The occurrence of left ventricular hypertrophy in normotensive individuals in a community setting in North-East Trinidad

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Objective: The purpose of this study is to determine primarily the occurrence of left ventricular hypertrophy (LVH) in normotensive Trinidadians.

Design and methods: Enrolment into the study required participants to have normal blood pressure (≤140/90) using the JNC 7 (The Seventh Report of the Joint National Committee on Prevention, Detection, Evaluation, and Treatment of High Blood Pressure) classification, free of type 2 diabetes, as well as no existing LVH. Upon entry into the study, participants were first screened for LVH using a standard 12-lead electrocardiogram (ECG), using the Sokolow–Lyon index and the Cornell index. ECHO was used to confirm or refute the diagnosis of LVH.

Results: A total of 209 patients met the criteria for entry into the study. Of these, 63.6% had LVH using Cornell criteria and 68.2% using LVH by Sokolow–Lyon criteria. Subsequently, ECHO confirmed the diagnosis in 2.9% using American Society of Echocardiography criteria and 1.5% using World Health Organization criteria. Thus the estimated prevalence of LVH in normotensive individuals was approximately 3%.

Conclusion: The estimated prevalence of LVH in normotensive individuals appears to be relatively high if an ECG is the single investigation performed, which is common in our setting and may also be common in the developing world. However, using ECHO, the prevalence of LVH approaches a value similarly reported in the literature. Therefore, these findings raise two important issues: 1) the use of criteria such as the Cornell and Sokolow–Lyon voltage criteria established in the developed world from populations of vastly different ethnic backgrounds may not be widely applicable, and 2) all individuals suspected of having LVH should have an ECHO.

Keywords: hypertension, normotensive, echocardiography, Sokolow–Lyon

Introduction

Left ventricular hypertrophy (LVH) occurs when left ventricular mass (LVM) exceeds the normal range, and it is a substantial independent predictor of cardiovascular morbidity and mortality. LVH can be physiological, occurring in athletes or adaptive to systemic hypertension, or aortic valve stenosis. It may occur in infiltrative systemic disease such as amyloidosis or lysosomal storage disease (Aderson–Fabry disease) or as familial hypertrophic cardiomyopathy. In a small proportion of individuals, LVH may occur without hypertension or any other recognized underlying pathology. We refer to this subset of LVH as LVH in normotensive individuals.

LVH is a silent epidemic in North America, affecting more than 16% of the US adult population, and it is noted to be a progressive disorder. It increases morbidity by increasing the risk of arrhythmias, ischemic heart disease, and myocardial infarction.
In addition, LVH is independently associated with an increased risk of stroke, sudden cardiac death, and heart failure. Carluccio et al reported that in 111 patients with acute myocardial infarction, 45% with LVH developed further cardiac complications compared with 6.9% without LVH. In 1995, Liao et al reported that patients with LVH and normal coronaries had a mortality of 16.1%. However, among those with single vessel disease and no LVH, the mortality was 10.9%. LVH more than doubled the risk of a fatal event, and for every 100 deaths, LVH independently accounted for 37. The presence of LVH was associated with a mortality risk of 50% greater than coronary artery disease. Scuteri et al in 2009 noted that increase in LVM is one-third of individuals over the age of 70.

In Trinidad and Tobago, cardiovascular disease is the leading cause of death, with hypertension being a significant contributor. In 2001, hypertension alone, without any other comorbidity accounted for 4.2% of total all-cause mortality.

Apart from hypertension, the Framingham Heart Study in 1990 identified age, body mass index, and both visceral and free fat mass, as factors contributing to LVH. The prevalence of LVH increases with age, affecting more than one-third of individuals over the age of 70. Patrick et al reported in 1995 that 27.4% of men and 9.7% of women in Tobago had LVH. He further showed that mean LVM increased progressively across 10-year age groups between 25 and 64 years in both sexes.

Diabetes and ethnicity have also been found to be associated with LVH. In 1998, Sato et al reported an increase in LVM in normotensive Type 1 diabetic patients. In 2003, Taskiran et al showed that in normotensive type 1 diabetic patients with autonomic neuropathy, LVM index (LVMI) was higher than healthy subjects.

As far back as 1986, Radice et al demonstrated that normotensive adolescents with a family history of hypertension had an LVM significantly greater than normotensive adolescents with normotensive parents. In 1990, Ali and colleagues further demonstrated that 12.2% of normotensive adolescents who had a parent with hypertension, developed LVH as compared with 2% whose parents were normotensive.

The purpose of this study is to determine primarily the prevalence of LVH in normotensive Trinidadians as well as to measure the proportion of various predisposing factors to cardiovascular disease among healthy individuals in the community.

Method

We used a cross-sectional study design. Participants were recruited from four Primary Health Care Facilities in North-East Trinidad. Enrolment into the study required participants to have normal BP (≤140/90) at the time of entry, using the JNC 7 (The Seventh Report of the Joint National Committee on Prevention, Detection, Evaluation, and Treatment of High Blood Pressure) guidelines and free from type 2 diabetes World Health Organization (WHO) criteria. In addition participants were required to have no previous history of a diagnosis of hypertension including hypertension in pregnancy, no current or previous history of being treated for hypertension, and age > 16 years. Participants who satisfied these criteria were further screened to exclude existing hypertension defined as a systolic BP ≥ 140 mmHg and a diastolic BP ≥ 90 mmHg, JNC 7 classification. Participants were also screened for pre-existing LVH due to any underlying pathological conditions such as diabetes, aortic stenosis, or alcoholic cardiomyopathy, by enquire and clinical records. Thus, only normotensive nondiabetic patients with no prior medical evidence of LVH were entered into the study.

Data were collected by asking the participants to complete a self administered pretested structured questionnaire containing 19 items. Clinical data collected include measurement of BP at the time of entry into the study. BP was measured initially in the sitting position using a mercury column sphygmomanometer using a standard technique: on the left arm of each subject using a cuff of appropriate size at the level of the heart. The cuff pressure was inflated 30 mmHg above the level of which the radial pulse disappeared and deflated slowly at a rate of 2 mmHg/second and the reading recorded to the nearest 2 mmHg. The first (appearance) and the fifth (disappearance) Korotkoff sounds were recorded as indicative of the systolic BP and the diastolic BP respectively. In cases where the two readings differed by over 10 mmHg, a third reading was made and the three readings averaged.

In addition, height (m) and weight (kg) were measured from which body mass index (BMI) (kg m⁻²) was calculated. Participants were classified as either overweight (BMI 25.0–29.9 kg m⁻²) or obese (≥30 kg m⁻²) according to WHO criteria.
Participants were then screened for LVH using a standard 12-lead electrocardiogram (ECG). Recordings were made with subjects lying flat in the supine position. The ECG machine was calibrated and tested prior to recording the ECG as outlined by the machine manual. The ECG coding quality assurance methods as outlined by the MONICA study was adopted for use in the study. As such, each ECG was coded by two independent coders, each unaware of the results of the other. Further, the codes established by the two coders separately were compared. However, if only one coder was available, the ECGs were subsequently re-coded after an interval of usually 24–48 hours. In the case of a discrepancy between the two coders, a third senior coder re-coded the ECG without knowing the codes of the two other coders. After having established his/her own code, the three coders discussed the findings and made a final decision.

We used both the Sokolow–Lyon index [S in V1 + R in V5 or V6 (whichever is larger) ≥35 mm] and the Cornell index [R in AVL + S in V3, ≥24 mm for males and >20 mm for females], to establish LVH by ECG. Participants who had elevated S and R waves due to arrhythmias, ventricular ectopic beats, ventricular extrasystole, myocardial ischemia, acute myocardial infarction, left bundle branch block, acute pericarditis, hyperkalemia, pulmonary embolism, and digoxin toxicity were not considered as having pure LVH and were therefore eliminated from the analysis.

M-mode echocardiograms were recorded on strip-chart paper with the subject in a partial left decubitus position. M-mode recordings were made with the ultrasound beam at or just below the tips of the mitral valve leaflets. We used conventional LVM calculations based on linear measurements derived from two-dimensional targeted M-mode 27 echocardiography normalized for body height to the allometric power of 2.7 (LVM = LVM/height²⁷). LVMI was measured by echocardiography normalized for body height to the allometric power of 2.7 (LVM = LVM/height²⁷). LVH was defined as a LVMI >50 g/m² for men and >47 g/m² for women. In addition, the LVM/BSA ratio was calculated as defined by the ASE, which is considered a more accurate way of normalizing LVM to the individual’s BSA, with LVH defined as >90 gm² for women and >114 gm² for men.

Due to constraints of availability and cost, echocardiography was restricted to approximately 64% of subjects who had LVH as defined by both the Sokolow–Lyon’s and Cornell criteria.

Data were analyzed using SPSS version 16. Ethical approval for this study was obtained from The Ethics Committee of The University of The West Indies.

**Results**

A total of 224 patients were recruited for the study. In order to ensure that all patients who entered the study did not have hypertension BP was assessed at the time of entry into the study. Consequently, 15 patients met the criteria for hypertension for the first time (mean systolic BP [MSBP] = 161 ± 17, mean diastolic BP [MDBP] = 97 ± 17 mmHg, and mean arterial pressures [MAP] ranging from 111 to 133.3 mmHg). In other words, while these patients gave no previous history of hypertension or were not receiving antihypertensive medications, BP at the time of entry into the study was elevated. Several factors may have contributed to this finding including white coat hypertension, unrecognized hypertension, or the use of drugs. In fact, four patients admitted to the use of cigarettes, five to the use of alcohol, three cocaine, and one the use of marijuana, a total of 14 of the 15 patients. In addition, seven reported a family history of high BP. Consequently, they were all eliminated from the analysis.

Of the 209 participants who were eligible for analysis, there were more females (124, 59.3%) than males 85 (40.7%), and the mean age of females (36, SD ± 13.8 years) was marginally higher than males (34, SD ± 13.5 years), a detail of the characteristics of the sample are listed in Table 1. The proportion of LVH found based upon the four criteria used, two for ECG and two for echocardiogram are listed in Table 2.
Table 1 Characteristics of study sample

<table>
<thead>
<tr>
<th>Sample characteristic</th>
<th>n (%)</th>
<th>Sample characteristic</th>
<th>n (%)</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>Gender</strong></td>
<td></td>
<td><strong>Smoking</strong></td>
<td></td>
</tr>
<tr>
<td>Male</td>
<td>85 (40.7%)</td>
<td>Current smoker</td>
<td>19 (9.1%)</td>
</tr>
<tr>
<td>Female</td>
<td>124 (59.3%)</td>
<td>Nonsmoker</td>
<td>130 (62.2%)</td>
</tr>
<tr>
<td>Total</td>
<td>209 (100%)</td>
<td>Previous smoker</td>
<td>60 (28.7%)</td>
</tr>
<tr>
<td><strong>Age range</strong></td>
<td></td>
<td><strong>Alcohol consumption</strong></td>
<td></td>
</tr>
<tr>
<td>17–26</td>
<td>33 (15.8%)</td>
<td>Yes</td>
<td>106 (50.7%)</td>
</tr>
<tr>
<td>27–36</td>
<td>15 (7.8%)</td>
<td>No</td>
<td>103 (49.3%)</td>
</tr>
<tr>
<td>37–46</td>
<td>20 (9.5%)</td>
<td>Total</td>
<td>209 (100%)</td>
</tr>
<tr>
<td>47–56</td>
<td>11 (5.2%)</td>
<td></td>
<td></td>
</tr>
<tr>
<td>&gt;56</td>
<td>5 (2.4%)</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Total</td>
<td>85 (40.7%)</td>
<td>&lt;3 days/month</td>
<td>39 (18.7%)</td>
</tr>
<tr>
<td><strong>Ethnicity</strong></td>
<td></td>
<td>3 × per week</td>
<td>38 (18.2%)</td>
</tr>
<tr>
<td>SEA</td>
<td>39 (18.7%)</td>
<td>&gt;3 × per week</td>
<td>40 (19.1%)</td>
</tr>
<tr>
<td>Africans</td>
<td>37 (17.7%)</td>
<td>Total</td>
<td>209 (100%)</td>
</tr>
<tr>
<td>Other</td>
<td>09 (4.3%)</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Total</td>
<td>85 (40.7%)</td>
<td>&lt;18.5</td>
<td>14 (6.7%)</td>
</tr>
<tr>
<td>BMI</td>
<td>124 (59.3%)</td>
<td>18–24.9</td>
<td>82 (39.5%)</td>
</tr>
<tr>
<td></td>
<td></td>
<td>25–29.9</td>
<td>70 (33.3%)</td>
</tr>
<tr>
<td></td>
<td></td>
<td>30–35</td>
<td>38 (18.1%)</td>
</tr>
<tr>
<td></td>
<td></td>
<td>&gt;35</td>
<td>5 (2.4%)</td>
</tr>
<tr>
<td><strong>BMI</strong></td>
<td></td>
<td>Total</td>
<td>209 (100%)</td>
</tr>
</tbody>
</table>

| Abbreviations: SEA, South East Asian; BMI, body mass index. |

Using the Sokolow–Lyon criteria, 27 participants had LVH. However, 5 of the 27 participants had ST segment elevation in addition to LVH and were consequently eliminated from analysis. Hence, only 22 (10.5%) patients had LVH using Sokolow–Lyon criteria (95% confidence interval [CI], 10.1–10.9), with a higher proportion among males (17/85, 20.0%) than females (5/124, 4.0%). The range of Sokolow–Lyon scores were 38.3–52.9 mm, with a mean of 40.3 (SD ± 4.7) mm.

Using Pearson’s correlation method, we attempted to correlate Sokolow–Lyon’s scores with LVM and found a correlation coefficient of 0.32 (P = 0.07), and with LVM/BSA, found a correlation coefficient of 0.37 (P = 0.045).

Similarly, using Cornell Voltage criteria, 11 (5.26%), (95% CI, 4.98–5.54) patients had LVH, again with a higher proportion in males (6/85, 7.0%) than females (5/124, 4.0%). The range for Cornell criteria for men was 29–40 mm with a mean of 32.5 (SD ± 4.03) mm, whereas for women was 21–42 mm with a mean of 26 (SD ± 8.3). Using Pearson’s correlation method, we attempted to correlate Cornell scores with LVM and found a correlation coefficient of 0.53 (P = 0.047), and with LVM/BSA, found a correlation coefficient of 0.62 (P = 0.021).

Of the individuals who tested positive by the Sokolow–Lyon criteria, 15/22 (68.2%) were randomly chosen for confirmatory echocardiograms of whom 4/15 (26.7%) tested positive for LVH using the ASE criteria and 2/15 (13.4%) tested positive for LVH using the WHO criteria.

Extrapolating these results to all 22 subjects with LVH using ASE criteria, the estimated proportion of LVH in normotensive individuals is 2.9% (95% CI, 2.77–3.23). However, using WHO guidelines the estimated proportion of confirmed LVH was 1.5% (95% CI, 1.34–1.66).

Among individuals who tested positive using the Cornell voltage criteria, 7/11 (63.6%) were randomly chosen for confirmatory echocardiograms, of whom 4/7 (57.1%) tested positive for LVH using the ASE criteria and 2/7 (28.6%) tested positive for LVH using the WHO criteria.

Extrapolating these results to all 11 subjects with LVH using ASE criteria, the estimated proportion of LVH in

Table 2 The proportion of LVH found using ECG and echocardiographic criteria

<table>
<thead>
<tr>
<th>Criteria</th>
<th>Male</th>
<th>Female</th>
<th>Total</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>ECG</strong></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Sokolow–Lyon</td>
<td>17 (20.0%)</td>
<td>5 (4.0%)</td>
<td>22 (10.5%)</td>
</tr>
<tr>
<td>Cornell</td>
<td>6 (7.0%)</td>
<td>5 (4.0%)</td>
<td>11 (5.3%)</td>
</tr>
<tr>
<td><strong>Echocardiogram</strong></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>ASE</td>
<td>3 (20.0%)</td>
<td>1 (6.7%)</td>
<td>4 (26.7%)</td>
</tr>
<tr>
<td>WHO</td>
<td>1 (6.7%)</td>
<td>1 (6.7%)</td>
<td>2 (13.3%)</td>
</tr>
</tbody>
</table>

| Abbreviations: ASE, American Society of Echocardiography; ECG, electrocardiogram; LVH, left ventricular hypertrophy; WHO, World Health Organization. |
normotensive individuals is 2.9% (95% CI, 2.77–3.23). However, using WHO guidelines, the estimated prevalence is 1.5% (95% CI, 1.34–1.66).

Discussion
The main finding of this study was an estimated proportion of LVH in normotensive individuals of 2.9%. Patrick and colleagues reported in 1995 a proportion of 27.4% in men and 9.7% in women using electrocardiographic (Sokolow–Lyon) criteria only.15 Our findings appear markedly lower, which may be attributed to more rigorous evaluation of LVH, ie, the use of echocardiograms as well as the use of other criteria apart from the Sokolow–Lyon criteria. In addition, these values were reported among participants with both diabetes and hypertension. A review of the literature revealed two other studies of LVH in patients without hypertension or any pathological conditions associated with LVH: the first was reported by Antoniucci et al, in 1997, who reported a prevalence of 2.7%–3.2% adjusted for height, age, BMI, and BSA.6

The other in 2007, by Ruiz-Nodar et al who showed that in a group of elderly normotensive subjects with a mean age of 72, the prevalence of LVH ranged from 30% by Cornell–Penn criteria to 14% by Devereaux criteria.31

Another important finding was although 60% of patients had increased LVM by echocardiographic evaluation when normalized for BSA according to ASE specifications and height according to WHO specifications, only four patients had LVH. Of these four participants, 20% (3/15) were overweight as opposed to 6.7% (1/15) who had a normal weight. Similarly de Simone et al reported in 1994 that 14% of normotensive individuals who were overweight had LVH, compared with 5% who were normal weight.32 Our findings raise an issue for further investigation in normotensive patients: will reduction in BP below currently recommended targets reduce LVH.

The traditional Sokolow–Lyon voltage and the Minnesota code high QRS amplitude (WHO Monica study) are prone to errors by extra cardiac factors such as body weight, obesity, breast tissue, pulmonary conditions, chest size and shape in relation to cardiac size, QRS axis and ventricular conduction problems. Cornell voltage criterion is less vulnerable to such factors.33 In our setting based on a higher Pearson’s correlation (Sokolow–Lyon r = 0.366, Cornell r = 0.618), we provide further support for this position. Sokolow and Lyon introduced their LVH criterion in 1949.32 Many studies have shown the sensitivity of this criterion is low, and it is perplexing that the Sokolow–Lyon’s LVH criterion continues to be used in the clinical evaluation of patients, especially in the developing world with limited resources for more advanced evaluations. Rodrigues et al in 2008 showed the classic Sokolow–Lyon Rappaport and Cornell Voltage criteria showed a low performance in relation to LVH, as confirmed by ECHO,33 our results support these findings as only 4 out of the 15 participants with LVH by Sokolow–Lyons criteria had confirmed LVH by echocardiography.

Ali and colleagues in 1990 showed that 12.2% of normotensive adolescents who had a parent with hypertension developed LVH.30 Our findings provide further evidence of a possible familial link between parental hypertension and the development of LVH, as 100% of participants with echocardiographic confirmation of LVH gave a family history of high BP.

A major limitation of the study was the use of two-dimensional echocardiography, which is less accurate and has greater test–retest variability than newer methods such as cardiac magnetic resonance imaging or real-time three-dimensional echocardiography, which are unavailable on the island. The number of confirmed participants with LVH by echocardiographic was small hence risk factor analyses could not be conducted. In addition the availability and cost of echocardiograms restricted the number of participants who could be further assessed.

We recommend further studies with larger sample sizes with the capacity for all participants to be evaluated further using echocardiography. Risk factors associated with LVH should also be assessed. LVH in normotensive individuals is an important subset of cardiac patients, as Brown et al34 showed that individuals who were normotensive and had LVH had a greater risk of developing cardiac complications and as such should receive adequate follow up.

In conclusion, we provide evidence of the existence of LVH in normotensive individuals in Trinidad with a proportion of 2.9%. LVH appears to be relatively high if an ECG is the single investigation performed, which is common in our setting and may also be common in the developing world. However using echocardiography, the prevalence of LVH approaches a similar value as reported in the literature. Therefore our study confirms: 1) the use of the Cornell and Sokolow–Lyon voltage criteria is applicable, and 2) all individuals suspected of having LVH should have a confirmatory echocardiogram as part of their management.

Disclosure
The authors report no conflicts of interest in this work.


28. Lang RM, Bierig M, Devereux RB, et al. 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(12):1440–1463.


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