Correlation of left atrial strain with left ventricular end-diastolic pressure in patients with normal left ventricular ejection fraction

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Abstract
Left ventricular diastolic dysfunction (LVDD) remains challenging to be assessed by echocardiography. We sought to explore the relationship between left atrial strain and left ventricular (LV) diastolic function in patients with normal left ventricular ejection fraction (LVEF) by invasive left-heart catheterization. 55 consecutive individuals with LVEF > 50% underwent LV catheterization. Standard transthoracic echocardiography was performed during 12 h before or after the procedure. Left atrial (LA) strain were obtained by speckle tracking echocardiography. When LVEF ≥ 50%, the group with elevated left ventricular end-diastolic pressure (LVEDP) (n = 35) showed decreased left atrial reservoir strain (LASr) (35.2 ± 7.7% vs 21.3 ± 7.2%, p < 0.001), left atrial conduit strain (LASct) (17.6 ± 6.3% vs 11.9 ± 4.1%, p < 0.001), left atrial contraction strain (LAScd) (16.6 ± 7.2% vs 9.5 ± 5.0%, p < 0.001) and increased E/e’ ratio (8.9 ± 2.6 vs 10.1 ± 3.5, p = 0.17). LVEDP negatively correlated with LASr (R = 0.662, p < 0.001), LASct (R = 0.575, p < 0.001) and LAScd (R = 0.456, p < 0.001), but not with E/e’. LASr, LASct and LAScd were all independent predictors of elevated LVEDP (p < 0.05), with a higher C-statistic for the model including LASr (0.95, 0.86 and 0.93 respectively). The area under the curve (AUC) for LASr is 0.914 (cutoff value is 26.7%, sensitivity is 90%, specificity is 82.9%). In patients with normal LV ejection fraction, left atrial strain presented good correlation with LVEDP, and LASr was superior to LASct and LAScd to predict LVEDP. LA strain demonstrated better agreement with the invasive reference than E/e’.

Keywords Left ventricular diastolic dysfunction · Left atrial strain · HFpEF

Introduction
Left ventricular diastolic dysfunction is an independent predictor of all-cause mortality in the general population, even in the preclinical stage [1], and evidence of LVDD is required for the diagnosis of heart failure with preserved ejection fraction (HFpEF) [2, 3]. So, it is increasingly important to evaluate LVDD accurately in routine clinical practice. Elevated left ventricular (LV) filling pressures are the main physiologic consequence of LV diastolic dysfunction [4].

Although invasive methods are considered the “gold standard” for evaluating left ventricular filling pressures and LV diastolic function, echocardiography is routinely used as a noninvasive alternative.

The 2016 ASE/SCAI guidelines streamline the use of four variables into a single algorithm to assess LV diastolic function, while accuracy will be affected in the presence of pulmonary arterial hypertension, severe tricuspid valve lesions, and low right atrial and right ventricular filling pressure etc. [4]. An accurate assessment of left ventricular diastolic function by transthoracic echocardiography is still needed. Speckle tracking is applied to directly reflect the intrinsic deformation of left atrium, which has a relatively independent load environment and geometric model and is less affected by the load change [5, 6]. It is suggested that left atrial strain should be used in diagnosis of LVDD [7]. The previous study indicated that LA strain correlates well with LVEDP, as well as pulmonary capillary wedge pressure [7–10]. While LASct and LAScd are rarely mentioned in
previous studies, cut-off value for LA strain is still undefined by invasive gold reference. Our study intended to explore the correlation between LA strain and LV diastolic dysfunction in patients with normal LVEF.

**Methods**

**Population**

Between June 2018 and November 2019, we prospectively studied 55 patients with LVEF ≥ 50% (mean age, 63 years old, 38 men [69.1%]) referred for left heart catheterization, and echocardiography was completed within 12 h before or after catheterization. Patients with confirmed or suspected coronary artery disease underwent left heart catheterization. Those with Atrial fibrillation, ST-segment elevation myocardial infarction (STEMI), non-ST-segment elevation myocardial infarction (NSTEMI), moderate or greater tricuspid regurgitation, moderate or greater mitral regurgitation, any mitral or aortic stenosis, prosthetic valves, hemodynamic instability or poor echocardiography imaging were excluded. The study was approved by the institutional enrolling board of the first affiliated hospital of Soochow university.

**Echocardiography**

Echocardiographic measurements were performed by two board-certified echocardiographers blinded to the LVEDP who finalized each measurement by consensus. The two-dimensional echocardiographic imaging of all subjects was performed by GE Vivid E9 and GE Vivid E95 equipment (Norway) 2.5 MHz transducer. The LA dimension was measured in the parasternal long-axis view at the ventricular end-systole. Mitral diastolic inflow was interrogated using pulsed-wave Doppler from the apical four-chamber view at the level of the mitral leaflet tips. Mitral early diastolic peak (E-wave) and late peak (A-wave) velocities and E/A ratio were measured. In the apical two and four-chamber views including the entire left atrium, LAV was determined using biplane Simpson method at end-systolic frames just before mitral valve opening (maximal left atrial volume, LAVmax), at end-diastolic frames coincided with the R-wave on the electrocardiogram (minimal left atrial volume, LAVmin) and during mid-diastole phase which is before echocardiographic P-wave (left atrial volume before A wave, LAVpre-A). Values were then indexed to the body surface area (LAVmax, LAVmin and LAVpre-A). left atrial ejection fraction (LAEF) were calculated by LAVmax minus LAVmin. In addition, the septal mitral annulus early (E’) velocity was measured by tissue Doppler imaging, and the E/E’ ratio was calculated using a cutoff value > 15 to represent elevated LV filling pressure. LVEF was measured using Simpson’s method, which was used as a standard index of global LV systolic function. All echocardiographic measurements used in the analysis were averaged from 3 heart beats.

**LA longitudinal strain analysis**

The two-dimension strain analysis package available on the Echo PAC work station (GE Healthcare) was used to measure the LA strain and strain rate. Similar to the left ventricle, the complete myocardial region of interest (ROI) of the LA is defined by the endocardial border. An adjustable ROI with a default width of 3 mm is recommended. The user can adjust the size and shape to include the thickness of LA wall and to avoid including the pericardium. The software divided the atrial endocardium into six segments, poorly displayed segments were automatically rejected by the software and excluded from the analysis. The global longitudinal strain and strain rate curves were generated by the software for each apical view. The operator could repeat the imaging or change software parameters such as the width of the region of interest and the smoothing functions to obtain satisfactory tracking. Apical four and two-chamber views were obtained in three consecutive heart beats, and electrocardiograph was recorded in the meantime. The LA strain is defined as the absolute strain value in LA three phases, which includes reservoir strain in systole (LASr), conduit strain in early diastole (LASct) and contraction strain in late diastole (LAScd). LV end-diastole was defined as initial zero reference point. The respective strains are LASr, calculated as difference between onset of filling and end-diastole; LAScd, calculated as difference between onset of atrial contraction (before start of Doppler A-wave) and onset of filling; LASct, calculated as difference between end-diastole and onset of atrial filling. The global strain was calculated as the average of both respective values form both views. (Fig. 1).

**Invasive LV pressure measurements**

A 6F pigtail catheter was placed in the left ventricle to obtain the invasive LV pressure. Excluding the unstable baseline of LV filling pressure, LVEDP was measured by mean values of 3–5 cardiac cycles. Image I was used to measure LVEDP at the beginning of QRS on Electrocardiography. A LVEDP value > 16 mmHg was defined as elevated LV filling pressure [3, 11]. LVEDP values were measured by two investigators blinded to echocardiographic data.

**Statistical analysis**

Analyses were performed using the SPSS 25.0. Continuous variables which are normally distributed are presented as mean ± SD. Variables that are not normally distributed
will be presented as median with inter-quartile ranges (IQR = 25th–75th percentile). T test and Mann–Whitney U test was used for comparison of groups. Categorical variables are expressed as absolute numbers and respective percentages, categorical variables were compared with chi squared tests. LA strain and LV diastolic dysfunction parameters such as septal e’ velocity, lateral e’ velocity, E/e’, E/A, LAEF, tricuspid regurgitation velocity, LAVI and LVEDP were evaluated using univariate logistic regression analysis. Differences with p values less than 0.05 were considered significant. The independent predictive value of echocardiographic variables was evaluated in three different multivariate models. Predictors of elevated LVEDP with p value ≤ 0.12 in the univariate analysis and age, were included in multivariate models. In our three models, LASr LASct and LAScd were analyzed separately due to their multicollinearity, with other control variables the same. The C-statistic was calculated in each model, in order to allow comparison between them. Receiver operating characteristics (ROC) curves were created to compare the performance of multiple variables in determining increased LVEDP.

Results

Characteristics of population

Patients’ clinical characteristics and echocardiographic measurements are shown in Table 1. LVEDP was elevated in 35 patients (63.6%), and normal in 20 (36.4%) patients. The study population (n = 55) has a mean age of 63 ± 10.0 years, among which 17 (31%) were female. Most characteristics are similar in both groups, such as age, diabetes, CAD etc. Significant differences between the two groups were observed in LA strain (LASr, LASct, LAScd). Vital signs were not significantly different when echocardiography and catheterization were performed.
Table 1  Characteristics of patients in two groups defined by LVEDP

| Variable                          | LVEDP ≤ 16 mmHg (n = 20) | LVEDP > 16 mmHg (n = 35) | p   |
|-----------------------------------|---------------------------|--------------------------|-----|
| Male                              | 15 (75%)                  | 23 (66%)                 | 0.48|
| Age                               | 63.8 ± 8                  | 62.4 ± 11.0              | 0.63|
| BMI                               | 24.1 ± 3.1                | 24.5 ± 3                 | 0.83|
| HR (beats/min)                    | 63.6 ± 10                 | 68.1 ± 9                 | 0.10|
| Medical history                   |                           |                          |     |
| HTN                               | 12 (60%)                  | 29 (83%)                 | 0.06|
| Diabetes                          | 4 (20%)                   | 10 (29%)                 | 0.50|
| CAD                               | 9 (45%)                   | 11 (31%)                 | 0.32|
| COPD                              | 1 (5%)                    | 1 (3%)                   | 0.69|
| CKD (stage ≥ 3) or ESRD           | 1 (5%)                    | 1 (3%)                   | 0.69|
| Medications before catheterization|                           |                          |     |
| Diuretic                          | 2                         | 3                        | 1.00|
| Calcium blocker                   | 5                         | 15                       | 0.19|
| BB                                | 6                         | 17                       | 0.18|
| ACEI                              | 5                         | 8                        | 0.86|
| ARB                               | 2                         | 3                        | 1.00|
| Angiographic findings             |                           |                          |     |
| Stenosis                          |                           |                          |     |
| None significant stenosis         | 9                         | 21                       | 0.28|
| < 50% of diameter                 | 3                         | 5                        | 1.00|
| > 50% of diameter                 | 6                         | 4                        | 0.18|
| Total occlusion                   | 2                         | 5                        | 0.97|
| Numbers of vessels with significant stenosis > 50% |   | | |
| 1                                 | 4                         | 6                        | 1.00|
| 2                                 | 3                         | 1                        | 0.26|
| 3                                 | 1                         | 2                        | 1.00|
| Vessels with stenosis > 50%       |                           |                          |     |
| LM                                | 1                         | 1                        | 1.00|
| LAD                               | 7                         | 7                        | 0.34|
| LCX                               | 3                         | 1                        | 0.26|
| RCA                               | 1                         | 3                        | 1.00|
| Echocardiographic parameters      |                           |                          |     |
| Mitral E (cm/s)                   | 66.1 ± 10.3               | 68.8 ± 14.7              | 0.47|
| Mitral A (cm/s)                   | 76.8 ± 17.8               | 87.2 ± 21.6              | 0.07|
| E/A ratio                         | 0.8 (0.7–0.8)             | 0.8 (0.6–0.9)            | 0.19|
| Septal e’ velocity (cm/s)         | 6.7 ± 2.0                 | 6.2 ± 1.9                | 0.356|
| Lateral e’ velocity (cm/s)        | 9.4 ± 2.5                 | 8.6 ± 3.1                | 0.385|
| E/e’ ratio                        | 8.5 (6.9–10.8)            | 9.9 (8.1–11.1)           | 0.24|
| TR peak velocity (m/s)            | 2.3 (2.1–2.5)             | 2.3 (2.1–2.6)            | 0.85|
| LVEF (%)                          | 66.7 ± 8.9                | 64.6 ± 7.2               | 0.95|
| LAVimax (ml/m²)                   | 26.9 ± 9.5                | 27.7 ± 9.6               | 0.89|
| LAVimin (ml/m²)                   | 10.6 (7.6–16.9)           | 10.8 (7.7–13.2)          | 1.00|
| LAVi pre-A (ml/m²)                | 17.8 ± 6.8                | 19.8 ± 6.7               | 0.54|
| LAEF (%)                          | 54.5 ± 8.1                | 55.3 ± 10.3              | 0.772|
| LASr (%)                          | 35.2 ± 7.7                | 21.3 ± 7.2               | < 0.001|
| LASct (%)                         | 17.6 ± 6.3                | 11.9 ± 4.1               | < 0.001|
| LAScd (%)                         | 14.7 (12.2–19.7)          | 9.6 (5.8–12.4)           | < 0.001|

Data are expressed as numbers or as mean ± SD. See text for details

LVEDP left ventricular end-diastolic pressure, HR heart rate, CAD coronary artery disease, CKD chronic kidney disease, COPD chronic obstructive pulmonary disease, ESRD end-stage renal disease, HTN hypertension. BB β-blocker, ACEI angiotensin-converting enzyme, ARB angiotensin II receptor blocker, LM left main coronary artery, LAD left anterior descending coronary artery, LCX left circumflex coronary artery, RCA right coronary artery, TR tricuspid regurgitation, LVEF left ventricular ejection fraction, LAVi left atrium volume index, LAEF left atrial ejection fraction, LASr LA reservoir strain, LASct LA conduit strain, LAScd LA contraction strain
Correlation between echocardiography assessment and LVEDP

The LA strain and LVEDP are correlated negatively: LASr \( (p < 0.001, R = 0.662) \), LASct \( (p < 0.001, R = 0.575) \), LAScd \( (p < 0.001, R = 0.456) \). No significant correlation was found between the LVEDP and E/e′ \( (p = 0.493) \). Figure 2 is scatter diagram of LA strain and E/e′. LASr has a stronger correlation with LVEDP, compared to other variables observed in our study.

Table 2 Multivariate regression analysis to identify predictors for elevated LVEDP

| Model | Variables | Univariate analysis | Multivariate analysis | C-statistic |
|-------|-----------|---------------------|-----------------------|-------------|
|       |           | OR (95% CI) p       | OR (95% CI) p         |             |
| 1     | LASr     | 0.76 (0.67–0.89) <0.001 | 0.75 (0.64–0.88) 0.001 | 0.95        |
|       | Age      | 0.98 (0.92–1.04) 0.42 | 0.93 (0.81–1.06) 0.29 |
|       | Mitral A | 1.02 (1.00–1.06) 0.12 | 1.06 (0.99–1.13) 0.12 |
|       | HR       | 1.07 (1.00–1.14) 0.06 | 1.04 (0.93–1.17) 0.51 |
|       | HTN      | 0.36 (0.10–1.27) 0.11 | 0.22 (0.03–1.73) 0.15 |
| 2     | LASct    | 0.78 (0.65–0.92) 0.003 | 0.72 (0.57–0.90) 0.004 | 0.86        |
|       | Age      | 0.98 (0.92–1.04) 0.42 | 0.95 (0.86–1.04) 0.27 |
|       | Mitral A | 1.02 (1.00–1.06) 0.12 | 1.05 (0.99–1.11) 0.09 |
|       | HR       | 1.07 (1.00–1.14) 0.06 | 1.04 (0.95–1.14) 0.42 |
|       | HTN      | 0.36 (0.10–1.27) 0.11 | 0.18 (0.04–0.92) 0.04 |
| 3     | LAScd    | 0.76 (0.64–0.90) 0.001 | 0.71 (0.57–0.89) 0.003 | 0.93        |
|       | Age      | 0.98 (0.92–1.04) 0.42 | 0.93 (0.81–1.06) 0.27 |
|       | Mitral A | 1.02 (1.00–1.06) 0.12 | 1.03 (0.98–1.08) 0.21 |
|       | HR       | 1.07 (1.00–1.14) 0.06 | 1.08 (0.98–1.20) 0.11 |
|       | HTN      | 0.36 (0.10–1.27) 0.11 | 0.39 (0.07–2.12) 0.27 |

CI confidence interval, HR heart rate, LASr LA reservoir strain, LASct LA conduit strain, LAScd LA contraction strain, HTN hypertension

Fig. 2 Scatter plots of variables with correlation to LVEDP, respectively LASr, LASct, LAScd and E/e′ ratio.
The predictive values of LAS\textsubscript{r}, LAS\textsubscript{ct} and LAS\textsubscript{cd} were evaluated in three different multivariate logistic regression analyses (Table 2), which contained the same variables (age, Mitral A, HR and hypertension), differing only by control variables. The LAS\textsubscript{r}, LAS\textsubscript{ct} and LAS\textsubscript{cd} were independent predictors for LVEDP > 16 mmHg in their respective models. The model shows that LAS\textsubscript{r} has a higher C-statistic when compared to the model with LAS\textsubscript{ct} and LAS\textsubscript{cd}.

**The accuracy of LVEDP > 16 mmHg predicted by LA strain**

Figure 3 shows that LA strain has good diagnostic accuracy for LVEDP > 16 mmHg. The area under the curve (AUC) for LAS\textsubscript{r} is 0.914 (cutoff value is 26.7%, sensitivity is 90%, specificity is 82.9%). For LAS\textsubscript{ct}, the AUC is 0.769 (cutoff value is 12.0%, sensitivity is 85%, specificity is 57%). The AUC of LAS\textsubscript{cd} is 0.844 (cutoff value is 11.0%, sensitivity is 90.0%, specificity is 68.6%), which shows that LAS\textsubscript{r} can predict LVEDP better than LAS\textsubscript{ct} and LAS\textsubscript{cd}.

**Discussion**

Left atrial strain presents significant feasibility and reproducibility, which has recently emerged as a powerful assessment in evaluation of left ventricular diastolic dysfunction [4]. We aim to evaluate the diagnostic accuracy of LA strain for LVDD in patients with normal LVEF by gold standard reference. In this study we demonstrated that left atrial strain was significantly impaired in the group with elevated LVEDP in patients with LVEF ≥ 50%. LAS\textsubscript{r} showed a stronger correlation with LVEDP than with LAS\textsubscript{ct}, LAS\textsubscript{cd} and other conventional diastolic echocardiography parameters. LA reservoir strain at cut-off of 26.7% predicted LVEDP > 16 mmHg
with 90% sensitivity and 82.9% specificity, which performed a higher diagnosis accuracy than LASct, LAScd and E/e’.

In the early phase of LVDD, reduced ventricular compliance and elevated LVEDP result in decreasing passive early transmitral diastolic flow, then atrial pump function enhances to compensate LV filling. As left ventricular diastensibility declines further, atrial pressure rises to maintain cardiac output, which may blunt the compliance of the LA. LA function is impaired as a consequence of chronically elevated LVEDP and decreased LA compliance [12]. This may be the potential mechanism for the inverse correlation between LA strain and LVEDP.

Multiple recent studies have demonstrated the correlation between LA strain and hemodynamic parameters [8, 9], but for patients with LVEF ≥ 50%, investigations with sufficient sample size about correlation between LA strain and LVEDP obtained by invasive gold standard reference, are still needed, and a definitive cut-off value for abnormal LA strain is still undefined.

Our study is consistent with the findings observed by Cameli and colleagues. They found that LASr was strongly negatively correlated with pulmonary capillary wedge pressure (PCWP) in patients with heart failure, which performed excellent sensitivity and specificity of 100% and 93% in assessing PCWP [9]. However, LVEDP > 16 mmHg was defined as LVDD instead of PCWP ≥ 18 mmHg in our study, which may recognize the patients at early stage of LVDD, because elevated PCWP only occurs at a decompensated LA when LVEDP increases significantly. Previous study showed that PCWP frequently underestimates LVEDP, and it only had a moderate ability to discriminate patients with normal or elevated LVEDP [13].

Amita Singh et al. have reported that LASr could predict elevated filling pressure with high accuracy and identified a cutoff value for LASr [8], while we have a larger sample size which may provide a more convincing evidence for cut off value of LA strain in patients with normal LVEF. Besides LASr, our study also demonstrated the correlation between LASct, LAScd and LVEDP. In our study LA strain presented a more sensitive diagnostic value for LVDD than conventional diastolic echocardiography assessments such as E/e’, which indicates that LA strain can recognize patients with LVDD even at early stage.

Poor correlation between E/e’ and LVEDP was observed in this study, which seems to contradict previous reports that E/e’ correlated with PCWP or LVEDP significantly [9, 10, 14, 15]. Possible explanations are as follows. Firstly, Previous study showed that E/e’ between 8 and 15 predicts LV Filling Pressure with poor accuracy [16]. However, 34 (61.8%) patients’ E/e’ ratio is between 8 and 15 in our study. Secondly, the accuracy of E/e’ ratio is uncertain in patients with advance decompensated HF, presence of cardiac resynchronization therapy, mitral annular calcification, surgical rings, prosthetic valves, several mitral regurgitations, atrioventricular or intraventricular conduction delay and symptomatic hypertrophic cardiomyopathy [4, 14, 17].

Our study showed that LAVI correlated with LVEDP poorly. LA dilatation usually implies the chronic, long-term elevation of LA pressure, and LAV can be normal in the early phase of LVDD [18, 19]. Many studies have suggested that LA strain may be superior to LAVI in evaluating LVDD even in the absence of LA enlargement [5, 6], which indicated that LA strain has better sensitivity. Moreover, LAV rarely normalizes with the reduction of LV filling pressure, and strong correlation of reduced LV filling pressure and improved LA strain demonstrates that LA strain may evaluate LA function remodeling better than LA size [20].

Current echocardiographic parameters show moderate sensitivity in evaluating LVDD, with the presence and severity of LVDD failing to be determined for a significant proportion of patients. LA strain measured by speckle tracking echocardiography can reclassify patients even in the early phase of LVDD. LA strain also has advantages when diagnosing LVDD in mitral prosthesis, mitral annular calcification, bundle branch block, post-cardiac surgery and AF. LA strain, with high reproductivity and feasibility, is suggested to be incorporated into routine evaluation of LVDD, especially for patients who are classified in the indeterminate LVDD group using the conventional recommended algorithm.

Limitations

The sample size in this study is limited, so selection bias still exists, however increasing the number of individuals will reduce the selection bias. The study was conducted on patients from only one medical center, which might impact the population selected and, consequently, the variables analyzed. Left atrium imaging is difficult to obtain and the image definition is poor in some cases, and some echocardiographic images have poor repeatability in left atrial strain.

Conclusion

For patients with normal LV ejection fraction, left atrial strain presented better correlation with LVEDP than E/e’. Moreover, LASr presented a better diagnostic value in predicting LVEDP than LASct and LAScd. It is suggested that left atrial strain should be used in diagnosis of LVDD.

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Compliance with ethical standards

Conflict of interest The authors declare that they have no conflict of interests.

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