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***Retrospective Study***

**Prognostic value of left atrial size in hypertensive African Americans undergoing stress echocardiography**

KhemkaA *et al*. Prognostic value of LA in AA

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**Abstract**

BACKGROUND

Left atrial (LA) enlargement is a marker of increased risk in the general population undergoing stress echocardiography. African American (AA) patients with hypertension are known to have less atrial remodeling than whites with hypertension. The prognostic impact of LA enlargement in AA with hypertension undergoing stress echocardiography is uncertain.

AIM

To investigate the prognostic value of LA size in hypertensive AA patients undergoing stress echocardiography.

METHODS

This retrospective outcomes study enrolled 583 consecutive hypertensive AA patients who underwent stress echocardiography over a 2.5-year period. Clinical characteristics including cardiovascular risk factors, stress and echocardiographic data were collected from the electronic health record of a large community hospital. Treadmill exercise and Dobutamine protocols were conducted based on standard practices. Patients were followed for all-cause mortality. The optimal cutoff value of antero-posterior LA diameter for mortality was assessed by receiver operating characteristic analysis. Cox regression was used to determine variables associated with outcome.

RESULTS

The mean age was 57 ± 12 years. LA dilatation was present in 9% (54) of patients (LA anteroposterior ≥ 2.4 cm/m2). There were 85 deaths (15%) during 4.5 ± 1.7 years of follow-up. LA diameter indexed for body surface area had an area under the curve of 0.72 ± 0.03 (optimal cut-point of 2.05 cm/m2). Variables independently associated with mortality included age [*P* = 0.004, hazard ratio (HR) 1.34 (1.10-1.64)], tobacco use [*P* = 0.001, HR 2.59 (1.51-4.44)], left ventricular hypertrophy [*P* = 0.001 , HR 2.14 (1.35-3.39)], Dobutamine stress [*P* = 0.003, HR 2.12 (1.29-3.47)], heart failure history [*P* = 0.031, HR 1.76 (1.05-2.94)], LA diameter ≥ 2.05 cm/m2 [*P* = 0.027, HR 1.73 (1.06-2.82)], and an abnormal stress echocardiogram [*P* = 0.033, HR 1.67 (1.04-2.68)]. LA diameter as a continuous variable was also independently associated with mortality but LA size ≥ 2.40 cm/m2 was not.

CONCLUSION

LA enlargement is infrequent in hypertensive AA patients when traditional reference values are used. LA enlargement is independently associated with mortality when a lower than “normal” threshold (≥ 2.05 cm/m2) is used.

**Key Words:** Mortality; Hypertension; African American; Left atrial enlargement; Stress echocardiography

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**Core Tip:** In hypertensive African American patients referred for stress testing, left atrial (LA) enlargement was infrequent when using the established references values for the general population. Indexed LA Antero-posterior diameter has a superior area under the curve compared to LA diameter alone for discrimination of survivors and non-survivors. LA enlargement is an independent predictor of mortality on long-term follow-up when assessed as a continuous variable or when using a lower reference value derived from our population.

**INTRODUCTION**

Left atrial (LA) enlargement is a known predictor of adverse cardiovascular events including atrial fibrillation, stroke and heart failure[1-3]. Hypertension can induce left ventricular (LV) remodeling resulting in increased LV mass and concentric hypertrophy, both of which are associated with LA enlargement[4,5].In the general population of patients undergoing stress echocardiography, LA enlargement [defined by an anteroposterior (AP) dimension ≥ 2.4 cm/m2] has been shown to be predictive of myocardial infarction and death[6,7]. LA enlargement has also been shown to be predictive of an abnormal stress echocardiogram[8]. African Americans (AA) have a high burden of cardiovascular disease as well as risk factors including hypertension and diabetes mellitus. Although morbidity and mortality risk in this population is well established, pharmacotherapy is less commonly utilized and AA have higher mortality relative to other ethnicities[9]. The prognostic value of LA enlargement in AA patients undergoing stress echocardiography is less clear. Additionally, it is unclear if reference values for LA enlargement established in white populations should be applied in AA. In African American patients, LA remodeling appears reduced relative to whites even when controlling for risk factors such as obesity, age, increased LV mass, and hypertension[10-12].The lower incidence of atrial fibrillation in AA may be attributed to their smaller LA size and may be related more to inter-racial differences in anteroposterior diameter rather than volume[10,13,14]. Despite less LA remodeling, AA are at increased for cardiovascular events and mortality compared to white patients[15]. However, LA size may also have prognostic value in this racial group[7,16]. The purpose of this study was to assess the prognostic value of LA size in hypertensive AA patients undergoing stress echocardiography and to determine a threshold value of LA enlargement associated with mortality.

**MATERIALS AND METHODS**

The Indiana University Institutional Review Board approved this study. The study population comprised of 583 consecutive AA patients with a history of hypertension referred for stress echocardiography at an urban community hospital in Indianapolis over a 2.5-year period.

***Clinical characteristics***

Clinical characteristics were extracted from the electronic health record. Patients were considered to have a smoking history if they were currently using tobacco or were a former smoker. Hypercholesterolemia was defined as total cholesterol greater than 200 mg/dL or if the patient was receiving lipid-lowering therapy. Obesity was defined as a body mass index ≥ 30 kg/m2. Patients were considered to have a history of coronary artery disease if they previously suffered a myocardial infarction, underwent a revascularization procedure, or had at least 50% diameter stenosis in one or more major epicardial coronary arteries by angiography. A history of heart failure was noted if there was a previous hospitalization for heart failure or a clinical diagnosis made in an outpatient setting with ongoing medical treatment for heart failure.

***Two-dimensional echocardiographic measurements and stress echocardiography***

LA diameter was measured as the maximum end-systolic anterior-posterior diameter in the parasternal long- or short-axis views. LA enlargement was defined as a dimension ≥ 2.4 cm/m2 when indexed to body surface area (BSA) based on studies in the general population[7,17]. LA volume index was not routinely assessed because only a minority of subjects had apical views visualizing the entire LA. In a small subset of patients with apical views that included the entire left atrium, LA volume index was also measured using the biplane Simpson’s method. LV diameters and wall thickness were obtained in the parasternal long- or short-axis views at the level of the mitral leaflet tips. LV mass was calculated using linear measurements with the following formula:

LV mass = 0.8 × {1.04 [(LVIDd + PWTd + SWTd)3 – (LVIDd)3]} + 0.6 g where:

LVIDd = maximum internal diameter at end-diastole.

PWTd = end-diastole posterior wall thickness.

SWTd = end-diastole septal wall thickness.

Left ventricular hypertrophy (LVH) was defined as an LV mass indexed to BSA greater than or equal to 96 g/m2 for women and 116 g/m2 for men[18,19]. Relative wall thickness (RWT) was calculated by the formula (2 × PWTd/LVIDd). LVH was further differentiated into concentric and eccentric hypertrophy if RWT was > 0.42 or ≤ 0.42, respectively. Concentric remodeling was defined as a normal LV mass with RWT > 0.42. Ejection fraction was calculated with either the area length method or with the modified Simpson’s method for patients with regional wall motion abnormalities.

Treadmill exercise was performed with protocols chosen based on the patient’s age and expected exercise ability. Standard end-points were used[20]. The Dobutamine protocol was conducted with a step-wise infusion using previously described methods and endpoints[21]. Images were obtained in the apical four- and two-chamber views and parasternal long- and short-axis views at rest, low-dose (5-10 µg/kg/min), peak dose and recovery in patients undergoing Dobutamine stress. Baseline and immediate post-stress images were obtained in patients undergoing exercise. Experienced echocardiographers blinded to the clinical data and follow-up interpreted the stress echocardiograms. An abnormal stress echocardiogram was defined by the presence of resting or stress-induced wall motion abnormalities in one or more of 16 myocardial segments[22].

***Follow-up***

Follow-up data was obtained retrospectively by review of the electronic health records and the Social Security Death Index database[23].The end-point for the study was all-cause mortality.

***Statistical analysis***

Continuous variables were reported as mean ± SD. Patient groups were compared using the Student *t*-test for continuous variables and Chi-square test for categorical variables. A two sided *P*-value < 0.05 was considered significant. Receiver-operating characteristic (ROC) curve analysis was used to determine the best cut-point of LA diameter for predicting mortality. The area under the curve (AUC) was calculated for both LA diameter and LA diameter indexed to BSA. The difference between the two AUC values was compared using the correlated area test statistic. Kaplan-Meier analysis of survival was performed using the best cut-point from ROC analysis. Cox proportional hazards model was used to assess predictors of mortality. Variables with *P* value < 0.05 were included in a multivariate analysis employing a forward conditional method. LA diameter was tested on multivariate analysis both as a continuous variable and as a categorical variable using the cut-point of 2.4 cm/m2 previously established in the general population and the best cut-point determined from ROC analysis in our study population. The relationship between LA diameter index and LA volume index was assessed by linear regression.

Statistical analysis was performed using SPSS version 18 (SPS, Chicago, IL, United States) and the software package ROCKIT[24].

**RESULTS**

***Patient characteristics***

Table 1 shows the clinical and stress echocardiographic characteristics of the patient population. Of the 583 patients, 32% were male and the mean age was 57 ± 12 years. A history of heart failure or coronary atherosclerosis was present in 11% and 19%, respectively. Ninety percent of the patients were on anti-hypertensive therapy and the mean resting systolic blood pressure was 140 ± 17 mmHg. An abnormal stress echocardiogram was noted in 17% of patients. Eleven percent had an ejection fraction less than 50%. LVH was present in 25% and concentric remodeling was present in 52%. Only 9% of the study population had an elevated LA diameter index, using the cut-point of 2.4 cm/m2 as defined in the general population.

***LA size and mortality***

During follow-up of 4.5 ± 1.7 years (max 6.9 years), 85 patients (15%) died. ROC analysis showed that LA diameter referenced to body surface area had a larger AUC compared to LA diameter alone (AUC of 0.72 ± 0.03 *vs* 0.66 ± 0.03, *P* = 0.002) for distinguishing survivors and those who died. Figure 1 demonstrates a plot of sensitivity and specificity of LA diameter for death during follow-up at 0.05 cm/m2 intervals. LA size above the reference value (2.4 cm/m2) had sensitivity and specificity for mortality during follow-up of 24% and 93%, respectively. The best cut-point for predicting death during follow-up (maximum of sensitivity and specificity) was 2.05 cm/m2, which produced a sensitivity and specificity of 61% and 72%, respectively.

***LA size and survival***

Figure 2 shows a Kaplan-Meier analysis of cumulative survival using the best cut-point of 2.05 cm/m2. Overall, survival was 92% if LA diameter index was ≤ 2.05 cm/m2 and 72% if LA diameter index was > 2.05 cm/m2 (*P*-value < 0.001).

***Predictors of mortality***

Table 2 shows univariate predictors for all-cause mortality. There were six independent predictors of mortality by multivariate analysis using the reference cut-point for LA enlargement (2.4 cm/m2, see Table 3). These included age, smoking history, heart failure, the need for Dobutamine stress, an abnormal stress echocardiogram, and LVH (Chi-square 102). LA enlargement was not a predictor. In a second multivariate analysis using the ROC defined cut-point of 2.05 cm/m2 for LA size, LA enlargement was found to be an additional independent predictor (Chi-square 107). A third multivariate analysis considering LA diameter index as a continuous variable rather than a categorical variable also found LA diameter index to be independently predictive in addition to the other six predictors (Chi-square 109).

***Comparison of LA anteroposterior diameter and la volume index***

In the 57 patients (10%) in whom LA volume index could be assessed, the *R*-value for the correlation of LA diameter index and LA volume index was 0.76. Fourteen subjects (25%) were identified as having LA enlargement by volume index based on a cut-point of 34 mL/m2 established in the general population[6].

**DISCUSSION**

Our study had three main findings. LA enlargement in the AP dimension was infrequent in AA with hypertension using reference values established in the general population. LA diameter indexed for BSA had a superior AUC to LA diameter alone. LA size was an independent predictor of mortality on long-term follow-up when assessed as a continuous variable or using the cut-point of 2.05 cm/m2 for enlargement derived from our population but not when using the cut-point of 2.4 cm/m2 derived from the general population.

***LA remodeling in AA***

In this study, only 9% of hypertensive AA were found to have LA enlargement. This is a lower than expected frequency of LA enlargement when compared to the general population. Among a broad sample of the Framingham study used to validate reference values of LA diameter, 22% of men and 29% of women had LA diameters that exceeded reference limits[2]. Compared to the Framingham study cohort, our population had a higher prevalence of hypertension (100% *vs* 33%), heart failure (11% *vs* 1%), and older age (mean age 57.4 years *vs* 50.8 years), which are all variables associated with LA enlargement[3,25,26]. Multiple studies have shown a higher prevalence of LA enlargement in white patients with hypertension (weighted average 37.3%)[27-30].

AA have a higher burden of hypertension and cardiovascular mortality with lower rates of pharmacologic interventions[9]. However, when traditional reference values for LA size are used, AA patient mortality risk may be underappreciated. Several studies have shown reduced LA remodeling in AA patients. In a cohort of men with hypertension (58% AA), investigators found that as age increased white patients had a greater mean LA diameter than AA patients[12]. Similarly, in a cohort of 3882 elderly subjects, AA men had significantly smaller mean LA diameter (1.9 mm LA dimension)[11]. Additionally, in a study evaluating the effect of race on the prevalence of atrial fibrillation, AA subjects were demonstrated to have significantly smaller LA diameters (2 mm smaller AP LA dimension)[10].

A more recent evaluation of 129 AA compared with 326 whites showed that in the presence of hypertension, the former had significantly smaller LA size despite similar ventricular relative wall-thickness, diastolic function, and 6-min walk test[31]. Why LA remodeling might be reduced in AA remains unclear although there is speculation that genetic and environmental factors influence the structure of the hearts of AA patients compared to hearts of white patients. Badertscher *et al*[31] found that AA have lower levels of collagen 1 telopeptide and higher levels of collagen 1 propeptide suggesting that different collagen homeostasis may contribute to atrial remodeling. While AA have a similar average LV mass index as whites, they have significantly smaller LV cavities and thicker LV walls, with a high percentage demonstrating the “concentric remodeling” pattern of cardiac structure[33-35]. This pattern was seen in a majority of our population with 52% displaying concentric remodeling.Similar genetic and environmental factors that produce differences in LV remodeling may also contribute to race related differences in LA remodeling. Gottdiener *et al*[12] proposed the possibility that in parallel with an increased LV wall thickness, there might also be a similar increase in LA wall thickness, which might reduce wall compliance and the resultant cavity size of the LA.

An additional possibility is that reduction of anterior-posterior LA dimension in AA is due to differences in chest and mediastinal structures rather than a consequence of true differences in LA remodeling. The LA is a relatively low-pressure chamber and its size and configuration is influenced by its surrounding structures. Manolio *et al*[11] reported that racial differences in LA dimensions were partially mitigated when accounting for chest dimensions and spirometric lung volumes. Given the close proximity of the ascending aorta to the LA, enlargement of the aortic root might limit the ability of the LA to expand in the antero-posterior direction. AA patients are known to have a higher than expected prevalence of aortic regurgitation, which was independently predicted by aortic root size[35]. In the small cohort of patients in our study who had LA volume measurements, the proportion (25%) that had enlargement remained lower than expected.

***Prediction of mortality by LA dimension***

From ROC analysis, the optimal cut-point for an abnormal LA diameter that predicts mortality in AA was well within the normal reference range. In contrast, the guidelines-defined cut-point had very low sensitivity for predicting mortality in our study population. While LA dilation is infrequent in AA, LA diameter does hold prognostic significance in this population when a lower threshold for abnormal is used.

LA diameter indexed to BSA improved prediction of mortality over LA diameter alone. Indexing of echocardiographic measurements to BSA is currently recommended by the American Society of Echocardiography but it has been argued that correcting for body size inappropriately “forgives” for obesity[36]. Our population included a large proportion of AA females, a population known to have a high prevalence of obesity[37]. Forty-five percent of our population was obese. Therefore, use of indexed LA diameters raises the potential of overcorrection for obesity in our study. However, we found that indexed LA diameter had superior prognostic value over LA diameter alone suggesting that the correction is appropriate in our population. To our knowledge, this is the first study to demonstrate superiority of indexed LA diameter over LA diameter alone.

***Comparative long term prognostic value of la dimension***

Our data found LA diameter index to be an independent predictor of all-cause mortality in addition to heart failure, age, smoking history, LVH, an abnormal stress echocardiogram, and the requirement for Dobutamine stress. Our study demonstrated that LA diameter predicted long-term outcome as survival curves continued to separate at 6 years of follow-up. Similar to our finding, data from the Framingham study found LA diameter to be predictive of death during 8 years of follow-up[2]. Within an AA cohort of the Atherosclerosis Risk in Communities study, those with the highest quintile of LA diameter had a higher risk of mortality during a median follow-up of 9.8 years[38].Several investigations have suggested that LA enlargement serves as a marker of chronic diastolic dysfunction over time and thus accounts for the accumulated risk of elevated cardiac filling pressures for cardiovascular events[39].Our results suggest different reference values are needed for AA patients to accurately evaluate their cardiovascular risk. This may also improve treatment in hypertensive AA patients which may translate to decreased mortality.

***Limitations of this study***

The primary limitation of this study is our use of LA diameter as opposed to LA volume index. LA volume is currently recommended by the American Society of Echocardiography as the most accurate measure of true LA size[6]. Unfortunately, majority of the patients in our study had truncated apical images utilized for stress echocardiography so we were unable to derive information on LA volume except in a minority of patients. In the small subset of patients there was a reasonable correlation between LA diameter and volume index. While LA volume is clearly a more accurate measure of true LA size, LA volume may be only marginally superior at identifying cardiovascular disease[26,40]. For patients undergoing stress echocardiography, LA diameter index has shown to offer adequate prognostic value and is probably acceptable for those with difficult visualization of the complete LA[7].

An additional limitation of our study was the large percentage of female subjects. Sixty-eight percent of our population was female. How this might affect the applicability of our data for predicting mortality in AA men is unknown, but previous data have suggested that indexing for body size nearly completely accounts for gender differences in LA dimensions[41].

**CONCLUSION**

LA enlargement is infrequent in AA with hypertension referred for stress testing when using the established references values for the general population. Indexed LA AP diameter has a superior AUC to LA diameter alone for discrimination of survivors and non-survivors. LA enlargement is an independent predictor of mortality on long-term follow-up when assessed as a continuous variable or when using a cut-point derived from our population.

**ARTICLE HIGHLIGHTS**

***Research background***

African Americans (AA) have higher cardiovascular (CV) risk factors including hypertension and mortality compared to other races. Left atrial (LA) size has shown prognostic value in white patients.

***Research motivation***

Prior research has suggested AA have smaller LA volumes and standard references values may not apply.

***Research objectives***

We investigated the prognostic value of LA size in hypertensive AA patients undergoing stress echocardiography.

***Research methods***

In this retrospective cohort study, we evaluated 583 consecutive AA patients with a history of hypertension referred for stress testing and evaluated LA diameter in the Antero-posterior window.

***Research results***

LA dilatation was present in 9% (54) of patients [LA anteroposterior (AP) ≥ 2.4 cm/m2]. There were 85 deaths (15%) during 4.5 ± 1.7 years of follow-up. LA diameter indexed for body surface area had an AUC of 0.72 ± 0.03 (optimal cut-point of 2.05 cm/m2). Variables independently associated with mortality included age (*P* = 0.004), tobacco use (*P* = 0.001), left ventricular hypertrophy (*P* = 0.001), need for pharmacologic dobutamine stress (*P* = 0.003), heart failure history (*P* = 0.031), LA diameter ≥ 2.05 cm/m2 (*P* = 0.027), and an abnormal stress echocardiogram (*P* = 0.033). LA diameter as a continuous variable was also independently associated with mortality but LA size ≥ 2.40 cm/m2 was not.

***Research conclusions***

LA enlargement is infrequent in AA with hypertension referred for stress testing when using the established references values for the general population. Indexed LA AP diameter has a superior prognostic value to LA diameter alone for discrimination of survivors and non-survivors. LA enlargement is an independent predictor of mortality on long-term follow-up when assessed as a continuous variable or when using a cut-point derived from our population.

***Research perspectives***

References values for LA size in AA patients may need to be adjusted to more accurately reflect CV risk and which may translate to more aggressive pharmacologic management.

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**Footnotes**

**Institutional review board statement:** In accordance with 45 CFR 46.101(b) and/or IU HRPP Policy, the above-referenced protocol is granted exemption. Exemption of this submission is based on your agreement to abide by the policies and procedures of the Indiana University Human Research Protection Program (HRPP) and does not replace any other approvals that may be required.

**Informed consent statement:** This was a retrospective study that the IRB deemed as exempt and so we did not need informed consent forms signed by patients.

**Conflict-of-interest statement:** The authors have nothing to disclose.

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**Figure Legends**



**Figure 1 Comparison of sensitivity and specificity of left atrial diameter index cut-points.** Sensitivity, specificity, and the summation of sensitivity and specificity are plotted for left atrial diameter index at 0.05 cm/m2 intervals. Sensitivity is shown in blue, specificity is shown in red, and the summation of the two is shown in green. The reference upper and lower limits of normal (2.4 cm/m2 and 1.5 cm/m2, respectively) are indicated with dashed lines. The optimal cut-point that maximizes sensitivity and specificity was 2.05 cm/m2 and is indicated with an asterisk (\*) on the graph.



**Figure 2 Kaplan Meier curve.** Cumulative survival estimates are compared between patients with left atrial (LA) diameter index values above (blue line, LA index < 2.05) and below (red line, LA index ≥ 2.05) the optimal cut-point that maximizes sensitivity and specificity for death. LA: Left atrial.

**Table 1 Baseline clinical and echocardiographic characteristics**

|  |  |
| --- | --- |
| **Clinical** | **Echocardiographic** |
| Age (yr) | 57 ± 12 | Ejection fraction (%) | 59 ± 10 |
| Male  | 32% | Reduced EF | 11% |
| Tobacco | 60% | LA Diam (cm) | 3.7 ± 0.6 |
| Family history of CAD | 34% | LA Diam index (cm/m2) | 1.9 ± 0.4 |
| Hyperlipidemia | 50% | Abn LA Diam index | 9% |
| Diabetes mellitus | 38% | LV mass (g) | 172 ± 59 |
| Obesity | 45% | LV mass index (g/m2) | 89 ± 29 |
| CAD | 19% | LV hypertrophy | 25% |
| Heart failure | 11% | Relative wall thickness | 0.51 |
| Atrial fibrillation | 3% | LV remodeling pattern |  |
| CKD (GFR < 60) | 17% | Normal geometry | 22% |
| Systolic BP (mmHg) | 140 ± 17 | Concentric remodeling | 52% |
| Hypertensive therapy | 90% | Concentric hypertrophy | 21% |
| Diuretic | 56% | Eccentric hypertrophy | 4% |
| Calcium channel blocker | 33% | Dobutamine study | 40% |
| ACE-I/ARB | 55% | Abnormal stress echo | 17% |
| Beta-blocker  | 46% |  |  |

Data are presented as the mean value + standard deviation or percent baseline prevalence. CAD: Coronary artery disease; CKD: Chronic kidney disease; GFR: Glomerular filtration rate; BP: Blood pressure; ACE-I: Angiotensin converting enzyme inhibitor; ARB: Angiotensin II receptor blocker; EF: Ejection fraction; LA: Left atrial; LV: Left ventricular.

**Table 2 Univariate predictors of all-cause mortality**

|  |  |  |  |  |
| --- | --- | --- | --- | --- |
| **Characteristic** | **Event** | **No event** | **Hazard ratio (95%CI)** | ***P* value** |
| Clinical |  |  |  |  |
| Age  | 62.9 | 56.5 | 1.04 (1.02-1.06) | < 0.001 |
| Male sex | 40% | 32% | 1.43 (0.92-2.20] | 0.109 |
| Tobacco | 79% | 57% | 2.55 (1.51-4.29) | < 0.001 |
| Fam. History of CAD | 28% | 35% | 0.73 (0.45-1.28) | 0.194 |
| Hyperlipidemia | 48% | 51% | 0.91 (0.59-1.40) | 0.662 |
| Diabetes mellitus | 52% | 36% | 1.77 (1.16-2.71) | 0.008 |
| Obesity | 32% | 47% | 0.56 (0.35-0.88) | 0.010 |
| CAD | 33% | 17% | 2.15 (1.37-3.38) | 0.001 |
| Heart failure  | 27% | 9% | 3.44 (2.13-5.56) | < 0.001 |
| Atrial fibrillation | 7% | 2% | 2.70 (1.18-6.19) | 0.019 |
| CKD (GFR < 60) | 29% | 15% | 2.37 (1.48-3.78) | < 0.001 |
| Systolic BP (mmHg) | 141.5 | 140.0 | 1.01 (0.999-1.02) | 0.438 |
| Echocardiographic |  |  |  |
| Reduced EF | 25% | 9% | 2.89 (1.76-4.72) | < 0.001 |
| Abn. LA index (2.40 cut-point) | 24% | 7% | 3.16 (1.91-5.22) | < 0.001 |
| Abn. LA index (2.05 cut-point) | 61% | 28% | 3.35 (2.17-5.18) | < 0.001 |
| LV hypertrophy | 52% | 21% | 3.62 (2.36-5.54) | < 0.001 |
| Relative wall thickness | 0.51 | 0.51 | 0.94 (0.19-4.57) | 0.941 |
| LV diastolic diameter | 4.54 | 4.34 | 1.57 (1.15-2.13) | 0.005 |
| LV systolic diameter | 3.24 | 2.98 | 1.54 (1.20-1.99) | 0.001 |
| Fractional shortening | 0.30 | 0.32 | 0.05 (0.00-0.57) | 0.017 |
| IV septum thickness | 1.20 | 1.11 | 2.51 (1.19-5.33) | 0.016 |
| LV post. wall thickness | 1.13 | 1.07 | 2.48 (1.04-5.92) | 0.040 |
| Dobutamine study | 68% | 35% | 3.55 (2.25-5.60) | < 0.001 |
| Abnormal stress | 35% | 14% | 2.76 (1.77-4.31) | < 0.001 |

CAD: Coronary artery disease; CKD: Chronic kidney disease; GFR: Glomerular filtration rate; BP: Blood pressure; EF: Ejection fraction; Abn. LA index: Abnormal left atrial diameter indexed to body surface area (cm/m2); LV: Left ventricular; IV: Intraventricular; Post.: Posterior; See text for further explanation of variables.

**Table 3 Multivariate predictors of all-cause mortality**

|  |  |  |
| --- | --- | --- |
|  | **Reference cut-point** **for Abn. LA Diam** | **Best cut-point** **for Abn. LA Diam** |
|  | **Chi-square 102** | **Chi-square 107** |
| **Predictor**  | **Wald**  | **HR (95%CI)** | ***P* value** | **Wald**  | **HR (95%CI)** | ***P* value**  |
| Age (per 10 yr) | 11.4 | 1.40 (1.15-1.71) | 0.001 | 8.2 | 1.34 (1.10-1.64) | 0.004 |
| Tobacco  | 11.9 | 2.61 (1.51-4.49) | 0.001 | 11.9 | 2.59 (1.51-4.44) | 0.001 |
| Heart failure | 6.4 | 1.92 (1.16-3.20) | 0.012 | 4.7 | 1.76 (1.05-2.94) | 0.031 |
| LVH  | 17.6 | 2.54 (1.64-3.93) | <0.001 | 10.5 | 2.14 (1.35-3.39) | 0.001 |
| Abnormal stress  | 5.9 | 1.79 (1.12-2.86) | 0.015 | 4.6 | 1.67 (1.04-2.68) | 0.033 |
| Dobutamine study | 8.5 | 2.09 (1.27-3.43) | 0.004 | 8.9 | 2.12 (1.29-3.47) | 0.003 |
| LA index ≥ 2.40  | –  | –  | NS  | 1  | 1 | 1 |
| LA index ≥ 2.05  | 1  | 1 | 1 | 4.9 | 1.73 (1.06-2.82) | 0.027 |

Column (Reference cut-point for Abn. LA Diam) demonstrated a multivariate analysis using the reference cut-point of 2.40 cm/m2 for defining an abnormal left atrium and column (Best cut-point for Abn. LA Diam) demonstrates a multivariate analysis using the best cut-point of 2.05 cm/m2. ABN: Abnormal; DIAM: Diameter, LVH: Left ventricular hypertrophy; LA index: Left atrium diameter index (cm/m2). 1Variable not included in calculation.