RESEARCH Open Access

Ambulatory hypertension diagnosed by 24-h mean ambulatory versus day and night ambulatory blood pressure thresholds in children: a cross-sectional study

Ajay P. Sharma^{1,2*}, Luis Altamirano-Diaz^{2,3}, Mohamed Mohamed Ali⁴, Katryna Stronks⁴, Amrit Kirpalani^{1,2}, Guido Filler^{1,2} and Kambiz Norozi^{2,3,5}

Abstract

Background: The agreement between the commonly used ambulatory blood pressure (ABP) thresholds to diagnose ambulatory hypertension in children (patient's 24-h mean ABP classified by 24-h 95th ABP percentile threshold, American Heart Association [AHA] threshold, or patient's day and night mean ABP classified by day-night 95th ABP percentile thresholds) is not known. We evaluated the agreement among 24-h ABP threshold, AHA threshold, and day-night ABP thresholds to diagnose ambulatory hypertension, white coat hypertension (WCH) and masked hypertension (MH).

Methods: In a cross-sectional study design, we analyzed ABP recordings from 450 participants with suspected hypertension from a tertiary care outpatient hypertension clinic. The American Academy of Pediatrics thresholds were used to diagnose office hypertension.

Results: The 24-h ABP threshold and day-night ABP thresholds classified 19% ABP (95% confidence interval [CI], 0.15–0.23) differently into ambulatory normotension/hypertension (kappa [κ], 0.58; 95% CI, 0.51–0.66). Ambulatory hypertension diagnosed by 24-h ABP threshold in 27% participants (95% CI, 0.22–0.32) was significantly lower than that by day-night ABP thresholds in 44% participants (95% CI, 0.37–0.50; P < 0.001). The AHA threshold had a stronger agreement with 24-h ABP threshold than with day-night ABP thresholds for classifying ABP into ambulatory normotension/hypertension (k 0.94, 95% CI 0.91–0.98 vs. k 0.59, 95% CI 0.52–0.66). The diagnosis of ambulatory hypertension by the AHA threshold (26%; 95% CI, 0.21–0.31) was closer to that by 24-h ABP threshold (27%, P = 0.73) than by day-night ABP thresholds (44%, P < 0.001). Similar agreement pattern persisted among these ABP thresholds for diagnosing WCH and MH.

Conclusions: The 24-h ABP threshold classifies a lower proportion of ABP as ambulatory hypertension than daynight ABP thresholds. The AHA threshold exhibits a stronger agreement with 24-h ABP than with day-night ABP thresholds for diagnosing ambulatory hypertension, WCH and MH. Our findings are relevant for a consistent interpretation of hypertension by these ABP thresholds in clinical practice.

¹ Division of Nephrology, Department of Paediatrics, University of Western Ontario, London, ON, Canada Full list of author information is available at the end of the article



^{*}Correspondence: ajay.sharma@lhsc.on.ca

Keywords: Pediatrics, Blood Pressure, Hypertension, White coat hypertension, Masked hypertension, Ambulatory Blood Pressure Monitoring

Background

The American Heart Association (AHA) and European Society of Hypertension recommend the use of ambulatory blood pressure monitoring (ABPM) in clinical practice to diagnose ambulatory hypertension [1, 2]. Ambulatory normotension or hypertension status helps to classify office blood pressure (OBP) into white coat hypertension (WCH) and masked hypertension (MH) [1–5]. In children, patient's 24-h/day/night mean ambulatory blood pressure (ABP) values are interpreted by the age-, sex-, and height-specific 24-h/day/night 95th systolic/diastolic ABP percentiles [6, 7].

Based on the assumption that 24-h ABP represents both day and night ABP, ambulatory hypertension in children is commonly diagnosed by either patient's 24-h mean ABP exceeding the 24-h 95th ABP percentile (24-h ABP threshold) or patient's mean day/night ABP exceeding the day/night 95th ABP percentiles, respectively (daynight ABP thresholds) [8–13]. The AHA guidelines have recommended a modified version of 24-h 95th ABP percentile (24-h 95th ABP percentile along with 24-h ABP load \geq 25%) [1, 3]. However, the agreement between 24-h ABP, day-night ABP, and AHA thresholds to diagnose ambulatory hypertension is not known.

Based on the observations that an individual can have a disconnect in the diagnosis of day and night ambulatory hypertension based on the corresponding day/night 95th ABP percentiles [14–18], we hypothesized that 24-h ABP threshold may diagnose ambulatory hypertension differently than day-night ABP thresholds. Therefore, our primary objective was to compare the diagnosis of ambulatory hypertension by the 24-h ABP and day-night ABP thresholds, as well to evaluate their relative agreement with the AHA threshold to diagnose ambulatory hypertension. Our secondary objective was to assess the agreement among these ABP thresholds to diagnose WCH and MH.

Methods

This was a single center, retrospective cross-sectional study performed after an approval by the Research Ethics Board of the University of Western Ontario (No. 16143E and 116,082). The study involved the retrospective review of existing clinical data and was therefore exempted from the need for an individual informed consent. The records of children who underwent ABPM at a tertiary care outpatient hypertension clinic (London, ON, Canada) were collected. The data was collected between January 2003

and December 2008 (n = 159) as a part of previous studies and recently between January 2018 and September 2020 (n=291) [19–23]. The participants were referred to our outpatient clinic at London Health Sciences Center because of suspected hypertension based on OBP assessments by the primary health care providers. During both the study periods, there was a uniformity in the protocol regarding offering ABPM to all patients older than 5 years and for evaluating secondary hypertension (the fourth report guidelines [24] during the first study period and similar recommendations by the American Academy of Pediatrics [AAP] guidelines during the second period [3]). The participants with an inadequate ABPM or missing OBP recordings were excluded. Anthropometric measurements including age, weight, and height were collected. Body mass index (BMI) was calculated as the weight (kg)/height (cm)2. BMI percentiles were calculated based on the Centers for Disease Control and Prevention reference intervals (overweight, 85th-95th percentiles; obese, > 95th percentile) [25].

Office blood pressure measurement

We performed OBP measurements by the methodology recommended by the fourth report and the AAP guidelines [3, 24]. In brief, a trained nurse measured OBP in a quiet state with child seated for 3 to 5 min, back supported, using an appropriate-sized cuff selected according to child's upper right arm. OBP was initially measured by a calibrated oscillometric device (V 100; Dinamap, Tampa, FL, USA) [26]. If oscillometric OBP measurements remain elevated (≥ 90th percentile), auscultatory OBP measurements were performed using a calibrated aneroid sphygmomanometer [3, 24]. An average of last two auscultatory OBP measurements was used to diagnose office hypertension [3, 24].

Ambulatory blood pressure measurement

A 24-h ABPM was performed with an oscillometric ambulatory BP monitors (model 90,207; Spacelabs Inc., Redmond, WA, USA) [6, 27]. A trained nurse chose an appropriate-sized cuff and conducted the ABPM as per the fourth report and AAP guidelines [3, 24]. The cuff was placed on the nondominant arm, with ABP recordings planned for every 20 min during day and 30 min during night [1, 3]. The participants were instructed to continue with their regular daily activity, to avoid strenuous exercise, and to maintain a wake-sleep log for defining day and night ABP [28]. The adequacy of ABP

recordings was established based on minimum one reading per hour during day and nighttime, and more than 40 readings in 24-h [3]. The 24-h, day and night systolic and diastolic ABP on each patient were analyzed by the respective 24-h/day/night 95th ABP percentile references, recommended by Wühl et al. [6] and endorsed by the AAP guidelines [3].

Outcomes

Our primary outcome was to compare ambulatory hypertension by the 24-h, day-night, and AHA ABP thresholds. Our secondary outcome was to evaluate WCH and MH by these ABP thresholds.

Definitions

Office hypertension as per the AAP guidelines: patient's systolic or diastolic OBP \geq age-, sex-, and height-specific 95th systolic/diastolic OBP percentiles or OBP \geq 130/80 for patients \geq 13 years [3].

Ambulatory hypertension as per 24-h ABP threshold: patient's 24-h mean systolic or diastolic ABP \geq 24-h age-, sex-, and height-specific 95th systolic/diastolic ABP percentiles [6].

Ambulatory hypertension as per day—night ABP thresholds: patient's day or night mean systolic or diastolic ABP \geq corresponding day/night age—, sex—, and height—specific 95th systolic/diastolic ABP percentiles [6].

Ambulatory hypertension as per the AHA threshold: patient's 24-h mean systolic or diastolic ABP \geq 24-h age-, sex-, and height-specific 95th systolic/diastolic ABP percentiles and 24-h systolic or diastolic ABP load \geq 25% [1, 3]. A 24-h systolic/diastolic ABP load was calculated as the percentage of ABP measurements higher than the 24-h mean 95th systolic/diastolic ABP percentiles [6].

Statistical methods

Normally distributed continuous variables were reported as mean (standard deviation, SD), otherwise as median (interquartile range, IQR). Categorical variables were reported as frequency and percentage. Continuous variables were compared with the parametric unpaired t-test or the nonparametric Mann–Whitney U-test, as appropriate. Categorical variables were compared with chi-square test. Systolic and diastolic OBP z-scores and percentiles were calculated based on the methodology recommended by the AAP guidelines [3, 29]. The 24-h, day and night systolic/diastolic ABP z-score, and 95th ABP percentiles were calculated using Box-Cox transformations with age-, and sex-specific estimates of the

distribution median, coefficient of variation, and degree of skewness recommended by Wuhl et al. [6]. The agreement between the ABP thresholds was calculated by the proportion of ABP classified similarly (accuracy) into ambulatory normotension/hypertension and by using the kappa (κ) statistics [30]. Adolescents were defined as those with age \geq 13 years, as recommended by the AAP guidelines [3]. Accuracy and kappa statistics were calculated on MedCalc ver. 18.11 (MedCalc Software, Ostend, Belgium). All other statistical analysis was performed IBM SPSS ver. 25.0 (IBM Corp., Armonk, NY, USA).

Results

Patient characteristics

In the initial screening, 544 participants who had ABPM studies during the recruitment period met the inclusion criteria. Ninety-four participants were excluded for the following reasons: 53 had less than two OBP recordings and 41 had an inadequate ABPM. Four hundred and fifty eligible participants aged 5 to 18 years with complete ABPM and OBP recordings were included in this analysis. The study sample included 59% adolescents ≥ 13 years, 59% males, and 55% overweight/obese participants. Each participant was included with a single ABPM recording in the analysis. The 24-h ABP threshold diagnosed ambulatory hypertension in 122 (27%; 95% confidence interval [CI], 0.22-0.32) and day-night ABP thresholds in 197 participants (44%; 95% CI, 0.37-0.50). Ambulatory hypertension by 24-h ABP threshold and day-night ABP thresholds did not differ in age, sex, BMI z-score, overweight/obese participants, office hypertension, primary hypertension, and blood pressure medication intake; however, ambulatory hypertension by day-night ABP thresholds had more adolescents and ambulatory hypertension by 24-h ABP threshold had higher systolic and diastolic ABP z-scores and ABP loads (Table 1) [3, 6]. As the data was included from two different time periods, we compared the patient characteristics over the two periods, which did not show a significant difference in first versus second period in age (mean \pm SD, 12.98 \pm 3.95 years vs. 13.06 \pm 3.37 years; P=0.83), males (60% vs. 59%; P=0.11), BMI z-score (median [IQR], 1.10 [0.26–2.02] vs. 1.23 [0.12–1.99]; P=0.70) and overweight/obese (54% vs. 56%; P=0.10) distribution.

Diagnosis of ambulatory hypertension, WCH, and MH by 24-h ABP vs. day-night ABP thresholds

The 24-h ABP threshold and day-night ABP thresholds demonstrated only a moderate agreement (κ , 0.58; 95% CI, 0.51–0.66) by classifying 81% ABP (95% CI, 0.72–0.89) similarly into ambulatory normotension/hypertension. As a result, 24-h ABP threshold diagnosed fewer

Sharma et al. Clinical Hypertension (2022) 28:34 Page 4 of 9

Table 1 Patient characteristics

Characteristic	Entire group (n = 450)	(A) Ambulatory hypertension by 24-h mean ABP threshold (n = 122)	(B) Ambulatory hypertension by day-night ABP thresholds (n = 197)	P-value (A vs. B)
Age (yr)	13.03 ± 3.58	12.68±3.73	13.27 ± 3.54	0.15
≥ 13	267 (59)	59 (48)	120 (61)	0.02
Male sex	267 (59)	75 (61)	114 (58)	0.59
Overweight/obese	248 (55)	70 (57)	116 (59)	0.72
BMI z-score	1.21 (0.16 to 2.00)	1.46 (0.38 to 2.18)	1.32 (0.31 to 2.05)	0.50
Office hypertension	57 (36)	93 (76)	131 (66)	0.06
OBP systolic z score	1.68 (0.33 to 2.30)	2.28 (1.51 to 3.00)	2.09 (1.25 to 2.78)	0.22
OBP diastolic z-score	0.60 (-0.05 to 1.41)	1.04 (0.37 to 1.64)	0.99 (0.33 to 1.53)	0.62
Primary hypertension	352 (78)	109 (89)	174 (88)	0.78
Secondary hypertension	98 (22)	13 (11)	23 (12)	0.78
No BP medication	322 (72)	102 (84)	164 (83)	0.81
BP medication	128 (28)	20 (16)	33 (17)	0.81
ABP systolic z-score	0.32 (-0.60 to 1.18)	1.94 (1.27 to 2.66)	1.29 (0.51 to 2.16)	< 0.001
ABP systolic load (%)	21.42 (7.14 to 44.44)	60.85 (41.40 to 80.00)	42.31 (22.47 to 65.63)	< 0.001
ABP diastolic z-score	0.13 (-0.69 to 1.18)	1.78 (1.07 to 2.65)	1.23 (0.40 to 2.03)	< 0.001
ABP diastolic load (%)	15.38 (6.67 to 34.20)	41.55 (28.20 to 60.47)	34.62 (16.95 to 51.06)	< 0.001

Data are presented as mean \pm standard deviation, number (%), or median (interquartile range). Office hypertension was diagnosed by the American Academy of Pediatrics thresholds [3]. 24-h mean ABP threshold: 24-h mean ABP \geq 24-h 95th ABP percentile according to the ABPM references [6]; Day-night ABP thresholds: mean day ABP \geq day 95th ABP percentile [6] or mean night ABP \geq night 95th ABP percentile [6]

ABP ambulatory blood pressure, BMI body mass index, OBP office blood pressure, BP blood pressure

ambulatory hypertension (27%; 95% CI, 0.22–0.32) than day-night ABP thresholds (44%; 95% CI, 0.37–0.50; P < 0.001) (Table 2, Fig. 1) [6]. Fewer ambulatory hypertension by 24-h ABP threshold translated into more WCH (33%, 95% CI 0.28–0.39 vs. 25%, 95% CI 0.20–0.30; P < 0.001) and fewer MH (6%, 95% CI 0.04–0.09 vs. 14%, 95% CI 0.11–0.18; P < 0.001) by 24-h ABP than day-night ABP thresholds (Table 2, Fig. 1) [6]. The pattern of ambulatory hypertension, WCH, and MH diagnosis by the 24-h ABP and day-night ABP thresholds remained consistent across the age, sex, BMI, primary hypertension, and no BP medication intake (Table 2) [6].

Agreement between 24-ABP and day ABP vs. night ABP thresholds

Despite the fact that 24-h ABP threshold reflects the mean of day and night ABP thresholds, only a moderate agreement between the 24-h and day-night ABP thresholds led us to look at the individual association between the 24-h ABP threshold and day ABP vs. night ABP threshold. We found that 24-h ABP threshold had a stronger agreement with day ABP (91%; 95% CI, 0.82–1.06) than night ABP (78%; 95% CI, 0.70–0.87; P<0.001) to classify ABP into ambulatory normotension/hypertension (Table 3) [6]. Therefore, the diagnosis of ambulatory hypertension was closer between 24-h ABP and day ABP threshold (difference, 3.70%; 95% CI, 1.97%–9.34%; P=0.20) than between 24-h ABP and night ABP

threshold (difference, 11%; 95% CI, 4.9%–17%; P<0.001). The agreement pattern between 24-h ABP and day ABP versus night ABP threshold remained consistent across the age, sex, BMI, primary hypertension, and no BP medication intake (Table 3) [6].

Agreement between the AHA threshold and 24-h ABP vs. day-night ABP thresholds

The AHA threshold demonstrated a strong agreement with 24-h ABP threshold (κ , 0.94; 95% CI, 0.91–0.98), and the two thresholds classified 98% ABP (95% CI, 0.89–1.07) similarly into ambulatory normotension/hypertension (Table 4) [3, 6]. In contrast, the AHA threshold showed only a moderate agreement with day-night ABP thresholds (κ , 0.59; 95% CI, 0.52–0.66), with a relatively lower proportion of ABP classified similarly by the two ABP thresholds (80%; 95% CI, 0.72–0.89) (Table 4) [3, 6]. The stronger agreement between the AHA and 24-h ABP threshold than with the day-night ABP thresholds remained consistent in sub-group analysis on adolescent, male, overweight-obese, primary hypertension, and no BP medication intake (Table 4) [3, 6].

The stronger agreement between the AHA and 24-h ABP thresholds translated into AHA threshold diagnosing ambulatory hypertension closer to 24-h ABP threshold (26%; 95% CI, 0.21–0.31 vs. 27%; 95% CI, 0.22–0.32; P=0.73) than day-night ABP thresholds (44%; 95% CI, 0.37–0.50; P<0.001) (Fig. 1). Similarly, the diagnosis of

Sharma et al. Clinical Hypertension (2022) 28:34 Page 5 of 9

Table 2 Diagnosis of ambulatory hypertension, white coat hypertension, and masked hypertension based on 24-h mean ABP and day-night ABP thresholds

Variable	24-h mean ABP threshold (%; 95% CI)	Day-night ABP thresholds (%; 95% CI)	Difference (%; 95% CI)
Ambulatory hypertension			
Entire group ($n = 450$)	27 (0.22 to 0.32)	44 (0.37 to 0.50)	17 (10.8 to 23)*
Age \ge 13 yrs. ($n = 267$)	22 (0.16 to 0.28)	45 (0.37 to 0.53)	23 (15.1 to 30.5)*
Male sex ($n = 267$)	28 (0.22 to 0.35)	43 (0.35 to 0.51)	15 (6.9 to 22.8)*
Overweight/obese ($n = 248$)	28 (0.22 to 0.35)	47 (0.38 to 0.56)	19 (10.5 to 27.1)*
Primary hypertension ($n = 352$)	31 (0.25 to 0.37)	49 (0.42 to 0.57)	18 (10.8 to 24.9)*
Secondary hypertension ($n = 98$)	13 (0.07 to 0.22)	23 (0.14 to 0.35)	10 (0.8 to 20.7)
No BP medication ($n = 322$)	32 (0.25 to 0.38)	51 (0.43 to 0.59)	19 (11.4 to 26.3)*
On BP medication ($n = 128$)	16 (0.09 to 0.24)	26 (0.17 to 0.36)	10 (0.01 to 19.8)
White coat hypertension			
Entire group ($n = 450$)	34 (0.28 to 0.39)	25 (0.20 to 0.30)	9 (2.1 to 13.9)*
Age \ge 13 yr ($n = 267$)	35 (0.28 to 0.43)	23 (0.18 to 0.30)	12.0 (4.3 to 19.5)*
Male sex ($n = 267$)	33 (0.27 to 0.41)	26 (0.20 to 0.33)	70 (-0.7 to 14.6)
Overweight/obese ($n = 248$)	37 (0.30 to 0.46)	25 (0.19 to 0.32)	12 (3.9 to 19.9)*
Primary hypertension ($n = 352$)	34 (0.29 to 0.41)	25 (0.20 to 0.31)	9 (2.3 to 15.6)*
Secondary hypertension ($n = 98$)	28 (0.18 to 0.41)	23 (0.14 to 0.35)	5 (-7.2 to 17)
No BP medication ($n = 322$)	36 (0.29 to 0.43)	25 (0.20 to 0.31)	11 (3.9 to 18)*
On BP medication ($n = 128$)	27 (0.19 to 0.38)	23 (0.15 to 0.33)	4 (-6.6 to 14.5)
Masked hypertension			
Entire group ($n = 450$)	6 (0.04 to 0.09)	14 (0.11 to 0.18)	8 (4.1 to 12)*
Age \ge 13 yr ($n = 267$)	6 (0.03 to 0.10)	17 (0.12 to 0.23)	11 (5.6 to 16.5)*
Male sex $(n = 267)$	6 (0.03 to 0.10)	14 (0.10 to 0.19)	8 (2.9 to 13.2)*
Overweight/obese ($n = 248$)	6 (0.03 to 0.10)	12 (0.08 to 0.17)	8 (2.9 to 13.2)*
Primary hypertension ($n = 352$)	6 (0.04 to 0.09)	15 (0.11 to 0.20)	9 (4.5 to 13.6)*
Secondary hypertension ($n = 98$)	6 (0.02 to 0.13)	11 (0.05 to 0.20)	5 (-3.1 to 13.4)
No BP medication ($n = 322$)	8 (0.05 to 0.12)	17 (0.13 to 0.22)	9 (3.9 to 14.1)*
On BP medication ($n = 128$)	1 (0.00 to 0.05)	7 (0.03 to 0.14)	6 (1.1 to 11.8)*

Office hypertension was diagnosed by the American Academy of Pediatrics thresholds [3]. 24-h mean ABP threshold: 24-h mean ABP \geq 24-h 95th ABP percentile according to the ABPM references [6]; Day-night ABP thresholds: mean day ABP \geq day 95th ABP percentile [6] or mean night ABP \geq night 95th ABP percentile [6] ABP ambulatory blood pressure, CI confidence interval, BP blood pressure, ABPM ambulatory blood pressure monitoring

*P < 0.05

WCH (34%; 95% CI, 0.29–0.40) and MH (6%; 95% CI, 0.04–0.09) by the AHA threshold was not significantly different than WCH (33%; 95% CI, 0.28–0.39; P=0.75) and MH (6%; 95% CI, 0.04–0.09; P>0.999 by the 24-h ABP threshold (Fig. 1). However, the diagnosis of WCH (25%; 95% CI, 0.20–0.30) and MH (14%; 95% CI, 0.11–0.18) by the day-night ABP thresholds was significantly different than WCH (difference, 9%; 95% CI, 3%–14.9%; P<0.001) and MH (difference, 8%; 95% CI, 4.1%–12%; P<0.001) by the AHA threshold (Fig. 1).

Discussion

The choice of an ABP threshold can influence the diagnosis of ambulatory hypertension, consequently that of WCH/MH [8]. In clinical practice, 24-h ABP, day-night

ABP, and AHA thresholds are commonly used to diagnose ambulatory hypertension [8–13]. We found that 24-h ABP threshold diagnosed significantly fewer ambulatory hypertension than day-night ABP thresholds, which reflected in the difference in the diagnosis of WCH and MH by these ABP thresholds. In addition, the AHA threshold diagnosed ambulatory hypertension, WCH, and MH closer to 24-h ABP threshold than day-night ABP thresholds.

Given the paucity of literature on the agreement between these commonly used ABP thresholds, our findings become novel and important for a consistent interpretation of ambulatory hypertension by these ABP thresholds in clinical practice. We found that 24-h ABP and day-night ABP thresholds classified ABP similarly

Sharma et al. Clinical Hypertension

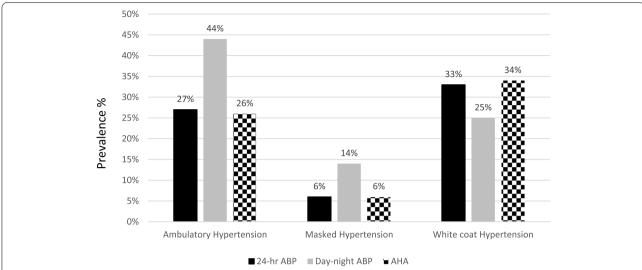


Fig. 1 Ambulatory hypertension (AH), masked hypertension (MH), and white coat hypertension (WCH) as per the 24-h ambulatory blood pressure (ABP), day-night ABP and American Heart Association (AHA) thresholds. The 24-h ABP threshold diagnosed fewer AH (27% vs. 44%; P < 0.001), MH (6% vs. 14%; P < 0.001), and more WCH (33% vs. 25%; P < 0.001) than day-night ABP thresholds. The 24-h ABP and AHA thresholds diagnosed AH (27% vs. 26%; P = 0.73), MH (6% vs. 6%; P > 0.999), and WCH (33% vs. 34%; P = 0.75) quite similarly, whereas the day-night ABP and AHA thresholds diagnosed AH (44% vs. 26%; P < 0.001), MH (14% vs. 6%; P < 0.001), and WCH (25% vs. 34%; P < 0.001) differently

into ambulatory normotension/hypertension in 81% participants, which can be explained by the fact that patient's 24-h ABP is expected to represent an average of the day and night ABP measurements. However, the noteworthy observation from our analysis was that 24-h ABP

threshold diagnosed 17% fewer ambulatory hypertension than the day-night ABP thresholds (27% vs. 44%). As a result, 24-h ABP threshold diagnosed more WCH and fewer MH than the day-night ABP thresholds, which remained consistent across subgroups based on age,

Table 3 Agreement between 24-h mean ABP and day versus night ABP for diagnosing ambulatory normotension/hypertension

Variable	Accuracy (%) ^{a)} (95th CI)	Kappa (95% CI)
Day ABP threshold		
Entire group ($n = 450$)	91 (0.82–1.06)	0.77 (0.70–0.83)
Age \geq 13 yr ($n = 267$)	91 (0.80–1.03)	0.73 (0.63–0.83)
Male sex $(n=267)$	92 (0.81–1.05)	0.82 (0.74–0.89)
Overweight/obese ($n = 248$)	91 (0.79–1.03)	0.77 (0.69–0.86)
Primary hypertension ($n = 352$)	90 (0.80–1.01)	0.77 (0.69–0.84)
Secondary hypertension ($n = 98$)	93 (0.75–1.15)	0.71 (0.50–0.93)
No BP medication ($n = 322$)	90 (0.79–1.01)	0.76 (0.68–0.84)
On BP medication ($n = 128$)	94 (0.78–1.12)	0.76 (0.60-0.93)
Night ABP threshold		
Entire group ($n = 450$)	78 (0.70–0.87)	0.52 (0.44–0.60)
Age \geq 13 yr ($n = 267$)	73 (0.63–0.84)	0.39(0.29-0.50)
Male sex $(n=267)$	78 (0.68–0.90)	0.57(0.47-0.68)
Overweight/obese ($n = 248$)	79 (0.69–0.91)	0.55(0.45-0.66)
Primary hypertension ($n = 352$)	76 (0.67–0.85)	0.49 (0.40-0.58)
Secondary hypertension ($n = 98$)	88 (0.71–1.09)	0.60 (0.39-0.81)
No BP medication ($n = 322$)	75 (0.66–0.85)	0.48 (0.39–0.58)
On BP medication ($n = 128$)	87 (0.72–1.05)	0.59 (0.41-0.76)

²⁴⁻h mean ABP threshold: 24-h ABP ≥ 24-h 95th ABP percentile according to the ABPM references [6]; Day ABP threshold: mean day ABP ≥ day 95th ABP percentile [6]; Night ABP threshold: mean night ABP ≥ night 95th ABP percentile [6]

ABP ambulatory blood pressure, CI confidence interval, BP blood pressure, ABPM ambulatory blood pressure monitoring

a) The proportion of ABP classified similarly by both the ABP thresholds into ambulatory normotension/hypertension

Sharma et al. Clinical Hypertension (2022) 28:34 Page 7 of 9

Table 4 Agreement between the AHA threshold and the 24-h mean ABP threshold versus day-night ABP thresholds for diagnosing ambulatory normotension/hypertension

Variable	AHA and 24-h mean ABP threshold		AHA and Day-night ABP thresholds	
	Accuracy (%) ^{a)} (95th CI)	Kappa (95% CI)	Accuracy (%) ^{a)} (95th CI)	Kappa (95% CI)
Entire group ($n = 450$)	98 (0.89–1.07)	0.94 (0.91–0.98)	80 (0.72–0.89)	0.59 (0.52–0.66)
Age \geq 13 yr ($n = 267$)	98 (0.86-1.10)	0.94(0.89-0.99)	76 (0.65–0.87)	0.49(0.39-0.58)
Male sex $(n=267)$	97 (0.85-1.09)	0.93(0.88-0.98)	83 (0.72-0.95)	0.64(0.55-0.73)
Overweight/obese ($n = 248$)	97 (0.85–1.10)	0.93 (0.89-0.98)	79 (0.69–0.91)	0.58(0.49-0.67)
Primary hypertension ($n = 352$)	97 (0.87-1.08)	0.93 (0.90-0.97)	78 (0.69–0.88)	0.56 (0.48-0.64)
Secondary hypertension ($n = 98$)	98 (0.81-1.21)	1.00 (1.00-1.00)	89 (0.72-1.10)	0.66 (0.48-0.85)
No BP medication ($n = 322$)	97 (0.86-1.08)	0.93 (0.89-0.97)	77 (0.69–0.87)	0.54 (0.46-0.63)
On BP medication ($n = 128$)	100 (0.83–1.18)	1.00 (1.00-1.00)	89 (0.74–1.07)	0.69 (0.54-0.84)

24-h mean ABP threshold: 24-h mean ABP \geq 24-h 95th ABP percentile according to the ABPM references [6]. Day-night ABP thresholds: day ABP \geq 95th day ABP percentile [6] or mean night ABP \geq 95th night ABP percentile [6]. AHA threshold: 24-h systolic or diastolic ABP \geq 95th ABP percentile and 24-h systolic or diastolic ABP load: proportion of systolic or diastolic ABP readings higher than 24-h mean systolic or diastolic 95th ABP percentile

AHA American Heart Association, ABP ambulatory blood pressure, CI confidence interval, BP blood pressure, ABPM ambulatory blood pressure monitoring

sex, BMI, primary hypertension, and no BP medication intake. Our finding on the discrepancy in the diagnosis of ambulatory hypertension by the 24-h and day-night ABP thresholds remains consistent with the adult literature that demonstrated a difference in ambulatory hypertension by the 24-h ABP threshold versus 24-h ABP threshold combined with day and night ABP thresholds [31, 32]. Based on these observations, it can be concluded that 24-h ABP and day-night ABP thresholds diagnose ambulatory hypertension differently, which can impact the determination of WCH and MH by these thresholds.

The difference in the diagnosis of ambulatory hypertension by 24-h ABP versus day-night ABP thresholds can be explained by a relatively stronger agreement between 24-h ABP and day ABP than between 24-h ABP and night ABP threshold. We found that 24-h ABP and day ABP thresholds classified 91% ABP similarly into ambulatory normotension/hypertension, which was higher than 78% ABP classified similarly by the 24-h and night ABP thresholds. It is tempting to speculate that the stronger agreement between the 24-h ABP and day ABP threshold than with the night ABP threshold results from a relatively higher representation of day ABP on 24-h ABPM, which can be explained by the standard recommendations of measuring day ABP more frequently, every 20 to 30 min, than more spread out night ABP recordings every 30 to 40 min [1, 3, 24].

Given the fact that the AHA guidelines have recommended a different ABP threshold than 24-h and day-night ABP thresholds to diagnose ambulatory hypertension, we evaluated the agreement between the AHA threshold and 24-h ABP/day-night ABP thresholds. We found that the AHA threshold had a stronger agreement

with 24-h ABP threshold than with day-night ABP thresholds. The AHA and 24-h ABP thresholds classified 90% ABP similarly into ambulatory normotension/ hypertension, which was higher than 80% ABP classified similarly by the AHA and day-night ABP thresholds. The stronger agreement between the AHA and 24-h ABP thresholds persisted regardless of age, sex, BMI, primary hypertension, or no antihypertensive medication intake. The stronger agreement between the AHA and 24-h ABP than day-night ABP thresholds can be explained by the fact that the AHA threshold is based on 24-h mean 95th ABP percentile along with ABP load≥25% estimated by 24-h mean 95th ABP percentile. As a result, it can be concluded that day-night ABP threshold and AHA threshold cannot be interchangeably used to diagnose ambulatory hypertension in clinical practice.

Strengths of our study included the use of a standardized methodology for undertaking ABPM and using ABPM references [6] recommended by the AHA guidelines [1, 3]. The use of OBP references to diagnose office hypertension was consistent with that recommended by the AAP guidelines [3]. Our study limitations include retrospective study design and unavailability of hypertension-induced target-organ damage assessment. Though there is paucity of data on the outcome-based studies with the AHA threshold, 24-h ABP 95th percentile has been shown to have a strong association with hypertension-induced target organ damage [33, 34]. The use of OBP measurements from a single visit may potentially influence the estimation of WCH and MH in our analysis. However, OBP measurements should not change our observations on ambulatory hypertension and the relative pattern of

a) The proportion of ABP classified similarly by both the ABP thresholds into ambulatory normotension/hypertension

interaction among the ambulatory thresholds to diagnose WCH/MH. Despite the fact that we included participants from two different time periods to enhance the statistical power of our analysis, a consistency over the two time periods in regards to evaluating secondary etiologies of hypertension [3, 24], machines used for OBP/ABP measurements, OBP/ABP thresholds for data interpretation, and no difference in age, sex, BMI z-score, overweight-obese distribution should minimize a potential cofounding effect of using data from two different time periods. A relatively larger representation of primary hypertension and those not on a blood pressure medication limits the generalizability of our findings to those with secondary hypertension and on blood pressure medications. Predominant Caucasian ethnicity limits the applicability of our observations to other ethnicities. In view of the tertiary care setting of our study, our results should be generalized to a primary care population with caution.

Conclusions

Our study addresses the important question of an appropriate diagnosis of ambulatory hypertension on 24-h ABP monitoring, and consequently that of WCH and MH, by the commonly used ABP thresholds in children. Our findings suggest that 24-h ABP threshold has a strong agreement with the AHA threshold to diagnose ambulatory hypertension, WCH and MH. However, notably both 24-h ABP and AHA thresholds diagnose ambulatory hypertension, WCH, and MH differently than the day-night ABP thresholds. Given the importance choosing an appropriate ABP threshold for diagnosing ambulatory hypertension, our findings are relevant for an appropriate interpretation of ambulatory hypertension, WCH and MH by these commonly used ABP thresholds in clinical practice and for research purposes.

Abbreviations

AAP: American Academy of Pediatrics; ABP: Ambulatory blood pressure; AHA: American Hypertension Association; ABPM: Ambulatory blood pressure monitoring; BMI: Body mass index; CI: Confidence interval; IQR: Interquartile range; MH: Masked hypertension; OBP: Office blood pressure; SD: Standard deviation; WCH: White coat hypertension.

Acknowledgements

We acknowledge Carlos Reis, RN for his meticulous assistance in organizing the ARPM recordings

Authors' contributions

For the manuscript, APS conceived the idea, conducted the statistical analysis, prepared the initial draft, and supervised overall development of the manuscript. Authors MA and KS contributed in drafting the manuscript and data collection. The authors KN, AK, GF and LAD revised the manuscript and played an important role in interpreting the results. The publication has been approved by all co-authors. The corresponding author has had full access to

the data in the study and final responsibility for the decision to submit for publication.

Funding

None.

Availability of data and materials

The datasets generated and analyzed during the current study are not publicly available due to patient's confidentiality but are available from the corresponding author on reasonable request.

Declarations

Ethics approval and consent to participate

Ethics approval was obtained from the Research Ethics Board of the University of Western Ontario (No. 16143E and 116082). The study involved the retrospective review of existing clinical data and was therefore exempted from the need for an individual informed consent.

Consent for publication

Not applicable.

Competing interests

The authors declare that they have no competing interests.

Author details

¹Division of Nephrology, Department of Paediatrics, University of Western Ontario, London, ON, Canada. ²London Health Sciences Centre, London, ON, Canada. ³Division of Cardiology, Department of Paediatrics, University of Western Ontario, London, ON, Canada. ⁴University of Western Ontario, London, ON, Canada. ⁵Department of Paediatric Cardiology and Intensive Care, Medical School Hannover, Hannover, Germany.

Received: 17 September 2021 Accepted: 21 June 2022 Published online: 15 November 2022

Reference

- Flynn JT, Daniels SR, Hayman LL, Maahs DM, McCrindle BW, Mitsnefes M, et al. Update: ambulatory blood pressure monitoring in children and adolescents: a scientific statement from the American heart association. Hypertension. 2014;63:1116–35.
- Lurbe E, Agabiti-Rosei E, Cruickshank JK, Dominiczak A, Erdine S, Hirth A, et al. 2016 European society of hypertension guidelines for the management of high blood pressure in children and adolescents. J Hypertens. 2016;34:1887–920.
- Flynn JT, Kaelber DC, Baker-Smith CM, Blowey D, Carroll AE, Daniels SR, et al. Clinical practice guideline for screening and management of high blood pressure in children and adolescents. Pediatrics. 2017;140:e20171904.
- Krmar RT. White-coat hypertension from a paediatric perspective. Acta Paediatr. 2019;108:44–9.
- Kirk V, Midgley J, Giuffre M, Ronksley P, Nettel-Aguirre A, Al-Shamrani A. Hypertension and obstructive sleep apnea in Caucasian children. World J Cardiol. 2010;2:251–6.
- Wuhl E, Witte K, Soergel M, Mehls O, Schaefer F. German working group on pediatric hypertension. Distribution of 24-h ambulatory blood pressure in children: normalized reference values and role of body dimensions. J Hypertens. 2002;20:1995–2007.
- Soergel M, Kirschstein M, Busch C, Danne T, Gellermann J, Holl R, et al. Oscillometric twenty-four-hour ambulatory blood pressure values in healthy children and adolescents: a multicenter trial including 1141 subjects. J Pediatr. 1997;130:178–84.
- Sorof JM, Poffenbarger T, Franco K, Portman R. Evaluation of white coat hypertension in children: importance of the definitions of normal ambulatory blood pressure and the severity of casual hypertension. Am J Hypertens. 2001;14(9 Pt 1):855–60.

- Stabouli S, Kotsis V, Toumanidis S, Papamichael C, Constantopoulos A, Zakopoulos N. White-coat and masked hypertension in children: association with target-organ damage. Pediatr Nephrol. 2005;20:1151–5.
- Shah S, Swartz S, Campbell J, Srivaths PR. Ambulatory blood pressures and central blood pressures are associated with cardiovascular morbidity in adolescent and young adult patients receiving chronic hemodialysis. Pediatr Nephrol. 2019;34:1261–8.
- Höcker B, Weber LT, John U, Drube J, Fehrenbach H, Klaus G, et al. Steroid withdrawal improves blood pressure control and nocturnal dipping in pediatric renal transplant recipients: analysis of a prospective, randomized, controlled trial. Pediatr Nephrol. 2019;34:341–8.
- Suláková T, Feber J. Should mean arterial pressure be included in the definition of ambulatory hypertension in children? Pediatr Nephrol. 2013;28:1105–12.
- Koshy S, Macarthur C, Luthra S, Gajaria M, Geary D. Ambulatory blood pressure monitoring: mean blood pressure and blood pressure load. Pediatr Nephrol. 2005;20:1484–6.
- Li Y, Wang JG. Isolated nocturnal hypertension: a disease masked in the dark. Hypertension. 2013;61:278–83.
- Bromfield SG, Shimbo D, Bertoni AG, Sims M, Carson AP, Muntner P. Ambulatory blood pressure monitoring phenotypes among individuals with and without diabetes taking antihypertensive medication: the Jackson heart study. J Hum Hypertens. 2016;30:731–6.
- Wijkman M, Länne T, Engvall J, Lindström T, Ostgren CJ, Nystrom FH. Masked nocturnal hypertension: a novel marker of risk in type 2 diabetes. Diabetologia. 2009;52:1258–64.
- 17. Fujita H, Matsuoka S, Awazu M. Masked isolated nocturnal hypertension in children and young adults. Pediatr Cardiol. 2018;39:66–70.
- Seeman T, Palyzová D, Dusek J, Janda J. Reduced nocturnal blood pressure dip and sustained nighttime hypertension are specific markers of secondary hypertension. J Pediatr. 2005;147:366–71.
- Sharma A, Altamirano-Diaz L, Grattan M, Filler G, Sharma AP. Comparative analysis of American heart association and European society of hypertension ambulatory blood pressure thresholds for diagnosing hypertension in children. Kidney Int Rep. 2020;5:611–7.
- Sharma AP, Mohammed J, Thomas B, Lansdell N, Norozi K, Filler G. Nighttime blood pressure, systolic blood pressure variability, and left ventricular mass index in children with hypertension. Pediatr Nephrol. 2013;28:1275–82.
- 21 Sharma AP, Norozi K, Grattan M, Filler G, Altamirano-Diaz L. Diagnosis of pediatric hypertension: European society of hypertension-recommended 24-hour vs. 24-hour-day-night ambulatory blood pressure thresholds. Am J Hypertens. 2021;34:198–206.
- Sharma AP, Mohammed J, Thomas B, Singh RN, Filler G. Using simplified blood pressure tables to avoid underdiagnosing childhood hypertension. Paediatr Child Health. 2015;20:297–301.
- Sharma AP, Altamirano-Diaz L, Mohamed Ali M, Stronks K, Kirpalani A, Filler G, et al. Diagnosis of hypertension: ambulatory pediatric American heart association/European Society of Hypertension versus blood pressure load thresholds. J Clin Hypertens (Greenwich). 2021;23:1947–56.
- 24. National High Blood Pressure Education Program Working Group on High Blood Pressure in Children and Adolescents. The fourth report on the diagnosis, evaluation, and treatment of high blood pressure in children and adolescents. Pediatrics. 2004;114(2 Suppl 4th Report):555–76.
- Centers for Disease Control and Prevention (CDC). National Center for Health Statistics: 2000 CDC growth charts for the United States: methods and development. Atlanta: CDC; 2002. Available from: https://www.cdc. gov/growthcharts/cdc_charts.htm. [cited 2020 Feb 02].
- 26. Lee CG, Park HM, Shin HJ, Moon JS, Hong YM, Kim NS, et al. Validation study of the Dinamap ProCare 200 upper arm blood pressure monitor in children and adolescents. Korean J Pediatr. 2011;54:463–9.
- Belsha CW, Wells TG, Bowe Rice H, Neaville WA, Berry PL. Accuracy of the SpaceLabs 90207 ambulatory blood pressure monitor in children and adolescents. Blood Press Monit. 1996;1:127–33.
- Jones HE, Sinha MD. The definition of daytime and nighttime influences the interpretation of ABPM in children. Pediatr Nephrol. 2011;26:775–81.
- Rosner B, Cook N, Portman R, Daniels S, Falkner B. Determination of blood pressure percentiles in normal-weight children: some methodological issues. Am J Epidemiol. 2008;167:653–66.
- Landis JR, Koch GG. The measurement of observer agreement for categorical data. Biometrics. 1977;33:159–74.

- Asayama K, Thijs L, Li Y, Gu YM, Hara A, Liu YP, et al. Setting thresholds to varying blood pressure monitoring intervals differentially affects risk estimates associated with white-coat and masked hypertension in the population. Hypertension. 2014;64:935–42.
- de la Sierra A, Vinyoles E, Banegas JR, Segura J, Gorostidi M, de la Cruz JJ, et al. Prevalence and clinical characteristics of white-coat hypertension based on different definition criteria in untreated and treated patients. J Hypertens. 2017;35:2388–94.
- Maggio AB, Aggoun Y, Marchand LM, Martin XE, Herrmann F, Beghetti M, et al. Associations among obesity, blood pressure, and left ventricular mass. J Pediatr. 2008;152:489–93.
- 34. McNiece KL, Gupta-Malhotra M, Samuels J, Bell C, Garcia K, Poffenbarger T, et al. Left ventricular hypertrophy in hypertensive adolescents: analysis of risk by 2004 national high blood pressure education program working group staging criteria. Hypertension. 2007;50:392–5.

Publisher's Note

Springer Nature remains neutral with regard to jurisdictional claims in published maps and institutional affiliations.

Ready to submit your research? Choose BMC and benefit from:

- fast, convenient online submission
- $\bullet\,$ thorough peer review by experienced researchers in your field
- rapid publication on acceptance
- support for research data, including large and complex data types
- gold Open Access which fosters wider collaboration and increased citations
- maximum visibility for your research: over 100M website views per year

At BMC, research is always in progress.

Learn more biomedcentral.com/submissions

