Early Detection of Left Ventricular Subclinical Systolic Dysfunction in Hypertensive Patients: Speckle Tracking at Rest and after Dobutamine Stress Echocardiography Study

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Abstract

Background: Hypertension is one of the major cardiovascular risk factors that may result in heart failure. Two-D transthoracic echocardiography (2-D TTE) recognizes reduction in left ventricular systolic function in late stages of the disease course, so early detection is of paramount importance.

Aim of the work: Our study intended to evaluate feasibility of speckle-tracking echocardiography (STE) to provide additional perceptions for early detection of hypertension induced left ventricular systolic dysfunction.

Patient and Methods: We enrolled 40 hypertensive patients, 21 women (52.5%) their mean age was 54.55±9 years and 20 normotensive age and gender matched control individuals (10 women, their mean age was 53.20±12.06 years). All had normal left ventricular systolic function by 2-D TTE. STE performed at rest and low dose dobutamine. Patients with significant ischemic or valvular heart disease as well as atrial fibrillation, conduction abnormalities and diabetes excluded.

Results: Systolic blood pressure was significantly higher in the hypertensive group patients (138.50±10.27 mmHg vs. 120.50±7.76 mmHg, respectively; P = <0.001). Compared to control group, hypertensive patients experienced significantly impaired global longitudinal STE at rest (-18.08±1.63% versus 20.50±1.52% respectively; P<0.001) and at low-dose dobutamine (-19.11±1.75% versus 22.61±1.88% respectively; P<0.008).

Conclusion: Speckle tracking echocardiography increases the sensitivity in detecting subclinical cardiac involvement in early stages of hypertension compared to conventional 2-D echocardiography.

Keywords: Hypertension; Ventricular Function; Global Longitudinal Strain; Dobutamine Echocardiography.

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Introduction

Hypertension is a prevalent and well recognized cardiovascular risk factor, which may lead to left ventricular (LV) systolic impairment through chronic pressure overload. LV hypertrophy (LVH) is a compensatory process in response to increased wall stress. However, this initially useful adaptive mechanism later becomes ‘a pathological change’ in the myocardium. Whilst LVH has been shown to be a powerful independent predictor for cardiovascular morbidity and mortality it can occur late in the course of the disease.1

Multiple recent studies have shown that LV ejection fraction (LVEF) lacks accuracy and sensitivity in detecting early subclinical impairment. In contrast, several studies using two-dimensional (2D) speckle-tracking echocardiography (STE) have demonstrated that despite normal LVEF, many patients in different clinical settings have longitudinal systolic dysfunction of the left ventricle. Thus, systolic analysis of the left ventricle using global longitudinal systolic strain has been suggested as a new standard assessment for global LV systolic function.2

Dobutamine Stress echocardiography was used in our study to assess the contractile reserve (CR) of hypertensive patients which is thought to represent an early manifestation of LV systolic dysfunction.4
Our study designated to detect the value of assessment of left ventricular function by measuring global longitudinal strain at rest and after dobutamine stress for detection of early subclinical left ventricular systolic dysfunction in patients with systemic hypertension.

Patient and Methods

We enrolled 40 known hypertensive patients and 20 normotensive age and gender matched control individuals admitted at Al-Hussain University Hospital, Cairo, between June 2019 and April 2020. The hypertensive group of patients had been selected after elective invasive assessment of their coronaries for another reason than the research, coronary angiograms proved evidence of no or non-significant coronary artery disease. An informed consent obtained from every patient after explanation of the research objectives and the purpose of this study.

Exclusion criteria: Patients with significant coronary artery disease, defined as >50% stenosis in any major epicardial coronary artery or its major branches on coronary angiography. Previous myocardial infarction or coronary revascularization. Non-sinus rhythm as atrial fibrillation. Left bundle branch block. Coexisting significant left sided valvular heart disease (more than mild in severity). Diabetes mellitus.

The following data were collected for each patient:

- Clinical Data: Patients’ data as gender, age, weight, height, hypercholesterolemia, smoking status, family history of ischemic heart disease and drug history. Clinical data as heart rate, systolic and diastolic blood pressures at rest and at low dose dobutamine. Beta-blockers held 48 hours before the echocardiographic study.

- ECG: Standardized 12 lead ECG recorded in every participant in the study. Systematic interpretation and recording of ECG data accomplished for documentation of rhythm, heart rate, presence or absence of left ventricular hypertrophy, conduction disturbances or manifestation of ischemic heart disease.

- Echocardiography: All subjects were examined at rest in the left lateral decubitus position to obtain adequate images in different standard views with a ultrasound machine using. Echocardiographic data were collected as follow: (A) Resting conventional 2D echocardiography: A comprehensive transthoracic echocardiogram with appropriate 2D, color, and Doppler imaging was performed. End-diastolic and end-systolic dimensions of the LV, end-diastolic thickness of inter-ventricular septum and LV posterior wall, LV Ejection fraction (derived from linear measurements obtained from 2D images), LV wall thickness, LV mass were measured. (B) Dobutamine stress echocardiography: Dobutamine stress echocardiography was performed in all participants according to standard protocol. Intravenous dobutamine was infused in incremental doses starting at five µg/kg/min. The dose then increased to 10µg/kg/min and 20µg/kg/min at three 3.14±0.39 cm respectively; P = 0.386) (Figure 3). min interval, these doses were chosen as they have been previously demonstrated to be safe and effective in detecting contractile reserve, without affecting heart rate, blood pressure, or loading conditions 6. Standard parasternal long- and short-axis views, and apical four-, two-, and three-chamber views of the left ventricle obtained at rest and at the end of each infusion stage 6. Visual wall motion analysis was performed by an experienced investigator using the American Society of Echocardiography’s 17-segment model in blinded fashion. (C) Two-dimensional speckle tracking global longitudinal strain: The speckle tracking strain analyses was performed on grey scale images of the left ventricle. Peak global systolic longitudinal strain was measured from the 18 segment measurements (six segments from each of the apical four-, two-, and three-chamber views) at rest and at low dose dobutamine stress. During strain analysis, the endocardial border was manually traced at end-systole, and the width of the region of interest was manually adjusted to include the entire myocardial wall thickness. Three cardiac cycles were analysed and the measurements were averaged.

Statistical analysis:

Results of the present study were statistically analysed using SPSS 25 (IBM, USA). Data were represented as median (interquartile range) or number and percentage. Numerical data were compared using Mann-Whitney U test while categorical data were compared using Fisher exact test or Chi-square test as appropriate. ROC curve was used to evaluate the performance of different tests differentiate between certain groups. The level of significance was taken at P value < 0.050 is significant, otherwise is non-significant.

Results

The resting mean SBP and DBP was significantly higher in the patients group compared to the control group either at rest (SBP; 138.50±10.27 mmHg vs. 120.50±7.76 mmHg, respectively; P = <0.001, DBP; 84.13±8.54 mmHg vs. 77.75±6.17 mmHg respectively; P = 0.004), or at low dose dobutamine (SBP; 140.50±9.18 mmHg vs. 121.75±10.29 mmHg, respectively; P = <0.001, DBP; 83.63±9.47 mmHg vs. 76.75±5.91 mmHg respectively; P = 0.004). The mean HR didn’t differ significantly between the patients and control groups either at rest (75.60±10.30 BPM vs. 74.90±8.04 BPM respectively; P= 0.477), or at low dose dobutamine (77.30±11.76 BPM vs. 76.00±6.98 BPM, respectively;P= 0.350), (Figure 1, 2).

Both the mean of the inter-ventricular septum (IVS) and LV posterior wall (LVPW) thickness were significantly higher in the patients group compared to the control group (IVS;1.20±0.13 cm vs. 0.90±0.17 cm respectively; P<0.001, LVPW; 1.22±0.10 cm vs. 0.89±0.15 cm respectively; P=0.001), with no difference in the mean of LVIDd and LVIDs between the patients group compared to control group (LVIDd; 4.68±0.54 cm vs. 4.83±0.46 cm, respectively; P=0.297, LVIDs; 3.07±0.50 cm vs. 3.07±0.45 cm respectively).

There was no significant difference in the mean LVEF between the patients group and the control
group either at rest (61.82±5.08 % and 61.43±4.41% respectively; P = 0.771) or at low dose dobutamine (65.46±5.08 % and 64.35±5.68% respectively; P = 0.311), while LV GLS was significantly lower in the patients group compared to the control group at rest (-18.08±1.63 % vs. -20.50±1.52 % respectively; 
P <0.001) and at low dose dobutamine (-19.11±1.75 % vs. -22.61±1.88% respectively; P =0.008) (Tables 1, 2).

Figure (1): Comparison between patients and control according to clinical characteristics at rest.

Figure (2): Comparison between patients and control according to clinical characteristics at low dose dobutamine.
**Figure (3):** Comparison between patients and control according to resting echocardiographic linear dimensions of LV.

Table (1): Comparison between patients and control according to resting echocardiographic characteristics of LV systolic function.

<table>
<thead>
<tr>
<th>LV systolic function</th>
<th>Patients (n=40)</th>
<th>Control (n=20)</th>
<th>t-test</th>
<th>p-value</th>
</tr>
</thead>
<tbody>
<tr>
<td>EF%</td>
<td>Mean±SD</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Range</td>
<td>61.82±5.08</td>
<td>61.43±4.41</td>
<td>0.085</td>
<td>0.771</td>
</tr>
<tr>
<td>GLS %</td>
<td>Mean±SD</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Range</td>
<td>-18.08±1.63</td>
<td>-20.50±1.52</td>
<td>30.921</td>
<td>&lt;0.001**</td>
</tr>
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</table>

P-value>0.05 NS; **p-value <0.001 HS  
T-Independent Sample t-test;

Table (2): Comparison between patients and control according to echocardiographic characteristics of LV systolic function at low dose dobutamine.

<table>
<thead>
<tr>
<th>Low Dose Dob Echo</th>
<th>Patients (n=40)</th>
<th>Control (n=20)</th>
<th>t-test</th>
<th>p-value</th>
</tr>
</thead>
<tbody>
<tr>
<td>EF%</td>
<td>Mean±SD</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Range</td>
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<td>64.35±5.68</td>
<td>1.020</td>
<td>0.311</td>
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<tr>
<td>GLS</td>
<td>Mean±SD</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Range</td>
<td>-19.11±1.75</td>
<td>-22.61±1.88</td>
<td>8.682</td>
<td>0.008*</td>
</tr>
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</table>

P-value>0.05 NS; *p-value <0.05 S  
T-Independent Sample t-test;
Discussion
The current study conducted to assess the value of measuring GLS at rest and at low dose dobutamine to detect subclinical LV systolic dysfunction in patients with systemic hypertension. This prospective observational study conducted from June 2019 to April 2020 and included 60 individuals, 40 patients were hypertensives and 20 normotensives control group. Matle, et al.7, Imbalzano, et al.8 and Hensel et al.9 were concerned with this issue in their studies. Matle et al.7 enrolled 129 patients in their study, 73 were hypertensives and 56 as a healthy control group with an average age 60.0 ± 8.7 and 56.7 ± 10.2 respectively. with no significant differences between the 2 groups according to either the age (P = 0.051) or sex (P =0.294 ). Imbalzano et al.4, included in 102 patients their study, 51 patients with isolated hypertension (33 males, mean age 56.5 ± 14 years) and 51 age and gender-matched healthy subjects (32 males, mean age 52 ± 13 years). According to the presence or absence of LVH, patients were classified as LVH(+) and LVH(−), respectively. Hensel et al9, enrolled 46 hypertensives with an average age 46.6 ± 14.4, and 46 healthy controls with an average age 44 ± 22.5, with no significant differences between the 2 groups according to either the age or sex. We used normotensive patients with comparable cardiovascular risk factors and we excluded patients with diabetes, significant coronary artery disease, non-sinus rhythm, left bundle branch block, coexisting significant left sided valvular heart disease (more than mild in severity), previous myocardial infarction, or coronary revascularization, therefore were able to examine GLS in a more homogenous patient population, where the only major factor adversely affecting LV deformation was hypertension. Matle et al.7, Imbalzano, et al.8, and Hensel et al9 also excluded these patients. Imbalzano, et al.9 excluded also dyslipidemic patients. In our study all patients underwent resting general echocardiographic examination and 2D STE GLS measurement then at low dose dobutamine 2D STE GLS and EF were measured, which is similar to that was done by Matle, et al. 7 Hensel, et al.9, measured 2D STE GLS and EF at three different levels of physical challenge, in the resting stage, after cycling at a level of 50Watts of resistance for two minutes, and at 150 Watts resistance. Imbalzano, et al.7, examined their patients in the resting state only.

The mean HR did not differ significantly between the patients and control groups either at rest (65.3 ± 10.5 BPM vs. 74.90±0.04 BPM respectively; P = 0.477), or at low dose dobutamine (77.3±11.76 BPM vs. 64.2 ± 9.08BP, respectively; P= 0.350) which is in agreement with Matle et al.7 who revealed at rest (75.60±10.50 BPM vs.74.90±0.04 BPM respectively; P= 0.525) and at low dose dobutamine (64.3 ± 11.1BP vs. 63.0 ± 9.0 BPM, respectively; P = 0.477) and also concordant with Imbalzano et al.8 who reported the mean HR didn’t differ significantly between the patients and control groups either hypertensives with LVH (66.6±9.7 BPM vs. 70±13BPM, respectively; P= NS) or hypertensives without LVH (68.8±9.2 BPM vs. 70±13BPM, respectively; P=NS). Hensel et al.9, reported the mean HR didn’t differ significantly between the patients and control groups either at rest (69.3 ±11.8BPM vs. 72.4 ±15.1 BPM, respectively; P= NS) or at 50 watt stress (110.7 ±11.6 BPM vs. 103 ±10.2 BPM, respectively; P= NS) but at 150 watt stress mean HR was significantly higher in the control group compared to the hypertensive group (140.2 ±18.1BPM vs. 124.5 ±15.8 BPM, respectively; P= 0.00).

Our study showed that both the mean of the inter-ventricular septum (IVS) and the left ventricular posterior wall (LVPW) thickness were significantly higher in patients compared to the control group (IVS;1.20±0.13 cm vs. 0.90±0.17 cm respectively<0.001, LVPW; 1.22±0.10 cm vs. 0.89±0.15 cm respectively; P<0.001) which is consistent with Matle et al.7 who revealed (IVS;1.1±0.2 vs. 1.0±0.2 and its p-value was 0.003, LVPW; 1.1 ± 1 vs. 1.0 ± 2, respectively and its p- value was 0.001) and consistent with Imbalzano et al.8 showed that both the mean of the inter-ventricular septum and LV posterior wall thickness were significantly higher in the hypertensive group with LVH compared to the control group (IVS;14±2.9 mm vs. 10.2±1mm respectively; P<0.001, LVPW; 12±1.1 mm vs. 7.4±0.8mm respectively; P<0.001) but no significant difference between the control group and the hypertensive group without LVH. Hensel et al.9, reported that there was significant difference between the hypertensive group and the control group as regard to IVS (IVS; 1 ±0.2 cm vs. 0.9 ±0.2 cm respectively<0.016) but no significant difference as regard to LVPW (1.1 ±0.2cm vs. 1 ±0.2cm respectively; P=NS).

Our study demonstrated that there was with no significant difference in the mean of LVIDd and LVIDs between the patients group compared to the control group (LVIDd; 4.68±0.54 cm vs. 4.83±0.46 cm, respectively; P = 0.297, LVIDs; 3.07±0.50 cm vs. 3.14±0.39 cm respectively; P = 0.386) which is in line with Matle et al.7. (LVIDd; 4.6±0.5 cm vs. 4.6±0.4 cm, respectively; P = 0.9, LVIDs; 2.6±0.4 cm vs. 2.9±0.6 cm respectively; P = 0.119). Hensel et al.9, reported that LVIDd significantly higher in the patients group compared to the control group (LVIDd; 5.4±0.5 cm vs 4.3±0.5 cm, respectively; P = 0.011) while LVIDs did not differ significantly (LVIDs; 3.1 ±0.5cm vs 3 ±0.5 cm respectively; P=NS).

Our study showed statistically significant mean of patients group compared to control group according to LV mass (213.5±47.88 g vs. 150.87±38.11g respectively<0.001) and LVMi (118.96±26.77 g/m2 vs. 81.43±21.70 g/m2 respectively; P <0.001) which is consistent with Hensel et al.9, who reported significant increase mean of patients group compared to control group according to LV mass, (188.4 ±59.4 g vs. 148.8 ± 44.2 g respectively; P<0.001) LVMi, (96.8 ±30.6 g/m2 vs. 81.1 ±21.3 g/m2 respectively; P <0.02), also Imbalzano et al.8 showed significantly higher mean LVMi of patients with LVH group compared to either control group (121.8 ±16 g vs. 63.6 ± 21 g respectively; P= 0.001) or patients without LVH (120.8 ±16 g vs. 81 ± 14.5 g respectively =0.001) also LVMi significantly higher in patients group without LVH compared to the control group (81 ± 14.5g vs. 63.6 ± 21 g respectively; P=0.01).
Our study showed that no significant difference in the mean LVEF between the patients group and the control group either at rest (61.82±5.08 % and 61.43±4.41 % respectively; P = 0.771) or at low dose dobutamine (65.46±5.08 % and 64.35±5.68% respectively; P = 0.311) which is concordant to that reported by Matle et al., who reported at rest (64.5 ± 6.0% and 64.5 ± 6.6% respectively; P = 0.986) and at low dose dobutamine (72.1 ± 6.5% and 71.2 ± 7.0% respectively; P = 0.424). Imbalzano et al. 8, reported no significant difference in the mean resting LVEF between the control group and the hypertensive group either hypertensives with LVH (63±5.9% and 59±8 % respectively; P =NS), or hypertensives without LVH (63±5.9% and 60±7% respectively; P = NS).

Our study revealed LV GLS was significantly lower in the patients group compared to the control group either at rest (-18.08±1.63 % vs. -20.50±1.52%), or at low dose dobutamine (-19.11±1.75 % vs. -22.61±1.88% respectively; P =0.008), which is concordant with Matle et al. 7 who revealed at rest (-17.1± 1.8% vs. -19.4± 1.5% respectively; P < 0.001) and at low dose dobutamine (-18.1± 2.3% vs. -22.6± 2.4%, respectively and its p-value was < 0.001) and also concordant with Imbalzano et al. 8, who revealed resting LV GLS significantly lower in hypertensive patients (65±5.9% and 64±6% respectively; P <0.01). Hensel et al.9, revealed LV GLS was significantly lower in the patients group compared to normotensive group, either hypertensives without LVH (-18±1.9% vs. – 20.4±2.5%, respectively; P = 0.02) or hypertensives with LVH (–15.9±3.3% vs. –20.4±2.5percentage. Respectively; P <0.001), also LV GLS significantly lower in hypertensive patients with LVH compared to hypertensive patients without LVH (–15.9±3.1percentage vs. -18±1.9%, respectively; P = 0.01). Hensel et al.9, revealed LV GLS was significantly lower in the patients group compared to the control group either at rest (-17.8±2.8 % vs. -20.5±2.3 % respectively; P <0.001) or at 50 watt stress (-18.63±2.4 % vs. -20.93±2.5 % respectively; P <0.001) or at 150 watt stress (-19.1±2.4 % vs. -22.13±3.1 % respectively; P <0.001).

Conclusion

Hypertensive patients have impaired left ventricular global longitudinal strain at rest and at low dose dobutamine despite normal resting left ventricular systolic function assessed by conventional 2D-Echocardiography.

Speckle tracking echocardiography increases the sensitivity in detecting subclinical cardiac involvement in early stages of hypertension compared to conventional 2-D echocardiography.

Global longitudinal strain measurement using speckle-tracking echocardiography is advisable to detect subclinical left ventricular systolic dysfunction in hypertensive patients, however, large-scale study is highly recommended to indorse these consequences.

References