Prevalence of left atrial enlargement and its risk factors in general Chinese population

Qiaoyun Ou, Yintao Chen, Shasha Yu, Xiaofan Guo, Huijie Zhao and Yingxian Sun*

Abstract

Background: Left atrial enlargement (LAE) has been proven to be significantly related to stroke and cardiovascular diseases. In China, few studies related to LAE have been conducted, especially in the general population. To the best of our knowledge, our study is the first to explore the prevalence of LAE and associated risk factors in the general Chinese population.

Methods: This study included a total of 11,956 subjects aged ≥35 years. All participants completed a questionnaire, and underwent complete physical examination, electrocardiogram (ECG) and echocardiogram. We defined LAE as a LA diameter exceeding 4.0 cm in men and 3.8 cm in women. We conducted a multivariable logistic regression analysis and a linear regression analysis to identify independent factors of LAE.

Results: The overall prevalence of LAE was 6.43 % for subjects aged over 35 years. The prevalence of LAE was 6.78 % in women and 6.02 % in men. The major risk factors of LAE were female sex (odds ratio [OR]: 1.229, β: 0.197), advancing age (OR: 1.015, β: 0.036), high systolic blood pressure (OR: 2.331, β: 0.185), high body mass index (BMI) (OR: 3.956, β: 0.373), diabetes (OR: 1.498, β: 0.030), high left ventricular myocardial index (OR: 1.003, β: 0.073), and low left ventricular ejection fraction, low heart rate, and low estimated glomerular filtration rate. Additionally, the association between BMI and LAE was the most obvious.

Conclusions: Female sex, advancing age, high systolic blood pressure, high BMI, diabetes, high left ventricular myocardial index, low estimated glomerular filtration rate, low left ventricular ejection fraction, and low heart rate were proven to be risk factors of LAE.

Keywords: Left atrial enlargement, Prevalence, Risk factors, General Chinese population

Background

Several studies have shown that alterations in the left ventricular structure and function are associated with an increased risk of cardiovascular morbid and fatal events [1–3]. Recently, left atrial enlargement (LAE) has been proven to be significantly related to stroke and cardiovascular diseases [4–7]. Furthermore, some studies have found that LAE may be a more sensitive indicator of cardiovascular diseases, compared with left ventricular remodeling [8, 9]. Therefore, the study of the prevalence of LAE and its risk factors is becoming increasingly important. However, very few studies conducted on the general population have focused on LAE. At present, almost all of the research is aimed at patients with high blood pressure (BP), or those who are hospitalized. Moreover, no research studies on the general Chinese population have focused on LAE. To the best of our knowledge, our study is the first to explore the prevalence of LAE and its risk factors in the general Chinese population, including over 10,000 participants. The aims of our study were as follows: (1) to investigate the prevalence of LAE in a rural Chinese population consisting of a general adult population rather than patients that were indicated to undergo echocardiographic examination for different reasons, (2) and to explore independent risk factors of LAE.

Methods

Study population

From January 2012 to August 2013, a representative sample of participants aged ≥35 years was selected to
characterize the prevalence and risk factors in rural areas of Liaoning Province, located in China. The study adopted a multi-stage, stratified, random-cluster sampling scheme. In the first stage, three counties (Dawa, Zhangwu and Liaoayang County) were randomly selected from Liaoning province. In the second stage, one town was randomly selected from each county (for a total of three towns). In the third stage, 8–10 rural villages from each town were randomly selected (for a total of 26 rural villages). Participants who were pregnant or had malignant tumors or mental disorders were excluded from the study, comprising a potential pool of 14,016 people. Of these, 11,956 participants agreed and completed the present study, yielding a response rate of 85.3 %. In this report, we used only the data from participants who completed the study, which provided a final sample size of 10,574 subjects (4768 men and 5806 women). For the present analysis, participants with atrial fibrillation showed by Electrocardiogram that made diastolic function and diameter assessment unreliable with conventional Doppler technique were also excluded.

The study was approved by the Ethics Committee of China Medical University (Shenyang, China). All procedures were performed in accordance with ethical standards. Written consent was obtained from all participants after they had been informed of the objectives, benefits, medical items and confidentiality agreement regarding their personal information. For participants who were illiterate, we obtained written informed consent from their proxies.

Data collection and measurements
Data were collected during a single visit to the clinic by cardiologists and trained nurses using a standard questionnaire in a face-to-face interview. Before the survey was performed, we invited all eligible investigators to attend an organized training session. The training included the purpose of this study, how to administer the questionnaire, the standard method of measurement, the importance of standardization and the study procedures. A strict test was administered after this training, and only those who scored perfectly on the test were accepted as investigators in this study. During data collection, our inspectors gave some further instructions and support. Data regarding the demographic characteristics, lifestyle risk factors, dietary habits, family income and family history of chronic diseases were obtained during the interview using the standardized questionnaire. The study was guided by a central steering committee with a subcommittee for quality control.

According to American Heart Association protocol, BP was measured three times at 2-min intervals after at least 5 min of rest using a standardized automatic electronic sphygmomanometer (HEM-907; Omron), which had been validated according to the British Hypertension Society protocol [10]. The participants were advised to avoid caffeinated beverages and exercise for at least 30 min before the measurement. During the measurement, the participants were seated with their arms supported at the level of the heart. The mean of three BP measurements was calculated and used in all analyses. Weight and height were measured to the nearest 0.1 kg and 0.1 cm, respectively, with the participants wearing light-weight clothing and without shoes. Body mass index (BMI) was calculated as the weight in kilograms divided by the square root of the height in meters. Fasting blood samples were collected in the morning after at least 12 h of fasting. Blood samples were obtained from an antecubital vein into Vacutainer tubes containing ethylenediaminetetraacetic acid (EDTA). Fasting plasma glucose (FPG), total cholesterol (TC), low-density lipoprotein cholesterol (LDL-C), high-density lipoprotein cholesterol (HDL-C), triglycerides (TGs) and other routine blood biochemical indexes were analyzed enzymatically using an autoanalyzer. All laboratory equipments were calibrated, and blinded duplicate samples were used for these analyses.

Echocardiographic measurements
Echocardiograms were obtained using a commercially available Doppler echocardiograph (Vivid, GE Healthcare, United States), with a 3.0-MHz transducer. The transthoracic echocardiogram examinations included M-mode, two-dimensional, spectral and color Doppler, and were performed with subjects resting in the supine position. Echocardiogram analyses and readings were performed by three doctors specialized in echocardiography. Consultations were made to two other specialists if any questions or uncertainty arose. The parasternal acoustic window was used to record two-dimensional and M-mode images of the left ventricle (LV) internal diameter, wall thickness, aortic root and left atrium (LA). The apical acoustic window was used to record 4- and 5-chamber images. Color Doppler recordings were adopted to identify valvular regurgitation. The correct orientation of imaging planes and Doppler recordings were verified using previously described procedures [11, 12]. The left ventricular end-diastolic dimension (LVIDd) was obtained in the LV minor axis at end-diastole, internal dimensions and interventricular septal thickness (IVST) and posterior wall thickness (PWT) were measured at end of the diastole and systole according to the recommendations of the American Society of Echocardiography [13, 14]. The LV mass was estimated by Devereux’s formula $0.8 \times [1.04 \times \left(\text{LVIDd} + \text{PWT} + \text{SWT}\right)^3 - \text{LVIDd}^3] + 0.6 \text{ g} [15]$ and normalized to body surface area (BSA). Left ventricular ejection fraction (LVEF) was measured from the four-chamber apical projection by the area product $\times$ the ventricular length. Two-dimensional guided
M-mode measurements of the LA posteroanterior dimension were performed from the parasternal long-axis view according to the standards of the American Society of Echocardiography.

Definitions
Left atrial enlargement was defined as an LA diameter exceeding 4.0 cm in men and 3.8 cm in women [16]. LAE as the index by body surface area (iLAE) was defined as indexed LA diameter exceeding 2.3 cm/m2 in both sexes [16]. Higher left ventricular myocardial index (LVMI) was defined as LVMI exceeding 115 g/m2 in men and 95 g/m2 in women [16]. According to the Standards of the American Society of Echocardiography [17], left atrial enlargement was defined as an LA diameter exceeding 4.0 cm in men and 3.8 cm in women [16]. According to the World Health Organization (WHO) criteria [18], left atrial enlargement was defined as indexed LA diameter exceeding 2.3 cm/m2 in men and 2.3 cm/m2 in women according to the American Society of Echocardiography [17].

Statistical analysis
Descriptive statistics were calculated for all the variables, including continuous variables (reported as mean values and standard deviations) and categorical variables (reported as numbers and percentages). The differences between the LAE and non-LAE groups were evaluated using the Student’s t-test, analysis of variance, non-parametric test or the χ2 test, as appropriate. Multivariate logistic regression analyses and linear regression analysis were used to identify independent factors of LAE, and odds ratios (ORs). Linear correlation coefficient β and corresponding 95% confidence intervals (CIs) also were calculated. All the statistical analyses were performed using SPSS version 17.0 software (SPSS Inc, Chicago, Illinois, USA), and P values less than 0.05 were considered statistically significant.

Results
Background characteristics
Patient baseline characteristics are shown in Table 1. A total of 10,574 participants (4768 men and 5806 women) were included in the study. The mean age was 53.81 years. The subjects in the LAE group were older and included a higher proportion of women than the non-LAE group (P < 0.001 for both). Participants with LAE had higher BP, higher body mass index (BMI), higher prevalence of diabetes mellitus, higher TG, higher TC, higher LVMI, lower left ventricular ejection fraction, and lower estimated glomerular filtration rate (eGFR) than those in the non-LAE group (P < 0.001).

Prevalence of LAE in different sexes and ages
The overall prevalence of LAE was 6.43% for subjects aged over 35 years. The prevalence of LAE was 6.78% in women and 6.02% in men. For men, the prevalence of LAE increased with advancing age (age 35–44 years: 3.32%; age 45–54 years: 5.31%; age 55–64 years: 6.95%; and age ≥65 years: 9.31%). For women, the prevalence of LAE was 3.72, 5.03, 9.17, 11.11%, respectively. As for the total population, the prevalence was 3.55, 5.44, 8.85, 11.37% respectively (Fig. 1).

Factors associated with LAE
Table 2 shows multivariable logistic regression analysis of risk factors related to LAE. The prevalence of LAE was higher for women than men (OR: 1.229, 95% CI: 1.003–1.507). Advancing age (OR: 1.015, 95% CI: 1.005–1.026), high systolic blood pressure (SBP) (OR: 2.331, 95% CI: 1.889–2.876), high diastolic blood pressure (DBP) (OR: 1.24, 95% CI: 1.016–1.513), high BMI (OR: 3.956, 95% CI: 3.208–4.879), high prevalence of diabetes (OR: 1.498, 95% CI: 1.19–1.886), high LVMI (OR: 1.003, 95% CI: 1.002–1.004), and low LVEF, lower heart rate, and lower eGFR were found to be the major risk factors of LAE. After discussing the association between men and women, respectively, no obvious correlation was found for DBP and LAE. The associations between diabetes, eGFR and LAE were significant in men (P < 0.05 vs. P < 0.05), but not in women (P = 0.263 vs. P = 0.751). Additionally, we tested the association by linear regression analysis (Table 3). Female sex, advancing age, high SBP, high BMI, high fasting plasma glucose, high LVMI, low eGFR, low LVEF, and low heart rate were proven to be highly significant risk factors of LAE. Additionally, the association between BMI and LAE was shown to be the most obvious.

Discussion
Our study found that the prevalence of LAE was 6.43% in the total population, 6.78% in women, and 6.02% in men. A survey found that the prevalence of LAE defined according to LA diameter in an urban population in
Poland was 15.7%. As for the indexed LA diameter, the prevalence was 8.8% [22]. It was found that their participants were much older (mean age of 63.0 years) and obese (mean BMI of 27.9 kg/m²), which may be the reason for the higher LAE prevalence among them than in our population. Stritzke et al. reported that the prevalence of LAE consists of 9.8% the German residents [23]. However, they defined LAE as \(iLA \geq 35.7\) and \(\geq 33.7\) ml/m² in men and women, respectively. Thus, we suspect that the differences in prevalence between their study and ours can be attributed to the different population studied, different age levels, different BMI levels, different BP levels, and different definitions of LAE. Further, compared to Europeans and Americans, Asians tend to have smaller, thinner bodies. Furthermore, our research population was much larger. Thus, it is understandable that the prevalence of LAE in our study was relatively low.

In addition, we also found that the prevalence of LAE as the index by BSA was higher than that defined by absolute LA diameter, especially among the women population. After indexed by body surface area, the prevalence of \(iLA\) was 15.35% in total population, 20.58% in female, and 9.09% in male. This finding was different from some studies in Europe and America [22]. The reason for this difference is that Asians tend to have smaller, thinner bodies, compared to Europeans and Americans, especially in elder women. Their BSA was smaller, so when LAE was defined as LA diameter normalized to BSA >2.3 cm/m² in both sexes, the prevalence of LAE in Asians became higher than Europeans and Americans, and the prevalence of \(iLA\) was significantly higher in females than in males. A study in Japan confirmed the results to

### Table 1 Characteristics of the study population

| Characteristics          | Total       | LAE          | Non-LAE      | \(P\)  |
|--------------------------|-------------|--------------|--------------|--------|
| Gender, female           | 5991 (54.16 %) | 406 (57.10 %) | 5585 (53.96 %) | 0.104  |
| Age, y                   | 53.81 ± 10.52 | 57.86 ± 10.40 | 53.53 ± 10.47 | <0.001 |
| Current smoking          | 3890 (35.17 %) | 194 (27.29 %) | 3696 (35.71 %) | <0.001 |
| Current drinking         | 2476 (22.39 %) | 151 (21.24 %) | 2325 (22.46 %) | 0.448  |
| SBP, mmHg                | 141.63 ± 23.36 | 158.18 ± 25.64 | 140.5 ± 22.76 | <0.001 |
| DBP, mmHg                | 81.97 ± 11.70  | 86.87 ± 12.52  | 81.63 ± 11.49  | <0.001 |
| HR, beats minute         | 71.66 ± 12.3   | 71.03 ± 11.90  | 71.70 ± 12.33  | 0.165  |
| BMI, Kg/m²               | 24.78 ± 3.66   | 27.54 ± 3.77   | 24.60 ± 3.58   | <0.001 |
| Diabetes mellitus        | 1148 (10.38 %)  | 133 (18.71 %)  | 1015 (9.81 %)  | <0.001 |
| HDL-C, mol/L             | 1.41 ± 0.38    | 1.32 ± 0.32    | 1.41 ± 0.38    | <0.001 |
| LDL-C, mol/L             | 2.93 ± 0.82    | 3.08 ± 0.85    | 2.92 ± 0.82    | <0.001 |
| TG, mmol/L               | 1.63 ± 1.47    | 1.89 ± 1.42    | 1.62 ± 1.48    | <0.001 |
| TC, mmol/L               | 5.24 ± 1.09    | 5.38 ± 1.11    | 5.23 ± 1.08    | <0.001 |
| eGFR, ml/min/1.73 m²     | 92.81 ± 15.87  | 88.02 ± 18.51  | 93.14 ± 15.62  | <0.001 |
| Hemoglobin, g/L          | 138.6 ± 18.74  | 138.28 ± 24.50 | 138.63 ± 18.29 | 0.638  |
| LVDD, cm                 | 4.71 ± 0.43    | 5.07 ± 0.55    | 4.68 ± 0.41    | <0.001 |
| IVST, cm                 | 0.89 ± 0.26    | 1.01 ± 0.38    | 0.88 ± 0.24    | <0.001 |
| PWT, cm                  | 0.87 ± 0.26    | 0.96 ± 0.36    | 0.86 ± 0.25    | <0.001 |
| LVMI, g/m²               | 86.83 ± 59.31  | 110.44 ± 87.80 | 85.21 ± 56.48  | <0.001 |
| LVEF, %                  | 62.97 ± 3.81   | 61.33 ± 4.77   | 63.09 ± 3.71   | <0.001 |
| E/A                      | 1.06 ± 4.28    | 0.91 ± 0.74    | 1.07 ± 4.42    | 0.339  |

Abbreviations: LAE left atrial enlargement, Non-LAE non-left atrial enlargement, BMI body mass index, SBP systolic blood pressure, DBP diastolic blood pressure, TC total cholesterol, TG triglycerides, LDL-C low density lipoprotein cholesterol, HDL-C high density lipoprotein cholesterol, eGFR glomerular filtration rate, HR heart rate, LVDD left ventricular end-diastolic dimension, IVST interventricular septal thickness, PWT posterior wall thickness, LVEF left ventricular ejection fraction

Note: data are expressed as mean ± standard deviation or n (%)
Table 2 Multivariable logistic regression analysis of risk factors related to left atrial enlargement

| Characteristics | Total OR (95 % CI) | Male OR (95 % CI) | Female OR (95 % CI) |
|-----------------|-------------------|-------------------|---------------------|
| Gender, female  | 1.229 (1.003–1.507) | 1.007 (0.992–1.022) | 1.017 (1.002–1.032) |
| Age             | 1.015 (1.005–1.026) | 1.033 (0.977–0.988) | 0.978 (0.968–0.988) |
| Current smoking | 0.741 (0.607–0.905) | 0.741 (0.570–0.963) | 0.730 (0.533–0.998) |
| Current drinking| 1.274 (1.001–1.62)  | 1.291 (0.987–1.688) | 1.308 (0.715–2.393) |
| SBP ≥ 140       | 2.331 (1.889–2.876) | 2.673 (1.901–3.758) | 2.033 (1.551–2.664) |
| DBP ≥ 80        | 1.24 (1.016–1.513)  | 1.302 (0.966–1.754) | 1.161 (0.888–1.518) |
| HR              | 0.980 (0.972–0.987) | 0.979 (0.967–0.990) | 0.978 (0.968–0.988) |
| BMI ≥ 30        | 3.956 (3.208–4.879) | 4.947 (3.53–6.932)  | 3.414 (2.61–4.467)  |
| Diabetes mellitus| 1.498 (1.19–1.886)  | 2.042 (1.443–2.889) | 1.192 (0.877–1.518) |
| TC              | 0.953 (0.880–1.033) | 0.977 (0.860–1.11)  | 0.931 (0.839–1.033) |
| TG              | 1.029 (0.977–1.083) | 0.973 (0.899–1.054) | 1.076 (1.002–1.154) |
| eGFR            | 0.991 (0.985–0.997) | 0.979 (0.969–0.989) | 0.999 (0.991–1.007) |
| Anemia          | 1.253 (0.981–1.6)   | 1.222 (0.771–1.938) | 1.326 (0.990–1.774) |
| LVMI            | 1.003 (1.002–1.004) | 1.002 (1.000–1.003) | 1.004 (1.003–1.006) |
| LVEF            | 0.928 (0.911–0.940) | 0.939 (0.913–0.966) | 0.919 (0.896–0.942) |
| E/A             | 0.892 (0.703–1.133) | 1.017 (0.934–1.107) | 0.523 (0.348–0.788) |

Table 3 Linear regression analysis of risk factors related to left atrial enlargement

| Characteristics | Total β (95 % CI) | Male β (95 % CI) | Female β (95 % CI) |
|-----------------|------------------|------------------|-------------------|
| Gender, female  | 0.197 (–0.174, –0.139) | 0.000 | 0.004 (0.0005, 0.003) |
| Age             | 0.036 (0.0005, 0.002) | 0.002 | 0.017 (–0.0006, 0.002) |
| Current smoking | 0.001 (–0.015, 0.016) | 0.897 | 0.001 (–0.020, 0.021) |
| Current drinking| 0.030 (0.009, 0.047) | 0.003 | 0.037 (0.008, 0.049) |
| Mean SBP        | 0.185 (0.003, 0.004)  | 0.185 | 0.176 (0.002, 0.003) |
| Mean DBP        | –0.045 (–0.002, –0.0006) | 0.034 | –0.018 (–0.002, 0.001) |
| HR              | –0.122 (–0.005, –0.003) | 0.000 | –0.138 (–0.005, –0.004) |
| BMI             | 0.373 (0.038, 0.042)  | 0.000 | 0.380 (0.038, 0.044) |
| FPG             | 0.030 (0.003, 0.012)  | 0.0005 | 0.037 (0.002, 0.015) |
| TC              | –0.019 (–0.014, –0.0004) | 0.038 | –0.007 (–0.012, –0.008) |
| TG              | 0.014 (–0.001, 0.008)  | 0.146 | –0.011 (–0.009, –0.004) |
| eGFR            | –0.029 (–0.001, –0.0002) | 0.005 | –0.050 (–0.002, –0.0005) |
| Hemoglobin      | –0.059 (–0.002, –0.0008) | 0.000 | –0.039 (–0.0001, 0.0002) |
| LVMI            | 0.073 (0.0004, 0.0007) | 0.000 | 0.075 (0.0003, 0.0006) |
| LVEF            | –0.038 (–0.006, –0.002) | 0.000 | –0.013 (–0.004, 0.001) |
| E/A             | –0.007 (–0.009, 0.003) | 0.392 | –0.002 (–0.007, 0.006) |

some extent [24]: Although the LA volume was similar between males and females, the LA volume indexed by BSA was significantly higher in females than in males. Our study results indicated that advancing age and female sex were significantly associated with LAE, which has been reported in previous studies [9, 25]. In the case of advancing age, there are a few possible causes: valvular degeneration, cardiac systolic or diastolic dysfunction, hypertension among other factors. Compared to men, the higher prevalence of LAE in women may be ascribed to several factors, including higher systolic (143.4 vs. 139.9 mmHg) and mean arterial pressure (103.5 vs.
100.3 mmHg), and higher LVMI (91.85 vs. 81.77 g/m²), among other factors.

In our research, high SBP was significantly associated with LAE in different adjusted models. After adjusting for sex, DBP was not found to have an obvious correlation with LAE, which was consistent with many other studies [23]. Stritzke et al. found that LA pressure load in hypertensive individuals resulted in an increase in LA size [23]. Additionally, the association between BMI and LAE was shown to be the most obvious in our study, which was consistent with many other studies [9, 23]. Especially, Stritzke et al. reported that obesity appears to be the most important risk factor for LAE in the general population [23]. We also found an important result: diabetes and lower eGFR were positively associated with LAE in the general population, which was also consistent with previous studies [27, 28]. However, in our study, the association between diabetes, eGFR and LAE was only observed in men but not in women, after adjusting for sex.

Actually, previous studies confirmed that higher LVMI was associated with left atrial size [26, 27]. Our study further validates that affirmation. Appleton et al. showed that increased LVMI was related to increased LV stiffness and increase in LV filling pressure, which leads to LA enlargement [25]. Furthermore, our finding that heart rate was inversely related to LAE was consistent with previous studies [26, 28].

Limitations
The present study had several limitations. First, our study was a cross-sectional study, which restricted the interpretation of the observed associations in terms of causality. Second, recent studies have shown that LA volume may be more accurate for the definition of LAE than LA diameter; therefore, the lack of LA volume data is a limitation of this study. However, the simple linear measurement is more common and convenient in daily clinical practice. Additionally, we consider that LA diameter is still helpful in identifying high-risk individuals.

Conclusion
Our population-based study indicated that the latest prevalence of LAE in rural areas of China is not as high as that in European and American countries. The prevalence was similar between males and females, overall the women was slightly higher. After indexed by BSA, the prevalence of LAE became higher, and it was significantly higher in females than in males. Female sex, advancing age, high SBP, higher BMI, diabetes, higher LVMI, lower eGFR, lower LVEF, and lower HR were found to be risk factors of LAE. Further prospective studies are required to verify these findings.

Ethics approval and consent to participate
The study was approved by the Ethics Committee of China Medical University (Shenyang, China). All procedures were performed in accordance with ethical standards. Written consent was obtained from all participants after they had been informed of the objectives, benefits, medical items and confidentiality agreement regarding their personal information. For participants who were illiterate, we obtained written informed consent from their proxies.

Consent for publication
Consent.

Availability of data and materials
All of data and materials are availability.

Competing interests
The authors declare that they have no competing interests.

Authors' contributions
QO collected the data, and analyzed and prepared the first draft of the manuscript; XG was involved in revising the manuscript critically for important intellectual content; SY and YC participated in analysis and interpretations of the data; HZ collected the data; YS conceived the study design, reviewed the manuscript and serves as guarantor for the contents of this paper. All authors read and approved the final manuscript.

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References
1. Levy D, Garrison RJ, Castelli WP, Savage DD, Kannel WB. Factor analysis of echocardiographic measurements. The Framingham Heart Study. J Chronic Dis. 1986;39:317-324.
2. Sundström J, Lind L, Arnlöv J, Zethelius B, Andrén B, Lithell HO. Echocardiographic and electrocardiographic diagnoses of left ventricular hypertrophy predict mortality independently of each other in a population of elderly men. Circulation. 2001;103:2946–51.
3. Li X, Li R, Yu W, Shi H, Wei L. Characteristics of coronary microvascular lesions in autopsy elderly with hypertensive left ventricular hypertrophy. Chin Med J. 2002;115:658–63.
4. Kizer JR, Bella JM, Palmieri V, Liu JE, Best LG, Lee ET, Roman MJ, Devereux RB. Left atrial diameter as an independent predictor of first clinical cardiovascular events in middle-aged and elderly adults: The Strong Heart Study (SHS). Am Heart J. 2006;151:412–8.
5. Tang RB, Yan XL, Dong JZ, Kalifa J, Long DY, Yu RH, Bai R, Kang JP, Wu JH, Sang CH, Ning M, Jiang CX, Salim M, Ma CS. Predictors of recurrence after a repeat ablation procedure for paroxysmal atrial fibrillation: role of left atrial enlargement. Europace. 2014;16:1569–74.
6. El Aouar LM, Meyerfreud D, Magalhães P, Rodrigues SL, Baldo MP, Brasil Y, El Aouar SM, El Aouar NA, Mill JG, Campos Filho O. Relationship between left atrial volume and diastolic dysfunction in 500 Brazilian patients. Arq Bras Cardiol. 2013;101:52–8.
7. Pierdomenico SD, Pierdomenico AM, Di Carlo S, Di Tommaso R, Cucurullo F. Left atrial enlargement and risk of ischemic stroke in elderly treated hypertensive patients. Am J Hypertens. 2014;27:1179–84.
8. Shigematsu Y, Nitohitsu S, Ogimoto A, Ohtsuka T, Okayama H, Higaki J. The influence of insulin resistance and obesity on left atrial size in Japanese hypertensive patients. Hypertens Res. 2009;32:500–4.

9. Su G, Cao H, Xu S, Lu Y, Shuai X, Sun Y, Liao Y, Li J. Left atrial enlargement in the early stage of hypertensive heart disease: a common but ignored condition. J Clin Hypertens. (Greenwich). 2014;16:192–7.

10. O’Brien E, Petrie J, Litter W, de Swiet M, Padfield PL, O’Malley K, Jamieson M, Altman D, Bland M, Atkins N. The British Hypertension Society protocol for the evaluation of automated and semi-automated blood pressure measuring devices with special reference to ambulatory systems. J Hypertens. 1990;8:607–19.

11. Devereux RB, Roman MJ. Evaluation of cardiac and vascular structure by echocardiography and other noninvasive techniques. In: Laragh JH, Brenner BM, editors. Hypertension: pathophysiology, diagnosis, treatment. 2nd ed. New York: Raven; 1995. p. 1969–85.

12. Sahn DJ, De Maria A, Kisslo J, Weyman AE. The Committee on M-mode Standardization of the American Society of Echocardiography: recommendations regarding quantitation in M-mode echocardiography: results of a survey of echocardiographic measurements. Circulation. 1978;58:1072–83.

13. Schiller NB, Shah PM, Crawford M, DeMaria A, Devereux R, Feigenbaum H, Gutgesel H, Reichek N, Sahn D, Sahn JJ. The American Society of Echocardiography Committee on Standards, Subcommittee on Quantitation of Two-Dimensional Echocardiograms: recommendations for quantitation of the left ventricle by two-dimensional echocardiography. J Am Soc Echocardiogr. 1989;2:358–67.

14. Roman MJ, Devereux RB, Kramer-Fox R, O’Loughlin J. Two-dimensional echocardiographic aortic root dimensions in children and adults: biologic determinants and normal limits. Am J Cardiol. 1986;54:507–12.

15. Devereux RB, Alonso DR, Lutas EM, Gottlieb GJ, Campo E, Sachs I, Reichek N. Echocardiographic assessment of left ventricular hypertrophy: comparison to necropsy findings. Am J Cardiol. 1986;57:450–8.

16. Lang RM, Bierig M, Devereux RB, Flachskampf FA, Foster E, Pellikka PA, Picard MH, Roman MJ, Seward J, Shanewise JS, Solomon SD, Spencer KT, Sutton MS, Stewart WJ. Recommendations for Chamber Quantification: a report from the American Society of Echocardiography’s Guidelines and Standards Committee and the Chamber Quantification Writing Group, developed in conjunction with the European Association of Echocardiography, a branch of the European Society of Cardiology. J Am Soc Echocardiogr. 2005;18:140–63.

17. Chobanian AV, Bakris GL, Black HR, Cushman WC, Green LA, Izzo Jr JL, Jones DW, Materson BJ, Oparil S, Wright JT Jr, Roccella EJ. The seventh report of the Joint National Committee on Prevention, Detection, Evaluation, and Treatment of High Blood Pressure: the JNC 7 report. JAMA. 2003;289:2560–72.

18. Hypertension control. Report of a WHO Expert Committee. World Health Organ Tech Rep Set 1996; 862:1–83.

19. Expert Panel on Detection, Evaluation, and Treatment of High Blood Cholesterol in Adults. Executive summary of the third report of The National Cholesterol Education Program (NCEP) Expert Panel on Detection, Evaluation, and Treatment of High Blood Cholesterol in Adults (Adult Treatment Panel III). JAMA. 2001;285:2486–97.

20. Wang CN, Liang Z, Wei P, Liu P, Yu JX, Zhang DM, Ma FLl. Changes in dietary patterns and certain nutrition-related diseases in urban and rural residents of Jiangsu Province, China, during the 1990s. Biomed Environ Sci. 2002;15:271–6.

21. Cuspidi C, Negri F, Lonati L, Muisenas ML, Capra A, Milan A, Sala C, Danzi GB, Longo M, Morganti A. Prevalence and correlates of echocardiographic left atrial enlargement in hypertensive outpatients in clinical practice. Clin Exp Hypertens. 2011;33:328–35.

22. Kopchek G, Sobier B, Podolec M, Walgowa M, Brzazda M, Zarecka J, Loster B, Nessler J, Pajak A, Podolec P. The prevalence of abnormal echocardiographic findings in a sample of urban adult population. Kardiol Pol. 2014;72:42–9.

23. Stritzke J, Markus MR, Duderstadt S, Lieb W, Luchner A, Dittrich A, Keil U, Hense HW, Schunkert H. The aging process of the heart: obesity is the main risk factor for left atrial enlargement during aging. J Am Coll Cardiol. 2009; 54:1602–9.

24. Wang SK, Yue H, Zou YB, Sun K, Fu CY, Wang H, and Hui RT. Prevalence and risk factors for left ventricular hypertrophy and left ventricular geometric abnormality in the patients with hypertension among Han Chinese. Chin Med J. 2012;125(1):21–6.

25. Katayama T, Fujisawa N, Tsuuya Y. Factors contributing to left atrial enlargement in adults with normal left ventricular systolic function. J Cardiol. 2010;55:196–204.