	-0.031 (-0.107 to -0.013)
Masked nighttime hypertension	0.664 (0.621 to 0.706)	0.689 (0.648 to 0.730)	0.689 (0.648 to 0.730)	0.646 (0.603 to 0.689)	0.025 (-0.010 to 0.034)	0.025 (-0.010 to 0.037)	-0.018 (-0.095 to 0.009)
Masked 24-hour hypertension	0.696 (0.652 to 0.739)	0.732 (0.691 to 0.772)	0.730 (0.690 to 0.771)	0.662 (0.618 to 0.707)	0.036 (-0.010 to 0.044)	0.035 (-0.025 to 0.051)	-0.033 (-0.112 to 0.001)
Participants taking antihypertensive medication							
Any masked hypertension	0.697 (0.641 to 0.754)	0.689 (0.632 to 0.747)	0.690 (0.632 to 0.747)	0.664 (0.606 to 0.723)	-0.008 (-0.056 to 0.026)	-0.008 (-0.050 to 0.031)	-0.033 (-0.082 to 0.006)
Masked daytime hypertension	0.682 (0.636 to 0.728)	0.721 (0.680 to 0.763)	0.721 (0.679 to 0.762)	0.651 (0.606 to 0.697)	<b>0.039</b> ( <b>0.041 to 0.095</b> )	<b>0.039</b> ( <b>0.043 to 0.094</b> )	-0.031 (-0.061 to 0.022)
Masked nighttime hypertension	0.664 (0.621 to 0.706)	0.689 (0.648 to 0.730)	0.689 (0.648 to 0.730)	0.646 (0.603 to 0.689)	<b>0.025</b> ( <b>0.013 to 0.064</b> )	<b>0.025</b> ( <b>0.016 to 0.053</b> )	-0.018 (-0.050 to 0.028)
Masked 24-hour hypertension	0.696 (0.652 to 0.739)	0.732 (0.691 to 0.772)	0.730 (0.690 to 0.771)	0.662 (0.618 to 0.707)	<b>0.036</b> ( <b>0.031 to 0.076</b> )	<b>0.035</b> ( <b>0.034 to 0.071</b> )	-0.033 (-0.068 to 0.005)

Statistically significant differences versus ASCVD risk score are shown in bold.

ASCVD: Atherosclerotic cardiovascular disease

CI: Confidence interval

\*Adjusted for age, sex, body mass index, less than high school education, and estimated glomerular filtration rate <60 ml/min/m<sup>2</sup>

†Comparing clinic SBP and DBP to 10-year predicted ASCVD Risk Score

‡Comparing clinic SBP to 10-year predicted ASCVD Risk Score

§Comparing clinic DBP to 10-year predicted ASCVD Risk Score

SBP: Systolic blood pressure

DBP: Diastolic blood pressure

**SUPPLEMENTAL FIGURES**

**Supplemental Figure 1.** Prevalence of 10-year predicted atherosclerotic cardiovascular disease risk categories among participants with and without masked hypertension stratified by those not taking antihypertensive medication (upper panel) and those taking antihypertensive medication (lower panel). The prevalence data, depicted here, are presented in Supplemental Table 5.

