The Role of Blood Pressure in Carotid Plaque Incidence: Interactions With Body Mass Index, Age, and Sex-Based on a 7-Years Cohort Study

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Background: Although high blood pressure (BP) is a risk factor for carotid plaque, its long-term prognostic value might be underestimated due to its confounding interactions with BMI, age, and gender. Therefore, we conducted a 7-year prospective cohort study to evaluate the prognostic value of BP for the incidence of carotid plaque.

Methods: The subjects enrolled in 2011 were free of carotid plaque at baseline and were followed up in 2018. Multivariate Cox proportional-hazards models were used to evaluate the association between BP and carotid plaque incidence.

Results: During the follow-up study, the incidence of carotid plaque was 36.5%. The significant positive linear trend showed that subjects with higher BP levels at baseline were more likely to develop carotid plaques at the end. Especially in the female subpopulation, after confounders being adjusted, the carotid plaque was associated with higher BP (adjusted HR 1.52, 95% CI 1.02–2.26), pulse pressure (PP) (adjusted HR 1.15, 95% CI 0.76–1.75), and mean arterial pressure (MAP) (adjusted HR 1.44, 95% CI 1.00–2.08). The adjusted HRs of hypertension, PP, and MAP (HR 27.71, 95% CI 2.27–338.64; HR 14.47, 95% CI 1.53–137.18; HR 9.97, 95% CI 1.29–77.28) were significantly higher after the potential antagonistic interactions between BP categorical indicators and age being adjusted, respectively.

Conclusion: High BP indicators might be associated with higher HRs of carotid plaque after adjusting interactions between BP indicators and BMI, age, and gender, which suggests that the incidence of carotid plaque in female adults with high BP indicators might increase significantly with the increase of age.

Keywords: carotid arteries, blood pressure, interaction, cohort studies, age
INTRODUCTION

Approximately one-third of the adult population in China has been diagnosed with carotid atherosclerosis, leading to a heavy economic and social burden (Clarke et al., 2017). Prevention of carotid plaque is crucial because carotid plaque is associated with an increased risk of cardiovascular disease (Naqvi and Lee, 2014; Franceschini et al., 2018; Sillesen et al., 2018). The risk factors for carotid plaque include age, sex, hypertension, diabetes, and dyslipidemia (Noflatscher et al., 2018; Zhao and Hatsukami, 2018).

Hypertension is the leading modifiable risk factor for cardiovascular diseases including carotid plaque (Hellings et al., 2010; Franceschini et al., 2018; Sillesen et al., 2018). Inadequate awareness, improper therapeutic and poor controlling of hypertension might increase the risk of cardiovascular diseases (Zhao and Hatsukami, 2018). In a series of cohort studies and relatively small clinical trials studies, hypertension has been proposed as a risk factor for carotid plaque (Clarke et al., 2017; Steffen et al., 2018; Wang et al., 2018; Zhao and Hatsukami, 2018; GBD 2013 Mortality and Causes of Death, 2015). However, applying a single indicator of blood pressure (BP) to assess the risk of carotid plaque limits the ability in the evaluation of the prediction efficiency of different BP indicators, such as pulse pressure (PP), mean arterial pressure (MAP), and isolated systolic hypertension. Furthermore, the interactions between BP indicators and other cardiovascular risk factors for the carotid plaque have not been systematically analyzed (Yang et al., 2013; Clarke et al., 2017). Furthermore, this interaction may be involved in the formation of carotid plaque (Rovella et al., 2018) and the observational studies of these potential interactions were still unclear.

In this 7-year follow-up cohort study, we aimed to explore the possible interactions between BP categorical indicators and age, BMI, and gender in the associations of BP and incidence of carotid plaque.

MATERIALS AND METHODS

Subjects
A total of 2,410 subjects who were free of carotid plaque, coronary heart disease, stroke, or heart failure at baseline were recruited...
TABLE 1 | Baseline characteristics of subjects.

|                      | All (n = 1177) | Men (n = 555) | Women (n = 622) | p   |
|----------------------|----------------|--------------|-----------------|-----|
| Age (year), median (IQR) | 51.00 (41.00–61.00) | 49.00 (40.00–58.50) | 53.00 (43.00–62.00) | <0.001 |
| Abdominal circumference (cm), median (IQR) | 87.00 (81.00–93.00) | 87.00 (82.00–93.00) | 86.00 (80.00–92.00) | <0.001 |
| SBP (mmHg), median (IQR) | 118.00 (109.00–129.00) | 119.00 (110.00–128.00) | 118.00 (107.00–129.00) | 0.429 |
| DBP (mmHg), median (IQR) | 75.00 (70.00–80.00) | 75.00 (71.00–82.00) | 74.00 (69.00–79.00) | <0.001 |
| Marital status [n (%)] |                |              |                 |     |
| Married               | 858 (72.9%) | 244 (43.9%) | 614 (98.7%) |     |
| Widowed/divorced      | 319 (27.1%) | 311 (56.0%) | 8 (1.2%) |     |
| TC (mmol/L), median (IQR) | 4.90 (4.32–5.49) | 4.71 (4.14–5.21) | 5.02 (4.49–5.66) | <0.001 |
| TG (mmol/L), median (IQR) | 1.14 (0.81–1.57) | 1.24 (0.88–1.71) | 1.06 (0.77–1.43) | <0.001 |
| LDL-C (mmol/L), median (IQR) | 2.94 (2.40–3.45) | 2.86 (2.36–3.37) | 2.98 (2.42–3.54) | 0.012 |
| HDL-C (mmol/L), median (IQR) | 1.37 (1.18–1.58) | 1.25 (1.08–1.42) | 1.50 (1.30–1.66) | <0.001 |
| FPG (mg/dL), median (IQR) | 95.40 (88.20–102.60) | 95.40 (88.20–102.60) | 93.6 (88.20–102.60) | 0.327 |
| BMI (kg/m²) [n (%)] |                |              |                 |     |
| <18.5                | 615 (52.2%) | 245 (44.1%) | 370 (59.4%) | <0.001 |
| 18.5–24.0            | 445 (37.8%) | 248 (44.6%) | 197 (31.6%) |     |
| >24.0                | 117 (9.6%) | 62 (11.1%) | 55 (8.8%) | <0.001 |
| Hypertension [n (%)] |                |              |                 |     |
| No                   | 897 (76.2%) | 465 (66.4%) | 432 (67.0%) | 0.756 |
| Yes                  | 280 (23.7%) | 90 (33.6%) | 190 (30.5%) |     |
| Dyslipidemia [n (%)] |                |              |                 |     |
| No                   | 847 (71.9%) | 397 (71.5%) | 450 (72.3%) | 0.944 |
| Yes                  | 330 (28.0%) | 158 (28.4%) | 172 (27.7%) |     |
| FPG [n (%)]          |                |              |                 |     |
| Normal               | 786 (66.7%) | 369 (66.4%) | 417 (67.0%) |     |
| High-normal          | 187 (15.8%) | 91 (16.4%) | 96 (15.4%) |     |
| IFG                  | 89 (7.5%) | 40 (7.2%) | 49 (7.8%) |     |
| Diabetes             | 115 (9.7%) | 55 (9.9%) | 60 (9.6%) |     |

Values are n (%) or median (IQR).
SBP, systolic blood pressure; DBP, diastolic blood pressure; TC, total cholesterol; TG, triglyceride; LDL-C, low-density lipoprotein cholesterol; HDL-C, high-density lipoprotein cholesterol; BMI, body mass index; FPG, fasting plasma glucose; IFG, impaired fasting glucose; IQR, interquartile range.

from the Department of Healthcare of the Fourth Medical Center of the People’s Liberation Army General Hospital. The age of the study population was ranging from 35 to 74 years. Subjects with insufficient follow-up information and those who could not be contacted were excluded from the study. The baseline survey was conducted at 2011 with annual follow-up during physical examination, and the follow-up endpoint in this study was at 2018. Subjects lost to follow-up and those for whom relevant data were missing were not included in the data analyses (Figure 1). All subjects provided the written informed consent upon enrollment to the study. This study was approved by the ethics committee of Fourth Medical Center of the People’s Liberation Army General Hospital.

**Measurement of Carotid Plaque**

Carotid artery ultrasonography was performed using a B-mode ultrasound system (EPIQ7 or HD7XE, Philips, Amsterdam, Netherlands) with a linear array transducer (7.5–12 MHz). The examination included bilateral longitudinal and transverse scans of the distal segments of the common carotid artery, the bifurcation segments of the common carotid artery, and the proximal segments of the internal and external carotid arteries. We measured the intima-media thickness (IMT) and plaque formation in the wall of the eight segments of vessels mentioned above. The presence of plaque was defined as one of the following: IMT ≥ 1.5 mm, a focal structure (≥0.5 mm) protruding into the arterial lumen, or the surrounding IMT ≥ 50% (Collaborators, 2017; Dong et al., 2019).

Carotid plaque was defined as the development of at least one plaque in the eight previously plaque-free arterial segments during reexamination in 2018. To test for inter-observer reliability, plaque measurements in 25 patients were repeated after 1 week by two independent technicians and the results showed high reliability (intraclass correlation coefficients were 0.94).
Subjects were given a physical examination in the medical insurance hospital in May every year. The subjects were at a fasting state in the morning and BP was measured twice with a 3-minute interval by trained nurses using digital automatic monitors (HBP-1300; Omron, Netherlands), and an average measurement. Pre-hypertension was defined as SBP of 120–139 mmHg and DBP of 80–89 mmHg were defined as normal BP, while those with SBP ≥140 mmHg and/or DBP ≥90 mmHg (China Hypertension Prevention Control Guidelines Revision Committee, 2019).

To better understand the role of BP in carotid plaque incidence, we analyzed a range of BP categorical indicators, including hypertension, PP, isolated systolic hypertension, and MAP. Hypertension was defined as SBP >140 mmHg and/or DBP >90 mmHg according to an average BP of two measurements. Pre-hypertension was defined as SBP of 120–139 mmHg or DBP of 80–89 mmHg and normotensive subjects as SBP <120 mmHg or DBP <80 mmHg. Isolated systolic hypertension was defined as SBP ≥140 mmHg and DBP <90 mmHg. PP was categorized into three subgroups according to the tertile points such as T1 (<38 mmHg), T2 (38 mmHg–47 mmHg), and T3 (>47 mmHg), and MAP were divided into tertiles such as ≤86 mmHg, 86–94 mmHg, and >94 mmHg.

Fasting venous blood samples were drawn and assayed for glucose, total cholesterol (TC), triglyceride (TG), low-density lipoprotein cholesterol (LDL-C), high-density lipoprotein cholesterol (HDL-C), and other biochemical parameters using an automatic analyzer (COBAS-701, Roche, Mannheim, Germany). Body mass index (BMI) was calculated as weight (kg) divided by height squared (m²). Individuals were classified as underweight (BMI <18.5 kg/m²), normal (BMI 18.5–23.9 kg/m²), or overweight (BMI ≥24.0 kg/m²) based on the BMI limits defined for Chinese adults (Department of Disease Control, Ministry of Health of the People’s Republic of China, 2006). According to their fasting plasma glucose (FPG) levels, subjects were classified as normal FPG (<99.0 mg/dl), high-normal FPG (99.1–108.2 mg/dl), impaired FPG (108.3–125.9 mg/dl), or diabetes (≥126.0 mg/dl) (Ding et al., 2015).

### Covariants

| Variable | Male carotid plaque (n = 555) | Male carotid plaque (n = 622) |
|----------|-------------------------------|-------------------------------|
|          | Incidence (% 95% CI) | Total person-years | Incidence rate per 100 person-years (% 95% CI) | Incidence (% 95% CI) | Total person-years | Incidence rate per 100 person-years (% 95% CI) |
| Hypertension |                                |                               |                                |                                    |
| No (n = 893) | 159 17.8 (16.0–20.0) | 6,251                          | 2.5 (2.3–2.7) | 104 11.6 (9.6–13.6) | 6,251                          | 1.7 (1.7–1.7) |
| Yes (n = 284) | 52 18.3 (14.4–22.2) | 1,988                          | 2.6 (2.4–2.8) | 115 40.5 (34.6–46.4) | 1,988                          | 5.8 (5.6–6.0) |
| Blood pressure |                                |                               |                                |                                    |
| Normal (n = 529) | 77 14.5 (10.6–18.4) | 3,703                          | 2.1 (1.9–2.3) | 47 8.9 (8.7–9.1) | 3,703                          | 1.3 (1.1–1.5) |
| High-normal (n = 364) | 82 22.5 (18.6–26.4) | 2,548                          | 3.2 (3.1–3.4) | 57 15.6 (11.7–19.5) | 2,548                          | 2.2 (2.0–2.4) |
| Hypertension (n = 284) | 52 18.3 (14.4–22.2) | 1,988                          | 2.6 (2.4–2.8) | 115 40.5 (34.6–46.4) | 1,988                          | 5.8 (5.6–6.0) |
| Pulse pressure |                                |                               |                                |                                    |
| T1 (n = 415) | 52 12.5 (8.6–16.4) | 2,905                          | 1.8 (1.6–2.0) | 42 10.1 (8.1–11.2) | 2,905                          | 1.4 (1.2–1.6) |
| T2 (n = 394) | 79 20.1 (16.2–24.0) | 2,758                          | 2.9 (2.7–3.1) | 55 14.0 (10.1–17.9) | 2,758                          | 2.0 (1.8–2.2) |
| T3 (n = 368) | 80 21.7 (17.8–25.6) | 2,576                          | 3.1 (2.9–3.3) | 122 33.2 (29.3–37.1) | 2,576                          | 4.7 (4.5–4.9) |
| Isolated systolic hypertension |                                |                               |                                |                                    |
| No (n = 1086) | 192 17.7 (15.6–19.7) | 7,602                          | 2.5 (2.5–2.5) | 175 16.1 (14.1–18.1) | 7,602                          | 2.3 (2.3–2.3) |
| Yes (n = 91) | 19 20.9 (13.1–28.7) | 637                           | 3.0 (2.6–3.4) | 44 48.4 (38.6–58.2) | 637                           | 6.9 (6.5–7.3) |
| Mean arterial pressure |                                |                               |                                |                                    |
| T1 (n = 401) | 53 13.2 (9.3–17.1) | 2,807                          | 1.9 (1.7–2.1) | 49 12.2 (8.3–16.1) | 2,807                          | 1.7 (1.5–1.9) |
| T2 (n = 397) | 66 16.6 (12.7–20.5) | 2,779                          | 2.4 (2.2–2.8) | 72 18.1 (14.2–22.0) | 2,779                          | 2.6 (2.4–2.8) |
| T3 (n = 379) | 92 24.3 (20.4–28.2) | 2,663                          | 3.5 (3.3–3.7) | 98 25.9 (22.0–29.8) | 2,663                          | 3.7 (3.5–4.1) |
| Total | 211 38.0 (34.1–41.9) | 3,885                          | 5.4 (5.2–5.4) | 219 35.2 (31.3–39.1) | 3,885                          | 5.0 (4.8–5.2) |

CI, confidence interval; T1, in the first tertile; T2, in the second tertile; T3, in the third tertile.

Statistical Analyses

All data analyses were conducted with SPSS 24.0 (IBM, Armonk, NY, USA). Continuous data were presented as median and
interquartile range (IQR), while categorical data as the number of cases and percentage (%). The normality of continuous variables was tested using the Shapiro–Wilk normality test. Inter-group differences in skewed data, such as sex-related variables, were assessed for significance using the Kruskal–Wallis test. Differences in categorical variables were assessed using the chi-squared test and $p < 0.05$ was considered statistically significant.

Multivariate Cox proportional-hazards models were built using the “enter” method for variable selection (Ding et al., 2015). In Model 1, age, dyslipidemia, FPG category, and BMI category were adjusted. Model 2 was adjusted for the same variables in Model 1 and the interaction between age and BP categorical indicators. Model 3 was adjusted for the same variables as Model 1 and the interaction between BMI and hypertension. $E$-value was reported in the sensitivity analysis, which is related to the potentially subject to unmeasured confounding (VanderWeele and Ding, 2017).
TABLE 3 | Analysis of variables that may affect the association between hypertension and risk of carotid plaque.

| Variables | Men Model 1 | Men Model 2 | Men Model 3 | Women Model 1 | Women Model 2 | Women Model 3 |
|-----------|-------------|-------------|-------------|---------------|---------------|---------------|
|           | HR (95% CI) | HR (95% CI) | HR (95% CI) | HR (95% CI)   | HR (95% CI)   | HR (95% CI)   |
| Hypertension | No 1 (0.78–1.56) | 1.05 (0.56–1.98) | 1.40 (1.03–1.89) | 1.40 (1.03–1.89) | 8.80 (1.46–52.98) | 1.55 (1.06–2.28) |
|            | Yes 1.10 (0.78–1.56) | 0.88 (0.13–6.13) | 1.05 (0.56–1.98) | 1.40 (1.03–1.89) | 8.80 (1.46–52.98) | 1.55 (1.06–2.28) |
| Age (year) | 1.04 (1.02–1.05) | 1.04 (1.02–1.05) | 1.04 (1.02–1.05) | 1.06 (1.04–1.07) | 1.07 (1.05–1.09) | 1.06 (1.04–1.07) |
| BMI (kg/m²) | <18.5 1.27 (0.95–1.71) | 1.23 (0.89–1.72) | 0.90 (0.66–1.22) | 0.89 (0.66–1.21) | 1.04 (0.68–1.60) |
|            | 18.5–24.0 1.15 (0.71–1.87) | 1.24 (0.70–2.20) | 1.32 (0.87–2.00) | 1.22 (0.60–1.87) | 1.50 (0.71–3.18) |
| Dyslipidemia | No 1.01 (0.74–1.36) | 1.00 (0.74–1.36) | 1.14 (0.86–1.52) | 1.11 (0.83–1.48) | 1.15 (0.87–1.53) |
| FPG         | Normal 1.06 (0.65–1.36) | 1.00 (0.74–1.36) | 1.14 (0.86–1.52) | 1.11 (0.83–1.48) | 1.15 (0.87–1.53) |
|            | High-normal 0.85 (0.57–1.25) | 0.84 (0.57–1.24) | 0.99 (0.69–1.44) | 0.99 (0.68–1.43) | 1.00 (0.69–1.45) |
|            | IFG 0.96 (0.58–1.58) | 0.95 (0.57–1.56) | 0.92 (0.58–1.46) | 0.94 (0.59–1.48) | 0.92 (0.58–1.46) |
|            | Diabetes 1.13 (0.75–1.70) | 1.15 (0.76–1.75) | 1.14 (0.76–1.73) | 1.05 (0.71–1.56) | 1.06 (0.72–1.57) | 1.06 (0.71–1.58) |

Model 1, adjusts for hypertension, age, dyslipidemia, BMI, and FPG categories. Model 2, adjusts for these variables and for the interaction between age and hypertension. Model 3, adjusts for the same variables as Model 1 and the interaction between BMI and hypertension.

RESULTS

Baseline Characteristics of Subjects
A total of 2,410 subjects were recruited in 2011, and the last follow-up was conducted in 2018 including 1,177 subjects (555 men and 622 women), of which 430 new cases of carotid plaque were found. The median age of the subjects was 51 years (IQR 41–61) ranging from 35 to 74 years. The proportion of women was slightly higher than that of men (52.85% vs. 47.15%). Abdominal circumference is 87 (82–93) in men vs. 86 cm (80–92 cm) in women. BMI (18.5–24.0 kg/m²) is 44.6% in men vs. 31.6% in women. DBP level is 75 mmHg (71–82 mmHg) in men vs. 74 mmHg (69–79 mmHg) in women (Table 1).

Incidence of Carotid Plaque and Association With Baseline BP
After analyzing 8,239 person-years of follow-up, we identified 430 new-onset carotid plaque (36.5%; 95% CI 33.8–39.3%). The incidence was 5.22% per 100 person-years (95% CI 5.02–5.44%). The incidence of carotid plaque was significantly higher among male subjects and subjects whose baseline BP categorical indicators were higher (Table 2).

Incidence of Carotid Plaque and Its Association With BP Levels
In the analysis which takes BP (SBP, DBP, PP, and MAP) as continuous variables, the incidence of carotid plaque increased among male subjects with the increase of age-adjusted BP indicators. These associations were found in female subjects except the PP indicator (Figure 2).

Incidence of Carotid Plaque and Association With BP Categorical Indicators
Multivariate Cox proportional hazards models were generated to calculate hazard ratios (HRs) of carotid plaque incidence in men and women according to different BP indicators, after adjusting for age, dyslipidemia, BMI, and FPG categories, and significant interactions between BP and age and BMI. Among women, the adjusted HRs for the carotid plaque were 8.80 (95% CI 1.46–52.98; Model 2) and 1.55 (95% CI 1.06–2.28; Model 3) in the presence of hypertension relative to normal or high-normal BP, while Model 1 indicated a much smaller HR (1.40, 95% CI 1.03–1.89). These adjusted HRs of women were higher than that of men (Table 3). Among women, the adjusted HRs for carotid plaque were 1.98 (95% CI 1.21–3.24; Model 3) in the presence of hypertension relative to normal BP. The adjusted HR for carotid plaque was 27.71 (95% CI 2.27–338.64; Model 2) in the presence of hypertension relative to normal BP. Among men, the adjusted HRs in models 1–3 were not statistically significant (Table 4).

We analyzed the incidence of carotid plaque according to PP tertiles. Among women, the adjusted HRs for subjects in the third tertile were 14.47 (95% CI 1.53–137.18; Model 2) and 1.83 (95% CI 1.01–3.32; Model 3), relative to the first tertile, and Model
TABLE 4 | Analysis of variables that may affect the association between BP (normal, high-normal, and hypertension) and risk of carotid plaque.

| Variables                        | Men                      | Women                     |
|----------------------------------|--------------------------|---------------------------|
|                                  | Model 1 | Model 2 | Model 3 | Model 1 | Model 2 | Model 3 |
| Hypertension                     |         |         |         |         |         |         |
| Normal BP                        | 1       | 1       | 1       | 1       | 1       | 1       |
| High-normal BP                   | 1.04 (0.76–1.41)         | 1.15 (0.29–4.62)          | 1.16 (0.84–16.17) | 1.34 (0.96–1.87) | 14.70 (2.32–93.29) | 1.53 (0.93–2.54) |
| Hypertension                     | 1.14 (0.72–1.80)         | 1.28 (0.13–12.75)         | 2.44 (0.80–73.84) | 1.52 (1.02–2.26) | 27.71 (2.27–338.64) | 1.98 (1.21–3.24) |
| Age (year)                       | 1.04 (1.02–1.05)         | 1.04 (1.02–1.06)          | 1.04 (1.02–1.05) | 1.06 (1.04–1.07) | 1.08 (1.06–1.11) | 1.06 (1.02–1.07) |
| BMI (kg/m²)                      | <18.5  | 1       | 1       | 1       | 1       | 1       |
|                                 | 18.5–24.0 | 1.27 (0.95–1.71)         | 1.27 (0.94–1.71) | 1.29 (0.91–1.82) | 0.94 (0.69–1.27) | 0.89 (0.66–1.21) | 1.21 (0.66–2.20) |
|                                 | >24.0   | 1.15 (0.71–1.87)         | 1.15 (0.71–1.87) | 1.24 (0.63–2.45) | 1.33 (0.87–2.01) | 1.24 (0.81–1.90) | 2.09 (0.28–15.40) |
| Dyslipidemia                     | No      | 1       | 1       | 1       | 1       | 1       |
|                                 | Yes     | 1.01 (0.75–1.37)         | 1.01 (0.75–1.39) | 1.01 (0.75–1.37) | 1.21 (0.91–1.60) | 1.17 (0.89–1.55) | 1.13 (0.851–1.51) |
| FPG                              | Normal  | 1       | 1       | 1       | 1       | 1       |
|                                 | Upper range of normal    | 0.85 (0.58–1.26)          | 0.85 (0.58–1.26) | 0.85 (0.57–1.25) | 1.02 (0.70–1.47) | 1.00 (0.69–1.46) | 1.01 (0.70–1.47) |
|                                 | IFG     | 0.96 (0.58–1.58)         | 0.96 (0.58–1.59) | 0.95 (0.58–1.58) | 0.94 (0.59–1.48) | 0.96 (0.61–1.51) | 0.93 (0.59–1.47) |
|                                 | Diabetes| 1.14 (0.76–1.72)         | 1.14 (0.76–1.72) | 1.12 (0.75–1.69) | 1.06 (0.72–1.57) | 1.08 (0.73–1.59) | 1.05 (0.71–1.57) |

Model 1, adjusting for hypertension, age, BMI, dyslipidemia, and FPG categories. Model 2, adjusting for hypertension, age, BMI, dyslipidemia, FPG categories, and interaction of age by hypertension. Model 3, adjusts for the same variables as Model 1 and the interaction between BMI and hypertension.

BMIs, body mass index; BP, blood pressure; FPG, fasting plasma glucose; IFG, impaired fasting glucose; HR, hazard ratio; CI, confidence interval.

1 gave a smaller HR of 1.15 (95% CI 0.76–1.75). Among men, the adjusted HRs for subjects in tertiles in Models 1–3 were not statistically significant and lower than that of women (Table 5).

When we analyzed the incidence according to MAP tertiles, among women, the adjusted HRs for subjects in the third tertile were 9.97 (95% CI 1.29–77.28; Model 2) and 2.05 (95% CI 1.25–3.35; Model 3), relative to the first tertile, and Model 1 gave a smaller HR of 1.44 (95% CI 1.00–2.08). Among men, the adjusted HRs for subjects in tertiles in Models 1–3 were not statistically significant and lower than that of women (Table 6). Isolated systolic hypertension was not significantly associated with carotid plaque after adjustment.

Sensitivity Analysis
E-value was calculated to estimate the potential effects of unmeasured significant confounding factors on the carotid plaque. When E-value was more than two, considerable unmeasured significant confounding factors could be needed to negate the existing adjusted HRs. In the sensitivity analysis, after adjusted for interactions between BP and BMI and age, all the E-value were more than two, which indicated the current associations tended to be more stable.

DISCUSSION
This cohort study evaluated the associations between BP indicators and the incidence of carotid plaque in Chinese adults. The findings showed that higher BP indicators might be associated with higher HRs of carotid plaque after adjusting for interactions between BP indicators and parameters such as BMI, age, and gender. These associations were significant in female subjects.

It is well-established that age, obesity, and hypertension are independent risk factors for carotid plaque (Genuth et al., 2003; Mahmoudi et al., 2011; Schwartz et al., 2016; Colafella and Denton, 2018). It is clear that there is an association between hypertension and the incidence of carotid plaque, but few studies have found that whether interactions among hypertension and its important influencing factors such as age, BMI, gender, and also whether the interactions could increase their influence on the onset of carotid plaque (Rovella et al., 2018). In this study, we observed that interactions of BP indicators with age or BMI were associated with the carotid plaque incidence in female adults. After the confounding factors and interactions of BP indicators with age and BMI being adjusted, high BP indicators significantly elevated the HRs of carotid plaque, while the HRs were lower if we adjusted only for confounding factors. Results were similar when we estimated the influence of PP and MAP tertiles on the incidence of carotid plaque. This suggests that age and BMI may act as both confounders and effect modifiers to influence the HRs of carotid plaque in female population. In addition, obesity was found to interact with age, sex, and BP on the incidence of carotid plaque, and the risk of carotid plaque due to overweight or obesity after adjusting for these interactions was substantially higher than that of when interactions were not controlled (Clarke et al., 2017). This might suggest that, after interactions being adjusted, the
Another potential explanation is gender differences in the sex hormones estrogen and testosterone and in the sex chromosome complement (Schwartz et al., 2016). In addition, it could be due to the sex differences in lifestyle. Unhealthy lifestyle habits such as smoking, drinking, and staying up late are typically more common in men (Liu and Li, 2015; Rosendorff et al., 2015). The incidence of carotid plaque might be attributed to the fact that the effect of an unhealthy lifestyle on the carotid plaque was more than that of BP indicators. Future work should explore some of these hypotheses in detail to clarify the apparent sex differences in how BP influences the incidence of carotid plaque.

There is a significant difference in BP changes both in men and women in the whole life cycle. Among middle-aged and elderly women, due to hormone levels, obesity and aging, the average BP level increased faster, and they were more likely to suffer from hypertension. In addition, women might be more likely to be obese during and after menopause. Moreover, obesity increases BP in men and women, but women have higher BP. Therefore, middle-aged and elderly women are more likely to suffer from obesity, hypertension, and the combined effects of these factors multiply increase the risk of cardiovascular diseases (Ventura-Clapier et al., 2017; Colafella and Denton, 2018).

This study is a 7-year follow-up with a large sample size, which allowed us to analyze 430 cases of new-onset carotid plaque. It is of great significance to explore the influencing factors of carotid plaques formation since it can provide important evidence for the primary prevention of cardiovascular diseases.
TABLE 6 | Analysis of variables that may affect the association between mean arterial pressure (MAP) tertiles and risk of carotid plaque.

|                  | Men Model 1 | Men Model 2 | Men Model 3 | Women Model 1 | Women Model 2 | Women Model 3 |
|------------------|-------------|-------------|-------------|---------------|---------------|---------------|
|                  | HR (95% CI) | HR (95% CI) | HR (95% CI) | HR (95% CI)   | HR (95% CI)   | HR (95% CI)   |
| MAP T1           | 1           | 1           | 1           | 1             | 1             | 1             |
|                  | 0.99 (0.69–1.42) | 0.64 (0.13–3.10) | 1.22 (0.72–2.06) | 1.17 (0.80–1.69) | 4.16 (0.51–34.07) | 1.39 (0.84–2.30) |
| T2               | 1.17 (0.82–1.67) | 1.75 (0.37–8.42) | 1.17 (0.67–2.02) | 1.44 (1.00–2.08) | 9.97 (1.29–77.28) | 2.05 (1.25–3.35) |
| T3               | 1.04 (1.02–1.05) | 1.04 (1.01–1.06) | 1.04 (1.02–1.05) | 1.06 (1.05–1.07) | 1.08 (1.05–1.11) | 1.06 (1.04–1.07) |
| Age (year)       | 1.04 (1.02–1.05) | 1.04 (1.01–1.06) | 1.04 (1.02–1.05) | 1.06 (1.05–1.07) | 1.08 (1.05–1.11) | 1.06 (1.04–1.07) |
| BMI (kg/m²)      | 1           | 1           | 1           | 1             | 1             | 1             |
| <18.5            | 1.26 (0.94–1.68) | 1.27 (0.94–1.71) | 1.52 (0.87–2.64) | 0.92 (0.68–1.25) | 0.89 (0.66–1.21) | 1.44 (0.81–2.56) |
| 18.5–24.0        | 1.14 (0.70–1.84) | 1.15 (0.70–1.86) | 0.90 (0.21–3.83) | 1.32 (0.87–2.01) | 1.28 (0.84–1.96) | 3.24 (0.44–24.13) |
| >24.0            | 1.00 (0.74–1.36) | 0.99 (0.73–1.35) | 1.01 (0.74–1.37) | 1.21 (0.91–1.60) | 1.18 (0.89–1.56) | 1.18 (0.89–1.57) |
| Dyslipidemia     | 1           | 1           | 1           | 1             | 1             | 1             |
| No               | 0.85 (0.58–1.25) | 0.86 (0.58–1.26) | 0.84 (0.57–1.25) | 1.02 (0.70–1.47) | 0.99 (0.69–1.44) | 1.02 (0.70–1.48) |
| Yes              | 0.97 (0.59–1.60) | 0.96 (0.58–1.58) | 0.96 (0.58–1.59) | 0.93 (0.59–1.47) | 0.94 (0.60–1.49) | 0.94 (0.59–1.49) |
| FPG              | 1           | 1           | 1           | 1             | 1             | 1             |
| Normal           | 1.14 (0.76–1.71) | 1.13 (0.75–1.69) | 1.14 (0.76–1.71) | 1.09 (0.74–1.61) | 1.09 (0.73–1.61) | 1.09 (0.74–1.62) |
| Upper range of normal | 0.97 (0.59–1.60) | 0.96 (0.58–1.58) | 0.96 (0.58–1.59) | 0.93 (0.59–1.47) | 0.94 (0.60–1.49) | 0.94 (0.59–1.49) |
| IFG              | 1.14 (0.76–1.71) | 1.13 (0.75–1.69) | 1.14 (0.76–1.71) | 1.09 (0.74–1.61) | 1.09 (0.73–1.61) | 1.09 (0.74–1.62) |

Model 1, adjusting for MAP tertiles, age, BMI, dyslipidemia, and FPG categories. Model 2, adjusting for MAP tertiles, age, BMI, dyslipidemia, FPG categories, and interaction of age by MAP tertiles. Model 3, adjusts for the same variables as Model 1 and the interaction between BMI and MAP tertiles. T1, the first tertile; T2, the second tertile; T3, the third tertile. BMI, body mass index; MAP, mean arterial pressure; FPG, fasting plasma glucose; IFG, impaired fasting glucose; HR, hazard ratio; CI, confidence interval.

Another strength is that the diagnosis of carotid plaque was conducted using color ultrasonography to detect IMT of the bilateral carotid arteries at eight locations. Meanwhile, we proposed that high BP significantly increases the HRs of carotid plaque after adjusting for covariants and interactions between BP indicators and parameters such as BMI, age, and gender. The influence of hypertension on carotid plaque formation depends on age, sex, and BMI to some extent. Clinicians should pay more attention to BP management of elder and fatter women to prevent carotid plaque.

Limitations of this study are the lack of information on BP changes and follow-up. Therefore, this result should be interpreted with caution since baseline BP cannot capture fluctuations in BP during follow-up, so we were unable to explore the effects of ambulatory BP on the incidence of carotid plaque. In addition, a lack of information on the occurrence of carotid plaque during a 7-year follow-up may cause some bias. Moreover, we did not collect complete data on the use of antihypertensive or other drugs for cardiovascular disease, or on lifestyle factors that might influence the incidence such as smoking, drinking, and physical activity.

CONCLUSION

In this study, we find that high BP indicators might be associated with higher HRs of carotid plaque after the interactions being adjusted in female adults, suggesting that the incidence of carotid plaque in female adults with high BP indicators and higher age or BMI might significantly increase. This hypothesis, however, needs further verification.

DATA AVAILABILITY STATEMENT

The raw data supporting the conclusions of this article will be made available by the authors, without undue reservation.

ETHICS STATEMENT

The studies involving human participants were reviewed and approved by Ethics Committee of the fourth medical center of PLA General Hospital. The patients/participants provided their written informed consent to participate in this study. Written informed consent was obtained from the individual(s) for the publication of any potentially identifiable images or data included in this article.

AUTHOR CONTRIBUTIONS

JL and SW conceived the study and its design. JL, XM, X-LR, LY, HX, and ZL collected the data. JL, SW, XM, and X-LR managed, analyzed, and interpreted the data. All the authors have read and approved the final manuscript.
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Conflict of Interest: The authors declare that the research was conducted in the absence of any commercial or financial relationships that could be construed as a potential conflict of interest.

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