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Assessment of Left Ventricular Dyssynchrony after Permanent Cardiac Pacing by Using Two Dimensional Speckle Tracking Echocardiography

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Abstract

Background: Echocardiography is important in assessing left ventricular mechanical dyssynchrony (LVMD) and left ventricular (LV) function after implant of a permanent pacemaker. Global longitudinal strain (GLS) and Left ventricular dyssynchrony assessment enable clinicians to detect early signs of LV dysfunction after cardiac pacing. This study assessed LV mechanical asynchrony and LV function in different pacing modes after permanent cardiac pacing by using tissue Doppler imaging (TDI) and speckle tracking echocardiography (STE).

Patients and Methods: Seventy female (42) and male (28) patients were enrolled in this prospective observational case study (mean age 60.99 ± 13.77 years) at Mansoura Specialized Medical Hospital over a period of 1 year from April 2018 to April 2019. All the patients were assessed by thorough history taking, clinical examination, conventional Echocardiography, TDI and two-dimensional (2D) STE. Results: Regarding parameters assessed by STE, there was a significant decrease in global longitudinal strain GLS $(P = 0.034^*)$ while there was a significant increase in time to peak strain standard deviation (TP-SD) by STE $(P < 0.001^*)$. Also, there was significant decrease in GLS $(P < 0.001^*)$ and significant increase in TP-SD by STE $(P = 0.001^*)$ in dual chamber pacemaker (DDD) group. Similarly, there was a significant decrease in GLS $(P < 0.001^*)$ and a significant increase in TP-SD by STE $(P < 0.001^*)$ in ventricular demanding pacing (VVI) group. Conclusion: The results show that GLS by 2D STE can help in detection of subclinical left ventricular dysfunction (LVD) after permanent pacemaker implantation before appearance of clinical symptoms. In addition, cardiac pacing can still lead to LVMD irrespective of the degree of the pacing mode. However, this may need upgrading later on to cardiac resynchronization therapy (CRT).

Keywords: LVMD, GLS, speckle tracking echocard-iography

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Introduction

The choice of an optimum pacing mode for the management of low heart rate is essential. In presence of disease in sino-atrial node dual chamber pacing is of choice [1]. The various pacing modes can lead to several problems due to some electrical changes leading to difference between timing in contraction of both ventricles in the heart which may cause cardiac failure and sudden cardiac death [2]. The artificial cardiac pacing via pacemaker alters the normal heart electrical activation when compared to the native activation resulting from atrioventricular (AV) nodal and His bundle system stimulation [3]. The pacing at the right ventricle causes abnormal contraction and decreased pump function. Right ventricular apical pacing can initiate asynchrony among LV interventricular septum (IVS) and free wall and also between right ventricle (RV) and LV [4]. It has also been showed that the existence of ventricular asynchrony is accompanied by a high risk of cardiac morbidity and death [5].

Echocardiography is essential in evaluating left ventricular asynchrony and function. Currently, realtime three dimensional echocardiography (RT3DE) and tissue Doppler imaging are of great sensitivity and frequently utilized procedures in order to quantify left ventricular mechanical asynchrony [6]. Speckle tracking echo-cardiography is recent modality depends on 2D grey scale imaging which helps in assessing left ventricular asynchrony. Suffoletto et al. [7] described the use of speckle tracking radial strain in quantifying asynchrony, which is the time difference in peak antero-septum to posterior wall strain more than or equal to 130 ms. Moreover, global longitudinal strain can help the cardiac clinicians to find out early signs of left ventricular dysfunction. Defining mechanisms of asynchrony will be useful for pacemaker programing choices in order to prevent further dysfunction of left ventricular [8]. The aim of this study was to evaluate left ventricular mechanical dyssynchrony and left ventricular function in various pacing modes after permanent cardiac pacing by using tissue Doppler imaging and speckle tracking echocardiography and to correlate these changes with other clinical, electrocardiographic and echocardiographic data.

Subjects and Methods

Ethics Statement

All procedures were performed as recommended by the guidelines as regard conventional and speckle tracking Echocardiography. The study was explained to all patients and they gave oral informed consent. Besides, the study was approved by the Ethics Committee of the Faculty of Medicine, Mansoura University. Informed consent was obtained from each individual participant involved in this study. This study was conducted in accordance with the 1964 Declaration of Helsinki and its subsequent amendments.

Study Population

Seventy (n = 70) patients were enrolled in this prospective observational case study, (mean age 60.99 ± 13.77 years, 28 males and 42 females), at Mansoura Specialized Medical Hospital over a period of 1 year from April 2018 to April 2019. Patients were enrolled into 2 groups. The first group had 30 patients with implanted single chamber pacemaker (Group A) while the second group had 40 patients with implanted Dual chamber pacemaker (Group B).

Methodology

All patients were assessed by thorough history taking including; age and gender, risk factors for coronary artery disease; hypertension, diabetes, dyslipidemia. Clinical examination included: blood pressure (BP), pulse, general examination and local cardiac examination. Investigations included: 12 lead surface electrocardiography (ECG), Echocardiography, TDI and 2D speckle tracking echocardiography.

Echocardiography

All of the patients had echocardiography using the General Electric Vivid E9 XD clear Dimensions ultrasound system (GE Healthcare, USA) using the matrix M5Sc transducer. With more stress on the following pulsed wave Doppler (PWD), TDI, septal posterior wall motion delay (SPWMD) by M mode, aortic pre-ejection delay, inter-ventricular mechanical delay, tissue synchronization imaging (TSI) were done to assess septal to lateral delay, septal to posterior delay and SD-Ts. Similarly, 2D speckle tracking echocardiography was done to assess TP-SD and global longitudinal strain via the same machine.

Exclusion Criteria

Exclusion criteria include:

- Cases with bad echocardiography window, cases with manifestations of obvious heart failure (HF), preceding heart operation, dilated cardiomyopathy, prosthetic valves and cases with preceding coronary artery disease (CAD) identified by proof of left ventricular wall motion aberrations at the echo or pathological Q waves in ECG, previous percutaneous coronary intervention (PCI) or coronary artery bypass grafting (CABG).
- 2. Previous implanted permanent pacemaker or implantable cardioverter defibrillator (ICD).
- 3. Frequent premature ventricular complexes (PVCs) in ECG
- 4. Patients with terminal co-morbidities such as end stage malignancy, end stage renal or liver diseases.

Statistical Analysis

The clinical and laboratory data were recorded on an "Investigation report form." These data were tabulated, coded then analyzed using the computer program SPSS (Statistical package for social science) version 25 to obtain the descriptive data.

In turn, the descriptive statistics data were calculated for the anthropometric measurements and laboratory data in the form of: Mean ± Standard deviation (SD). Frequency (Number-percent)

Analytical Statistics

- In the statistical comparison between the different groups, the significance of difference was tested using the following test:
 - Student's t-test: Used to compare between mean of two groups of numerical (parametric) data. Inter-group comparison of categorical data was performed by using chi square test (X2-value).
 - Paired t-test was used for comparison within groups
 - Wilcoxon singed ranks test (Z) was used for comparison within group
 - A P value of < 0.05 was considered statistically significant and a P value of
 < 0.0001 was considered highly significant (HS) in all analyses.

Results

The study included 70 patients (mean age 60.99 \pm 13.77 years) who were divided into two groups. The first group A of patients (mean age 65.7 ± 12.3 years) undergone pacing mode single chamber pacemaker VVI while the second group B of patients $(57.45 \pm 13.89 \text{ years})$ had pacing mode dual chamber pacemaker. The mean demographic clinical characteristics and major risk factors of CAD of each group was assessed at 1 month and after 3 months of permanent pacemaker implantation (Table 1). The study recruited patients with the following risk factors of cardiovascular diseases (CVD). They included 11 patients (15.7%) who were smokers, 46 patients (65.7%) who were hypertensive, 9 patients (12.9%) who were dyslipidemic and 11 patients (15.7%) who were diabetic patients. Table 1 shows demographic clinical characteristics and major risk factors of CAD in the whole studied population while Table 2 demonstrates baseline criteria among the two studied groups.

| | All patients $(n = 70)$ | All patients $(n = 70)$ | |
|----------------------|-------------------------|-------------------------|--|
| | No | % | |
| Age | | | |
| <40y | 6 | 8.6% | |
| 40-60y | 28 | 40% | |
| >60y | 36 | 51.4% | |
| Mean \pm SD | 60.99 ± 13.77 | 7 | |
| Sex | | | |
| Male | 28 | 40% | |
| Female | 42 | 60% | |
| Hypertension | 46 | 65.7% | |
| Diabetes | 11 | 15.7% | |
| Smoking | 11 | 15.7% | |
| Dyslipidemia | 9 | 12.9% | |
| Beta blocker | 10 | 14.3% | |
| ACEI | 38 | 54.3% | |
| Diuretic | 31 | 44.3% | |
| Aspirin | 13 | 18.6% | |
| Warfarin | 4 | 5.7% | |
| Congested neck veins | 18 | 25.7% | |
| Edema L.L | 29 | 41.4% | |
| Dyspnea | 18 | 25.7% | |

Table 1. Baseline demographic, clinical characteristics and major risk factors of CAD among the 70 patients employed in the study

Abbreviations; ACEI: angiotensin converting enzyme inhibitor, L.L: lower limb.

SD: Standard Deviation.

Data are expressed in number abs as percentages.

Table 2. Baseline, clinical characteristics and major risk factors of CAD among the two studied groups (VVI versus DDD group) of patients

| | Group B (n = 40) | | Group A (n = 30) | | χ2 | Р |
|----------------------|---------------------|-------|---------------------|-------|-------|--------|
| | No | % | No | % | | |
| Hypertension | 27 | 67.5% | 19 | 63.3% | 0.132 | 0.716 |
| Diabetes | 8 | 20% | 3 | 10% | 1.294 | 0.255 |
| Smoking | 10 | 25% | 1 | 3.3% | 6.076 | 0.014* |
| Dyslipidemia | 6 | 15% | 3 | 10% | 0.383 | 0.536 |
| Betablocker | 8 | 20% | 2 | 6.7% | 2.489 | 0.115 |
| ACEI | 24 | 60% | 14 | 46.7% | 1.228 | 0.268 |
| Diuretic | 17 | 42.5% | 14 | 46.7% | 0.121 | 0.728 |
| Aspirin | 10 | 25% | 3 | 10% | 2.551 | 0.110 |
| Warfarin | 0 | 0% | 4 | 13.3% | 5.657 | 0.017* |
| Congested neck veins | 10 | 25% | 8 | 26.7% | 0.025 | 0.875 |
| Edema L.L | 18 | 45% | 11 | 36.7% | 0.491 | 0.484 |
| Dyspnea | 12 | 30% | 6 | 20% | 0.897 | 0.343 |

Abbreviations; LL: lower limb, ACEI: angiotensin converting enzyme inhibitor, N: number, P: probability value, *: significant value> 0.05, X2: chi square, %: percentage.

Data are expressed as number and percentages with $\chi 2$ and p values *P < 0.05.

| | 1 month | 3 month | t | Р |
|--------------------------------|--------------------|--------------------|--------|----------|
| | (n = 70) | (n = 70) | t | |
| E DT | 230.41 ± 79.65 | 228.31 ± 84.53 | 0.317 | 0.752 |
| E/e' ratio | 11.07 ± 4.4 | 9.24 ± 3.65 | 9.844 | < 0.001* |
| Tei index | 0.69 ± 0.2 | 0.78 ± 0.22 | 5.113 | < 0.001* |
| LAD | 3.97 ± 0.57 | 4.03 ± 0.56 | 3.308 | 0.001* |
| LVEDD | 5.45 ± 0.61 | 5.67 ± 0.66 | 8.345 | < 0.001* |
| LVESD | 3.75 ± 0.59 | 3.95 ± 0.66 | 7.415 | < 0.001* |
| EF | 60.36 ± 8.39 | 56.3 ± 8.6 | 10.339 | < 0.001* |
| FS | 32.53 ± 5.74 | 29.63 ± 5.74 | 8.887 | < 0.001* |
| APED | 136.56 ± 27.97 | 151.11 ± 25.44 | 6.729 | < 0.001* |
| IVMD | 30.04 ± 21.78 | 36.61 ± 21.58 | 4.416 | < 0.001* |
| SPWMD by M Mode | 79.43 ± 36.6 | 93.71 ± 46.88 | 3.698 | < 0.001* |
| SD Ts by TSI | 53.89 ± 19.14 | 59.8 ± 18.37 | 3.632 | 0.001* |
| Septal posterior delay by TSI | 30.41 ± 74.11 | 25.7 ± 84.14 | 0.404 | 0.688 |
| All segments Max delay by TSI | 164.13 ± 54.2 | 174.99 ± 50.83 | 2.262 | 0.027* |
| Septal to lateral delay by TSI | 23.63 ± 77.53 | 15.59 ± 88.07 | 0.634 | 0.528 |
| GLS by STE | -15.96 ± 7.29 | -14.25 ± 3.4 | 2.160 | 0.034* |
| TP SD by STE | 35.04 ± 20.47 | 47.94 ± 23.92 | 5.522 | < 0.001* |

Table 3. Comparative analysis of echocardiographic parameters after 1 and 3 months of permanent pacemaker implantation among the whole studied population of 70 patients.

Abbreviations; N: number, T: student t-test, P: probability value, *: significant value< 0.05, DT: deceleration time, LAD: left atrial dimension, LVEDD: left ventricular end diastolic dimension, LVESD: left ventricular end systolic dimension, EF: ejection fraction, FS: fractional shortening, APED: aortic pre-ejection delay, IVMD: inter-ventricular mechanical delay, SPWMD: septal posterior wall motion delay, TSI: tissue synchronization imaging, SD-Ts: standard deviation of time to peak systolic velocity, GLS: global longitudinal strain, STE: speckle tracking echocardiography, TP SD: standard deviation of time to peak strain.

Data are presents as mean \pm SD with t and P values; *P < 0.001.

Table 4. Comparative analysis of different echocardiographic parameters after 1 and 3 months of pacemaker implantation in DDD group comprising of 40 patients

| DDD | 1 month | 3 month | Т | Р |
|--------------------------------|--------------------|--------------------|-------|-----------|
| | (n = 40) | (n = 40) | - | • |
| E DT | 215.35 ± 69.74 | 225.28 ± 75.44 | 1.280 | 0.208 |
| E/e' ratio | 10.23 ± 3.85 | 8.58 ± 2.99 | 6.570 | < 0.001** |
| Tei index | 0.73 ± 0.22 | 0.78 ± 0.21 | 3.668 | 0.001** |
| LAD | 3.85 ± 0.52 | 3.92 ± 0.56 | 2.421 | 0.020* |
| LVEDD | 5.47 ± 0.62 | 5.72 ± 0.66 | 6.520 | < 0.001** |
| LVESD | 3.81 ± 0.63 | 4.01 ± 0.71 | 5.965 | < 0.001** |
| EF | 58.93 ± 8.61 | 54.75 ± 8.84 | 7.318 | < 0.001** |
| FS | 31.35 ± 5.74 | 28.7 ± 5.77 | 5.239 | < 0.001** |
| APED | 133.63 ± 28.71 | 148.25 ± 27.01 | 5.689 | < 0.001** |
| IVMD | 28.68 ± 22.72 | 36.35 ± 23.78 | 5.476 | < 0.001** |
| SPWMD by M Mode | 73.75 ± 31.78 | 89.5 ± 44.43 | 3.348 | 0.002* |
| SD Ts by TSI | 53.9 ± 19.16 | 59.4 ± 18.45 | 2.460 | 0.018* |
| Septal posterior delay by TSI | 42.35 ± 70.63 | 28.35 ± 84.52 | 0.976 | 0.335 |
| All segments Max delay by TSI | 163.5 ± 53.47 | 172.1 ± 47.04 | 1.279 | 0.208 |
| Septal to lateral delay by TSI | 13.8 ± 74.53 | 3.48 ± 83.31 | 0.600 | 0.552 |
| GLS by STE | -15.41 ± 3.88 | -14.08 ± 3.93 | 6.058 | < 0.001* |
| TP SD by STE | 37.7 ± 21.04 | 49.25 ± 25.75 | 3.719 | 0.001* |

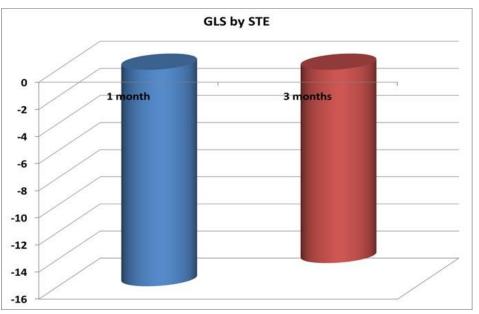
Abbreviations; N: number, T: student t-test, P: probability value, *: significant value> 0.05, DT: deceleration time, LAD: left atrial dimension, LVEDD: left ventricular end diastolic dimension, LVESD: left ventricular end systolic dimension, EF: ejection fraction, FS: fractional shortening, APED: aortic pre-ejection delay, IVMD: inter-ventricular mechanical delay, SPWMD: septal posterior wall motion delay, TSI: tissue synchronization imaging, SD-Ts: standard deviation of time to peak systolic velocity, GLS: global longitudinal strain, STE: speckle tracking echocardiography, TP SD: standard deviation of time to peak strain. Data are expressed mean \pm SD and with T and P values; *P < 0.05 and **P < 0.001.

| Group A | $ \begin{array}{c} 1 \text{ month} \\ (n = 30) \end{array} $ | | | Р |
|--------------------------------|--|--------------------|--------|----------|
| E DT | 250.5 ± 88.45 | 232.37 ± 96.54 | 1.656 | 0.108 |
| E/e' ratio | 12.19 ± 4.88 | 10.12 ± 4.28 | 7.544 | < 0.001* |
| Tei index | 0.65 ± 0.17 | 0.78 ± 0.24 | 3.830 | 0.001* |
| LAD | 4.12 ± 0.59 | 4.17 ± 0.54 | 2.475 | 0.019* |
| LVEDD | 5.41 ± 0.61 | 5.62 ± 0.66 | 5.162 | < 0.001* |
| LVESD | 3.67 ± 0.54 | 3.87 ± 0.6 | 4.434 | < 0.001* |
| EF | 62.27 ± 7.82 | 58.37 ± 7.94 | 7.509 | < 0.001* |
| FS | 34.1 ± 5.42 | 30.87 ± 5.56 | 9.119 | < 0.001* |
| APED | 140.47 ± 26.95 | 154.93 ± 23.08 | 3.849 | 0.001* |
| IVMD | 31.87 ± 20.7 | 36.97 ± 18.64 | 1.735 | 0.093 |
| SPWMD by M Mode | 87 ± 41.54 | 99.33 ± 50.17 | 1.883 | 0.070 |
| SD Ts by TSI | 53.88 ± 19.44 | 60.33 ± 18.56 | 2.701 | 0.011* |
| Septal posterior delay by TSI | 14.5 ± 76.81 | 22.17 ± 84.93 | 0.394 | 0.697 |
| All segments Max delay by TSI | 164.97 ± 56.06 | 178.83 ± 56.09 | 2.037 | 0.051 |
| Septal to lateral delay by TSI | 36.73 ± 80.76 | 31.73 ± 92.99 | 0.263 | 0.794 |
| GLS by STE | -17.11 ± 2.57 | -14.15 ± 2.42 | 16.118 | < 0.001* |
| TP SD by STE | 31.51 ± 19.47 | 46.18 ± 21.53 | 4.109 | < 0.001* |

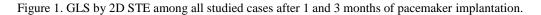
 Table 5. Comparative analysis of all echocardiographic parameters after 1 and 3 months of pacemaker implantation in group A of 30 patients

Abbreviations; N: number, T: student t-test, P: probability value, *: significant value< 0.05, DT: deceleration time, LAD: left atrial dimension, LVEDD: left ventricular end diastolic dimension, LVESD: left ventricular end systolic dimension, EF: ejection fraction, FS: fractional shortening, APED: aortic pre-ejection delay, IVMD: inter-ventricular mechanical delay, SPWMD: septal posterior wall motion delay, TSI: tissue synchronization imaging, SD-Ts: standard deviation of time to peak systolic velocity, GLS: global longitudinal strain, STE: speckle tracking echocardiography, TP SD: standard deviation of time to peak strain.

Data are expressed mean \pm SD and with T and P values; *P < 0.05 and **P < 0.00.



Data are expressed mean \pm SD.



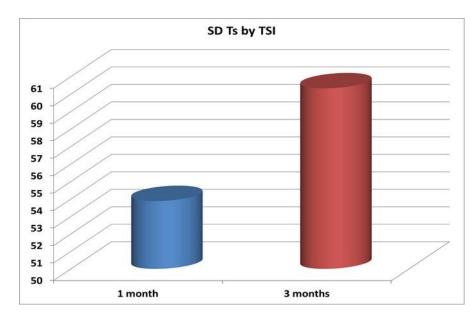


Figure 2. Standard deviation of time to peak systolic velocity by tissue synchronizing imaging after1 and 3 months of dual chamber pacemaker implantation.

Comparative analysis of echocardiographic parameters after 1 and 3 months of PPM implantation among the 70 patients.

Regarding diastolic function by pulsed wave Doppler PWD, the results show no significant change in E wave deceleration time (P = 0.752) while there was a significant decrease in E/e' ratio (P < 0.001*) and a significant increase in myocardial performance index Tei index (P < 0.001*).

Regarding internal dimensions and systolic function, the data show that there was significant increase in the following: left atrial dimension (LAD) (P = 0.001^*), left ventricular end systolic dimension LVESD (P < 0.001^*) and left ventricular end diastolic dimension LVEDD (P < 0.001^*), while there was significant decrease in ejection fraction EF (P < 0.001^*) and fractional shortening FS (P < 0.001^*).

Regarding parameters of LV dyssynchrony, the results reveal that there was a significant increase in aortic pre-ejection delay APED (P < 0.001^*), interventricular mechanical delay (P < 0.001^*) and septal posterior wall motion delay SPWMD by M mode (P < 0.001^*).

Regarding parameters assessed by tissue synchronization imaging TSI, the findings show that there was significant increase in the following: SD-Ts ($P = 0.001^*$) and all segments max delay ($P = 0.027^*$), while there was no significant change in

the following: septal posterior delay (P = 0.688) and septal to lateral delay (P = 0.528).

Regarding parameters assessed by STE, the results reveal that there was significant decrease in global longitudinal strain GLS (P = 0.034^*), while there was a significant increase in TP-SD by STE (P < 0.001^*).

The data further show that in both group A and group B studied groups there was a significant increase in most of LV dyssynchrony parameters and a significant decrease in LV GLS by 2D STE after 1 and 3 months of pacemaker implantation.

Discussion

The major finding of our study is that in short-term follow up of left ventricular function after permanent cardiac pacing. The results of this study have revealed that there was subclinical LV dysfunction by 2D STE in most of cases, either single or dual chamber pacing which needs close observation of such patients and long-term follow up for fear of pacing induced cardiomyopathy and beginning of heart failure symptoms appearance. Cardiac pacing is the single efficient management for manifested sinus node disorders and it can enhance manifested chronotropic incompetence [9]. Furthermore, several research studies have revealed enhancement in symptoms and function by cardiac pacing in cases with atrioventricular block [10].

Right ventricular (RV) pacing or left bundle branch block (LBBB) can lead to a left ventricular (LV) contraction shape which is dyssynchronous. Because of the out-of-phase contraction and relaxation of the LV septal and lateral walls, there is a loss of stroke work to internal power transmission from a contracting wall to the opposed wall that is in relaxation. Another result of the dyssynchronous LV contraction design is that the LV lateral wall contracts last resulting in overstretched and thus, accomplished a disproportionate fraction of the total stroke work [11].

The concept of 'pacing-induced' cardiomyopathy (CMP) has been applied to define the condition characterized by LV dilatation and hypokinesia which is frequently manifested with HF. In turn, this is accompanied with a high burden of RV pacing. However, pacing-induced CMP has not been especially involved in classification of cardiomyopathies [12].

Echocardiographic evaluation of LV dyssynchrony has been broadly utilized as it is noninvasive, extensively accessible and has no risk or complications. Currently, nearly all of the clinical evaluation modalities utilized tissue Doppler imaging (TDI) [13-14]. However, the most recent research study has employed the usage of speckle-tracking echocardiography (STE) and 3-dimensional (3D) echocardiography in tackling the clinical problems [15]. As such, this study evaluated LV mechanical dyssynchrony and LV function in different pacing modes after permanent cardiac pacing by using TDI and speckle tracking echocardiography. The study also showed that most of LV dyssynchrony parameters were affected (increased) after cardiac pacing whatever the pacing mode was either single or dual chamber pacing. In contrast, LV peak GLS was decreased after permanent pacemaker implantation after 1 and 3 months of implantation. The data also revealed that diastolic dysfunction seemed to be appeared among most of our patients.

The results of this study is in total agreement with those of Algazzar et al., [16] who found that mitral deceleration duration revealed marked changes among two modes of cardiac pacing in both group A and B, only at 6 month duration. Doppler designs to investigate of mitral influx was able to reveal the pressure gradient among the left atrium and LV. The trans-mitral speeds were closely correlated with preload and inversely correlated with ventricular relaxation.

The results also reveal a significant (P < 0.001^*) increase in myocardial performance index Tei index which is in agreement with Algazzar et al., [16] who found elevation in myocardial performance index (MPI) in the both groups with marked changes among them up to 6 months. In the present study, there was a significant increase in left atrial dimension (LAD) (P = 0.001^*), left ventricular end systolic dimension (LVESD) (P < 0.001^*) and left ventricular end diastolic dimension (LVEDD) (P < 0.001^*) while there was a significant decrease in ejection fraction EF (P < 0.001^*) and fractional shortening FS (P < 0.001^*).

Lieberman et al., [17] studied a group of cases with conserved LV ejection fraction (LVEF) and they found that RV apical pacing caused a modest reduction in LVEF, while the LV the measurements were still unaffected. In the present study, there was a significant increase in aortic pre-ejection delay APED (P < 0.001*), interventricular mechanical delay (P < 0.001*) and septal posterior wall motion delay SPWMD by M mode (P < 0.001*).

Alhous et al., [18] stated that in a study done on 25 cases, three showed SPWMD \geq 130 ms at starting point. With right ventricular apical pacing to elevate SPWMD, the cases with intra-ventricular dyssynchrony utilizing such approach was elevated to six. Also, right ventricular pacing led to elevated IVMD compared to starting point, and in five cases IVMD was \geq 40 ms. In the present study, there was a significant decrease in global longitudinal strain GLS (P = 0.034*) while there was significant increase in TP-SD by STE (P < 0.001*).

Moreover, our study revealed a significant increase in the following: SD-Ts ($P = 0.001^*$) and all segments with Max delay ($P = 0.027^*$). Our data are in complete agreement of those reported by Pastore et al., [19] who evaluated LV dyssynchrony by utilizing tissue Doppler echo at baseline and following at least one day of constant RV apical pacing. Employing total of 101 cases, their results showed marked LV dyssynchrony.

Dasgupta et al., [20] stated that myocardial deformation imaging may be a clinically useful tool for the prediction of a decline in LV systolic function following pacemaker implantation. Abnormalities in apical 4 chamber view (A4C) seem to appear before LVEF decline and as soon as 1-month post pacemaker implantation. On the contrary, El-Shabrawi et al., [21] found that chronic RV apical pacing in children after tetralogy of fallot repair was associated with better clinical status, preservation of RV systolic function, and prevention of progressive QRS prolongation.

In summary, it is of paramount importance to emphasize on the detection of subclinical left ventricular dysfunction after cardiac pacing which is so important on short and long term to recognize pacing- induced cardiomyopathy or ventricular dysfunction for possibility of upgrading conventional pacemaker to CRT to maintain LV function. Hence, the role of STE in measuring GLS became essential as it may detect LV dysfunction even with normal LVEF.

Conclusion

It is concluded that the new modalities in echocardiography as GLS and 2D STE can help in detection of subclinical left ventricular dysfunction after permanent pacemaker implantation before appearance of symptoms. Also, cardiac pacing can lead to LVMD whatever the pacing mode is which in turn may need later on upgrading to biventricular pacing.

Study Limitations

These include:

- 1. The small sample size was an important shortcoming of the current study. It should be noted that a limited number of patients met the inclusion criteria and were enrolled into the study. This small sample size could reduce the statistical power of our analyses. Short term follow up period.
- 2. Comparison of the different pacing modes only did not taking into consideration

different pacing sites (RV apical pacing; RV septal pacing either low, mid or high; His bundle pacing). Moreover, RT3DE and 3D STE needed for more evaluation. The need for correlation of study results regarding LV dyssynchrony with primary clinical end points as HF symptoms, the need for hospitalization and mortality.

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Ethical Compliance

The authors have stated all possible conflicts of interest within this work. The authors have stated all sources of funding for this work. If this work involved human participants, informed consent was received from each individual. If this work involved human participants, it was conducted in accordance with the 1964 Declaration of Helsinki. If this work involved experiments with humans or animals, it was conducted in accordance with the related institutions' research ethics guidelines.

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