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Racial Differences in Relation Between Carotid and Radial Augmentation Index

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Abstract

Background—Augmented central artery wave reflection is a cardiovascular disease risk factor. Augmentation index (AI) obtained from peripheral artery waveforms provides qualitatively similar information to AI from central artery waveforms. Little information is available, however, regarding the influence of racial difference in association between central and peripheral AI.

Methods—We studied 47 White adults (45±17 yr, 20 women) and 94 age-matched Asian adults (45±14 yr, 42 women).

Results—The White group was significantly taller than the Asian group, whereas there were no significant group differences in blood pressure and heart rate. Carotid and radial AI tended to be lower in White compared with Asian adults ($P<0.10$ for both). Such tendency disappeared when the difference in height was taken into account using ANCOVA ($P=0.84$ and $P=0.77$, respectively). Radial AI was strongly and positively correlated with carotid AI in White adults ($r=0.75$, $P<0.0001$) as well as in Asian adults ($r=0.82$, $P<0.0001$). The slope and intercept of linear regression line between radial and carotid AI of White adults were highly comparable with those of Asian adults.

Conclusion—AI in the conveniently located peripheral vasculature may provide a surrogate measure of central AI irrespective of difference in race (e.g., Asian vs. White populations).

Keywords

reflection wave; applanation tonometry; race; central blood pressure

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Introduction

Augmented wave reflection in the central artery is an important determinant of cardiovascular risk [1–3], and augmentation index (AI) is a well-established indicator of arterial wave reflection [4]. However, the acquisition of arterial pressure waveforms required for AI is difficult to perform in the central arteries (e.g., the aorta, carotid artery). Accordingly, many investigators have adapted the measurement of AI on the radial arteries and have used it to gain insight into carotid or aortic AI [5–7]. The available evidence is encouraging in this line of research, but it is not clear if the relation between radial artery AI and central artery AI is influenced by the race or ethnic status. This is critical in order for this technique to have a greater generalizability. Accordingly, the purpose of this study was to determine the influence of race (more specifically, Whites and Asians) on association between radial AI and carotid AI.

Methods

Subjects

Forty-seven white adults and 94 age-matched Asian adults (1 to 2 ratio match up of subject samples) who were free of overt cardiovascular diseases were studied after at least 4 hours of fasting and abstinence from caffeine. This study was reviewed and approved by the local Institutional Review Board, and all subjects signed the informed consent.

All measurements were performed after an abstinence of caffeine and a 3-hour fast. After resting for at least 15 min in a quiet, temperature-controlled room, carotid and radial arterial AI were measured in a random order by two different vascular testing devices. Carotid arterial waveforms were recorded on the automatic device (VP-2000, Colin Medical Technology, San Antonio, TX) in duplicate using an arterial applanation tonometry probe incorporating an array of 15 micropiezoresistive transducers on the left common carotid artery with a collar, which is able to hold the probe on the carotid artery by an optimal constant pressure. The systolic foot (diastolic pressure), shoulder, and peak (systolic pressure) of carotid arterial pressure waveforms were automatically detected by using algorithms based on a band-pass filtering (5–30 Hz) and fourth derivatives [8] (Figure 1). The band-pass filtering was performed in order to eliminate artifacts. Carotid AI was calculated as follows:

$$\text{Carotid AI(\%)} = \Delta P / \text{carotid PP} \times 100$$

where ΔP is augmentation of carotid systolic pressure about the first systolic shoulder, PP is pulse pressure [4]. The validity and reliability of carotid AI measurements with this automatic device have been established [9].

Radial AI was measured using an applanation tonometry-based automated radial AI measurement device (HEM-9010AI; Omron Healthcare, Kyoto, Japan). The arterial applanation tonometry probe, incorporating an array of 40 micropiezoresistive transducers, was held on the left radial artery while pressure against the radial artery was automatically modulated in order to obtain an optimal radial arterial waveform. Systolic foot (e.g., diastolic pressure) and the first and second systolic peaks were automatically detected using fourth derivatives for each radial arterial waveform and averaged. Radial AI was calculated as follows:

$$\text{Radial AI(\%)} = (\text{radial SBP}_2 - \text{DBP}) / (\text{radial SBP}_1 - \text{DBP}) \times 100$$

where SBP_1 and SBP_2 are the first and second systolic peaks of radial arterial pressure. The validity and reliability of detecting the first and second peak radial systolic pressures with the fourth derivative wave has been established (Figure 1) [10].

Unpaired t-test and ANCOVA were performed to compare variables of interest between groups. Univariate correlation analysis and Bland-Altman plots were used to assess relationships between variables of interest. The effect of the race on the relation between carotid and radial AI was assessed by a general linear regression model.

Results

White adults were significantly taller and heavier than Asian adults, whereas no significant group differences were observed in blood pressure and heart rate (Table 1). Carotid and radial AI tended to be lower in white compared with Asian adults ($P < 0.10$ for both). However, such tendency was disappeared when the difference in height was taken into account using ANCOVA ($P = 0.84$ and $P = 0.77$, respectively).

Radial AI was correlated with carotid AI in White adults ($r = 0.75$, $P < 0.0001$) and in Asian adults ($r = 0.82$, $P < 0.0001$) (Figure 2). The regression line for White adults ($y = 0.95x - 64.1$, black line) was almost superimposed on that for Asian adults ($y = 0.93x - 60.9$, gray line). As seen in Bland-Altman plots, the differences between radial and carotid AI were not different between White and Asian adults ($66.5 \pm 10.3\%$ vs. $67.3 \pm 14.1\%$, $P = 0.71$, Figure 2).

In the pooled subjects, general linear regression model revealed that radial AI ($\beta = 0.57$, $P < 0.0001$) and age ($\beta = 0.17$, $P < 0.01$) were significant independent correlates for carotid AI ($R^2 = 0.71$, $P < 0.0001$), whereas race, sex, height, body weight, body mass index, heart rate, and blood pressure were ejected from the model by not being independently significant.

Discussion

Increase in central artery wave reflection is gaining a reputation as a risk factor for cardiovascular disease. Because of its clinical importance, the development of promising noninvasive techniques to screen high-risk patients is desirable. We and others have reported that radial AI, which is easier to measure and implement, may be able to provide qualitatively similar information to carotid AI [5–7]. These relations have been observed in some populations, including general populations [5,6] and medicated hypertensive populations [7]. The results of the present study extend these previous by suggesting that these relations can be extrapolated to different racial populations, including Whites and Asians.

A cohort study composed of monozygotic and dizygotic female twin pairs indicated that AI has significant heritable component (approximately 37 % of total variation in AI) [11]. These results imply that genetic components might affect association between carotid and radial AI. Contrary to this hypothesis, radial AI was significantly correlated with carotid AI similarly in both White and Asian groups. Furthermore, the Bland-Altman plot revealed that differences between radial and carotid AI of White adults were highly comparable with those of Asian adults. General linear regression model revealed that radial AI and age were significant independent variables for carotid AI, whereas race, sex, height, body weight, body mass index, heart rate, and blood pressure were ejected from the model by not being independently significant. These results suggest that the race does not affect the relation between carotid and radial augmentation index.

To our knowledge, this is the first study to assess the effect of race on association between measures of arterial wave reflection evaluated from central and peripheral arterial pulse waves. In the present study, carotid and radial AI tended to be greater in Asian subjects than in White

subjects. This group difference was primarily due to the smaller height in Asian subjects compared with White populations as the group difference was disappeared when height was taken into account. AI, measure of central wave reflection, is influenced by various factors (i.e., sex, aging, height, arterial stiffness, heart rate, blood pressure) [6,12–15]. Since shorter height is associated with the smaller length from the heart to major wave reflection sites [16, 17] and earlier return of reflection waves from periphery to the heart, a person, who has short stature, tends to exhibit a greater AI. East-Asian is known to be smaller in height than Caucasian or other white populations [18,19], and this was reflected in our study sample.

There is an increasing recognition that central systolic and pulse pressure is more relevant than peripheral (brachial) measure for the prediction and pathophysiology of cardiovascular disease [20]. Several investigators indicated that the late systolic peak of peripheral (radial) pressure wave closely approximates central (aortic) systolic pressure [21–24]. Our present results extend these findings by showing that the association between central and peripheral AI is similar in different racial populations. Hence, peripheral AI reflects the ratio between central and peripheral pulse pressure [21]. It is important to emphasize that radial AI derives information about central wave reflection and central pulse pressure without using the general transfer function.

In conclusion, measurement of AI in the conveniently-located peripheral vasculature may provide a surrogate measure of central AI irrespective of difference in the race. Future cross-sectional studies comparing the Black adults (not studied in the present study) with other populations (Whites, Asians, etc.) as well as future prospective longitudinal studies to clarify the relation between radial AI and the morbidity and mortality of cardiovascular disease are warranted.

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References

1. London GM, Blacher J, Pannier B, Guerin AP, Marchais SJ, Safar ME. Arterial wave reflections and survival in end-stage renal failure. *Hypertension* 2001;38:434–438. [PubMed: 11566918]
2. Weber T, Auer J, O'Rourke MF, Kvas E, Lassnig E, Lamm G, Stark N, Rammer M, Eber B. Increased arterial wave reflections predict severe cardiovascular events in patients undergoing percutaneous coronary interventions. *Eur Heart J* 2005;26:2657–2663. [PubMed: 16183688]
3. Weber T, Auer J, O'Rourke MF, Kvas E, Lassnig E, Berent R, Eber B. Arterial stiffness, wave reflections, and the risk of coronary artery disease. *Circulation* 2004;109:184–189. [PubMed: 14662706]
4. Kelly R, Hayward C, Avolio A, O'Rourke M. Noninvasive determination of age-related changes in the human arterial pulse. *Circulation* 1989;80:1652–1659. [PubMed: 2598428]
5. Melenovsky V, Borlaug BA, Fetics B, Kessler K, Shively L, Kass DA. Estimation of central pressure augmentation using automated radial artery tonometry. *J Hypertens* 2007;25:1403–1409. [PubMed: 17563562]
6. Sugawara J, Komine H, Hayashi K, Maeda S, Matsuda M. Relationship between augmentation index obtained from carotid and radial artery pressure waveforms. *J Hypertens* 2007;25:375–381. [PubMed: 17211244]
7. Sugawara J, Komine H, Hayashi K, Yoshizawa M, Yokoi T, Maeda S, Tanaka H. Agreement between carotid and radial augmentation index: Does medication status affect the relation? *Artery Research* 2008;2:74–76.

8. Sugawara J, Hayashi K, Yokoi T, Cortez-Cooper MY, DeVan AE, Anton MA, Tanaka H. Brachial-ankle pulse wave velocity: an index of central arterial stiffness? *J Hum Hypertens* 2005;19:401–406. [PubMed: 15729378]
9. Cortez-Cooper MY, Supak JA, Tanaka H. A new device for automatic measurements of arterial stiffness and ankle-brachial index. *Am J Cardiol* 2003;91:1519–1522. [PubMed: 12804752]
10. Takazawa K, Tanaka N, Takeda K, Kurosu F, Ibukiyama C. Underestimation of vasodilator effects of nitroglycerin by upper limb blood pressure. *Hypertension* 1995;26:520–523. [PubMed: 7649592]
11. Snieder H, Hayward CS, Perks U, Kelly RP, Kelly PJ, Spector TD. Heritability of central systolic pressure augmentation: a twin study. *Hypertension* 2000;35:574–579. [PubMed: 10679500]
12. Hayward CS, Kelly RP. Gender-related differences in the central arterial pressure waveform. *J Am Coll Cardiol* 1997;30:1863–1871. [PubMed: 9385920]
13. Kelly RP, Millasseau SC, Ritter JM, Chowienzyk PJ. Vasoactive drugs influence aortic augmentation index independently of pulse-wave velocity in healthy men. *Hypertension* 2001;37:1429–1433. [PubMed: 11408390]
14. Smulyan H, Marchais SJ, Pannier B, Guerin AP, Safar ME, London GM. Influence of body height on pulsatile arterial hemodynamic data. *J Am Coll Cardiol* 1998;31:1103–1109. [PubMed: 9562014]
15. Wilkinson IB, MacCallum H, Flint L, Cockcroft JR, Newby DE, Webb DJ. The influence of heart rate on augmentation index and central arterial pressure in humans. *J Physiol* 2000;525(Pt 1):263–270. [PubMed: 10811742]
16. Latham RD, Westerhof N, Sipkema P, Rubal BJ, Reuderink P, Murgo JP. Regional wave travel and reflections along the human aorta: a study with six simultaneous micromanometric pressures. *Circulation* 1985;72:1257–1269. [PubMed: 4064270]
17. Murgo JP, Westerhof N, Giolma JP, Altobelli SA. Aortic input impedance in normal man: relationship to pressure wave forms. *Circulation* 1980;62:105–116. [PubMed: 7379273]
18. McDowell MA, Fryar CD, Ogden CL, Flegal KM. Anthropometric Reference Data for Children and Adults: United States, 2003–2006. *National Health Statistics Reports* 2008;10.
19. Ministry of Education C, Sports, Science and Technology of Japan: Physical Development and Health. 2007
20. Williams B, Lacy PS, Thom SM, Cruickshank K, Stanton A, Collier D, Hughes AD, Thurston H, O'Rourke M. Differential impact of blood pressure-lowering drugs on central aortic pressure and clinical outcomes: principal results of the Conduit Artery Function Evaluation (CAFE) study. *Circulation* 2006;113:1213–1225. [PubMed: 16476843]
21. Munir S, Guilcher A, Kamallesh T, Clapp B, Redwood S, Marber M, Chowienzyk P. Peripheral augmentation index defines the relationship between central and peripheral pulse pressure. *Hypertension* 2008;51:112–118. [PubMed: 17998476]
22. Pauca AL, Kon ND, O'Rourke MF. The second peak of the radial artery pressure wave represents aortic systolic pressure in hypertensive and elderly patients. *Br J Anaesth* 2004;92:651–657. [PubMed: 15003985]
23. Richardson CJ, Maki-Petaja KM, McDonnell BJ, Hickson SS, Wilkinson IB, McEniery CM. Comparison of estimates of central systolic blood pressure and peripheral augmentation index obtained from the Omron HEM-9000AI and SphygmoCor systems. *Artery Research* 2009;3:24–31.
24. Takazawa K, Kobayashi H, Shindo N, Tanaka N, Yamashina A. Relationship between radial and central arterial pulse wave and evaluation of central aortic pressure using the radial arterial pulse wave. *Hypertens Res* 2007;30:219–228. [PubMed: 17510503]

Carotid arterial pressure waveform**Radial arterial pressure waveform**

$$\text{Carotid AI (\%)} = \Delta P / \text{carotid PP} \times 100$$

$$\text{Radial AI (\%)} = P_2 / P_1 \times 100$$

Figure 1.

Augmentation index (AI) (*left*: carotid artery; *right*: radial artery). SBP=systolic blood pressure, DBP=diastolic blood pressure, PP=pulse pressure, P₁=first systolic peak – DBP, P₂=second systolic peak – DBP.

Figure 2.

(*Top*) Relation between carotid augmentation index (AI) and radial AI in Asian (closed circles) and White (open circles) adults. Asian (gray line): $y=0.93x-60.9$, $r=0.82$, $P<0.0001$; White (black line): $y=0.95x-64.1$, $r=0.75$, $P<0.0001$. (*Bottom*) Bland and Altman's Plots between carotid AI and radial AI in Asian and White adults. Mean differences were $66.5\pm10.3\%$ in Asian (gray lines) and $67.3\pm14.1\%$ in White (black lines).

Table 1

Physical characteristics

	Asian	White	P-value
N, male/female	52 / 42	27 / 20	n.s.
Age, yrs	45 ± 14	45 ± 17	n.s.
Height, cm	166 ± 8	171 ± 10	<0.01
Body mass, kg	63 ± 12	73 ± 15	<0.0001
BMI, kg/m ²	22.9 ± 3.0	25.1 ± 4.4	<0.001
Heart rate, bpm	61 ± 9	60 ± 10	n.s.
Systolic BP, mmHg	117 ± 10	120 ± 16	n.s.
Diastolic BP, mmHg	71 ± 8	69 ± 9	<0.10
Mean BP, mmHg	88 ± 8	88 ± 11	n.s.
Carotid AI, %	9.3 ± 18.0	3.4 ± 21.5	<0.10
Radial AI, %	75.9 ± 15.9	70.8 ± 17.0	<0.10
ΔAI, %	66.5 ± 10.3	67.3 ± 14.1	n.s.

Data are mean±SD. BMI = body mass index, BP = blood pressure, AI=augmentation index ΔAI = Radial AI – Carotid AI.